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THE COMMENTARY
THE COMMENTARY
EUROPEAN COMMUNITIES – MEASURES AFFECTING ASBESTOS AND ASBESTOS-CONTAINING PRODUCTS

I. INTRODUCTION

The present case represents one of the high profile disputes handled by a WTO panel and the Appellate Body, and is an important precedent on product differentiation and WTO compatibility of health measures. Canada challenged the French Government’s 1996 ban on the manufacture, sale and import of all forms of asbestos and products containing them. Although the toxicity of asbestos is widely recognised and, in the words of the Appellate Body, ‘the value pursued by the measure is both vital and important in the highest degree’\(^1\), Canada claimed that chrysotile asbestos was safe under properly controlled use and therefore should not be prohibited outright.

The dispute received massive publicity and was attentively watched by non-governmental organisations and other non-state actors. These groups also tried to present pertinent information to the panel and the Appellate Body, in addition to the Parties’ submissions, as to how the WTO adjudicating bodies should decide on certain factual and legal issues in question. The panel and the Appellate Body were anxious to please all involved interests, but their treatment of amicus curiae briefs caused harsh reactions among both the Member States’ official delegates to the WTO as well as non-governmental organisations and other representatives of the civil society.

Both the panel and the Appellate Body report break new legal ground in manifold ways. Not only was the panel, and subsequently also the Appellate Body, called upon for the first time to explicitly rule on a complainant’s allegation that the TBT Agreement was violated, but the present dispute also marks the first ever under the WTO regime in which an otherwise GATT-inconsistent measure was determined by a panel or the Appellate Body to be justified under Article XX(b) of the GATT 1994. Furthermore, the appeal was the first occasion for the Appellate Body to examine Article XXIII:1(b) of the GATT 1994 regarding so-called “non-violation” complaints, and it took the opportunity to make some general observations on the relationship between Article XXIII:1(a) and XXIII:1(b) of the GATT 1994 and the nature of non-violation complaints in general.

Arguably, the most profound finding of the Appellate Body in the present case was the introduction of fundamental human interests such as health risks into the ‘like product’ test pursuant to Article III:4 of the GATT 1994, thus reversing the panel’s finding on this point. Particularly with

\(^1\) Appellate Body report, para. 172.
II. FACTUAL BACKGROUND

As of 1 January 1997, the French Government put into force Decree No. 96-1133 which established a total ban on asbestos fibres and products containing asbestos.2 The Decree, in essence, consists of a prohibition of asbestos fibres and of products containing asbestos fibres (Article 1), coupled with limited and temporary exceptions from the prohibitions for certain ‘existing materials, products or devices containing chrysotile fibre’ (Article 2). The remaining operative provisions of the Decree contain additional rules governing the grant of an exception (Articles 3 and 4) and the imposition of penalties for violation of the prohibitions in Article 1 (Article 5). Furthermore, certain used ‘vehicles’ and ‘agricultural and forestry machinery’ are entirely excluded, until 31 December 2001, from certain aspects of the prohibitions pursuant to Article 1 (Article 7).

The French Government adopted the Decree against the backdrop of widespread recognition that asbestos is a highly toxic material which poses a significant threat to human health. Its carcinogenicity has been scientifically acknowledged for some time by relevant international bodies such as the WHO.3 Nevertheless, due to the asbestos fibres’ exceptional chemical features and characteristics, they have found manifold use in industrial applications. In particular, chrysotile (or white asbestos) has been exploited for commercial purposes.

III. DISPUTE SETTLEMENT PROCEDURE

In a communication dated 28 May 1998, Canada requested consultations with the European Communities pursuant to Article XXII of the GATT 1994, Article 11 of the Agreement on the Application of Sanitary and Phytosanitary Measures (SPS Agreement), and Article 14 of the Agreement on Technical Barriers to Trade (TBT Agreement), concerning the French Decree on the prohibition of asbestos and products containing asbestos. Consultations were held among the parties, but they failed to resolve the dispute satisfactorily. Accordingly, the Dispute Settlement Body (DSB) was informed on 8 October 1998.

Consequently, Canada formally requested the DSB to establish a panel pursuant to Article 6.2 of the DSU in order to examine the matter in the light of the GATT 1994, the SPS Agreement and the TBT Agreement. At

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2 Journal officiel, France, 26 December 1996.
3 Cf. panel report, para. 8.188, for an overview on the development of knowledge of the risks associated with asbestos.
its meeting held on 25 November 1998, the DSB established a panel, and the parties to the dispute agreed on the following composition thereof:

   Chairman:  Mr. Adrian Macey  
   Members:  Mr. William Ehlers  
               Mr. Ake Linden  

Brazil, the United States, and Zimbabwe reserved their rights as third-parties to the dispute in accordance with Article 10 of the DSU.

The panel substantively met with the parties on 1-2 June 1999 and 20 January 2000, respectively, and held a meeting with scientific experts on 17 January 2000. Its final report was submitted to the parties on 25 July 2000 and circulated to WTO members on 18 September 2000.

On 23 October 2000, Canada notified the DSB of its decision to appeal certain issues of law covered in the panel report pursuant to Article 16.4 of the DSU, and filed a Notice of Appeal with the Appellate Body pursuant to Rule 20 of the Working Procedure for Appellate Review. The members of the Appellate Body hearing the case were Florentino P. Feliciano (Presiding), James Bacchus and Claus-Dieter Ehlermann. Canada and the European Communities filed their appellant’s submissions on 16 and 21 November 2000, respectively, and their appellee’s submissions on 1 December 2000. Brazil and the United States again participated as third parties. In the absence of any objection by a participant or third participant, Zimbabwe was allowed to attend the oral hearing as a passive observer.


IV. CLAIMS

1) Before the panel

Canada requested the panel to address the following substantive claims and to rule that

i) the French Decree is a technical regulation within the meaning of Annex 1 to the TBT Agreement. As such, it is incompatible with paragraphs 1, 2, 4, and 8 of Article 2 of that Agreement;

ii) the French Decree is incompatible with Articles III:4 and XI of the GATT 1994;

iii) in the event that the panel is unable to find a violation of Article XXIII:1(a) of the GATT 1994, it nevertheless shall conclude that the provision of subparagraph (b) of that Article applies.
In addressing the substantive claims set forth by Canada, the European Communities submitted that

i) the French Decree does not fall under the TBT Agreement. If the panel nevertheless finds that the Decree is covered by that Agreement, the European Communities requested the panel to conclude that the Decree complies with the relevant provisions thereof;

ii) with respect to the GATT 1994, the panel shall confirm that either the French Decree does not establish less favourable treatment for imported products than for like domestic products within the meaning of Article III:4 of that Agreement or that the Decree is necessary to protect human health within the meaning of Article XX(b) of the GATT 1994;

iii) Article XXIII:1(b) of the GATT 1994 does not apply.

2) In appeal

Canada appealed the panel’s findings and conclusions on two grounds:

i) it argued that the panel erred in its definition of the term ‘technical regulation’ pursuant to Article 2 of the TBT Agreement and the Annex 1 thereto. Canada claimed that the Appellate Body should find that the French Decree as a whole falls within the scope of that Agreement, and subsequently determine that the French Decree is inconsistent with paragraphs 1, 2, 4, and 8 of Article 2 of that Agreement;

ii) it claimed that the French Decree is not justified under Article XX(b) of the GATT 1994. As part of this argument, it also asserted that the panel failed to make an objective assessment of the matter as required by Article 11 of the DSU.

The European Communities responded by asking the Appellate Body to reverse the panel report on two issues:

i) it argued that chrysotile asbestos fibres are not ‘like’ polyvinyl alcohol (‘PVA’), cellulose and glass fibres, and that cement-based products containing chrysotile asbestos fibres are not ‘like’ cement-based products containing PVA, cellulose and glass fibres. Consequently, the panel erred in its interpretation and application of the term ‘like products’ in Article III:4 of the GATT 1994, and the French Decree does not violate that Article;

ii) it appealed certain legal findings on Article XXIII:1(b) of the GATT 1994 but requested the Appellate Body to uphold the panel’s conclusion that Canada did not establish nullification or impairment of a benefit within the meaning of that provision.
V. FINDINGS OF THE PANEL

1) Preliminary issues: scientific experts and *amicus curiae* briefs

In the course of the procedure, the panel reached two decisions which it considered to be explicitly reproduced as an integral part of the findings:

Firstly, the panel decided to consult experts on an *individual* basis rather than establish an expert group in order to obtain information on the circumstances of exposure to chrysotile asbestos and the associated risks as well as on the effectiveness of a ‘controlled use’ of chrysotile. It made reference to the *EC – Hormones case* and interpreted Article 13 of the DSU to the effect that this Article does not prevent a panel from consulting experts individually and leaves the determination of whether it is necessary or appropriate to establish an expert group to the sound discretion of a panel. Moreover, the panel also elaborated on Article 14.2 of the TBT Agreement which provides for the establishment of a technical expert group if requested by a party or on the panel’s own initiative. That article constitutes *lex specialis* which prevails, in the case of a conflict, over similar provisions under the DSU. The panel concluded, based on the clear wording of both Article 13 of the DSU and Article 14.2 of the TBT Agreement, that neither of these provisions makes the establishment of such a group mandatory, and they can thus be seen as complementary. Therefore, it was entirely within the panel’s discretion to obtain additional scientific information by consulting experts individually rather than requesting a joint report on the various scientific or technical issues.

Secondly, the panel received five *amicus curiae* briefs submitted by sources other than the parties or third parties to the dispute. The parties were informed accordingly, and the European Communities formally incorporated by reference two of them into its own submission on an equal footing. The panel informed the parties that it will consider these two documents on the same basis as the other documents furnished by the European Communities in the present dispute, and gave Canada the opportunity to fully comment on their content. As regards the remaining three *amicus curiae* briefs, the panel decided not to take them into consideration.

2) TBT Agreement and the definition of ‘technical regulation’

The panel first addressed Canada’s claim that the French Decree is to be qualified as a ‘technical regulation’ pursuant to Annex 1 to the TBT Agreement. It thoroughly examined the exact meaning of the term ‘technical regulation’ and concluded that a measure constitutes a ‘technical regula-

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6 These two being *amicus* briefs submitted by the *Collegium Ramazzini* and *American Federa- tion of Labor and Congress of Industrial Organizations*, respectively.

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tion’ if i) the measure affects one or more given products; ii) the measure specifies the technical characteristics of the product(s) which allow them to be marketed in the Member State that took the measure; and iii) compliance is mandatory.7

Based on that definition, the panel went on to qualify the French Decree. It considered it useful, for practical reasons, to engage in a separate examination of the provisions specifically concerning the ban on asbestos and the provisions allowing certain limited exceptions. It concluded that

i) the part of the French Decree which generally prohibits asbestos and asbestos-containing products does not fall within the definition of a ‘technical regulation’ pursuant to the TBT Agreement. The panel based its finding on the view that the measure at issue did not define the characteristics of the products whose import is banned nor does it cover processes or methods for the production of asbestos or asbestos-containing products as required by its definition;

ii) the part of the French Decree which contains certain limited exceptions to the general ban on asbestos does fall within the scope of the definition of ‘technical regulation’. The panel referred to the exhaustive list in Article 2.II of the Decree which covers the identification of the materials, products or devices benefiting from the exceptions to the general ban. Moreover, that Article, supplemented by Article 3.I, also sets out the criteria for marketing the products identified in the Decree and not solely those for excluding products from the market. Since those marketing criteria relate to the characteristics of one or more given products or processes or production methods relating to them, the panel concluded that this part of the Decree is covered by the TBT Agreement.

However, Canada did not make any specific claims regarding the limited exceptions to the prohibition. The arguments brought forward by Canada related only to the part of the Decree containing the general ban which did, according to the panel, not qualify as ‘technical regulation’. Therefore, the panel concluded that there was no ground for further deliberations on the issue as to whether that part of the Decree was compatible with the TBT Agreement or not.

3) Violation of Article III:4 of the GATT 1994

By way of introduction, the panel examined the applicability of Article III:4 and Article XI:1 of the GATT 1994 to the French Decree. It drew attention to the Note Ad Article III which specifically covers a situation in

7 In footnote 49 to para. 8.64, however, the panel added that ‘these criteria are by no means exhaustive.’
which a law, regulation or requirement applies both to an imported product and to the like domestic product and is enforced in the case of the imported product at the time or point of importation. The panel established that the latter was in fact the case, making reference to the practice under GATT 1947, and concluded that Article III:4 applied to the French ban on asbestos imposed by the Decree.8

Then, the panel turned to Canada’s claim that asbestos and non-asbestos substitutes were ‘like products’ pursuant to Article III:4 of the GATT 1994. It based its analysis on the criteria as originally developed by the Border Tax Adjustment Working Party and ever since consistently applied by almost all panels and the Appellate Body in this respect. These criteria are the following: i) physical characteristics; ii) consumers’ tastes and habits; iii) the product’s end-uses in a given market; and, additionally, iv) tariff classification. When examining the properties, nature and quality criterion, the panel considered whether it should take into account the relevance of the risk of a product for human or animal health as argued by the European Communities. It rejected to do so since ‘introducing a criterion on the risk of a product into the analysis of likeness within the meaning of Article III would largely nullify the effect of Article XX(b).’9 Overall, the panel determined that chrysotile fibres, on the one hand, and PVA, cellulose and glass fibres, on the other hand, are similar in properties, nature and quality and have similar end-uses, and it concluded therefrom that these products are ‘like’ within the meaning of Article III:4 of the GATT 1994. This holds true, according to the panel, for these products taken separately, i.e. not incorporated in a product, as well as for products containing chrysotile or substitute fibres.

The panel went on to examine the second requirement set forth in Article III:4 and established that the Decree treated imported chrysotile and chrysotile-containing products de jure less favourably than domestic PVA, cellulose and glass fibre products. Overall, the panel thus arrived at the conclusion that Article III:4 of the GATT 1994 was violated.

4) Justification of the measure under Article XX(b) of the GATT 1994

In accordance with the consistent case law, the panel adopted a two-step approach to determine whether the French Decree was justified by an exception set forth in Article XX of the GATT 1994.

First, the panel established that the measure at issue fell within the range of policies designed to protect human life or health and thus was covered by paragraph (b). It based its finding on the statements of the con-

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8 In the light of the observation of the applicability of Article III:4 and the subsequent finding with regard to the violation of that Article by the Decree, the panel did not consider it necessary to further examine whether the ban was also covered by Article XI:1 of the GATT 1994.
9 Panel report, para. 8.130.
sulted scientists and statistics delivered by the relevant international bodies such as the WHO according to which the deadly and carcinogenic characteristics of asbestos are well and widely recognized. Consequently, it considered that ‘the evidence before it tends to show that handling chrysotile-cement products constitutes a risk to health rather than the opposite’, and concluded that the European Communities made a *prima facie* case for the existence of a health risk.\(^\text{10}\) The panel then turned to the question of whether the measure is ‘necessary’ to protect human life or health within the meaning of Article XX(b) and applied the traditional *necessity test* which requires that there be no *reasonably available* and less trade restrictive alternative measure to achieve the policy goal. The panel determined that, in the light of France’s public health objectives, the European Communities made a *prima facie* case for the non-existence of a reasonably available alternative to the total banning of chrysotile products. It dismissed Canada’s argument that controlled use of asbestos is both as effective and reasonably available as a total ban by particularly focusing at controlled use in the building sector and by ‘do-it-yourself enthusiasts’. Since Canada could not rebut the presumption established by the European Communities, the panel consequently held that the French Decree satisfied the conditions of paragraph (b) of Article XX of the GATT 1994.

Next, the panel proceeded to examine the requirements of the introductory clause of Article XX, the so-called *chapeau*, whose main purpose is the prevention of abuse of exceptions set forth in Article XX. The panel concluded that the *application* of the measure at issue constituted neither a ‘means of arbitrary or unjustifiable discrimination’ nor a ‘disguised restriction on international trade’. As regards the latter, it focused its attention, *inter alia*, on the term ‘disguised’ and held that ‘as far as the design, architecture and revealing structure of the Decree are concerned, we find nothing that might lead us to conclude that the Decree has protectionist objectives.’\(^\text{11}\)

In conclusion, the panel held that the French Decree met the requirements of Article XX(b) of the GATT 1994, and its violation of Article III:4 of the GATT 1994 was thus justified by that Article.

5) **Non-violation nullification or impairment under Article XXIII:1(b) of the GATT 1994**

At the outset, the panel dismissed two preliminary objections brought forward by the European Communities. First, it rejected the claim that the rules on non-violation nullification and impairment apply only to measures which *do not otherwise fall* under other provisions of the GATT 1994. Moreover, it dismissed the argument that, while it may be possible to have ‘legitimate expectations’ in connection with a purely ‘commer-

\(^\text{10}\) Panel report, paras. 8.193-194.

\(^\text{11}\) Panel report, para. 8.238.
cial’ measure, it is not conceivable to claim ‘legitimate expectations’ with respect to a measure taken in order to protect human life or health which can be justified under Article XX(b) of the GATT 1994.

The panel then turned to the substantive aspects of Canada’s claim. It recalled that, making reference to the Japan – Photographic Film and Paper case, the wording of Article XXIII:1(b) itself establishes three elements whose existence needs to be proven: i) the application of a measure by a Member State to the WTO; ii) the existence of a benefit accruing under the applicable agreement; and iii) the nullification or impairment of a benefit as a result of the application of the measure. The panel considered it appropriate, when examining the existence of a ‘benefit’, to focus on the question whether the French Decree could reasonably have been anticipated by the Canadian Government at the time that it was negotiating the various tariff concessions covering the products concerned. It concluded that, at the conclusion of the Uruguay Round, Canada could indeed have reasonably foreseen that France might, in the short term, adopt more restrictive measures on the use of asbestos, including the introduction of a total ban.

Therefore, the panel held that Canada did not establish the existence of nullification or impairment of a benefit within the meaning of Article XXIII:1(b) of the GATT 1994 as a result of the application of the French ban on asbestos products.

VI. FINDINGS OF THE APPELLATE BODY

1) Preliminary issue: amicus curiae briefs

After the panel had received five amicus curiae briefs in the proceedings before it, the Appellate Body anticipated the possibility that it might receive unsolicited submissions as well, and formally adopted, in order to facilitate the fair and orderly conduct of the appeal, an Additional Procedure pursuant to Rule 16(1) of the Working Procedures. It did so after consultations among all seven members of the Appellate Body and also taking into account the comments of the parties to the dispute on the issue. The Additional Procedure was explicitly put in place for the purpose of this appeal only, and was posted on the WTO website on 8 November 2000. In essence, it contained a ‘request for leave’ procedure for amicus briefs and listed a series of requirements to be met by the applicants in order to be granted leave to file a written brief.

Subsequently, the Appellate Body received seventeen applications requesting leave. The Appellate Body, however, denied leave in each case; six of the seventeen applications were received after the deadline as specific...
fied in paragraph 2 of the Additional Procedure, and the other eleven submissions failed to comply with all the requirements set forth in paragraph 3 of the Additional Procedure. The Appellate Body sent a copy of its decision denying application for leave to each applicant. However, it did not indicate what the concrete shortcomings of the respective applications were.

In sum, the Appellate Body did not take into consideration any other source than the submissions presented by the parties and third parties to the dispute.

2) TBT Agreement and the definition of ‘technical regulation’

The Appellate Body reversed the panel’s finding in a twofold manner: Firstly, it set out that the proper legal character of the French Decree cannot be determined unless the measure is examined as an integrated whole. Accordingly, an objective assessment needs to take into account both the prohibitive and the permissive elements that are part of the measure, and the provisions specifically concerning the ban cannot be considered separately from those governing the exception to use asbestos in certain situations.

Secondly, it turned to the definition of the term ‘technical regulation’ as stipulated in Annex 1 to the TBT Agreement and interpreted by the panel. It emphasised, *inter alia*, that ‘product characteristics’ may be prescribed or imposed in either a positive or a negative form; thus, a document may also require, negatively, that products must *not* possess certain ‘characteristics’. Moreover, it held, in contrast to what the panel found, that nothing in the text of the TBT Agreement requires that a ‘technical regulation’ must apply to ‘given’ products which are actually named, specified or otherwise expressly identified in the regulation. Based on these considerations, it ruled that the French Decree, although negatively formulated, effectively did describe certain objective features or ‘characteristics’ of all products, namely that they be free of asbestos. Accordingly, the measure at issue constitutes a ‘technical regulation’ under the TBT Agreement.

However, the Appellate Body decided not to address the claims brought forward by Canada and thus not to ‘complete the legal analysis’. It considered such an analysis inappropriate since it would have had to deal with various interpretative issues of first impression not yet adjudicated by a panel before. Moreover, it hinted that the facts on the record might in any case have been insufficient to correctly apply the relevant provisions of the TBT Agreement.

3) Violation of Article III:4 of the GATT 1994

The Appellate Body reversed the panel’s finding that chrysotile fibres and PVA, cellulose and glass fibres are ‘like products’ within the meaning of Article III:4 of the GATT 1994.
It approved the panel’s approach to base the analysis on the Working Party *Border Tax Adjustment* criteria but, at the same time, stated that the panel erred in not considering and weighing all the evidence and, in particular, in not comprehensively examining *all* four of the *Border Tax Adjustment* criteria. Fundamentally, the Appellate Body held, reversing the panel’s finding on that point, that considerations of health effects need to be taken into account in the analysis of the physical properties of a product for the purpose of ‘likeness’ under Article III:4 of the GATT 1994. It expressed its inability to ‘see how this highly significant physical difference cannot be a consideration in examining the physical properties of a product.’\(^\text{14}\) In sum, the Appellate Body reached the conclusion that

‘the Panel disregarded the quite different ‘properties, nature and quality’ of chrysotile asbestos and PCG fibres, as well as the different tariff classification of these fibres; it considered no evidence on consumers’ tastes and habits; and it found that, for a ‘small number’ of the many applications of these fibres, they are substitutable, but it did not consider the many other end-uses for the fibres that are different. Thus, the only evidence supporting the Panel’s finding of ‘likeness’ is the ‘small number’ of shared end-uses of the fibres.’\(^\text{15}\)

Likewise, the Appellate Body reversed the panel’s finding that products containing chrysotile and those containing substitute materials are ‘like products’ for the purpose of Article III:4 of the GATT 1994.

Subsequently, the Appellate Body considered it appropriate to complete the analysis and to examine, on the basis of the factual findings of the panel and the undisputed facts in the panel record, whether the products at issue are ‘like products’ pursuant to Article III:4. It eventually found that ‘all of this evidence is certainly far from sufficient to satisfy Canada’s burden’ of proving that the products at issue are ‘like products’.\(^\text{16}\) In conclusion, the Appellate Body determined that the French Decree was consistent with Article III:4 of the GATT 1994.

4) Justification of the measure under Article XX(b) of the GATT 1994

After having found that the French Decree did not violate Article III:4 of the GATT 1994, the Appellate Body could have decided not to rule on Canada’s appeal concerning Article XX of the GATT 1994, for reasons of judicial economy. Nevertheless, it addressed the requirements of para-

\(^\text{14}\) Appellate Body report, para. 114 (original emphasis).

\(^\text{15}\) Appellate Body report, para. 125.

\(^\text{16}\) Appellate Body report, para. 141; it is interesting to note that one member of the Division of the Appellate Body hearing the case made a concurring statement on the way the Appellate Body completed the analysis and weighed the evidence.
graph (b) of Article XX in turn, and particularly deliberated on the term ‘necessary’. Overall, it upheld the panel’s conclusion that the measure at issue was justified by Article XX(b) of the GATT 1994.

As part of the argument that the French Decree was not justified under Article XX(b), Canada also asserted that the panel failed to ‘make an objective assessment of the matter’ as required by Article 11 of the DSU. In essence, Canada’s claim concerned the selection and expertise of the appointed scientists and the credibility and weight the panel ascribed to different elements of evidence. The Appellate Body, however, found that ‘the panel’s appreciation remained well within the bounds of its discretion as the trier of facts’ and consequently dismissed Canada’s appeal on Article 11 of the DSU.

5) Non-violation nullification or impairment under Article XXIII:1(b) of the GATT 1994

The European Communities claimed that the panel erred in some of its legal interpretations on Article XXIII:1(b) of the GATT 1994. The Appellate Body, however, dismissed the appeal, and consequently upheld the panel’s findings that Article XXIII:1(b) may also apply to measures which fall within the scope of application of other provisions of the GATT 1994 as well as to measures which pursue health objectives and can be justified under Article XX(b) of the GATT 1994.

VII. LEGAL COMMENTARY

1) The issue of amicus curiae briefs

The handling by both the panel and the Appellate Body of the issue as to whether to take amicus curiae briefs formally into consideration or not seems, at first sight, to be in line with the current practice under WTO dispute settlement. In 1998, the Appellate Body fundamentally ruled in the US – Shrimps case that ‘a panel has the discretionary authority either to accept and consider or to reject information and advice submitted to it, whether requested by a panel or not.’ It based its ruling on the wording of Article 13 of the DSU. Likewise, the Appellate Body approved for itself, in the US – Shrimps and US – British Steel cases, the ‘legal authority under the DSU to accept and consider amicus briefs in an appeal in which we find it pertinent and useful to do so.’ In practice, however, such briefs have extremely rarely been accepted and are usually rejected; to

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17 Appellate Body report, paras. 162, 177.
date, only one independently submitted non-governmental brief has been formally taken into account by a panel as amicus curiae brief.20

Ever since, the issue has been highly controversial, and Member States to the WTO are deeply divided over non-state actors’ status before panels and the Appellate Body. The most serious of a range of systemic concerns is related to Member States’ rights and the government-to-government nature of WTO dispute resolution. Accepting and comprehensively taking into account non-governmental briefs confers ‘outsiders’ more access to WTO dispute adjudication than to Member States that must demonstrate at the outset of a dispute settlement proceeding that they have a ‘substantial interest in the matter’ in order to be granted participation as third parties.21

The Appellate Body’s decision to set up an Additional Procedure in the present dispute can be understood only against the backdrop of that controversy. The adoption of the Additional Procedure itself caused considerable turmoil among many delegates from both developed and developing countries, and an extraordinary meeting of the General Council was called to consider the legitimacy of the new filing procedure. Yet, the General Council refrained from reversing the policy but forwarded a stern warning to the Appellate Body urging it to exercise ‘extreme caution’ on the issue.22 The Procedure contained a list of stringent procedural requirements; of utmost significance, albeit drawn up in indeterminate language, is the criterion stipulated in paragraph 3(f) of the Procedure according to which an applicant must state ‘why it would be desirable (...) for the Appellate Body to grant the applicant leave to file a written brief’ and indicate ‘in what way the applicant will make a contribution to the resolution of this dispute that is not likely to be repetitive of what has already been submitted by a party or third party’. In essence, the Appellate Body denied leave in each case forwarding as its reasoning the following standard explanation to the respective applicants: ‘Your application (...) has been denied for failure to comply sufficiently with all the requirements set forth in paragraph 3 of the additional procedure.’ It is mere guessing that most of the applications were turned down on grounds that they could not prove a potentially substantial contribution to the process of factual and legal findings in addition to the parties’ submissions.

20 Namely in the Australia – Measures Affecting Importation of Salmon case, WT/DS18/RW, report of the implementation panel, para. 7.8. Moreover, in United States – Import Prohibition of Certain Shrimp and Shrimp Products, WT/DS58/AB/R, Appellate Body report, para. 83, the Appellate Body accepted for consideration an unsolicited brief which was not attached to any party’s submission.

21 Cf. Article 10 of the DSU.

The current WTO practice on *amicus curiae* briefs and the approach followed by the panel and the Appellate Body in the present case raise three issues which are of primary concern. Firstly, both the panel, as the trier of facts, and the Appellate Body essentially made use of the wide discretionary power which is conferred on them in the matter and seemed to remain well within the bounds of their discretion. Moreover, they were under no legal obligation to justify the rejection of all unsolicited pleadings. However, the *complete lack of any reasons* for the rejection in both the panel and the Appellate Body report is grist to the mill of those who accuse the WTO and its dispute settlement system of not abiding by fundamental legal principles such as transparency, predictability and consistency of the law. The intention of the Appellate Body originally was, by adopting the *Additional Procedure*, to guarantee the fair and orderly conduct of the handling of *amicus curiae* briefs, but it obviously failed to achieve that goal in turning down the applications without delivering sound statements of reasons. Being mindful of the immense workload of panels and the Appellate Body, and the large number of unsolicited submissions due to the politically sensitive nature of the dispute, the reasoning could well have been short in nature and termed in summary wording. Nevertheless, it is submitted that a sound reasoning would have needed to substantially address every individual submission and its content.

Secondly, panels and the Appellate Body have very occasionally accepted and considered unsolicited submissions without having provided generally valid and handy criteria as to the appropriate source and nature of information, or the circumstances under which panels and the Appellate Body may consider information ‘desirable’ and ‘useful’. The *Additional Procedure* was explicitly put in place for the purpose of this appeal only but quite bluntly reveals the current need for authoritative guidelines which generally outline the handling of *amicus curiae* briefs. The main goal is to set aside the present state of legal uncertainty and to thus ensure the promotion of procedural fairness and predictability. It is, however, unlikely that panels or the Appellate Body will ever be in a position to develop satisfactory rules of procedure on their own. Current disagreement on the issue has simply assumed too vast a proportion, and the value of placing *amicus curiae* briefs before WTO adjudicating bodies can arguably be settled in a mutually satisfactory manner only by the Member States themselves.

Thirdly, due to inaction by the Member States, or due to their inability to agree on a common position, panels and the Appellate Body may nevertheless show their tendency to somehow fill the legislative gap and develop their own guidelines. For the time being, it is most likely that both panels and the Appellate Body will continue their cautious approach

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2) TBT Agreement and the definition of ‘technical regulation’

Since the coming into force of the WTO to date, 33 notifications have been referred to the DSB in which the complainants explicitly mentioned the TBT Agreement and claimed, *inter alia*, an alleged violation of that Agreement. Usually, the complainants argued that the measure in question fell within the definition of a ‘technical regulation’ pursuant to Annex 1 and that the measure was inconsistent with Article 2 of that Agreement. However, in all disputes eventually brought before a panel, a violation of the TBT Agreement was no longer alleged at the panel stage, and neither a panel nor the Appellate Body had the opportunity to rule on issues governed by that Agreement. This matter of fact is striking since a number of previous cases involved measures which could unequivocally be qualified as ‘technical regulations’, at least by applying a wide definition, and an obvious reason for the parties’ reluctance to invoke provisions of the TBT Agreement is not easily discernible. Eventually, the panel, and subsequently the Appellate Body, were in the present case called upon for the first time to explicitly rule on a complainant’s allegation that the TBT Agreement was violated.

The adjudication by the panel and the Appellate Body of Canada’s claims attracted the attention of scholars and practitioners alike. Three main issues have primarily caught their interest: Firstly, the Appellate Body correctly held, reversing the panel’s finding on that point, that ‘product characteristics’ as stipulated in Annex 1.1 to the TBT Agreement may be prescribed or imposed in either a positive or a negative form, and, as a consequence, a document may also negatively require that products must not possess certain characteristics. Moreover, the Appellate Body made it clear that compliance with the TBT Agreement, and particularly with Article 2.9.2 thereof, requires identification of the product coverage of the technical regulation but does not mean that the products necessarily be named or otherwise explicitly identified in the regulation. The Appel-

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[^24]: According to the Appellate Body in the present case, para. 81, the provisions of the Tokyo Round *Agreement on Technical Barriers* ‘were also never the subject of even a single ruling by a panel’.

late Body based its finding on the grounds that ‘there may be perfectly sound administrative reasons for formulating a ‘technical regulation’ in a way that does not expressly identify products by name, but simply makes them identifiable – for instance, through the ‘characteristic’ that is the subject of regulation.’ That rationale is both cogent and self-explanining, and, by the same token, its soundness becomes apparent in the light of the facts and circumstances of the present case. It was arguably impossible for the French administering authority to name each and every product to which the ban on asbestos shall apply – it is simply all products, and the ‘characteristic’ which is the subject of the regulation is that they be free of asbestos. In conclusion, the Appellate Body’s interpretation of the term ‘technical regulation’ results in a broader scope of application of the TBT Agreement than the panel’s approach did.

Secondly, while the Appellate Body found that the French Decree constituted a technical regulation, it reserved its judgement on the Decree’s compatibility with the TBT Agreement ‘in the light of the novel character’ of the obligations stipulated in that Agreement and due to ‘insufficiency of the facts on the record’27. This outcome might be understandable in the light of the incomplete factual record before the Appellate Body. Nevertheless, it is not satisfactory from a due process point of view. Canada properly raised the claim of TBT-inconsistency of the French Decree before the panel, and it was also perfectly legitimate to appeal the panel’s finding on the issue. However, Canada never received a decision on that point. This constellation once again reveals the issue of whether the Appellate Body shall have the power to remand a matter to the original panel with instructions to consider the remaining claims in the light of the appellate findings. As the law stands at the moment, the Appellate Body lacks authority to do so. In previous appeals, the Appellate Body has on occasion completed the legal analysis itself, but it consistently insisted ‘that we could do so only if the factual findings of the panel and the undisputed facts in the panel record provided us with a sufficient basis for our own analysis’.28

Thirdly, the Appellate Body touched upon the relationship between the TBT Agreement and GATT 1994. It held that ‘the TBT Agreement imposes obligations on Members that seem to be different from, and additional to, the obligations imposed on Members under the GATT 1994.’29 It cautiously added that not all internal measures covered by Article III of the GATT 1994 necessarily fall under the scope of the TBT Agreement, but it did not further elaborate on the practical consequences of that finding. One reading of the Appellate Body’s statement, based on its wording, may be to conclude that the provisions of the two agreements operate

26 Appellate Body report, para. 70.
27 Appellate Body report, para. 82.
28 See, for an overview on the case law, Appellate Body report, para. 78 and fn. 48 and 49 thereto.
29 Appellate Body report, para. 80 (original emphasis).
concurrently, and thus to undermining the notion that the TBT Agreement should be considered \textit{lex specialis} to the GATT 1994.\textsuperscript{30} If that is the case, both the GATT 1994 and the TBT Agreement may apply to a single dispute provided that the measure at issue falls within the ambit of some provisions of both agreements. The most significant implication of such an interpretation would become apparent when looking at the main regulatory purpose of the TBT Agreement: It is not limited to non-discrimination, but also addresses excessive regulations and applies irrespective of a discriminatory impact. Thus, a complainant would be supposed to invoke the TBT Agreement rather than Article III of the GATT 1994 when a technical regulation does not explicitly discriminate against imported products but creates otherwise illegitimate obstacles to international trade.\textsuperscript{31} However, it seems too daring to definitely draw the conclusion of concurrent application at the moment, and only future practice in this respect will clarify the issue.

3) The ‘like product’ test pursuant to Article III:4 of the GATT 1994

Arguably the most profound finding of the Appellate Body in the present case was the introduction of fundamental human interests such as health risks into the ‘like product’ test pursuant to Article III:4 of the GATT 1994. The Appellate Body elaborated on the criteria for determining likeness at great length and fundamentally clarified the issue. According to it, the following is the current state of law:

At the outset, the Appellate Body placed particular emphasis on Article III:1 of the GATT 1994 as stating the ‘general principle’ which informs the specific obligations contained in Articles III:2 and III:4. It compared the two paragraphs and pointed out that the scope of ‘like’ in the latter is broader than in the former, thus assigning Article III:4 ‘a relatively broad product scope.’\textsuperscript{32} Subsequently, it turned to examine specifically the meaning of Article III:4. By way of introduction, it recalled that an assessment of likeness can appropriately be made only on a case-by-case basis and necessarily entails an ‘unavoidable element of individual, discretionary judgement’.\textsuperscript{33} The proper framework for the analysis is still provided by the approach developed by the \textit{Border Tax Adjustment Working Party} and consists of the traditional four criteria which are: i) physical characteristics; ii) consumers’ tastes and habits; iii) the product’s end-

\textsuperscript{30} Guidance in this respect may be provided by the elaboration on the relationship between the SPS Agreement and GATT 1994, see \textit{European Communities – Measures Concerning Meat and Meat Products (Hormones)}, WT/DS26/AB/R, WT/DS48/AB/R, panel report, paras. 8.34-45.


\textsuperscript{32} Appellate Body report, para. 100.

uses in a given market; and, additionally, iv) tariff classification. By doing so, the Appellate Body confirmed the case law as originally developed under GATT 1947 and consistently followed under the WTO ever since, at the same time implicitly rejecting the ‘aims-and-effect’ test again.

The Appellate Body assessed the significance of the Border Tax Adjustment criteria by determining that they are to be treated as simple tools to assist in the task of sorting and examining the relevant evidence and do not constitute a closed list. On the contrary, the adoption of the Border Tax Adjustment criteria does by no means ‘dissolve the duty or the need to examine, in each case, all of the pertinent evidence’. The Appellate Body noted that neither the text of Article III:4 nor the practice of panels to date suggests that any evidence should be excluded a priori from a panel’s examination of likeness, and explicitly acknowledged that evidence relating to health risks associated with a product may perfectly well be pertinent in such an examination. However, the Appellate Body recoiled from taking too revolutionary a step. Firstly, it continued to rely upon the Border Tax Adjustment criteria and considered it inappropriate to introduce a separate criterion; instead, it evaluated health risks under the two already existing criteria of physical properties and consumers’ tastes and habits. Secondly, albeit stating that the products at issue were characterized by a ‘highly significant physical difference’, the Appellate Body nevertheless continued to examine the remaining criteria of the Border Tax Adjustment Working Party. Even physically very different products may be determined ‘like’ if all of the evidence, taken together, leads to the conclusion that there is a competitive relationship between the products. Still, ‘a very heavy burden’ of proving the contrary is placed, according to the Appellate Body, on the complainant in such a constellation.

In sum, the Appellate Body considered fundamental human interests such as health risks inherent in a product relevant for the purpose of the ‘like product’ test pursuant to Article III:4 of the GATT 1994. At the same time, it appeared to uphold its economic approach to the likeness analysis; in essence, the test is a market assessment of the competitive relationship between products in the relevant marketplace, and all evidence affecting the products should be taken into account and weighed. This market-based attitude prompted one member of the Division hearing the appeal to make, for the first time in an Appellate Body report ever, a concurring statement. He submitted that ‘the definitive characterisation (of unlikeness between the products at issue) may and should be made even in the absence of evidence concerning the other two Border Tax Adjustment criteria’. He pointed out that he could not imagine what evi-

34 Panel report, paras. 8.112-113; Appellate Body report, para. 101.
35 Appellate Body report, para. 102 (original emphasis).
36 Appellate Body report, para. 118.
37 Appellate Body report, para. 152 (original emphasis).
evidence could have outweighed and set at naught the undisputed deadly nature of chrysotile asbestos fibres when compared to substitute fibres.

The impact of the Appellate Body’s finding on future practice is difficult to assess. The following two observations may provide a tentative outlook: Firstly, the strategy of dealing with health issues within Article III:4 of the GATT 1994 allows for enhanced product and regulatory differentiation and enables Member States to refine their policy options in this respect. In essence, it is an expression of deference to Member States. At the same time, however, enhanced product differentiation potentially affects market access rights of other Member States.

Secondly, it seems unclear to what extent characteristic features associated with a product other than health risks may be taken into account in future ‘like product’ cases. Asbestos fibres were inherent in the products in question, and its impact could thus be evaluated under the Border Tax Adjustment criteria of physical characteristics and consumers’ tastes and habits. The Appellate Body report does not reveal whether, and if so to what extent, policy objectives such as the protection of the environment, for instance, will be considered equally relevant for the analysis of likeness under Article III:4 of the GATT 1994 as the elimination of health risks posed by asbestos fibres was in the present case. It is unclear, to give a practical example, whether the Appellate Body’s findings would support the consideration of the level of air pollution caused by the use of two products in assessing whether such products are ‘like’ pursuant to Article III:4 of the GATT 1994.

Finally, the Appellate Body was aware of the fact that its interpretation of Article III:4 may imply less frequent recourse to Article XX(b) of the GATT 1994. It countered, thus refuting the panel’s argumentation on this point, that Article III:4 and Article XX(b) are distinct and independent provisions, and concluded that consideration of health risks for the purpose of determining likeness ‘does not deprive the exception in Article XX(b) of effet utile.’38 This statement is correct from the Appellate Body’s perspective on what the basic principles of the two articles are: Article III:4 guarantees a competitive relationship in the marketplace between like products by preventing protectionism in the application of regulatory measures, whereas evidence under Article XX(b) serves the different purpose of assessing whether a Member State has a sufficiently legitimate basis for adopting a WTO-inconsistent measure on the grounds of human health. In essence, this outcome is crucial from a practitioner’s perspective as the differentiation between products on the basis of health considerations under Article III:4 shifts the burden of proof from the Member State who restricts market access on health grounds to the member state who challenges the allegedly inconsistent restriction.

38 Appellate Body report, para. 115 (original emphasis).
4) Article XX(b) of the GATT 1994 and the protection of health

The present dispute marks the first ever under the WTO regime in which an otherwise inconsistent measure was determined by a panel or the Appellate Body to be justified under Article XX(b) of the GATT 1994. By doing so, the panel exempted the French Decree from the obligation of National Treatment, because it had first established that asbestos and substitute products were ‘like’. The Appellate Body, on the other hand, addressed Canada’s claim of error under Article XX in appeal even though it could be viewed as moot. Because the Appellate Body reversed the Panel’s interpretation of like product and found that the French decree did not violate GATT Article III, a determination as to whether the application of an exception to justify a violation would seem unnecessary. There was no violation of any WTO obligation. This is not the same as deciding whether to exercise judicial economy in a case where a panel or the Appellate Body find it appropriate to save resources by not finding multiple violations. Rather, in this case, the Appellate Body has taken the more dangerous step of deciding on legal matters and making important legal interpretations that are completely irrelevant to the resolution of the dispute.

By deciding Canada’s claim of error, the Appellate Body has opened itself to additional charges that it is engaging in “judicial activism” and exceeding its authority in reaching issues and making interpretations on highly controversial matters.39

The Appellate Body arrived at two conclusions of fundamental significance with respect to the protection of health and national policy determinations which involve societal value judgements as to whether to accept a risk, and, if not, what measure to select. Firstly, it noted that ‘it is undisputed that WTO Members have the right to determine the level of protection that they consider appropriate in a given situation.’40

The Appellate Body thus rejected the notion that a Member State’s freedom to establish the level of protection should be subject to considerations of proportionality. As a consequence, national authorities are free to set levels of protection as ambitious as they deem appropriate; they even can pursue a zero risk policy.41 This determination is not ripe for

39 Notably, Article 17.12 of the DSU directs the Appellate Body to address each of the issues raised in accordance with paragraph 6 during the appellate proceedings. Given that the Appellate Body has refused in the past to address issues on the basis of judicial economy, it would seem even more compelling for the Appellate Body to avoid making findings where the issues are completely irrelevant to the resolution of the dispute.
40 Appellate Body report, para. 168.
41 Cf. the Appellate Body’s ruling with respect to SPS measures in Australia – Measures Affecting Importation of Salmon, WT/DS18/AB/R, para. 199: ‘The determination of the appropriate level of protection, a notion defined in paragraph 5 of Annex A [to the SPS Agreement], ..., is a prerogative of the Member concerned and not of a panel or of the Appellate Body.’ (original emphasis).
judicial review due to policy considerations. Nevertheless, the Appellate Body’s statement has to be read in the light of the findings in the Korea – Beef case according to which a Member State will not easily persuade a panel that its level of protection is zero risk if its policy instrument appears to be structurally incapable of achieving that level.\footnote{Korea – Measure Affecting Imports of Fresh, Chilled and Frozen Beef, WT/DS161/AB/R, WT/DS169/AB/R, Appellate Body report, para. 178.}

Secondly, the Appellate Body dealt with Canada’s assertion that the panel erred in finding that ‘controlled use’ is not a reasonably available alternative to the French Decree. It recalled the case law on the term ‘necessary’ as stipulated in various paragraphs of Article XX of the GATT 1994, and again approved the test consistently applied by panels and the Appellate Body in this respect, namely whether there is a reasonably available alternative which is less restrictive of trade than the measure in question. Moreover, it made explicit reference to the Korea – Beef case in which it had stated that ‘the more vital or important (the) common interests or values’ pursued, the easier it would be to accept measures designed to achieve those ends as necessary.\footnote{Korea – Measure Affecting Imports of Fresh, Chilled and Frozen Beef, WT/DS161/AB/R, WT/DS169/AB/R, Appellate Body report, para. 162.} The Appellate Body then turned to the facts of the present case and stated that

\begin{quote}
‘the objective pursued by the measure is the preservation of human life and health through the elimination, or reduction, of the well-known, and life-threatening, health risks posed by asbestos fibres. The value pursued is both vital and important in the highest degree.’\footnote{Appellate Body report, para. 172.}
\end{quote}

This paragraph, read in the light of the cited statement in the Korea – Beef case, suggests that there may be divergent levels of scrutiny with which a panel needs to analyse whether a measure is ‘necessary’ or not. The degree of deference then could depend on the relative importance of the various objectives or interests at stake. In essence, it seems that the Appellate Body consequently followed its reasoning in the Korea – Beef case and established a more deferential approach towards Member States’ regulatory choices in the field of vital health interests. However, the actual engagement of the Appellate Body in the present case as to whether the French ban was necessary in order to achieve a policy of zero risk does not reveal such limited scrutiny. The Appellate Body simply concluded that ‘controlled use’ would not be an alternative measure to meet France’s chosen level of health protection.

The Appellate Body’s findings in this respect, and the refinement of the necessity test, raise the question of whether it is appropriate for panels and the Appellate Body to engage in a judgement on the ‘relative vital
importance’ of various legitimate interests. Do panels and the Appellate Body really have the authority to determine and classify as to whether certain rights and objectives count more than others? From a purely legal point of view, the legality of such a determination appears to be doubtful, at least. Moreover, panel and Appellate Body activism in deciding on the “relative importance” of various policy objectives may adversely affect legal certainty and predictability, because such assessments often involve a value-based weighting of societal, ethical and political considerations. It might be more adequate to predominantly vest the power to classify policy objectives in the competent national authorities and not to subject such determinations to panel and Appellate Body review. The traditional necessity test has the advantage that it does not necessarily require panels to engage in a politically sensitive judgement on the “relative importance” of various interests

5) Non-violation nullification or impairment under Article XXIII:1(b) of the GATT 1994

When addressing Canada’s claim that the application of the French Decree resulted in nullification or impairment of a benefit pursuant to Article XXIII:1(b), the panel turned to examine the existence of a ‘benefit’ because the qualification of the Decree as a ‘measure’ was not in dispute. In non-violation cases, this benefit has traditionally been the ‘legitimate expectations’ of improved market access opportunities after a tariff concession. The panel followed this line and focused on the requirement of non-foreseeability which is the most important element of legitimate recourse to non-violation protection; it is here that panels normally subject the complainant’s claims to a closer look. At stake was the question of whether the French Decree could reasonably have been anticipated by the Canadian Government at the time that it was negotiating the various tariff concessions covering the products concerned. The panel answered the question of foreseeability in the affirmative, thus dismissing Canada’s claim and basing its finding on the fact that ‘measures to limit the use of asbestos were being proposed at the international level, and countries at the same level of social and economic development as France had already banned the use of chrysotile asbestos at the end of the Uruguay Round.’

In addition to its deliberations on the substance of Canada’s claim, the panel made a noteworthy statement concerning the burden of proof applicable to non-violation complaints pursuant to Article XXIII:1(b) of the

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47 Panel report, para. 8.303; it is noteworthy that the European Communities itself issued as early as in 1990 Directive 90/394/EEC providing for the replacement of asbestos.
GATT 1994. After pointing to Article 26.1(a) of the DSU, which requires the complaining party to carry the burden of presenting ‘a detailed justification’ in support of any non-violation complaint, it held that

‘because of the importance conferred on them a priori by the GATT 1994, as compared with the rules governing international trade, situations that fall under Article XX justify a stricter burden of proof being applied in this context to the party invoking Article XXIII:1(b), particularly with regard to the existence of legitimate expectations and whether or not the initial Decree could be reasonably anticipated.’

Canada did not appeal this finding, nor any other made by the panel, so this legal interpretation has not yet been fully tested by the Appellate Body. It was only the European Communities that appealed the panel report on two preliminary issues, in essence repeating the argument that the French Decree fell outside the scope of application of Article XXIII:1(b).

At the outset, the Appellate Body noted that this appeal was the first case ever brought before it concerning Article XXIII:1(b) of the GATT 1994, and it thus considered it useful to make some general preliminary observations on the relationship between paragraphs (a) and (b) of that Article. It made explicit reference to the Japan – Photographic Film and Paper case and confirmed that ‘the remedy in Article XXIII:1(b) should be approached with caution and should remain an exceptional remedy.’

The Appellate Body then turned to the claims in appeal. The panel’s finding that Article XXIII:1(b) may also apply to measures that pursue health rather than commercial objectives, and that can be justified under Article XX(b) of the GATT 1994, caused quite a stir, and is likely not to be unanimously welcomed by commentators. However, the Appellate Body confirmed this ruling and based its view mainly on the text of Article XXIII:1(b). According to the Appellate Body, ‘the use of the word ‘any measure’ suggests that measures of all types may give rise to a cause of action under that provision’.

Moreover, it corroborated such a reading by adding that an attempt to draw the distinction suggested by the European Communities between so-called health and commercial measures would be very difficult in practice, and it could not see merit in the argument that previous case law would suggest that only ‘commercial’ measures can be subject to non-violation complaints.

This finding seems to be correct from a purely textual interpretation on Article XXIII:1(b) of the GATT 1994. Moreover, it is important to note

Panel report, para. 8.282.
Appellate Body report, para. 186, making reference to Japan – Measures Affecting Consumer Photographic Film and Paper, WT/DS44/R, panel report, para. 10.37. This panel report was not appealed.
Appellate Body report, para. 188.
that the Appellate Body was exclusively called upon to review the panel's threshold issue of the scope of application of non-violation complaints. Its mandate did not allow it to substantively address the European Communities additional argument that a Member State cannot have reasonable expectations of continued market access for products which are shown to pose a serious risk to human life or health. According to the Appellate Body, such an argument 'relates to the substance of a claim that has been determined to fall within the scope of application of Article XXIII:1(b) and, in particular, concerns the issue whether a 'benefit' has been 'nullified or impaired' by a measure restricting market access for products posing a health risk.'\(^{51}\) The European Communities did not appeal the panel's findings relating to the 'nullification or impairment' of a benefit through the frustration of reasonable expectations by application of the French Decree.

The Appellate Body's statement indicates, and rightly so, that the impact of this finding should not be overrated. In essence, it is limited to the procedural question of whether a measure which pursues health objectives and can be justified under Article XX of the GATT 1994 can at the same time be formally challenged in a non-violation case. In substance, however, such a measure consistent with Article XX is arguably unlikely to meet the requirements for a successful non-violation complaint. The chapeau of Article XX generally prevents the abuse of exceptions under Article XX and requires a Member State to act in good faith. The protection of good faith is also at the core of non-violation complaints.\(^{52}\) It would seem inconsistent for a panel to find that a Member State acted in conformity with the chapeau of Article XX (i.e., a finding of acting in good faith), and for the same panel to find frustration of legitimate expectations (i.e., a finding of not acting in good faith) under the doctrine of non-violation. Moreover, the participants in the Uruguay Round knew that the value of the concessions negotiated in that Round could be adversely affected by measures taken to pursue legitimate policy objectives as stipulated in Article XX of the GATT 1994.\(^ {53}\) It might be a difficult task for a complainant to present valid arguments to the effect that legitimate expectations in terms of a benefit reasonably expected to accrue to it shall prevail over a measure lawfully adopted to protect, for instance, human, animal or plant life or health.

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\(^{51}\) Appellate Body report, para. 190.


\(^{53}\) Cf. the European Communities' submission in appeal, Appellate Body report, para. 38.
THE PANEL REPORT
MEASURES AFFECTING ASBESTOS AND ASBESTOS–CONTAINING PRODUCTS

Report of the Panel

The report of the Panel on European Communities – Measures Affecting Asbestos and Asbestos–Containing Products is being circulated to all Members, pursuant to the DSU. The report is being circulated as an unrestricted document from 18 September 2000 pursuant to the Procedures for the Circulation and Derestriction of WTO Documents (WT/L/160/Rev.1). Members are reminded that in accordance with the DSU only parties to the dispute may appeal a panel report. An appeal shall be limited to issues of law covered in the Panel report and legal interpretations developed by the Panel. There shall be no ex parte communications with the Panel or Appellate Body concerning matters under consideration by the Panel or Appellate Body.

Note by the Secretariat: This Panel Report shall be adopted by the Dispute Settlement Body (DSB) within 60 days after the date of its circulation unless a party to the dispute decides to appeal or the DSB decides by consensus not to adopt the report. If the Panel Report is appealed to the Appellate Body, it shall not be considered for adoption by the DSB until after the completion of the appeal. Information on the current status of the Panel Report is available from the WTO Secretariat.
I. INTRODUCTION

1.1 In a communication dated 28 May 1998, Canada requested consultations with the European Communities (EC) pursuant to Article XXII of the General Agreement on Tariffs and Trade (GATT), Article 11 of the Agreement on the Application of Sanitary and Phytosanitary Measures (SPS Agreement) and Article 14 of the Agreement on Technical Barriers to Trade (TBT Agreement), concerning certain measures taken by France for the prohibition of asbestos and products containing asbestos (WT/DS135/1 – G/SPS/GEN/72 – G/TBT/D/15). Canada’s request states that these measures include, but are not limited to, Decree No. 96-1133 of 24 December 1996 (the “Decree”) banning asbestos, issued pursuant to the Labour Code and the Consumer Code, as amended. On 12 June 1998, Brazil asked to take part in the consultations because of its substantial trade interest (WT/DS/135/2).

1.2 In a communication dated 8 October 1998, Canada informed the Dispute Settlement Body (DSB) that the consultations held with the EC had failed to resolve the dispute satisfactorily. Consequently, Canada requested the DSB to establish a panel to examine the French measure concerning the prohibition of asbestos and products containing asbestos. In its communication, Canada claimed that the Decree, as well as any other measure which Canada might indicate, were inconsistent with Articles 2 and 5 of the SPS Agreement, Article 2 of the TBT Agreement, Articles III and XI of the GATT 1994, and, under Article XXIII:1(b) of the GATT 1994, nullified or impaired one or several advantages accruing to Canada directly or indirectly under the WTO Agreement or impeded the attainment of an objective of the Agreement owing to the fact that the banning of asbestos by France was applied whether or not it conflicted with the Agreement (WT/DS/135/3).

1.3 At its meeting held on 25 November 1998, the DSB established a panel pursuant to Canada’s request. At that meeting, the parties to the dispute agreed that the Panel should have the following terms of reference:

“To examine, in the light of the relevant provisions of the covered agreements cited by Canada in document WT/DS135/3, the matter referred to the DSB by Canada in that document and to make such findings as will assist the DSB in making the recommendations or in giving the rulings provided for in those agreements”.

1.4 On 29 March 1999, the parties to the dispute agreed that the Panel would be constituted as follows (WT/DS135/4):

Chairman: Mr. Adrian Macey;
Members: Mr. William Ehlers;
Mr. Åke Lindén.

1.5 Brazil, the United States and Zimbabwe reserved their rights as third parties to the dispute, in accordance with Article 10 of the Dispute Settlement Understanding.

1.6 The Panel met with the parties on 1 and 2 June 1999 and with third parties on 2 June 1999. A meeting with scientific experts was held on 17 January 2000. The second substantive meeting with the parties was held on 20 January 2000.

1.7 In a communication dated 27 September 1999, the Chairman of the Panel informed the DSB that the Panel was unable to present its report within the six-month period provided in Article 12.8 of the Memorandum of Understanding. The reasons for the delay are set out in document WT/DS/135/5. In two subsequent communications dated 7 March 2000 and 28 June 2000 respectively, the Chairman of the Panel informed the DSB that the Panel would require more time to complete its report (WT/DS135/6 and WT/DS135/7).

1.8 The Panel transmitted its interim report to the parties on 13 June 2000. Its final report was submitted to the parties on 25 July 2000.
II. FACTUAL ASPECTS

A. BASIC DATA CONCERNING ASBESTOS

2.1 Asbestos is a “fibrous mineral of hydrated silicates” which can be divided into two groups: amphiboles and serpentine. There are five varieties of asbestos within the amphibole group: anthophyllite, amosite (or brown asbestos), crocidolite (or blue asbestos), actinolite, and tremolite. The serpentine group comprises only chrysotile (or white asbestos). These varieties of asbestos have different physical and chemical properties.

2.2 It is mainly amosite, crocidolite and chrysotile which are exploited for industrial and commercial purposes. The special qualities of asbestos fibres (for example, resistance to very high temperatures and to different types of chemical attack) due to their particular physical and chemical properties have meant that they have been put to many uses, for example, for the manufacture of industrial and consumer products and in the building industry.

B. DECREE NO. 96-1133 OF 24 DECEMBER 1996 BANNING ASBESTOS

2.3 On 24 December 1996, the French Government adopted Decree No. 96-1133 banning asbestos, issued pursuant to the Labour Code and the Consumer Code (décret no. 96-1133 relatif à l’interdiction de l’amiante, pris en application du code de travail et du code de la consommation) (hereinafter “the Decree”). The Decree entered into force on 1 January 1997. The following are its principal provisions:

2.4 Article 1 provides for a ban on asbestos in the following terms:

“I. – For the purpose of protecting workers, […] the manufacture, processing, sale, import, placing on the domestic market and transfer under any title whatsoever of all varieties of asbestos fibres shall be prohibited, regardless of whether these substances have been incorporated into materials, products or devices.

II. – For the purpose of protecting consumers, […] the manufacture, import, domestic marketing, exportation, possession for sale, offer; sale and transfer under any title whatsoever of all varieties of asbestos fibres or product containing asbestos fibres shall be prohibited […]”

2.5 Article 2 of the Decree allows some exceptions to the ban in Article 1.

“II. – On an exceptional and temporary basis, the bans instituted under Article 1 shall not apply to certain existing materials, products or devices containing chrysotile fibre when, to perform an equivalent function, no substitute for that fibre is available which:

- On the one hand, in the present state of scientific knowledge, poses a lesser occupational health risk than chrysotile fibre to workers handling those materials, products or devices;

- on the other, provides all technical guarantees of safety corresponding to the ultimate purpose of the use […]”

2.6 The exceptions allowed are the subject of an exhaustive list prescribed by the French authorities and reviewed annually (Article 2.II). Any exception under Article 2 must be the subject of a declaration by the head of the business establishment, the importer or the party responsible for domestic marketing of a product covered by Article 2; the purpose of the declaration is to be able to determine that the activity to which it refers meets the criteria set out in the first paragraph of the Article, taking into account scientific and technological progress (Article 3).
Article 4 contains requirements on the levels of exposure to be respected when manufacturing or processing products covered by Article 2, as well as their labelling and marking. Article 5 lays down penalties (fines) for breaches of the Decree’s provisions.

III. ARGUMENTS OF THE PARTIES

A. CLAIMS BY THE PARTIES

3.1 In view of the facts and arguments put forward, Canada requests the Panel to find that:

(a) Decree No. 96-1133 of 24 December 1996 banning asbestos is incompatible with the Agreement on Technical Barriers to Trade insofar as it is a technical regulation that:

(i) Creates an unnecessary obstacle to international trade contrary to the provisions of Article 2.2;

(ii) is not based on effective and appropriate international standards – nor is it in compliance with them – contrary to the provisions of Article 2.4;

(iii) is not based on orders relating to chrysotile and chrysotile-containing products with respect to the product performance of chrysotile, contrary to the provisions of Article 2.8; and

(iv) violates the national treatment disciplines and the most-favoured-nation clause of Article 2.1.

(b) In addition, the Decree is incompatible with the GATT 1994 in that it:

(i) Creates a prohibition or a restriction on the import of chrysotile and chrysotile-containing products, contrary to the provisions of Article XI.1; and

(ii) favours the national industry of products like chrysotile fibre and chrysotile-cement products, contrary to the national treatment disciplines of Article III.4.

3.2 In the event that the Panel is unable to find a violation of Article XXIII.1(a) of the GATT 1994, Canada requests that the Panel find a violation of Article XXIII.1(b) of the GATT 1994.

3.3 In view of the foregoing, Canada requests that the Panel recommend to France that it make the Decree compatible with its obligations under the Agreement on Technical Barriers to Trade and the GATT 1994.

3.4 In view of the facts and legal arguments, the European Communities (hereinafter the “EC”) ask the Panel to confirm that, in light of the rules of the GATT 1994, Decree No. 96-1133 of 24 December 1996 banning asbestos:

(i) Should not be examined in relation to the scope of Article XI of the GATT 1994;

(ii) does not establish less favourable treatment for similar imported products than for domestic products, within the meaning of Article III.4 of the GATT 1994;
The EC also ask the Panel to find that:

(i) The Decree is not covered by the Agreement on Technical Barriers to Trade and that, in any case, it complies with the relevant provisions of that Agreement.

Finally, the EC further ask the Panel to find that:

(i) Article XXIII.1(b) of the GATT 1994 does not apply.

Consequently, the EC ask the Panel to reject all the arguments put forward by Canada.

B. FACTUAL ARGUMENTS

1. Introductory remarks

Canada recalls that, since 1 January 1997, France has prohibited the manufacture, processing, sale, importation, exportation, domestic marketing, possession for sale, offer and transfer of all varieties of asbestos fibres, regardless of whether these substances have been incorporated into materials, products or devices. There are four temporary exceptions to this general ban. The ban will become total on 1 January 2002. Canada is challenging the ban on chrysotile fibre and products containing it. Before the ban, France imported 20,000 to 40,000 tonnes of chrysotile fibre from Canada each year, equivalent to over two thirds of the total quantity imported into France. After the French Government announced its intention to ban asbestos in July 1996, imports of Canadian chrysotile dropped to under 15,000 tonnes. In 1997, the year the ban took effect, just 18 tonnes were imported. At present, for all practical purposes, they have disappeared.

Canada asserts that, unlike amphibole fibres – the asbestos most hazardous to health, which was previously used widely in France – chrysotile fibres can be used without incurring any detectable risk. Chrysotile fibres are found today in a limited number of products, where they are encapsulated in an inert matrix. These products do not pose any risk to businesses, the general public, or the environment. Exclusive use of chrysotile fibre and the adoption of effective methods for reducing dust creation are sufficient health-protection guarantees. Before banning asbestos, France had been applying controlled use practices. It is past uses, especially the spraying of brittle asbestos in fireproofing, which are the main causes of asbestos-related health problems in France. Given the long latency period between exposure to asbestos and the onset of any related diseases, workers who were victims of heavy exposure with virtually no protection 30 years ago are experiencing serious health problems today. People are currently being exposed to asbestos dust released by fireproofing that is disintegrating. The use of materials containing brittle asbestos was prohibited when the Decree that is being challenged by Canada was adopted.

According to Canada, the ban on asbestos does nothing to correct the problems resulting from past asbestos use. The ban was adopted at a time of heavy media coverage of diseases caused by uncontrolled use of asbestos in France. Alarmist campaigns condemned all forms of asbestos use and there was pressure on public officials to take action. Spurred by courts that were concurrently examining the liability of political leaders in the “tainted blood” affair, the French Government chose to prohibit chrysotile and all its uses in the hopes of assuaging public opinion, which had been badly shaken. The ban on asbestos is nothing but a political reaction on the part of the French Government to anti-asbestos propaganda. In many respects, the French reaction is identical to the reaction of the United States Environmental Protection Agency (EPA) in 1989, when it prohibited asbestos under pressure from panicked United States public opinion. Unable to justify its ban scientifically, the EPA had to reverse its decision in 1992 and acknowledge that modern products
containing chrysotile enclosed in a matrix of cement or resin do not pose any detectable risk to public health. Today, although amphiboles are prohibited in the United States, a number of products containing non-brittle chrysotile are permitted.

3.11 Canada notes that France claims that its measure is based on a report from the Institut National de la Science et de la Recherche Médicale (INSERM). Yet several experts who have analysed the report have sharply criticized the methods of the INSERM researchers. They have also very severely criticized the findings of their report. Those experts are of the opinion that the INSERM report is not a credible basis for justifying a total ban on all varieties and all uses of asbestos for public health purposes. In the justification of the measure given by Directorate General III (Industry) of the European Commission, it was admitted that application of a practice of controlled use of asbestos in industry enabled true control of the risk of diseases attributable to occupational exposure, in the case of workers involved in the mining and processing of chrysotile. The practice of controlled use also applies to other possible situations of exposure to asbestos. The ban on chrysotile involves using substitute products whose effects on human health are, by INSERM’s own admission, unknown. The use of substitute products is not covered by clearly-established standards, yet the INSERM researchers state how important it is to evaluate the potential risks involved. Permitting the use of these products without taking the requisite precautions may cause a repeat of the mistakes that were made at the time when the risks associated with asbestos use were unknown or poorly controlled. The undetectable risk from chrysotile is thus replaced by the unknown risk from substitutes. This results in inconsistencies in the regulation of potentially hazardous products in France.

3.12 Canada is not challenging the right of the Members of the WTO to take whatever measures are necessary to protect the health and safety of their populations. That right, however, must be exercised in compliance with the obligations that a Member has under the WTO Agreements. In that respect, France was not entitled to adopt a total ban on asbestos, with no distinctions concerning fibres and products, in the absence of scientific proof of the health risks posed by modern products containing chrysotile. Yet, one thing needs to be said: the total ban is both irrational and disproportionate considering the fact that the manufacture and use of modern chrysotile asbestos products do not pose any detectable health risks. In Canada’s view, the scientific data on which France has based its case do not justify such a radical measure as the ban on chrysotile fibres and the prohibition of all uses that can be made of them. Moreover, Canada affirms that the ban does nothing to correct the problem of past exposure to asbestos, nor does it solve the problem of managing the asbestos already in use in France. Ultimately, the total ban on chrysotile fibre and chrysotile-containing products is an excessive measure. Other measures that are less restrictive on international trade and thus compatible with France’s international obligations were available and would have enabled France to achieve its aim just as well as a ban. Accepting France’s approach would give each Member the option of completely banning potentially hazardous natural products rather than adopting a reasonable approach of risk management based on their use. In terms of international trade, the total ban on asbestos creates a barrier to importation of chrysotile fibre and chrysotile-containing products into the French market. It is also an internal measure that upsets the competitive relationship between chrysotile fibre and chrysotile-containing products and like products of French or foreign origin: it is therefore a discriminatory measure.

3.13 The European Communities respond that the Canadian statement that amphiboles are “the asbestos most hazardous to health” is only recognized for the risk of mesothelioma (as explained in the INSERM report): it is accepted, however, that chrysotile has a carcinogenic potential at least comparable to amphiboles for lung cancer, as claimed below by the EC. This confusion is systematic in the Canadian arguments, as though it was less serious and less dangerous for health to suffer lung cancer than mesothelioma. It is also inaccurate to state, as Canada does, that amphibole fibres “were used widely in France”. Between 1945 and 1988, some 97 per cent of the asbestos consumed by France was chrysotile asbestos and since 1988 it has represented the totality. Regarding the methods which Canada deems “effective for reducing dust creation”, the EC note that the dust
levels created through the use of some materials that comply with the standard ISO 7337 are very much higher than the limit values allowed in France and even than the limit values recommended by the WHO group mentioned by Canada. Canada confines its analysis to the processing of raw asbestos and deliberately disregards users of material containing asbestos working on building sites or carrying out servicing or maintenance work. The dust elimination techniques cited in the standard ISO 7337 have shown their ineffectiveness for this type of changeable and mobile work; the dust levels regularly measured greatly exceed the thresholds adopted by France and many countries for which it has been proved that there is an excessive risk.

3.14 According to the EC, it is incorrect to state, as Canada does, that the use of materials containing brittle asbestos was banned in France prior to 1996. Before 1996, France authorized the use of all types of brittle asbestos, except for certain specifically prohibited uses such as fireproofing. It cannot be asserted either that “The main causes of asbestos-related health problems in France are past uses, especially the spraying of brittle asbestos in fireproofing”. In fact, the increase in asbestos-related diseases precedes the practice of fireproofing: it started in the 1950s, whereas fireproofing only really started to be practised from the 1960s onwards; the very long latency period for cancer caused by asbestos means that cancer cases can only be attributed to fireproofing since the 1990s, whereas the mesothelioma mortality rate in France has for a long time shown a rapidly ascending curve. Furthermore, this increase in disease concerns workers in very different industrial sectors.

3.15 The EC point out that, when Canada claims that the prohibition imposed by France in summer 1996 was “a political reaction … to anti-asbestos propaganda”, its interpretation of the decisions by a country’s Government is wrong: the EC explain below all the restrictive measures taken by France for a long time and gradually. The statement does not take into account the fact that seven other European countries took an identical measure some years ago, without Canada accusing them of having taken their decisions for “political” reasons or attacking them, as far as the EC know.

3.16 The EC note that it also subjective to state that several experts who analysed the INSERM report “sharply criticized” the methods used and “very severely criticized” its findings. The report by the Expert Panel of the Royal Society of Canada appointed by the Canadian Government contains praise for the work of the INSERM experts, even if certain points are raised, as is the rule for complex scientific problems. Some pages of the report of the Royal Society of Canada formally contradict Canada’s statement: for example, pp. 4-7 of the English text of the report, which list the important points on which the Panel agrees with INSERM’s findings, and the comments on pp. 9-18. It seems that, on the major points of INSERM’s conclusions, the Canadian experts agree or make comments which are a matter of scientific discussion without questioning INSERM’s findings. The EC also note that the Panel of the Royal Society of Canada worked too quickly, as is recognized in several instances (see, for example, page 15 of the English text), that it was unable to reach a consensus (page 15 of the English text), and that it worked on the basis of an incomplete document whose translation had not been reviewed (page 1), which clearly accounts for certain incorrect interpretations due to misunderstandings. This last aspect calls for a comment: Canada in fact obtained a copy of the pre-publication text of the INSERM report, without making a request to INSERM or to the French Government; it translated this into English without having the translation revised by the authors, who were not informed of the existence of the Panel and still less of its composition. At no time (neither during the examination of the INSERM report nor afterwards) did the Government of Canada or the Royal Society of Canada request explanations or comments from the French experts who participated in the INSERM report, which would no doubt have helped to remove certain ambiguities. It is absolutely alien to the tradition of scientific discussion to proceed in this way: scientific debate is obviously necessary, but it is usually based on contradictory discussions to which each may contribute arguments and not on procedures from which one of the parties is excluded.9

3.17 The EC reject several of Canada’s statements concerning substitute products. It is incorrect to state that “the use of substitute products is not covered under clearly estab-
lished standards”. The products used as substitutes for asbestos differ according to the use to which they are to be put. They are all chemical products. As such, they are subject to the regulations applicable to chemical substances for the purposes of risk prevention, and where appropriate to the regulations on carcinogenic substances if there is an established or suspected risk of carcinogenicity. Regarding the Canadian statement that “The undetectable risk from chrysotile is thus replaced by the unknown risk from substitutes”, the EC point out that undetectable risk is not the same as absence of risk, contrary to what Canada seeks to claim. On this point, the INSERM report showed clearly and in detail that the low risks associated with low levels of exposure (either due to chrysotile or any other substance) are indeed undetectable for methodological reasons that are explained in detail. It is therefore quite unacceptable to try to “absolve” chrysotile on the pretext that if it is inhaled in very low doses there will be an undetectable risk: if one followed this reasoning, it could be concluded that no substance is carcinogenic on the pretext that the corresponding risks are undetectable at very low levels of exposure (for example, it is clear that the risk of cancer from tobacco is undetectable if one inhales tiny quantities of cigarette smoke). The truth is that the risks from chrysotile are not only detectable, but have been detected for a long time because they are so great if there is a high level of exposure; the EC assert that this is still the case today, even with “modern” products.

3.18 The EC also emphasize that the majority of substitute products for asbestos are substances which have been used regularly for other purposes for decades without any risk associated with their use being detected, unlike the scientifically proven risk associated with the use of asbestos. No substitute product for chrysotile in fibro-cement is recognized as carcinogenic at the international level. Some fibrous substitute products used in a number of very limited cases may be suspected of being carcinogenic, but in any case their carcinogenicity for humans has not been proven scientifically at the international level. This scale of risk has been known since June 1996, the date of the French decision. It cannot be asserted that there is an “absence of scientific proof of the health risks posed by modern products containing chrysotile”. The EC reject this statement, based on international authorities such as the WHO and the ILO. To France’s knowledge, the methods of manufacturing asbestos-cement have not developed to any significant degree for several years. Furthermore, the fibre emissions are much higher than the limit values allowed for the servicing and maintenance of buildings, which Canada does not mention. Consequently, the concept of “modern” products does not have any meaning. The aim of the French regulation contested by Canada is to prevent the extension of the existing risk by halting the dissemination of this product, which is scientifically recognized as dangerous. Other measures have been taken by the Government in order to deal with problems due to past exposure and the management of asbestos already in place.

3.19 The EC reject the Canadian statement that “Accepting France’s approach would give each Member the option of completely banning potentially hazardous natural products rather than adopting a reasonable approach of risk management based on their use”. Contrary to Canada’s assertion, the risks of chrysotile are not potential but proven, as the WHO declared in 1998. Moreover, the WTO Agreements give each Member the sovereign right to choose the level of protection it wishes to adopt. Confronted with such a diffuse risk that is impossible to control in such divergent exposed populations, no country can state that it has been able to establish a responsible approach to the management of a carcinogenic risk other than by replacing the incriminated substance. Contrary to Canada’s claim, the effect of the French measure is not to favour products like asbestos of French origin. France does not manufacture the substitute products mainly used to replace asbestos in asbestos-cement, but imports them from other countries. For example, PVA, an asbestos-cement substitute, is produced in only two plants worldwide, one in China and one in Japan. Moreover, substitute products are not like products because they are less dangerous and their chemical composition differs.

2. Economic and trade data

3.20 Canada indicates that, in 1997, global production of chrysotile was about 2 million tonnes. The Commonwealth of Independent States is the number one producer, fol-
owed by Canada. Then come China, Brazil and Zimbabwe. International trade in chrysotile is of special importance to Canada, which is the world’s number one exporter. Canadian exports to the five continents totalled 430,000 tonnes in 1997. Before the Decree took effect, more than two thirds of French imports of chrysotile came from Canada. The effect of the Decree was felt as of 1996 due to the announcement of the intention to ban asbestos: French imports of chrysotile from Canada dropped by more than half compared to 1995, from approximately 32,000 tonnes to about 14,000 tonnes. Once the ban took effect in early 1997, imports, for all practical purposes, disappeared. For 1997, imports were just 18 tonnes of Canadian chrysotile fibres. The Decree has eliminated the French market for chrysotile.

All Canadian mines are located in Quebec Province. With annual production valued at Can$ 225 million, the chrysotile mining industry currently provides about 1,300 direct jobs and as many indirect jobs in Quebec. Moreover, the chrysotile processing industry provides about 1,500 jobs in friction product, composite material and asbestos textile companies, mainly located in Quebec. More than 4,000 Canadian jobs depend directly or indirectly on the chrysotile industry.

3.21 Canada explains that there are up to 3,000 commercial applications for asbestos, the most important of which are the following: (i) as a reinforcement material for cement, plastic or rubber; (ii) as part of brake linings or clutches; (iii) in the form of spun fibres for the protection of insulating woven fabrics or cords; (iv) in the past, it was applied by spraying in order to give boats, structural beams, pipes and boilers fire-resistant finishes; (v) in the past, it was used as a thermal insulation on pipes and boilers. Throughout the world, the most important current application is the manufacture of chrysotile products. The worldwide production and use of amphiboles (amosite and crocidolite) comprise less than 3 per cent of the total amount of asbestos produced and continue to decline. Therefore, chrysotile is now the only type of asbestos used. Furthermore, modern asbestos products are not brittle and fibre emissions during their transportation, installation and use (including subsequent losses due to alteration and abrasion) have been reduced to an absolute minimum, unlike earlier products which released much larger amounts of fibres into the environment.

3.22 The European Communities explain that, in 1973, world production of asbestos reached a peak of 5.2 million tonnes. Since then, production has steadily declined, falling to 1.92 million tonnes in 1997, some 60 per cent of which is produced by Canada and Russia. Canada is a major producer of chrysotile asbestos. It consumes little asbestos and thus exports the bulk of its production. It should be noted that Canada produces and exports only chrysotile asbestos. Canada is thus exporting to other countries the “public health risk” associated with chrysotile. The EC emphasize that, since asbestos was first used for industrial purposes, some 95 per cent of the asbestos consumed worldwide has been chrysotile asbestos. Between 1945 and 1980, some 97 per cent of the asbestos consumed by France was chrysotile asbestos. From 1988, onwards, all the asbestos consumed in France has been chrysotile asbestos.

3.23 The EC indicate that, given the many uses of asbestos, the range of asbestos-containing products is extremely wide. Depending on their physical appearance, there are five main categories: (i) bulk asbestos: asbestos wadding for the thermal insulation of ovens, boilers, fire-doors, refrigerating equipment, flocked asbestos for the underside of concrete slabs and for steel frames for the fireproofing and soundproofing of buildings; (ii) asbestos sheets or boards: asbestos paper and board for thermal insulation, for the protection of welded joints (plumbing) and for the protection of work surfaces (glass industry), boards for the fitting of false ceilings, fireproof surfaces, partitions, etc.; (iii) textile asbestoses: asbestos cord (sealing of oven doors, laboratory applications), buffer strips (to protect against heat), fireproof coverings and curtains, air, gas and liquid filters, insulation of electrical equipment; (iv) asbestos incorporated in cement products (asbestos-cement): roof tiles, roof cladding, window sills, facing of buildings, interior partitions and false ceilings, other forms of panelling, flues, ventilation shafts, rainwater pipes, plant tubs and gardening equipment; (v) asbestos in binding or bonding agents (resins, bitumens): friction linings (brake linings, clutch linings, linings for presses, winches, gantries, lifts, engines), road
surfaces, flooring, decorative shingles, sealants, plaster-based finishes and coatings, glues and gums and asbestos-based paints. Asbestos-cement accounts for 90 per cent of all asbestos used.

3.24 As far as the EC know, the production methods for asbestos-cement have not developed significantly in recent years. Canada cannot therefore claim that “modern asbestos products are not brittle and fibre emissions during their transportation, installation and use ... have been reduced to an absolute minimum, unlike earlier products”. Furthermore, fibre emissions greatly exceed the limit values permitted for all building servicing and maintenance activities, which Canada does not mention.

3.25 Canada states that, since the pathogenic potential of amphiboles is greater than that of chrysotile – this is true both for mesotheliomas and for lung cancer – a review of how amphibole asbestos has been used in France in the past is necessary. Here again, the data advanced by the EC must be corrected. According to Canada, it is incorrect to state, as the EC claim, that from 1945 to 1988 only 3 per cent of asbestos used came from the amphibole family; this is in complete contradiction with the studies reviewed by INSERM, which reveal that of the samples taken after 1990 in buildings containing asbestos in France, 18 per cent of the samples contained amphibole fibres, including 13 per cent which contained only amphibole fibres. It is also incorrect to claim that the use of amphiboles in France ceased in 1988: the INSERM data contradict these claims by the EC, clearly stating that the use of amphiboles continued well beyond 1988, i.e. at least until 1992. Given the much higher pathogenic potential of amphiboles compared to chrysotile and the fact that amphiboles are present in nearly 20 per cent of more than 1,000 exposures to asbestos recorded and measured in France, it is quite plausible to attribute the problem of asbestos in France to past situations. While amphiboles have been banned in France since 1994, along with the use of friable materials, the significant risks associated with their presence remain. For instance, according to the WHO, building maintenance personnel are today exposed to high risks, not because of the current uses of chrysotile, but particularly because of the large quantities of “mixed in-place friable asbestos fibres”. Similarly, Canada seeks to refute the French argument that, since 95 per cent of the asbestos used in the world today is chrysotile asbestos, the proportion of diseases that is attributable to it is “quite indisputably” of the same magnitude. This reasoning, dubious at best, denies the entire heritage of past uses of amphibole fibres, the existence of a latency period, as well as the greater pathogenicity of amphibole asbestos.

3. French legislation in its context

3.26 Canada considers that special attention should be paid to social and political climate in which the French ban was adopted. This climate is very revealing as concerns the factors that motivated France’s action and, in large part, explains the severity of that action: the political need to assuage an alarmed French population – what France’s Minister for Education, Mr. Claude Allègre, then called “un phénomène de psychose collective” (mass psychosis). Alarmed political leaders were evidently afraid of being held criminally liable if they were not seen to be doing something about the problem. In recent years in France, media coverage of several cases of diseases linked to occupational exposure to asbestos has inflamed public opinion. That, obviously, contributed to the adoption of the Decree. Indeed, since the spring of 1995, the media in France have presented alarmist articles and reports on the risks of asbestos to a public that was already shaken by two other public health cases: the “mad cow” and “tainted blood” crises. Those controversies led to the examination of the liability of political leaders in criminal courts. In the summer of 1995, with public outcry centred on Jussieu University as backdrop, the French Government commissioned a report from INSERM. INSERM was asked to study the health effects of the main types of exposure to asbestos. In late 1995, France presented an oversight programme to reduce the risks associated with asbestos. A ban did not seem to be under consideration at that time. On 21 June 1996, INSERM submitted an executive summary of its report to the French Government. On 25 June 1996, the filing of a complaint by the Association nationale de défense des victimes de l’amiante (ANDEVA) (National Association for the Protection of Asbestos Victims), specifically accusing the Government of inaction, re-
portedly helped to move France towards the ban. Seven days later, on 2 July 1996, INSERM published the summary of its report on the health effects of the main types of exposure to asbestos. The next day, 3 July 1996, France announced its intent to declare a ban on asbestos.

3.27 Canada asks why the great haste? Why was the ban announced less than two weeks after the French Government first obtained the summary of the INSERM report? Is it credible that in only two weeks the French Government could review and analyse the scientific, social and economic considerations related to such an important decision? And all this some 16 months before the final report – as opposed to the executive summary – saw the light of day? And why was the ban announced only one week after a legal action for poisoning was filed? The timing suggests that the ban was politically motivated and that the INSERM report merely provided the *ex post facto* scientific rationale. On 31 July 1996, yet another complaint for asbestos poisoning was filed, this time against eight former French ministers. All this took place during the contemporaneous examination of French political leaders’ criminal liability in the tainted blood scandal. According to Canada, French political leaders were evidently under tremendous pressure to take action, any action, to be seen to be remedying a situation blown out of all proportion by the media. French parliamentarians have themselves acknowledged the pressure from public opinion and the media. In a joint report on asbestos, the Senate and the National Assembly stated: “It is because a definitive response was so late in coming that the asbestos issue is today drawing so much media attention and political opinion is being diverted.”

3.28 Canada notes that, in July 1996, thus at the same time as it was preparing to prohibit asbestos, the French Government called for research proposals aimed specifically at developing fibres to replace asbestos. This specific effort on the part of the French Government was designed to accelerate the conversion of industrial groups to substitute fibres and to enable France to become a leader in substitute fibres in the global market. According to Canada, it was in that climate, when a panic had overtaken public opinion and commercial interests in French industry were moving toward the development of substitute products and fibres, that the Decree banning asbestos was adopted.

3.29 The *European Communities* note that Canada criticizes the rapidity with which the French Government announced that it was going to ban asbestos after the publication of the executive summary of the INSERM report, as well as the time taken by the French authorities to publish the final report. According to the EC, these arguments show that, on the one hand, Canada appears to ignore what constitutes a risk assessment and, on the other, how much time it takes to produce a formal publication of the results of a risk assessment. It is not uncommon practice for governments to act on the basis of unpublished results of a risk assessment. It takes on average more than one, if not two, years before formal publication in the form of a book or otherwise because the documents need careful editing and all bibliographical references must be checked. This is normal procedure also for similar publications by the WHO, the IARC and other international bodies such as the Joint FAO/WHO Expert Committee on Food Additives (JECFA) or the Joint FAO/WHO Meeting on Pesticide Residues (JNPR), whose formal publications frequently take more than a year to be published. It would be unreasonable if governments were to be prevented from taking action quickly on the basis of the substantive results which they have in the form of an executive summary as protection of health should be given priority over formalities prior to publishing results. The EC also emphasize that the interim results and the executive summary were immediately made available to Canada for comments. The only criticism which Canada was able to make against the INSERM report is the critique made in the report by the Royal Society of Canada. But by looking at the text of that report, it can be seen that the authors’ criticisms were not all unanimous; moreover, this report, which criticizes meticulous scientific work by 11 scientists, was adopted within the space of two weeks. The EC also consider that Canada’s attempt to convey the impression that the Canadian scientists who reviewed the INSERM report are “better” than the French scientists who prepared the report is unacceptable. This suggestion has no foundation in fact nor in science. Canada is of course entitled to employ the scientists it wishes to assess...
asbestos-related risks in Canada in the past. The EC do not criticize the quality of their report. Nevertheless, the EC expect Canada to show equal respect for the French scientists who prepared the INSERM report, bearing in mind that this report conforms to the great majority (if not all) of the views of the scientific community on the risks posed by asbestos and that its findings have even been confirmed subsequently in reports by international bodies such as the WHO report 203.21

3.30 The EC state that, contrary to Canada’s assertions, France’s ban on asbestos is not an isolated and irrational act arising out of a wave of panic. During the 1970s and early 1980s, scientific evidence showed that all types of asbestos were likely to cause asbestosis, lung cancer and mesothelioma. In response to these public health concerns, many European countries began to introduce national legislation restricting, and ultimately prohibiting, the marketing and use of asbestos. Rules designed to control the use of asbestos in the workplace and ultimately to limit the release of asbestos into the atmosphere were also introduced. It cannot be asserted, as Canada does, that in spring 1995 French public opinion was shaken by the “mad cow” crisis: the events to which Canada refers occurred in 1996. Likewise, it is incorrect to state that “In late 1995, France presented an oversight program to reduce the risks associated with asbestos … [and that a] ban did not seem to be under consideration at that time.” A total ban was one of the measures envisaged by the French Government as of 1995. A decision was finally taken not because of the filing of complaints by a victims’ association but as a result of the findings in the INSERM report given to the ministers concerned in June 1996.

3.31 The EC point out that a number of countries have either introduced bans on asbestos or plan to do so. The health authorities of many more are looking in detail at the dangers which the various forms of asbestos represent. As early as 1983, Iceland introduced a ban (with some limited exceptions) on all types of asbestos (updated in 1996). In 1984, Norway introduced a ban (with some limited exceptions) on all types of asbestos (revised in 1991). Since 1989, Switzerland has in principle prohibited the use of asbestos, including chrysotile asbestos, in accordance with an amendment to the law on environmentally harmful substances. In only two types of situation may an exception be granted to this principle, subject to the express authorization of the Federal Office for the Protection of the Environment: (i) where technology has not yet identified an asbestos-free substitute and where the amount of asbestos used does not exceed what is necessary to attain the desired objective; or (ii) where the technical characteristics of the product or object are such that it is impossible to use alternative components which do not contain asbestos (Annex 3.3 to the Order of 11 January 1989 amending the Order of 9 June 1986). An exemption may also be granted “on grounds of national defence” for material used in carrying out defence-related tasks, although only with the agreement of several ministerial departments. In New Zealand, the Asbestos Regulations 1983 ban the use of asbestos in the construction of new buildings. The ban applies to all types of asbestos: chrysotile, crocidolite, tremolite, actinolite, anthophyllite and amosite. In January 1999, a new amendment to the Asbestos Regulations of 1983 was published, providing for: (i) a ban on the import of chrysotile asbestos; (ii) amendments to the law relating to the recycling of asbestos-containing products; (iii) more rigorous and more systematic health checks; and (iv) a higher level of protection for workers who handle asbestos. Since 1 January 1999, the Czech Republic no longer imports nor processes asbestos, in whatever form. Asbestos is on its list of dangerous (carcinogenic) substances. In Australia, asbestos legislation falls within the remit of each of the federal states. However, there is a government agency, the National Occupational Health and Safety Commission (NOHCS), which is responsible for promoting health and safety at work throughout the country. The research division of the NOHCS recently published a report on the health risks posed by the use of chrysotile asbestos.

3.32 The EC point out that, with a view to ensuring a high level of health protection in the European Community and preserving the unity of the single market, numerous legislative acts have been adopted at Community level since 1980. On 4 May 1999, it was decided to introduce a total ban on all types of asbestos with effect from 1 January 2005. As early as 1972, Denmark introduced a ban on applying asbestos by the spraying process.
and on using it for insulation. It would appear to have been the first such ban anywhere in the world. In 1986, Denmark placed a total ban on asbestos, allowing some limited exceptions up to 1993. In 1972, the United Kingdom banned imports of crocidolite (blue asbestos). This decision was supplemented by the setting of limit values for exposure to asbestos dust at the workplace. In 1975, Sweden banned the marketing and use of crocidolite, and in 1976 the use of asbestos-cement products. In 1986, it introduced a total ban on asbestos with a few exceptions. In 1977, France set initial limit values for asbestos dust and in 1978 it banned the spraying of asbestos fibres. In 1996, France imposed a total ban on asbestos, with some limited exceptions. In 1977, the Netherlands banned crocidolite and the use of asbestos for spraying. In 1991, it introduced a total ban on asbestos, with exceptions which applied until 1997. In 1990, Austria banned the use of chrysotile asbestos, with some limited exceptions. In 1992, Finland and Italy imposed a total ban on asbestos, with exceptions which applied until 1993. In 1993, Germany imposed a total ban on asbestos, with some limited exceptions. In 1998, Belgium imposed a total ban on asbestos, with some limited exceptions.

3.33 The EC recall that, at the beginning of the 1980s, it became clear that there was a need for harmonization at European Community level. Two key Community directives were adopted: Directive 83/477/EEC and Directive 83/478/EEC. Under Directive 83/477/EEC, the Member States of the European Community must require employers to assess the risks to workers of exposure to asbestos; they must also take the necessary preventive measures. The Directive prohibits the application of asbestos by means of the spraying process and sets limit values for exposure. It also lays down a range of preventive measures such as medical surveillance of workers and appropriate cleaning of the place of work. Lastly, it lays down specific measures to be taken to protect workers involved in removing asbestos. Directive 83/478/EEC laid down the first Community measures in respect of the marketing of asbestos. It prohibits the marketing and use of crocidolite (with some exceptions) and introduces compulsory labelling for all asbestos-containing products. Directive 85/610/EEC extends this prohibition to all other types of asbestos for certain specific uses. During the same period, Council Directives 78/319/EEC and 87/217/EEC were adopted, laying down measures to prevent and reduce environmental pollution, including the control of asbestos-containing waste. Both these Directives were updated by Directive 91/689/EEC.

3.34 The EC also recall that, at the beginning of the 1990s, a good deal of scientific evidence concerning the risks from asbestos became available and safer substitute products were developed to replace asbestos in many applications. European legislation evolved rapidly so as to achieve tighter control of the risks. In 1991, for example, all types of asbestos were classed as category I carcinogens under Directive 67/548/EEC. The Directive defines category I as comprising substances known to be carcinogenic to man. Then in 1991, Directive 91/382/EEC amending Directive 83/477/EEC on the protection of workers (see above), lowered the limit values for exposure to 0.6 fibres per cm$^3$ for chrysotile asbestos and 0.3 fibres per cm$^3$ for all other forms of asbestos. One year previously, Directive 90/394/EEC, which deals with the protection of workers from exposure to all carcinogens, had introduced the principle of replacement. Under this principle, employers are required, where this is technically possible, to replace carcinogens used at work with substances which are not dangerous or which are less dangerous. As regards the placing of asbestos on the market, Directive 91/659/EEC introduced a total ban on the marketing and use of all forms of asbestos fibre, except chrysotile asbestos, whose use was nonetheless prohibited for fourteen specific categories of product. Other chrysotile-containing products (e.g. asbestos-cement) remained outside the Directive’s scope of application.

3.35 The EC point out that, as there is sufficient scientific evidence to justify a ban, the Commission of the European Communities has decided to propose a directive banning the marketing and use of chrysotile asbestos, with one exception and some transitional arrangements. On 4 May 1999, the proposed directive received a favourable vote from the Member States meeting within the Standing Committee set up by Directive 76/769/EEC. It is due to be finally adopted by the Commission in the near future. Under the new directive, the ban on chrysotile asbestos is to be implemented throughout the European Union.

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by 1 January 2005 at the latest. Each Member State may choose the pace at which it wishes to move towards this harmonized position. Decisions will have to be adopted in the light of the situation within the domestic industry and in accordance with the legal procedures applicable in each country. The new directive would extend the current ban to all remaining applications of chrysotile asbestos, with one exception. Only diaphragms used for electrolysis in certain chlorine plants may be exempted from the prohibition. Diaphragms constitute a special case since they represent the only current application of chrysotile asbestos for which it is not technically possible to use a substitute product without creating a safety risk (i.e. the risk of an explosion). Diaphragms are not placed on the market. Under the directive, the exemption applied to diaphragms will be reviewed (on the basis of an independent scientific risk assessment), both as part of the general review of the directive in 2003 and specifically in 2008. The directive recognizes that scientific knowledge about asbestos and substitute products is continually developing. It therefore stipulates that the relevant scientific issues and derogations be reviewed before 2003, in order to decide whether further legislative measures are necessary in this field.

3.36 The EC note that, in France, the first preventive rules on health and safety at work go back to the beginning of the century. For about a decade, French legislation has been based on the European directives on health and safety at work. All regulatory provisions are the subject of consultations with the social partners, i.e. the representatives of employer and employee associations within the Senior Council for the Prevention of Occupational Hazards. In addition to specific legislation covering certain particularly harmful substances, France has general rules on health and safety (risk assessment, informing and protecting workers, etc.). As regards the prevention of chemical hazards, France has incorporated into its Labour Code both the EC Directive on chemical agents (Articles R231-55 et seq.) and EC Directive 90/394/EEC on carcinogens (Articles R231-56 et seq.). Since 1977, France has had specific legislation on asbestos, designed to protect workers from inhaling asbestos dust. Following the example of other European countries, this legislation was progressively tightened up as knowledge developed until, finally, it was decided to impose a ban in 1996. It should be pointed out that, in France, the number of illnesses linked to the inhalation of asbestos dust which are recognized as being of occupational origin and for which compensation is paid has been rising constantly for several years. For example, between 1985 and 1995, the number of asbestos-linked occupational diseases for which compensation was paid (mesothelioma, lung cancer, asbestosis, pleural abnormalities) increased fourfold. The cost of such illnesses ran to F1.08 billion in 1997, or almost 40 per cent of the compensation budget for all diseases recognized as being of occupational origin in France. By way of comparison, musculoskeletal problems, which are very widespread and are at the top of the list in numerical terms, account for only 18.8 per cent of the total financial cost of occupational diseases. This increase in diseases, together with the lower life expectancy of workers formerly employed in the asbestos industry, recently prompted the French Government to introduce an early retirement plan for employees who worked in factories manufacturing asbestos products.

3.37 The EC point that the situation in France until 1994 was the following: in 1997, the IARC classed all forms of asbestos (including chrysotile) as known carcinogens. It was then that the first specific regulatory provisions were adopted in France in order to limit the risk of exposure to asbestos dust. A series of texts produced by the Ministries with responsibility for health, labour, infrastructure and consumer issues followed, restricting the use of asbestos, either by prohibiting the use of the most harmful forms or by prohibiting its use in certain processes (spraying) or in the manufacture of certain products. In the field of health at work, French legislation has, since 1977, strictly regulated working conditions in the asbestos-processing industry, in particular by setting limit values for exposure. These have been lowered several times to comply with the requirements of Community Directives. In addition to having an asbestos-processing industry producing essentially asbestos-cement, France has used asbestos in numerous sectors including construction (flocking and lagging), the manufacture of everyday consumer goods (floor slabs, brake linings, textiles and boarding) and heavy industry (shipbuilding, metal working). Particular emphasis must be placed on the fact that many asbestos-containing products, especially semi-finished products, were accessible to the general public and sold in large retail...
outlets (sheets of asbestos-cement and asbestos board, asbestos yarn, heat-resistant gloves, ironing-board covers, etc.).

3.38 The EC explain that, over the years, a series of regulatory measures reduced the applications for which asbestos might be used. Two types of measure were adopted:

(a) Partial bans on processes or products: (i) in 1977, the Ministry of Housing and Planning and the Ministry of Health and Social Security prohibited asbestos flocking in residential premises; (ii) in 1978, the Ministry of Health prohibited in its turn the use of flocking in all buildings; (iii) in 1988, the Ministry of Economic Affairs restricted to a few products (notably asbestos-cement pipes and gaskets) the use of those forms of asbestos considered at the time to be the most harmful, and made labelling compulsory for all products containing other forms of asbestos; (iv) in 1994, the Ministry of Economic Affairs placed a permanent ban on the forms then considered to be the most harmful and prohibited the use of all other forms of asbestos, including chrysotile, in a wide range of products (toys, smoking accessories, paints and varnishes, filters, etc.);

(b) a regulatory framework for authorized activities: (i) in 1977, the Ministry of Labour laid down strict regulations on working conditions in the asbestos-processing industry; (ii) in 1987, the Ministry of Labour lowered the limit values for exposure; (iii) in 1990, the Ministry of Labour prohibited the recruitment of casual workers to carry out the removal of flocked asbestos or demolition work which would expose them to asbestos dust; (iv) in 1992, the Ministry of Labour again lowered the limit values for exposure and issued a reminder that applying asbestos by spraying was totally prohibited. It also laid down rules on procedures for removing flocked asbestos.

3.39 The EC also indicate that 1990-1995 marked the turning point. Following the adoption of the ILO recommendation in 1986 and from the early 1990s, it became apparent that the risks in the servicing and maintenance sector needed to be better assessed. A data bank (EVALUTIL) was set up which made it possible to assess exposure to asbestos among users of asbestos-containing products. This data bank highlighted the extremely high levels of exposure among certain construction workers (“exposure peaks” occurred during operations such as the cutting of flocked cladding containing 5 per cent chrysotile, the cutting of asbestos-lined fireproof doors, etc).

3.40 The EC note that, given the dangers of inhaling fibres from flocked asbestos used in the construction industry between 1950 and 1977, on 13 December 1989, the French Senior Council for Public Health called on local authorities to compile a register of all buildings, especially public buildings, in which flocked asbestos was present. The feasibility of compiling such a register was assessed by a municipal health and safety department in a pilot town between 1991 and 1994. The study came up against major difficulties linked to the fact that some companies no longer existed and in some cases records had not been kept. In the light of these disappointing results, on 15 September 1994, the Senior Council for Public Health expressed the desire to see legislation introduced on the monitoring of flocked asbestos so as to protect exposed populations. The legislation was to be based on an early warning threshold expressed in terms of the erosion of flocked surfaces and corresponding to a level of atmospheric dust in the building concerned of 5 fibres per litre (“f/l”) (which corresponds to the average asbestos pollution measured in the outside air) and on a 25 f/l threshold above which work would have to be carried out. In the period prior to the implementation of this legislation, the Ministries of Labour and Health sent two circulars to prefects, dated 15 September 1994 and 31 July 1995 respectively, setting out the thresholds, removal and analysis techniques and the procedures for removing flocked asbestos.
3.41 The EC point out that, at the end of 1994, the Ministry of Social Affairs (Directorate for Industrial Relations and Directorate-General for Health) convened a group of experts, whose discussions brought to light certain scientific data, on the one hand, and certain gaps in the existing legislation on the other. The experts’ findings were as follows:

(i) an increase in the number of cases of mesothelioma in France and the worrying forecasts in a study conducted by Julian Peto in the United Kingdom. The study predicted a growth in asbestos-related deaths which would total between 2,700 and 3,000 per year in the 2020s; 
(ii) the serious risk for workers engaged in the servicing or maintenance of asbestos-containing products or equipment. This applies in particular to workers involved in finishing work in the construction industry, such as plumbers or electricians, since their exposure at “peak” moments may be very high; 
(iii) the emergence of cases of mesothelioma at levels of exposure which were below limit values in force at the time under French legislation. On the other hand, it was felt that there was nevertheless a need to collect together all existing scientific knowledge on asbestos so as to shed light on the effect of low doses and on how harmful chrysotile asbestos actually was. In the light of these findings, French legislation on the protection of workers appeared limited. Firstly, the thresholds seemed to be too high and secondly, the provisions designed for industries and removal work where asbestos was known to be present were scarcely or not at all effective when it came to maintenance workers. This conclusion prompted the authorities to take decisive action to reinforce the existing preventive measures and to obtain more detailed information on the risks linked to asbestos.

3.42 The EC emphasize that, on 3 July 1995, the Senior Council for the Prevention of Occupational Hazards adopted the following guidelines:

(i) to improve scientific knowledge of risks (launching of the joint INSERM report); 
(ii) to enhance protection of workers, especially those engaged in servicing and maintenance work; 
(iii) to raise awareness among all those exposed to the dangers of asbestos; 
(iv) to improve compensation for asbestos-related occupational diseases; 
(v) to begin discussions on whether asbestos should be banned. Contrary to what is asserted in the Canadian submission, the possibility of a ban was not ruled out. In July 1995, therefore, the French Government decided to commission INSERM to carry out a detailed study of asbestos-related diseases, based on all internationally available research. In December 1995 it also adopted a general action plan comprising measures to combat asbestos-related hazards, taking account of problems in the areas of public health, the environment, the protection of workers and compensation for asbestos-related occupational diseases.

3.43 The EC point out that the implementation of the 1995 action plan presented by the Minister for Social Affairs made it possible, without waiting for the findings of the INSERM joint report, to introduce general legislation in order to monitor the situation as regards buildings, limit occupational exposure and improve compensation by reorganizing and reviewing the tables of occupational diseases caused by asbestos. This action plan was intended to meet growing concerns about servicing and maintenance workers in the construction sector, a risk on the subject of which Canada remains silent. The action plan comprised the following measures:

(i) owners of buildings were obliged to identify asbestos flocked surfaces and asbestos-containing insulation before 31 December 1999, or before the end of 1996 in the case of priority buildings, i.e. those housing young people or children (Decree No. 96-97 of 7 February 1996); 
(ii) reduction in the limit values for occupational exposure (“VLEP”) to asbestos to the lowest technically possible level (0.1 fibre/cm³); the difference in the limit values for “pure” chrysotile and amphibole forms of asbestos was to be abolished on 1 January 1998, although this would apply only to a very limited number of manufacturing industries (Decree 96-98 of 7 February 1996); 
(iii) strict regulatory provisions on asbestos removal work; works inspectors were entitled to close asbestos removal sites if they did not consider the protective measures to be adequate (legislative provision); a prohibition was placed on hiring casual workers to carry out work involving contact with asbestos; 
(iv) the preparation of specific preventive rules for servicing and maintenance work (Decree No. 96-98 of 7 February 1996); 
(v) the creation of Table 30 bis on the recognition of occupational diseases was created specifically to cover bronchial cancer caused by asbestos; this disease could now be recognized without requiring
the presence of medical “markers”. The necessity of these measures, taken during the first half of 1996, was confirmed by the INSERM report, which was sent to the Minister for Labour and Social Affairs at the end of June 1996. This report, which shed new light on the subject, prompted the French Government to take additional steps without delay, including banning asbestos with effect from 1 January 1997.

3.44 The EC indicate that, in the summer of 1995, contacts between the Canadian and French administrations took place. France was seeking more effective ways of managing the risk, which it already considered very serious, while Canada proposed setting up an international research centre on the prevention of respiratory diseases. Several meetings took place with the Canadian party immediately after publication of the INSERM report and prior to the issue of the Decree on a ban; at that time, Canada asked for a derogation. Several meetings took place between the French and Canadian parties following discussions between the relevant Canadian and French ministers and on the instructions of the Prime Minister, who had received a letter from Mr. Chrétien. These meetings took place between July 1996 (announcement of the decision to impose a ban) and December 1996 (the ban entered into effect).

3.45 The EC stress that the scientific discussions took place in an atmosphere of mutual respect. On 8 October 1996, the Quebec experts met the experts from INSERM and then on 9 October 1996 they met the adviser on asbestos to the French Minister for Social Affairs (Mr. Roigt), together with the competent person from the Industrial Relations Directorate. Moreover, on 29 October 1996, France arranged an entire day’s technical meeting between the Canadian-Quebec delegation and the French experts. During the day, the questions of the harmfulness of asbestos, the question of substitute fibres, the criteria on which France would allow exceptions to the ban, and the “safe” use of asbestos-cement were discussed. On that occasion, the experts from Canada and Quebec acknowledged the high quality of INSERM’s work and showed an interest in discussing a number of scientific points, as is customary among researchers. The EC point out that this oral recognition is in contrast with the criticism of the INSERM report expressed by Canada.

(a) As regards the harmfulness of asbestos, the Quebec delegation reported on the recently published study by Mr. Siemiatycki on women living in close proximity to asbestos mines (the study by Camus et al. had not yet been published when the INSERM report was produced). The French experts from INSERM pointed out that they were aware of the study and had examined it in the context of their joint report. In their view, it showed that the risk of bronchial cancer was practically nil at that level of exposure, but that there was a risk of contracting mesothelioma from the asbestos contained in the dust from the chrysotile mines, at levels of exposure accumulated over a lifetime of a few fibres per cm$^3$ per year. The French experts pointed out that this level was easily reached during the cutting of asbestos-cement and that no country had succeeded in containing this risk.

(b) As regards substitute fibres, the French party felt that further epidemiological data should be obtained, although no victims of the use of substitute products had so far been reported. Decisions were taken “on the basis of the knowledge available”. The Canada-Quebec delegation informed the Ministry of Labour and Social Affairs of its desire to be involved in INSERM’s work on the fibres used as substitutes for asbestos. Since the purpose of the report was to summarize all international studies on the subject, there was no objection to consulting Quebec scientists as part of the research. At the beginning of 1997, however, when INSERM looked for specialists in the field, its bibliographical search did not identify any Canadian experts (whether English or French speaking) with sufficient publications on the subject. In the end, two experts were proposed by Canada’s Health and Safety
Research Institute, and both Dr. Gibbs and J. Siematycki attended the hearing organized by INSERM.

(c) As regards exceptions to the ban, the French administration presented the draft decree and explained the circumstances in which an exemption might be granted: there had to be no substitute products performing an equivalent function to asbestos which were less harmful to workers and which offered equivalent technical guarantees of safety for users.

(d) Lastly, the issue of the “safe” use of asbestos-cement was discussed. The main argument put forward by Canada and by Quebec in favour of extending the list of exemptions was that asbestos can be used in a “safe” way. The French experts from the INRS (National Safety Research Institute) and the Caisse Nationale d’Assurance Maladie (National Sickness Insurance Scheme) presented findings based on measurements which were significantly different from those presented by Canada.

3.46 The EC point out that, for reasons of substance, the request that an exemption be granted for asbestos-cement piping was not admissible. The general ban laid down by the Decree allows for exemptions only when there is no substitute product which can perform an equivalent function and which: (i) presents less of a risk to workers; and (ii) offers all technical guarantees of safety for users. In the case in point, it was very easy to replace asbestos by a less dangerous product. All alternatives to asbestos-cement are less dangerous than chrysotile. In fact, all asbestos-cement can be replaced by products which show no sign of being carcinogenic, whether they are non-fibrous products (ductile iron, plastic) or fibrous products (cellulose, polyvinyl alcohol).

3.47 The EC recall that the ban on asbestos or any material or product containing it has been in force since 1 January 1997. It applies to both industrial and commercial use, i.e. manufacture, processing, possession with a view to selling, offering for sale, import, export or any form of transfer. However, there are a very limited number of strictly monitored exceptions to the general ban. If a temporary and limited exemption is to be granted, there must be no substitute product which could perform an equivalent function and which: (i) on the basis of current scientific knowledge, presents less of a risk to workers; and (ii) provides equivalent guarantees in terms of safety of use (braking performance, for example). Firms which manufacture asbestos-based products or import asbestos must make a declaration to the Ministry justifying the use of asbestos and reporting on the progress made as regards substitution. The administration then registers the declarations which it considers to be in compliance with the legislation. The exceptions are included in a list drawn up by ministerial order.

3.48 The EC indicate that asbestos was used in numerous industrial applications because of its physico-chemical properties: good mechanical resistance (especially to pressure), resistance to high temperatures and to chemical attacks (especially corrosion). Replacing it with a substitute material requires a strict approach applied on the basis of specifications setting out what is required of the substitute product in very specific conditions of use. Once the substitute product has been identified, it must not only be tested, but its reliability over time must also be checked. This can be done by an external body (national or international) on the basis of qualification and approval procedures which, in sectors such as the aeronautical, nuclear or chemical industries, are very long and complex. The sectors most affected by residual use of asbestos are the nuclear, chemical, petrochemical and aeronautical industries.

3.49 The EC emphasize that the Decree provides for an updating procedure designed to eliminate the remaining exceptions as rapidly as possible. The list of categories of exception laid down by ministerial order is therefore reviewed each year by the Senior Council for the Prevention of Occupational Hazards, which shortens the list in the light of technical developments.
(a) The companies registered as a result of the annual declarations made in order to qualify for exemptions from the ban on asbestos are questioned on the progress they have made towards the replacement of asbestos. The findings are compared with the data held by the experts from the INRS46 and the Caisse Nationale d’Assurance Maladie. Experts from co-signatory ministries are consulted (Directorate-General for Competition, Consumer Affairs and Fraud Prevention, Ministries of Industry, Infrastructure, Environment, and Agriculture, and the Directorate-General of Customs). On this basis, proposals for changes are sent for an opinion to the Senior Council for the Prevention of Occupational Hazards, which comprises employer and employee representatives as well as experts.

(b) For example, the following proposals were made at the end of 1997.

(i) As regards brake linings, substitute products have been developed for heavy industrial installations and equipment, ships and special vehicles including heavy land vehicles and vehicles weighing more than 3.5 tonnes, except for certain military vehicles for which the approval procedures are not yet complete. However, it seems that the brake linings used in aircraft braking systems can be replaced only in certain cases. Where asbestos performs the function of braking agent, tests seem to have been conclusive. However, if asbestos is used as a binder between the brake support and the braking component (generally a fibreglass compound or Teflon), no substitute product has yet been found. Likewise, brake linings for compressors and sliding-vane rotary vacuum pumps have to endure extreme pressures (over 300 bars) and temperatures (over 350°C). The substitute products tested to date have not proved satisfactory; often made of graphite, they are deformed by the combined action of pressure and temperature. Finally, it was proposed that only those exceptions be withdrawn which applied to heavy industrial installations and equipment, ships and floating structures and special non-military land vehicles weighing over 3.5 tonnes.

(ii) As regards gaskets, in sectors such as the aeronautical, nuclear and chemical industries or in specific applications, less progress has been made, given the multiplicity and complexity of technical constraints. Substitute products do not offer sufficient guarantees in terms of their resistance over time to corrosive fluids. Accordingly, it was proposed that the exemption, as set out in the relevant paragraph, be maintained in its entirety.

(c) Monitoring is organized as follows: the list of firms which have made a declaration in accordance with the legislation and the list of their customers are forwarded to the other ministerial departments concerned and to the inspection units so as to facilitate checks by customs authorities, by the Directorate-General for Competition, Consumer Affairs and Fraud Prevention, and by the Labour Inspectorate.

4. Circumstances of exposure to asbestos and asbestos-related pathologies

Canada states that, today, science generally recognizes few or no demonstrable effects of environmental asbestos on health, and concludes that, at worst, the risk is undetectable. Asbestos is used in a vast array of products, because of its very useful and often unique characteristics. During extraction and handling of the mineral, manufacturing and...
use of its products, as well as their final disposal, however, a certain number of asbestos fibres are released into the environment. Because the hazards resulting from high levels of exposure to asbestos fibres in certain work environments of the past are known, there is concern over the possible effects of any exposure to concentrations of asbestos. Today, chrysotile fibres are bonded to cement or other particulate or encapsulated materials in a plastic, cement, asphalt, or resin matrix. Brittle products and, to a negligible extent, non-brittle products, however, release a certain number of fibres. Canada considers that, in spite of an ever-increasing understanding of the importance and impact of the sources and characteristics of asbestos fibres on human health, much confusion and many misunderstandings exist regarding the various aspects of asbestos production, such as product manufacturing, the types of fibres used, their presence in the environment, and, especially, the health effects of exposure. Canada contends that, as far as the health effects of exposure to asbestos in the workplace are concerned, it is now recognized that in the past several lung diseases were caused by high levels of exposure to asbestos dust inhaled under various working conditions. In the early 1950s, there were many concerns about the link between exposure to asbestos dust in the workplace and certain diseases (asbestosis, lung cancer, and mesothelioma). In spite of the great contrasts between industrial exposure in the past, which obviously had certain effects on health, and exposure in the general environment today, fears of health risks remain excessive; and yet, the risk is undetectable for the public at large. In order to evaluate the health effects of exposure to asbestos, a considerable number of toxicological tests have been performed on animals. Canada asserts that currently most of the results indicate that only fibres of more than 5 micrometers are pathogenic, and that considerable amounts must be inhaled (or implanted in animals) to cause reactions, including cancer. The length of most fibres in the ambient air, however, is less than 5 micrometers.

According to Canada, it is inevitable that some asbestos fibres are released into the air, water, and soil due to the use of asbestos and asbestos products on a worldwide scale, but concentrations of asbestos in the environment were higher in the past, because of inadequate control measures. In addition to artificial sources, some natural sources (rock alteration) are a significant source of fibres in the ambient air; even water contains natural asbestos fibres. According to the WHO, the total amount of asbestos released by natural sources probably exceeds emissions from industrial sources. A large number of artificial asbestos sources have been studied but, in most cases, there did not seem to be significant fibre emissions. Some of these sources, however, have raised the most fears, for example, brake linings, factory emissions, mines, shipyards, and asbestos in buildings. Although there were some major industrial emissions in the past, a good share of current atmospheric emissions are not respirable, because of the relatively large size of most dust particles and other substances to which the fibres are normally linked. Canada considers that the strong bond between chrysotile and cement in chrysotile-cement products reduces the release of fibres into the air to a minimum. Furthermore, some asbestos products are coated, which makes it very difficult for the fibres to be released. Regarding brake linings, because of the high temperatures caused by friction, up to 99 per cent of the asbestos is physically and chemically transformed into inert and non-hazardous material. Moreover, the fibre remaining after this transformation is less than 5 micrometers in length, and, therefore, has no biological effects. One of the major problems that remain is the elimination of brittle asbestos products that were once used in buildings, especially if they contain amphiboles, as well as the demolition of buildings that may contain significant amounts of these products.

Concerning the concentrations of fibres in the environment, Canada notes that asbestos fibres are present in outdoor as well as indoor air. The average long-term concentration calculated for several years of exposure, taking into account indoor and outdoor air conditions, appears to be between 0.0002 and 0.001 /ml of more than 5 micrometers, with an average concentration of approximately 0.0005 /ml of air. Some of this asbestos comes from natural erosion. In urban areas, however, atmospheric asbestos comes mainly from the extensive use of this product. It seems that ambient air concentrations are currently lower than those observed a few years ago. Regarding exposure to asbestos from drinking
water, whether or not it is routed through asbestos-cement pipes, drinking water contains 200,000 to 2,000,000 fibres per litre. Canada considers that, in order to evaluate the health effects of asbestos in the environment, it is important to determine the consequences of human exposure, as well as possible inhalation or ingestion of these fibres. There is a lot of confusion regarding the inhalation of all types of particles that penetrate the airways. In Canada’s view, it must be recognized that only extremely fine particles can penetrate the deep areas of the lungs. Many inhaled particles are immediately exhaled, because they do not succeed in depositing themselves. Large particles that penetrate through the upper parts of the airways are handled by an extremely effective biological elimination system; the mucociliary mechanism traps the fibres and pushes them out of the airways. Canada considers that the health effects of ingested asbestos are practically nil. The results of animal studies on ingestion by feeding have been absolutely negative, as have the vast majority of human epidemiological studies conducted in areas where asbestos concentrations in water are naturally high due to the presence of asbestos in the rock over which the water flows.

3.53 Canada notes that the WHO clearly reaffirmed, particularly in a 1989 follow-up, that the use of chrysotile-cement pipes is completely safe for transporting drinking water. It is essential to distinguish clearly between the obvious health effects associated with very high exposure to asbestos in the workplace and the absence of demonstrable health effects linked to current environmental asbestos concentrations. Therefore, the WHO considers the risk of mesothelioma and lung cancer attributable to mass exposure to asbestos is so low as to be undetectable. According to Canada, long-term exposure at rates of around 0.0005 fibres of more than 5 micrometers per ml of air corresponds to a possible cancer risk equal, at the most, to the extremely low risks of “rare events,” for example, the risk of being killed by lightning or getting cancer from consuming meat grilled over charcoal, or by an increase in the exposure to cosmic rays caused by one transatlantic airline flight each year. Consequently, the risk of cancer due to the exposure to an environmental concentration of 0.0005 f/ml of more than 5 micrometers would be only about one in 100,000 (estimated lifetime risk). In other words, it is a rate that does not justify any additional measures. According to the French Académie nationale de médecine, “this figure, which equals zero based on the degree of biological uncertainty […] indicates that there is no demonstrable risk.” These estimated environmental risk values are based on workplace health data and on the use of a linear lung cancer model, as well as on an exponential model for mesothelioma. The estimated values are conservative for various reasons, particularly because of the fact that a large number of previous workplace exposure values underestimated the actual exposure conditions for different types and mixtures of asbestos, and because the model used assumed they would apply even to extremely low exposure values, which means that they do not consider the possibility of a “threshold” effect under which there would be no detectable effect. In reality, Canada concludes that the risk could be even lower because, now, the public is exposed mainly to chrysotile, which is less hazardous than amphiboles.

3.54 Canada points out that environmental risks have to be put into perspective. Generally, it is acknowledged that, of the risks to which our society is exposed, some are less serious than others. Often, however, many people have a poor appreciation of the relative value of these risks, even though they have been published for several years. The following table provides an overview of the risks of exposure to environmental asbestos concentrations in comparison with other risks. It shows that the estimated risk attributed to this exposure is equal to or lower than that of very rare events.
Canada emphasizes that, because of the extremely low risks linked to the current use of asbestos products, there is no reason to have to reduce the use of chrysotile or tighten restrictive measures if adequate restrictive measures have been taken. Such was the case in France in 1997, according to the French Senate and National Assembly. Despite the importance of applying general restrictive measures, there are currently three specific cases that require particular vigilance: (i) the presence and elimination of old asbestos products in buildings; (ii) the demolition of buildings containing significant amounts of asbestos; and (iii) the elimination of asbestos waste. In addition, particular attention must be paid to the hazards caused by amphiboles. Because of the relatively high risks, compared to those of chrysotile, which are linked to exposure to various types of amphiboles in the workplace, most countries, including Canada and France, have prohibited their use. Special restrictive measures were adopted for extraction and crushing, as these activities can produce large amounts of dust. Appropriate regulation of modern crushing procedures allows asbestos to be exploited with minimal emission of fibres into the environment. For this purpose, for example, effective air purification systems such as electrostatic precipitators, cyclones, or sleeve chambers are used. The uncontrolled use of certain types of asbestos and of certain work processes was responsible in the past for unacceptable emissions. Today, with the help of advanced technologies and the widespread use of wet processing, it is relatively easy to use chrysotile in such a way as to ensure the protection of workers’ health while avoiding possible air contamination.

EVALUATION OF LIFETIME RISKS (SELECTED CASES) *(according to data mainly from the United States)*

<table>
<thead>
<tr>
<th>Lifetime risk per 100,000 inhabitants</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Extremely high risk</strong></td>
</tr>
<tr>
<td>Smoking (all causes of mortality)</td>
</tr>
<tr>
<td>Smoking (cancer only)</td>
</tr>
<tr>
<td><strong>Very high risk</strong></td>
</tr>
<tr>
<td>Driving an automobile</td>
</tr>
<tr>
<td><strong>High risk</strong></td>
</tr>
<tr>
<td>Frequent air travel (deaths)</td>
</tr>
<tr>
<td>Cirrhosis of the liver, moderate alcohol consumption (deaths)</td>
</tr>
<tr>
<td>Pedestrians hit by automobiles (deaths)</td>
</tr>
<tr>
<td><strong>Moderate risk</strong></td>
</tr>
<tr>
<td>Low alcohol consumption (one beer per day) (cancer)</td>
</tr>
<tr>
<td>Death by drowning (all recreational activities)</td>
</tr>
<tr>
<td>Atmospheric pollution, U.S., benzopyrene (cancer)</td>
</tr>
<tr>
<td>Natural radiation at sea level (cancer)</td>
</tr>
<tr>
<td>Cosmic rays, frequent airline travel (cancer)</td>
</tr>
<tr>
<td><strong>Low risk</strong></td>
</tr>
<tr>
<td>Household accidents (deaths)</td>
</tr>
<tr>
<td>Secondhand smoke (cancer)</td>
</tr>
<tr>
<td>Diagnostic X-rays (cancer)</td>
</tr>
<tr>
<td>(Rate of risk which few people are willing to use their own resources to reduce, Royal Society, London.)</td>
</tr>
<tr>
<td><strong>Very low risk</strong></td>
</tr>
<tr>
<td>Natural radiation, people living in brick houses (cancer)</td>
</tr>
<tr>
<td>Smallpox vaccinations, by vaccination (deaths)</td>
</tr>
<tr>
<td>One transcontinental flight per year (deaths)</td>
</tr>
<tr>
<td>Saccharine, average consumption in the U.S. (cancer)</td>
</tr>
<tr>
<td>Consumption of Miami or New Orleans water (cancer)</td>
</tr>
<tr>
<td>(Rate of risk at which few people feel measures are necessary, in the absence of clear links to consumer products, Royal Society, London.)</td>
</tr>
<tr>
<td><strong>Extremely low risk</strong> (<em>rare events</em>)</td>
</tr>
<tr>
<td>One transcontinental flight per year, natural radiation (cancer)</td>
</tr>
<tr>
<td>Lightning (deaths)</td>
</tr>
<tr>
<td>Hurricane (deaths)</td>
</tr>
<tr>
<td>Consumption of one charcoal cooked steak per week (cancer)</td>
</tr>
<tr>
<td><strong>RISK DUE TO ENVIRONMENTAL ASBESTOS CONCENTRATIONS</strong> <em>(cancer)</em></td>
</tr>
<tr>
<td>(approximately 1 per 100 000 or less)*</td>
</tr>
<tr>
<td>(Acceptable risk, WHO, drinking water (cancer))</td>
</tr>
<tr>
<td>(Additional restrictive measures are certainly not justified, Royal Society, London, United Kingdom)</td>
</tr>
</tbody>
</table>
3.56 Regarding the limitation of asbestos emissions during construction activities, Canada notes that the use of modern chrysotile products prefabricated in the factory for the construction industry reduces the problems of environmental contamination to a minimum. In these chrysotile-cement construction materials, the asbestos fibres are strongly linked to a matrix. If it should, nonetheless, become necessary to cut them on the construction site, the use of tools that almost entirely eliminate emissions (low-speed saws, with water injection or equipped with suction units), and the wearing of a mask by the operator guarantee their safety. It is easy to obtain a copy of the codes of practice that employees must follow when handling modern chrysotile products.

3.57 Canada considers that, in general, the extremely low risks of exposure to asbestos in buildings have been accurately demonstrated. In some cases, however, the materials enclosing the sprayed asbestos in the buildings can deteriorate and pose a certain hazard to their occupants. In France, as in Canada and everywhere else, regulations provide that, beyond a certain concentration of airborne asbestos in buildings, corrective measures such as isolation, encapsulation, or even elimination must be considered. If a large amount of asbestos is present in an old building to be demolished, especially if amphiboles are involved, it may be advisable to eliminate these materials in advance, if possible. The procedures to be followed in these projects are well documented. Canada also considers that the elimination of most modern asbestos products should present only a slight difficulty if a building has to be demolished. It is important to use effective measures for handling, packaging, and transport, as well as for waste disposal. The procedures to be followed for these operations are well documented and often standardized. These measures, which are normally easy to implement, allow asbestos waste in the environment to be kept to a satisfactory minimum. Industrial waste containing asbestos is normally easy to handle and confine, because it is often wet and, therefore, does not produce powder. Furthermore, many industries today recycle their waste because this measure, in addition to being more economical, allows the waste to be disposed of effectively. In most countries, industries must comply with regulations limiting emissions of asbestos into the atmosphere.

3.58 To conclude, Canada contends that, in light of the available data, there is no scientific justification for the prohibition or reduction of the manufacturing or use of modern asbestos products. This conclusion is as valid for developed countries as it is for developing countries. It is enough to continue to take adequate restrictive measures, particularly for mineral extraction and crushing, the manufacturing and use of products, and the elimination of asbestos waste.

3.59 The European Communities respond that Canada presents the health effects of asbestos by distinguishing on the one hand the environmental risks and, on the other, certain asbestos-related occupations: mining and crushing, manufacturing of products, building and demolition of buildings, elimination of waste. Canada simply disregards the fact that once asbestos has been used in the construction of a building and until it is demolished, any installation containing asbestos is likely, during its life cycle, to require some form of plumbing, heating or electrical work carried out by professionals or amateurs. Regular servicing and maintenance work carried out by people who are unaware of whether the material on which they are working contains asbestos or not may, if asbestos is present, lead to extremely high peaks of exposure, significantly higher than the exposure limits, and which have been proven to be carcinogenic in a perfectly “detectable” way. According to the EC, the large majority of cancers due to asbestos are the result of such work (servicing and maintenance) on materials containing chrysotile. For many years, scientific bodies and governmental authorities disregarded this type of risk. In the early 1990s, scientists started to note an increase in cases of mesothelioma affecting professionals who had never worked in the asbestos-manufacturing industry. France, following the example of several other countries, became aware of the seriousness of the risk faced by these workers and individuals, exacerbated by the number of asbestos products freely on sale and frequently used. In such situations, controlled use practices, which the EC consider do not in any case eliminate the risk, are inapplicable. Persons liable to be exposed to the risk are unaware that they are working with products containing asbestos. They are therefore unable to protect themselves. Even if they did know, controlled use practices imply the use of
costly material and equipment and make even the simplest and quickest operations very complicated. Taking these elements into account, France decided, by means of the regulation contested by Canada, to halt the dissemination of such risks by banning all future uses of products containing asbestos.

3.60 The EC point out that Canada asserts that “today science generally recognizes few or no demonstrable effects of environmental asbestos on health”. If the idea of ambient air is limited to the usual levels of exposure in towns or buildings that have not deteriorated, the INSERM report reaches the same conclusion unambiguously in several places. But the French regulation is not solely intended to prevent the risk due to ambient exposure, i.e. exposure caused by living in a building containing asbestos. The French regulation calls for caution in this area, but the purpose of the contested ban is to prevent the extension of risks due to exposure, usually only of an occasional nature, for professional activities (servicing and maintenance in buildings/public works, for example) or leisure activities (handymen). The EC state that these risks have been scientifically proven both by metrological data concerning the concentration of asbestos fibres and by international epidemiological data.

3.61 The EC consider that the Canadian assertion that “Today chrysotile fibers are bonded” or “encapsulated” in other materials and therefore do not present any risk is misleading because health problems are not simply due to the presence of chrysotile fibres in materials, but to the fact that very high levels of fibres are released into the atmosphere when working with these materials, for example, when sawing, cutting, etc. This idea is often repeated in Canada’s arguments in order to make it appear that asbestos incorporated in various materials, including asbestos-cement, is not dangerous: the purpose of this assertion is to provide misleading reassurance, and it will not be systematically contested. Moreover, it should be noted that the manufacture of asbestos-cement has always required that the asbestos and other materials be bonded. In this connection, it cannot be stated, as Canada does, that there are “modern” manufacturing methods. To the knowledge of the EC, the production methods for asbestos-cement have not evolved to any significant degree for many years. It is also wrong to claim that “The strong bond between chrysotile and cement in chrysotile-cement products reduces the release of fibers into the air to a minimum”. This claim is incorrect when the product in chrysotile-cement is the subject of servicing or maintenance which requires drilling or cutting. In such cases, the release of dust into the air is several hundred times higher than the value limit allowed in France, and the United States, or even the limit recommended by the WHO and cited by Canada. The EC note that the risks of “asbestosis” have been known since the beginning of the century and the first regulations adopted to protect the health of workers from the dangers of asbestos were drawn up in England in 1931, as is recalled in the European submission. The risks of asbestos for health have thus been recognized for a very long time.

3.62 The EC point out that Canada cites very low risks for health at very low exposures (0.0005 f/ml). It should be emphasized, however, that: (i) these are the same values as those cited in the INSERM report for such levels of exposure, which Canada appears to disregard; (ii) these values were obtained using the same model as that used by INSERM, whose validity Canada nonetheless denies. If the model can be used by Canada to show that the risks are low at low levels of exposure, why would it not be valid for higher levels of exposure? The EC note that Canada does not provide any justification for this contradiction and adds that this model does not take into account “a threshold below which there is no effect”: this ignores the scientific consensus that there is no such threshold. The WHO report of 1988 on the dangers of chrysotile (cited by the EC but never by Canada, which only refers to the 1986 WHO report) recognizes that there is no threshold for chrysotile. The EC note that Canada acknowledges that it has banned the use of amphibole asbestos, like many other countries. Taking into account the equally dangerous threat of lung cancer from both chrysotile asbestos fibres and amphibole asbestos fibres and the theoretical possibility of so-called “safe” use, the Canadian ban on amphiboles alone appears inconsistent, unless it is justified by the fact that Canada does not produce amphibole fibres. It is difficult to understand why it should not be possible to ensure “safe” use of
fibres of the amphibole variety when Canada claims that this is possible for fibres of the chrysotile variety: the production methods and the conditions of use are nonetheless strictly identical in both cases.

3.63 According to the EC, Canada’s assertion that “It is relatively easy to use chrysotile in such a way as to ensure the protection of workers’ health” is a euphemism that hides the real problems faced in practice. On the one hand, such protection can only exist if the worker knows he is handling asbestos products. On the other, following the ISO 7337 standard recommendations means cumbersome and costly individual measures (practically turning oneself into a cosmonaut) without being certain that no fibre can get through the protection. Furthermore, the asbestos-cement market is very diffuse. It is utopian to imagine that all the public concerned could apply the Canadian safe use programme. France recognizes that this is the only way of limiting as far as possible the risks of such operations when the asbestos is already there and it has adopted very strict regulations to this effect. The EC nevertheless contest the fact that Canada seeks to impose the adoption of such measures on very large parts of the economic sectors. In addition to their lack of effectiveness, these measures have a significant impact on the costs of French companies and are unknown to diffuse users of asbestos materials (on building sites, for example): asbestos-cement products were marketed in France as materials that did not present any particular risk, as Canada intimates when it refers to “modern asbestos-cement materials” in which asbestos fibres are “encapsulated!” The EC point out that, as shown by the occupational disease statistics, it is equally impossible to ensure the effective protection of workers in the raw asbestos processing industry, even with sophisticated dust collection and ventilation systems.

3.64 The EC consider that the Canadian claim that “the elimination of most modern asbestos products should present only a slight difficulty if a building has to be demolished” is a euphemism. Asbestos products used in the building industry are fragile and inevitably break or dissolve, releasing fibres into the atmosphere. In addition, the need to remove asbestos before demolition involves huge expenditure because any removal of asbestos requires the observance of draconian technical regulations essential for the protection of the workers. Lastly, the EC reject the Canadian argument that “in light of the available data, there is no scientific justification for the prohibition or reduction of the manufacture and/or use of modern asbestos products. Chrysotile asbestos, like other types of asbestos, is dangerous for the health of workers and the population, no safe use allows the risk to be reduced. Scientifically, the French measure on a general ban is justified.

3.65 The EC state that, as the risks from exposure to asbestos were gradually identified and became the subject of preventive rules and regulations, the various types of exposure were listed and were placed in the following three categories: the first category includes occupational exposure of workers, including “primary” users and “secondary” users. “Primary” users are those working in the asbestos industry: mining of asbestos, manufacture of asbestos products (asbestos fibres, asbestos-cement, gaskets, asbestos boards for soundproofing or thermal insulation; since 1965, there has been no asbestos mining in France and less than 1,500 people were working in the asbestos-processing industry at the time the ban came into effect. “Secondary” users are in enterprises which use asbestos products (building industry, metalworking, shipbuilding, etc.): enterprises which, in the course of their work on buildings or plants, encounter materials containing asbestos (servicing and maintenance, electrical work, plumbing, etc.); enterprises engaged in the removal and/or containment of asbestos, particularly flocked asbestos; in France, the use of asbestos products involved several hundred thousand persons at the time of the ban and, at the present time, servicing, maintenance and do-it-yourself using asbestos products affect several million people; removal and containment of asbestos involves several hundred persons. The EC note that the Canadian guide on “controlled” use covers only the mining of asbestos and the processing of raw asbestos into asbestos products, i.e. it applies only to “primary” users. According to the EC, the second category includes para-occupational and domestic exposure, where people come into contact with workers in the first category. Many members of the general population encounter similar exposure to servicing and maintenance workers when engaged in do-it-yourself activities. Such individuals even
undertake the removal of asbestos on their own. They are therefore exposed without being aware of it, for asbestos is often undetectable, as the Canadian submission points out. Environmental exposures are in the third category and three types of source can be distinguished: (i) pollution from a natural source of geological origin; (ii) pollution from a particular “industrial” source; (iii) pollution from asbestos in buildings and various types of equipment. This breakdown reflects that generally found in other industrial countries, except for those countries where asbestos is mined, such as Quebec, where 1,300 persons fall into the category of “primary” users.

3.66 The EC recall that three diseases are caused by asbestos. Firstly, mesothelioma is a pleural cancer for which the only known cause is the inhalation of asbestos. Asbestos in all its forms (amphiboles or chrysotile) is the only known agent which can cause mesothelioma or cancer of the pleura. In this disease, liquid forms between the lung and the thoracic cavity (pleurisy) causing pain and breathlessness. Asbestos of the amphibole type and chrysotile asbestos both cause mesothelioma, although amphiboles would seem to be somewhat more carcinogenic (but only in the case of mesothelioma). This cancer occurs mostly as a result of occasional low-intensity exposure. After a long latency period (lasting for 30 years on average), this cancer enters the terminal phase, which lasts for one year on average. There is as yet no curative treatment with any effect. In 1996 alone, an estimated 750 persons died of mesothelioma in France. Secondly, cancer of the lung may be due to the inhalation of asbestos, but unlike mesothelioma there are other known triggers. The effects of lung cancer are quicker to make themselves felt than those of mesothelioma and both the symptoms and the outcome of the disease are widely known. The first symptoms include blood in the sputum, breathlessness and a general deterioration in the patient’s health. Only certain forms of cancer can be treated, depending on their location and histological nature. Both chrysotile asbestos and the amphibole varieties can cause this form of cancer, their carcinogenic impact being comparable. Asbestosis is a form of pulmonary fibrosis (or pneumoconiosis) arising from an accumulation of asbestos fibres, just as silicosis results from the inhalation of silica. The fibrosis takes the form of a scarring process which follows from the inflammatory reactions caused by the presence of asbestos fibres (or silica dust) in the pulmonary alveoli. The fibrous thickening of the thin alveolar wall prevents the circulation of oxygen. Once exposure has ceased, the disease either stabilizes or becomes progressively worse, with respiratory difficulties which may end in the patient’s death. No curative treatment is yet available. Asbestosis results from what is considered a high level of occupational exposure. It develops over an average period of seven to eight years. Currently there are some 150 cases of asbestosis per year in France; in the United States there were on average 733 cases per year between 1982 and 1993.

3.67 The EC explain that, in all, asbestos is responsible for some 2,000 deaths per year in France, including 750 deaths from mesothelioma. These figures are based on reliable and verified data. In the INSERM report (pages 172 to 180) a detailed description is given of the data in question, which are drawn from two major French studies and from cancer records. The figures have been fully borne out by the surveys conducted by the Programme National de Surveillance du Mésothéliome (PNSM) (National Mesothelioma Surveillance Programme), which was set up under the French Health Monitoring Authority in 1998. In France, asbestos-related occupational diseases account for almost 40 per cent of all social security expenditure in respect of occupational diseases, i.e. over F 1 billion per year.

3.68 The EC point out that one important point in this connection is that mesothelioma is regarded as a very reliable indicator of exposure to asbestos. Researchers plot the course of the disease against the use of asbestos since this technique yields important data for assessing the risk involved in various situations where people are exposed to asbestos. Unlike lung cancer, which is the other cancer caused by asbestos, mesothelioma has certain special characteristics: (i) apart from exposure to asbestos, no other causal factor present in the industrialized countries has been established or even seriously suspected. Interpreta-
in both men and women; whenever levels are found in excess of this frequency, they are regarded as a sure indicator of the presence of asbestos. It is for these reasons that mesothelioma is studied to ascertain the harmful effects of asbestos. But this approach, which is justified for methodological reasons, should not lead one to overlook the fact that most asbestos-related cancer deaths are attributable to lung cancer. According to the EC, it is generally recognized\(^7^\) that to obtain the total number of cancer deaths attributable to asbestos, the number of mesothelioma deaths has to be doubled at the very least.

3.69 The EC recall four important dates which mark the progress of international awareness of the health risks. In 1977, the WHO, acting through the IARC, recognized that all varieties of asbestos, including chrysotile, were carcinogenic, causing both lung cancer and mesothelioma, and classified them in group I (known carcinogens). In 1986, the ILO recommended in Convention No. 162 that legislators should make provision for the replacement of asbestos by less harmful materials or technologies, at the earliest opportunity. In 1996, the WHO recommended that asbestos, including chrysotile, should be replaced by harmless substitutes wherever possible. Lastly, in 1998, the WHO again drew attention to the carcinogenic effects of chrysotile, particularly as a cause of mesothelioma; it reiterated its call for the replacement of chrysotile and pointed out that numerous categories of worker were at risk in many areas.

3.70 The EC indicate that the diseases described above and caused by chrysotile asbestos (mesothelioma, lung cancer, asbestosis) are incurable. As it has been recognized for decades that all types of asbestos are life-threatening, measures to protect workers have been taken, and have been in place for a very long time in many countries. Since regulatory measures were first put in place in the United Kingdom in 1931, the ceilings set in many countries for exposure at work have been gradually lowered. The use of certain forms of asbestos has been banned in certain countries and that of any form of asbestos in others. In 1977, following a great many international scientific studies, the WHO listed asbestos in all its forms as a substance known to be carcinogenic to man.\(^7^8\) Chrysotile asbestos and amphibole asbestos, which cause identical fatal diseases, cannot be treated separately. According to the EC, the diseases identified must be dealt with in a consistent fashion. These two varieties of asbestos cause the same cancers and the exposure conditions are identical. The EC note that, logically, just as it recommends the “safe” use of chrysotile asbestos, Canada could equally well have promoted the “safe” use of amphibole asbestos rather than approving and encouraging its prohibition. It is difficult to understand how Canada can accept the grounds for government measures banning amphibole asbestos while denying their legitimacy in the case of chrysotile.

3.71 The EC point out that asbestos is the cause of an international public health problem which affects both France and Canada. Mortality due to mesothelioma has been increasing in men in industrialized countries by some 5 to 10 per cent per annum since the 1950s.\(^7^9\) A similar increase has also been seen in France.\(^8^0\) A recent study coordinated by J. Peto\(^8^1\) found that in seven European countries nearly 10,000 people had died from mesothelioma during the period 1990-1994. The most recent forecasts on mesothelioma mortality show that over the next few decades there will inevitably be hundreds of thousands of deaths caused by exposure to asbestos dust. Peto’s 1999 study, mentioned above, also looks at the evolution of pleural mesothelioma mortality in six Western European countries (Great Britain, France, Germany, Italy, the Netherlands, Switzerland) over the period 1995-2029. This study estimates the total number of deaths due solely to mesothelioma in these six countries at around 200,000. According to the EC, the number of deaths will at least double between the periods 1990-1994 and 2015-2019, when it will reach 6,700 deaths per annum. If these figures are extended to all the countries of Western Europe and deaths from lung cancer are added, by 2029 around 500,000 deaths from cancer will have been caused by exposure to asbestos.

3.72 The EC indicate that, in France, the number of mesothelioma deaths in 1996 is estimated at 750. Contrary to what Canada states,\(^8^2\) this figure of 750 estimated by INSERM rests on solid data. The data come from two major French studies based on actual cases of mesothelioma since 1979 in the geographical areas covered by the cancer registers in France.
The frequency of mesothelioma is steadily increasing; all the French registers listing cases of mesothelioma show that the increase observed is 25 per cent every three years. According to the EC, these data are fully confirmed by the first available data provided by the Programme National de Surveillance du Mésothéliome (PNSM), which was set up under the authority of the Réseau National de Santé Publique in 1998.83 A recently published study,84 similar to that by Peto et al., predicts that the annual number of deaths from mesothelioma in France will continue increasing until 2020, in all the hypotheses studied. By that date, there could be 1,040 annual deaths from mesothelioma among men; the total number of deaths from mesothelioma over the whole period 1996-2020 could be 20,000 deaths among men and 2,900 deaths among women. INSERM has estimated that around 1,200 deaths from lung cancer were caused by asbestos in France in 1996.85

3.73 The EC note that, in Canada, the promoter of the so-called “safe” use of chrysotile, the health situation is no better than in other countries. It is regrettable that Canada has never provided its own health statistics on certified deaths from mesothelioma, in spite of the repeated requests made during the WTO consultations in Geneva on 8 July 1998. Unlike most other industrial countries, which are concerned about the deleterious effects of asbestos on the health of their population, Canada has never, as far as the EC are aware, published any data on the incidence of mesothelioma over a long period among its own population, although such data are available. The table below thus had to be reconstituted by French experts from the raw world data published regularly by the IARC.86

### ANNUAL INCIDENCE OF MESOTHELIOMA IN CANADA AND QueBEC

<table>
<thead>
<tr>
<th>PERIOD</th>
<th>1978-92</th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Canada</td>
<td>Quebec</td>
<td></td>
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<td>M</td>
<td>F</td>
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<tr>
<td>1978-82</td>
<td>6</td>
<td>2</td>
<td>9</td>
<td>4</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1983-87</td>
<td>9</td>
<td>2</td>
<td>11</td>
<td>3</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1988-92</td>
<td>11</td>
<td>2</td>
<td>15</td>
<td>3</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

This table shows that in Canada, each year between 1978 and 1982, six men per million were affected by mesothelioma. Ten years later, more than 11 men per million per year were affected; in other words, the figure doubled in less than 15 years, which is a particularly high rate of growth. According to the EC, the situation in Quebec, the province which produces Canada’s asbestos, is even worse: the increase in the number of cases was of the same order, but the rates are consistently higher than in Canada as a whole (about 35 to 50 per cent higher, depending on the period). The fact that the rates for women have remained more or less stable (though markedly higher in Quebec) shows indisputably that the cause of these mesotheliomas is essentially work-related. Furthermore, the fact that the increase in the frequency of mesothelioma-type cancers can be seen throughout Canada, according to the EC, shows that the risk of death from chrysotile is not confined to the asbestos mining industry (which exists only in the province of Quebec), but that it affects all sectors of the economy. The EC also emphasize that Canada, which produces only chrysotile asbestos, has mainly used this type of asbestos.

3.74 The EC note that the number of mesotheliomas, and hence the number of asbestos-related cancers, correlates with the amount of asbestos imported. The more asbestos that is imported into a country, the more deaths there are from cancer caused by asbestos. Analysis of the data for ten Western countries87 shows a very clear and strong correlation between cases of mesothelioma and consumption of asbestos per inhabitant, measured by the amount of imports. A study was conducted where the rates of cancer in the ten countries were compared with the total amount of asbestos imported per inhabitant (the study analyses the statistical correlation between these two values). This correlation is extremely strong (the very revealing correlation coefficient is 0.70). According to the EC, the number of cases of cancer thus increases proportionally with the increase of imports of asbestos into each country. It is important at this stage to bear in mind that about 95 per cent of all the asbestos used in the world is chrysotile.
The EC stress that chrysotile is a known carcinogen causing lung cancer and mesothelioma. The fact that chrysotile is carcinogenic has long been internationally recognized. Since 1977, the IARC has classified both chrysotile and amphiboles as carcinogens endangering human health (i.e. category I). The IARC has recognized that chrysotile causes lung cancer and mesothelioma in numerous epidemiological studies of exposed workers. The WHO reaffirmed that chrysotile asbestos “has been associated with an increased risk of pneumoconiosis, lung cancer and mesothelioma in numerous epidemiological studies of exposed workers”. The WHO thus confirms the conclusions reached by the 1996 INSERM report. Still more recently, INSERM’s conclusions were again confirmed by the United Kingdom’s Health and Safety Commission (HSC), which considers all forms of asbestos dangerous as they can all cause mesothelioma, lung cancer and asbestosis. The EC note that Canada never cites these WHO and HSC reports, but on the other hand it does cite the recent publication by Camus et al. on cancer mortality among women living in the vicinity of Quebec’s chrysotile asbestos mines. This study does not reveal any risk of lung cancer among these women. Canada fails to mention, however, that other studies, carried out among workers in the Quebec chrysotile mines, had already shown that the dose observed, as compared with the effect noted, was the lowest of all the estimates published in the international scientific literature on the risk of lung cancer. The number of cancers observed among workers in the Quebec asbestos mines is in fact much smaller than that found in all other situations linked to asbestos. This was pointed out in the INSERM report.

The EC point out that the 1998 WHO report also explicitly noted that: “The exposure-response relationship between chrysotile and lung cancer risks appears to be 10-30 times higher in studies of textile workers than in studies of workers in mining and milling industries.” It should be noted that these risks were observed in an American textile factory which had imported chrysotile asbestos from the Quebec mines. According to the EC, there is an international consensus that there is no threshold of harmlessness for chrysotile. It is important to make the distinction between setting exposure ceilings and the existence of a threshold of harmlessness beneath which there would be no health risk. These are two different problems which do not have the same rationale. The exposure ceilings are set by the authorities, taking various factors into account, such as the technical possibility of obtaining sufficiently low levels of exposure in the environment or the technical means of measuring the actual levels in the air for control purposes. The concept of the threshold of harmlessness is a biological and medical one: the aim is to establish whether biological effects are caused by chrysotile below a given level of exposure. There is thus no contradiction between the fact that it is scientifically accepted that there is no biological threshold of harmlessness and the fact that, nonetheless, an exposure limit is set at a certain level that can be achieved and easily measured for control purposes. The 1998 WHO report carried out under the International Programme on Chemical Safety states that, for chrysotile: “No threshold has been identified for carcinogenic risks”. This international authority thus confirms the conclusions of the INSERM report on this point.

The EC point out that most malignant tumours have their origin in the transformation of a single cell, and a very low dose of a carcinogen is quite capable of inducing a transformational mutation in the genetic make-up of a cell. In epidemiology and the biological sciences it is now agreed that the most plausible model for carcinogenicity is one with no threshold value. According to the EC, the results of the most recent studies confirm this fact, while virtually all the references cited by Canada are from the 1980s. Thus, the French study by Iwatsubo et al. shows that asbestos has carcinogenic effects at levels well below those specified in earlier publications. Other recent studies analysed in the INSERM report show similar results (see pages 122 and 123 of the report).

In the EC’s view, the model which considers that the risk of cancer is directly proportional to the dose of asbestos inhaled is that which is usually adopted. The model of simple proportionality between the dose of asbestos and the risk of cancer is that which is generally accepted nowadays. In the case of exposure to asbestos, such a model, which sees the risk of cancer as directly proportional to the dose of asbestos inhaled, fits very
satisfactorily with direct epidemiological observations of high exposure (above 1 fibre/ml). As applied to low doses, this model is generally considered to be the most scientifically plausible. The EC point out that this model, which is strongly criticized by Canada, is that used by all the official committees of experts to date; it is also used in the HEI (Health Effects Institute) report so comprehensively cited by Canada. The EC stress that the values cited by Canada (0.0002 fibres/ml) in no way correspond to the cases covered by the French Decree challenged by Canada. Such excessively low values are found in the ambient air of towns and buildings: at this infinitesimal level, the risk is clearly undetectable, as the INSERM report states in several places (pages 145 and 146, and 224-230). The ban on asbestos, in France and in other countries, is not intended to eliminate the approximately 0.0002 fibres/ml which exist “naturally” in the air. The ban is intended simply to protect all the workers and users of asbestos who are often exposed to much higher values, which may reach tens of fibres per millilitre (i.e. several thousand times more) in the case of standard procedures for dealing with materials containing asbestos-cement. Chrysotile represents the vast majority of the asbestos used throughout the world. About 95 per cent of all the asbestos used since the end of the Second World War has been of the chrysotile variety and more than 80 per cent of this chrysotile is incorporated into asbestos-cement products. This explains, in particular, the very high incidence of mesothelioma among building workers. It is therefore quite indisputable, in the EC’s view, that it is indeed the chrysotile variety of asbestos which causes the vast majority of asbestos-related cancers. This has also been shown in recent publications.

3.79 The EC point out that chrysotile and asbestos-cement pose a threat to an extremely large section of the population. Production workers are not the only section of the population affected. The EC note that, when discussing exposure to asbestos, Canada refers only to workers in the production and processing of asbestos (“primary users”). However, such workers are relatively few in number compared with the huge group of “secondary users”. It is because these secondary users are very numerous and run significant risks that there is a very rapid rise worldwide in the mortality rates for cancers caused by asbestos. As the British Health and Safety Commission (HSC) points out, the risks for secondary users of asbestos have, in the past, been underestimated.

3.80 The EC note that the occupations and branches of industry affected have changed and diversified. Several studies show the scale of exposure to asbestos in all countries. They also show the huge variety of occupations and sectors that are affected. In the 1960s, the main occupations affected were those in the insulation industry, in the production and processing of asbestos, in the central heating industry and in the shipyards. In the 1980s and 1990s, in contrast, the highest incidence was found among those whose work involved materials containing asbestos. It was only in the 1990s that studies were published on mortality by type of occupational exposure to asbestos, raising awareness of the large-scale health problem which such exposure posed for many categories of worker. In Great Britain, the study published by Peto et al. in 1995 dealt with mesothelioma mortality in England and Wales for the years 1979-1980 and 1985-1990. The table below (taken from this study) shows the principal occupations in which deaths from mesothelioma have been observed; the percentages represent the distribution by occupation of all the deaths from mesothelioma that have occurred in England and Wales. It shows the variety and the relative importance of the various occupations exposed to asbestos to a significant degree. The types of work are classified in descending order of frequency of death from mesothelioma. The occupations shown in this table account for some 50 per cent of all the deaths from mesothelioma which occurred during the period studied.
3.81 The EC point out that the figures in the table are not estimates but death rates which have actually been recorded. The occupations at high risk of mesothelioma include welders, dockers, laboratory technicians, painters and decorators, plasterers, fitters, upholsterers, power station workers, etc. Thus, the building sector alone accounts for a quarter of all the deaths from mesothelioma which have occurred in England and Wales, a proportion considered by Peto et al. probably to be underestimated. When the numbers of deaths from mesothelioma are compared to the number of people who work in each occupation, it can be seen that the jobs which are proportionally most affected are sheet metal workers/boiler-makers (including workers in shipyards) and industrial coach-builders. They are followed by plumbers, carpenters and electricians. In France, a recent study showed that, depending on the generation, between 18 and 25 per cent of French men, i.e. millions of people, have been exposed to asbestos at least once in the course of their working life. Another French study (currently under way), conducted in six départements with a sample of men who had retired between 1994 and 1996, shows the extreme variety of job periods which entailed exposure to asbestos. The EC emphasize that many sectors are exposed to asbestos, and that they are not the traditional industries of asbestos mining and processing. Thus, 45 per cent of building and public works occupations are exposed to asbestos. More than 40 per cent of industrial production jobs involve exposure to asbestos, metallurgy and the machines and tools sector being particularly affected.

3.82 The EC state that the study carried out by Y. Iwatsubo on the French population confirms these data. This is one of the most important worldwide studies analysing the relationship between exposure to asbestos and the risk of mesothelioma; it covers the entire population of the country concerned. Although it had not been published at the time, this study was the subject of an analysis in the INSERM report (see pages 121 and 122). According to the EC, the very large scale of the study showed that there is a clear risk of cancer at levels of exposure that are lower than those previously recognized. This study shows that the vast majority of cases of cancer occur in workers who are “secondary users”. Great numbers of people are often employed in the areas in question, which explains the large number of cases of mesothelioma which occur. The study shows, for example, that exposure to asbestos took place in 54 per cent of job periods in the construction sector.

3.83 The EC consider that the principal data which have been presented illustrate the ubiquity of asbestos in the workplace which can, at sufficiently high levels of exposure, lead to numerous cases of fatal disease. As a rule, the very many categories of workers

<table>
<thead>
<tr>
<th>Occupation</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Builder</td>
<td>3.6</td>
</tr>
<tr>
<td>Painter and decorator</td>
<td>2.2</td>
</tr>
<tr>
<td>Building maintenance</td>
<td>2.2</td>
</tr>
<tr>
<td>Engineer - non specific</td>
<td>2.3</td>
</tr>
<tr>
<td>Machine operator</td>
<td>4.0</td>
</tr>
<tr>
<td>Electrician</td>
<td>3.6</td>
</tr>
<tr>
<td>Sheet metal worker/boiler-maker</td>
<td>2.3</td>
</tr>
<tr>
<td>Electric power station worker</td>
<td>0.4</td>
</tr>
<tr>
<td>Chemical engineer</td>
<td>0.4</td>
</tr>
<tr>
<td>Industrial coach-builder</td>
<td>0.8</td>
</tr>
<tr>
<td>Plasterer</td>
<td>0.6</td>
</tr>
<tr>
<td>Technician - non specific</td>
<td>0.5</td>
</tr>
<tr>
<td>Laboratory technician</td>
<td>0.5</td>
</tr>
<tr>
<td>Upholsterer</td>
<td>0.4</td>
</tr>
<tr>
<td>Metal sprayer</td>
<td>1.1</td>
</tr>
<tr>
<td>Boiler operator</td>
<td>0.9</td>
</tr>
<tr>
<td>Building foreman</td>
<td>0.9</td>
</tr>
<tr>
<td>Electrical engineer</td>
<td>0.9</td>
</tr>
<tr>
<td>Plasterer</td>
<td>0.6</td>
</tr>
<tr>
<td>Industrial designer</td>
<td>0.6</td>
</tr>
<tr>
<td>Engineer - non specific</td>
<td>2.3</td>
</tr>
<tr>
<td>Machine operator</td>
<td>4.0</td>
</tr>
<tr>
<td>Carpenter</td>
<td>5.7</td>
</tr>
<tr>
<td>Fitter</td>
<td>6.8</td>
</tr>
<tr>
<td>Dockers</td>
<td>1.5</td>
</tr>
<tr>
<td>Welder</td>
<td>1.6</td>
</tr>
</tbody>
</table>
affected handle materials containing asbestos sporadically, particularly asbestos-cement in building and public works occupations. These workers are often unaware of the risk they run. In fact, as Canada rightly and insistently notes, asbestos is undetectable when mixed with other materials, particularly cement. It is therefore not possible for the innumerable workers in all these sectors to be systematically informed of the risks they are taking when working with these materials. It is for this reason, above all, that there are also considerable risks to the general population such as do-it-yourself enthusiasts, who may often use products containing asbestos and may be exposed to significant quantities of asbestos dust when cutting, sanding, sawing and doing repairs of all kinds. It is therefore not entirely true, as Canada maintains, that there is no danger to the public from asbestos. According to the EC, Canada controversially finds that there is little difference between the undetectable risk associated with the levels of exposure in the ambient air - measured by the rate of asbestos fibres within a town or in the vicinity of a building - and the risks associated with the occasional, but sometimes high, levels of exposure which a very large section of a country’s population may face. Far from affecting only those working in asbestos production (mining and processing), the danger linked to inhaling asbestos at much higher levels than those cited by Canada is now particularly serious for users of products containing asbestos, whether they use such products in an occupational (textile workers, building workers etc.), para-occupational and/or domestic (do-it-yourself) context. In France, several hundred thousand or even millions of daily users of asbestos are therefore affected; they are never mentioned in Canada’s submission.

Canada responds that, even if the EC refuse to recognize this fact, there are significant physical and chemical differences between chrysotile and amphibole asbestos, distinctions which are reflected in the pathogenic potential of the two types of asbestos. These distinctions are crucial in this case since the current problem of asbestos in France is due essentially to past uses and to the use of amphibole fibres. According to Canada, the distinction between chrysotile and amphibole asbestos is also important by virtue of the fact that the extrapolations performed by INSERM to assess the risks associated with chrysotile asbestos are based on exposure to amphibole fibres, in proportions of up to 100 per cent and in circumstances which have nothing to do with the current uses of chrysotile asbestos. Today, 97 per cent of chrysotile fibres are used in high-density and non-friable materials. Construction materials and pipes based on asbestos-cement account for nearly 90 per cent of the international market. Friction products, for their part, account for approximately 7 per cent of the market, with the remaining 3 per cent consisting of various products such as seals, textiles and membranes. The debate must therefore focus on the current uses of chrysotile, i.e. essentially chrysotile-cement.

In simple terms, according to Canada, three characteristics of fibrous materials are currently recognized as being significant parameters which determine biological activity: “durability” (or biopersistence), “dimension,” and “dose.” In Canada’s view, these characteristics are all relevant in the assessment of the health risk, as WHO, INSERM and the EC recognize. It is important to highlight the lower biopersistence and pathogenicity of chrysotile fibres compared to amphibole fibres in order to demonstrate that the current health problems linked to asbestos are largely due to exposure to amphiboles. Canada emphasizes that the ban on current uses of chrysotile is not the solution to the health problems that France is experiencing today. The use of modern research techniques, in particular the mineral analysis of pulmonary tissues, sometimes called lung burden study, has made it possible to identify the “durability” factor as being a key parameter in the study of the pathogenicity of inhaled particles. This characteristic, which varies significantly from one particle to another and is probably linked to the chemical composition and crystalline configuration of the particles, will determine the extent of a basic biological phenomenon: biopersistence, i.e. the period during which the inhaled particles persist in the lungs and exert a harmful effect on the surrounding tissues before being eventually dissolved or eliminated.

Canada states that the recent studies which use both the fibre mass and the number of fibres as dose units confirm that amphiboles are more pathogenic than chrysotile. The WHO declares that the use of gravimetric data “may be misleading when comparing samples of chrysotile and amphibole asbestos, because the former may contain more than ten times more fibres per unit weight”. Thus, the studies demonstrating that
there is no distinction between chrysotile and amphiboles on a gravimetric basis mean in fact, according to the WHO, that amphiboles are more than ten times more dangerous than chrysotile, fibre per fibre. This is confirmed by INSERM, which reveals that the risk of mesothelioma is in fact ten times greater for amphiboles than for chrysotile. Several studies published in the early 1980s were conducted on samples of pulmonary tissues from workers whose deaths were thought to be linked to exposure to asbestos, compared to control subjects who had been exposed to various levels of urban pollution. The results showed that the quantities of amphiboles present in the pulmonary tissue of the experimental subjects were one hundred times higher than those found in the control subjects, but that the quantities of chrysotile were similar. Thus, the workers died from exposure to amphiboles, not to chrysotile.

3.87 Canada states that, according to the WHO and studies reviewed by it, the biopersistence of chrysotile is lower than that of amphiboles:

“The fractional deposition of chrysotile was lower than for amosite and crocidolite [...] The alveolar clearance of chrysotile was faster than that of crocidolite. [...] The retention of chrysotile, as measured a few days after the end of the 6-week exposure period, was only about one third that of amphiboles. [...] This difference in the lung clearance of chrysotile and amphibole fibres has been confirmed by several studies.”

3.88 Canada emphasizes that the half-life of amphibole fibres, although difficult to estimate, appears to be in the order of decades, while that of chrysotile fibres may be only a few months. The research of Dr. Bernstein, on the basis of the Interim Protocol for the Inhalation Biopersistence of Mineral Fibres of the EC, confirms, in a 1998 study, the higher biopersistence of amphiboles compared to chrysotile. Chrysotile alone, according to Dr. Bernstein, “would have little if any toxicological effect.” Even INSERM recognizes “[Tr.] the difference in carcinogenicity between the two types of fibres for mesothelioma.” According to INSERM: “[Tr.] experimental studies have shown that the biopersistence of chrysotile fibres was lower than that of amphiboles.” The IARC also recognizes that: “[I]n manufacturing and application industries mesotheliomas have been caused by exposure to crocidolite, and less frequently to amosite and chrysotile.” Finally, the IARC adds that when fibres are identified in the lungs, amphibole fibres are predominant.

3.89 Canada also recalls that Drs. Kumar, Cotran and Robbins state, in their authoritative medical textbook on pathology:

“The basis for the carcinogenicity of asbestos is still a mystery. Clearly, the physical form of the asbestos is critical; very nearly all cases are related to exposure to amphibole asbestos, which has long, straight fibers, and not to serpentine chrysotile.”

3.90 For his part, Prof. Sir Richard Doll states that:

“First, there is the difference between the effects of chrysotile and amphiboles which is so great in relation to mesothelioma that it is possible to argue that chrysotile does not cause mesothelioma at all.”

3.91 Canada emphasizes that the French Académie nationale de médecine stated, in 1996, that amphibole fibres are “[Tr.] currently considered the most dangerous” and that chrysotile fibres are “[Tr.] considered to pose little danger because of their spontaneous degradation in the human organism. [...] Chrysotile asbestos is a form of asbestos which has not caused mesotheliomas, except in cases of massive and prolonged exposure. This is apparently explained by its solubility in the organism.” Canada recalls that, in 1997, the French Ministry of Labour, through the report of the Groupe scientifique pour la surveillance des atmosphères de travail (Workplace air quality scientific monitoring group) (G2SAT), cited by the EC, was of the opinion that chrysotile, once in an acidic medium – the lungs for example – ceases to exhibit “virtually any carcinogenic activity”: 123
"[Tr.] It has been demonstrated that chrysotile is much more easily eliminated from the human lung than the other forms. Moreover, it ceases to exhibit virtually any carcinogenic activity (by intra-cavitary injection) after acid attack, which dissolves most of the magnesium."

3.92 Canada points out that chrysotile fibres are "curly" and downy whereas amphibole fibres are straight and rigid like needles. A 1998 WHO study notes that "Inhalation of respirable straight fibres [amphiboles] is reported to be associated with greater penetration to the terminal bronchioles than in the case of ‘curly’ fibres [chrysotile]." Once they have entered the respiratory tract, chrysotile asbestos fibres, because of their curly shape, are more easily cleared by the mucociliary process than straight and rigid amphibole fibres. Canada observes that, for chrysotile fibres which nonetheless manage to become lodged in the lungs, the macrophages are able to deal more easily with chrysotile fibres than amphibole fibres. In addition, amphibole fibres are much more resistant than chrysotile fibres in an acidic medium such as the lungs; they will therefore remain there longer than chrysotile fibres. According to the WHO, the lower biopersistence of chrysotile compared to amphiboles may be due in part to the fact that chrysotile fibres dissolve in an acidic medium such as the lungs, while amphibole fibres resist dissolution in this medium.

3.93 Canada states that it is a basic principle in toxicology that the health risks posed by a toxic agent are directly proportional to the duration of the contacts with the target organs. Thus, since chrysotile is less biopersistent than amphiboles, it logically follows that chrysotile is less pathogenic. In a study by Coffin et al. cited by the WHO in 1998:

"Large differences in the incidence of mesothelioma in intratracheal injection studies were demonstrated [...] crocidolite [was] 30-60 times more tumorigenic than chrysotile on fibre number basis."

3.94 Canada points out that, contrary to what the EC and the United States claim, the United States EPA states that studies demonstrate a difference in carcinogenicity between the types of asbestos fibres. The medical textbooks on pathology also indicate a clear distinction between amphiboles and chrysotile:

"It is important to make the distinction between various forms of amphiboles and serpentines, because amphiboles, even though less prevalent, are more pathogenic than the serpentine chrysotile, particularly with respect to induction of malignant pleural tumors (mesotheliomas). Indeed, some studies have shown the link is almost invariably to amphibole exposure.

This comes back to one of Canada’s main criticisms of the INSERM report, i.e. that the extrapolations performed by INSERM are based on data on exposure to amphiboles or to mixtures of fibres containing amphiboles. The danger of amphiboles cannot and must not be used to justify a ban on chrysotile, in light of the fundamental differences between the two types of fibres.

3.95 Canada asserts that convincing evidence of this distinction between the risk posed by chrysotile and the risk posed by amphiboles is the fact that the asbestos regulations in several industrialized countries make a clear distinction between the two types. Because of the greater risk involved, the regulations impose stricter exposure limits in the case of amphiboles than in the case of chrysotile. The following table indicates, for the sake of example, certain regulatory distinctions.

<table>
<thead>
<tr>
<th>Country</th>
<th>Amphiboles</th>
<th>Chrysotile</th>
</tr>
</thead>
<tbody>
<tr>
<td>France (1994)</td>
<td>0.3 f/ml</td>
<td>0.6 f/ml</td>
</tr>
<tr>
<td>European Communities</td>
<td>0.3 f/ml</td>
<td>0.6 f/ml</td>
</tr>
<tr>
<td>Canada</td>
<td>0.2 f/ml (crocidolite)</td>
<td>0.5 f/ml (amosite)</td>
</tr>
<tr>
<td>Japan</td>
<td>0.2 f/ml (amosite)</td>
<td>0.6 f/ml</td>
</tr>
<tr>
<td>Spain</td>
<td>0.5 f/ml</td>
<td>1 f/ml</td>
</tr>
<tr>
<td>United Kingdom</td>
<td>0.2 f/ml</td>
<td>0.4 f/ml</td>
</tr>
</tbody>
</table>
3.96 Canada notes that amphiboles have been banned in France since 1994, while chrysotile continued to be used there until 1997. Amphiboles have also been banned in the United Kingdom since 1992. Convention 162 and Recommendation 172 of the ILO both call for a ban on crocidolite, which they do not however impose for chrysotile. A WHO Committee which met in Oxford in 1989 also made the same distinction in its recommendations: "For crocidolite and amosite asbestos, on the basis of health, it is recommended that their use be prohibited as soon as possible." In the case of chrysotile, far from suggesting a ban, it suggests an exposure limit of 1 f/ml. The regulatory distinction recognizing the difference in pathogenicity is also found in the disputed Decree, which, for instance, makes an explicit distinction between amphiboles and chrysotile, permitting exceptions only for chrysotile and not for amphiboles. Canada points out that it has demonstrated that the pathogenic potential of amphiboles is much higher than that of chrysotile. Consequently, INSERM is taking the wrong approach by using exposure to amphiboles to determine the risks associated with chrysotile. It is the high pathogenicity of amphiboles – used more extensively in France than the EC would have one believe – which, with the use of friable materials, is the cause of the health problems observed today in the French population. Canada considers that serious questions must be raised as to the reasons which prompt the EC to ignore – indeed even to deny in the way they treat the available scientific data – the existence of all this evidence which establishes, beyond the shadow of a doubt, the significant difference in pathogenicity between chrysotile and amphiboles. Indeed, the latter are the basic cause of the French problem, while the former has no health effects when used in a context of controlled use.

3.97 Canada points out that the EC claim that there is no safe threshold of exposure to asbestos, whether of the amphibole or chrysotile variety. INSERM came to a different conclusion, at least with respect to asbestosis:

"[Tr.] Most of the epidemiological data gathered in exposed occupational populations suggest that clinically and/or radiologically confirmed asbestosis appears only following sufficiently high exposures [...]."

3.98 Canada recalls that the EC have adopted the position that human experience has not demonstrated the existence of a threshold of exposure in the case of lung cancer or mesothelioma below which exposure to asbestos dust poses no health hazard. On the contrary, human experience, via the epidemiological data, supports the idea of a threshold. According to Canada, it is wrong to claim that there is an "[Tr.] international consensus that there is no safety threshold for chrysotile." An EC study cited by WHO suggests the existence of a threshold:

"It is very likely that there is a practical level of exposure below which it will be impossible to detect any excess mortality or morbidity due to asbestos. [...] Thus, it is possible that there is a level of exposure (perhaps already achieved in the general public) where the risk is negligibly small." 

3.99 According to Canada, this appears also to be the position of the EC when they state that their main data "[Tr.] illustrate the ubiquitous nature of asbestos in the workplace which can, at sufficiently high exposure levels, cause numerous cases of fatal diseases." On the contrary, this statement presumes that no disease appears at low exposure levels to asbestos dust in the workplace. The existence of a threshold suggests the possibility of demonstrating that the effect does not manifest itself at or under an identified dose. However, the unequivocal scientific demonstration of a zero effect is impossible, as DG XXIV acknowledges.

3.100 Canada asserts that, when the available epidemiological data are insufficient to make it possible to determine the risks of cancer associated with low exposure to a toxic contaminant, a methodology called "risk analysis" is used. Quantitative risk analyses project the risks observed at high exposure (experimental or occupational studies) to low exposure possibly associated with unobservable low risks. Such projections outside the field of empirical observations must be based on mathematical models. Canada indicates that the
linear model is one of the possible risk projection models. This model is simple to calculate mathematically and statistically. It implies, however, that there is no exposure value, no matter how low, which is not associated with a certain level of risk, no matter how low. The uncertain choice of the best model has a major impact on the estimated risk. In the case of asbestos, the linear model is in fact used to extrapolate risks from workers heavily exposed in the past to the exposures 100,000 times lower of the general population and asbestos workers today. In statistics, any extrapolation far from the region of observations (observable values, available studies) is speculative and very risky.

3.101 According to Canada, the EC assert that the linear model is the most plausible by using the following fallacious argument: "[Tr.] This hypothesis of the linearity of the risk based on the level of exposure and the lack of a threshold are part of the currently accepted assumptions concerning carcinogens; for instance, it is included in the ILO encyclopedia. This hypothesis is therefore the most plausible." However, this does not mean that this is the most plausible model, but rather the simplest and most "conservative" model in that it projects higher risks in the region of low doses than the other mathematical models of carcinogenesis. The risk assessment agencies emphasize this distinction and this criterion. The EC confuse the lack of an identified threshold with the lack of a threshold. For instance, the WHO report on chrysotile asbestos notes that: "No threshold has been identified for carcinogenic risks." For Canada, this means simply that a specific threshold cannot be identified, if there is one. At no place in the WHO report or in any other risk assessment is it claimed that there is no threshold. On the contrary, this possibility is mentioned by the Health Effects Institute-Asbestos Review (HEI-AR). According to this group of international experts, which included experts in modelling of cancer risk such as J. Peto, D. G. Hoel and W. Nicholson, the linear dose-response model is not adopted for its validity but rather because it tends to over-estimate the actual risk. This model ignores the natural biological defence mechanisms against toxic invaders of the body, which, as Canada has previously demonstrated, are very effective in the case of chrysotile. With respect to the validity of the linear model in general, Prof. Doll wrote as follows:

“We have no real ground for postulating that a linear relationship for lung cancer can be extrapolated back to the levels of dose with which we are concerned in non-occupational settings.”

3.102 Canada points out that, similarly, Ames and Gold wrote: “linear extrapolation from the maximum tolerated dose in rodents to low-level exposure in humans has led to grossly exaggerated forecasts in mortality.” E. Fournier and M.-L. Efthymiou are even more severe. In their view “linear extrapolation to zero is an unscientific methodology whose social consequences are so immense that it warrants unconditional elimination.” Canada indicates that, once a cell has been altered carcinogenically, there are natural defence mechanisms of the organ or of the organism which seek to destroy or eliminate the invaders. The conventional models of carcinogenesis do not take this broader organic context into account. The EC and INSERM appear to confuse dose and exposure in their justification of the linear model. This model is applied to the ambient exposures of workers and of the general population, while it is justified in carcinogenesis on the basis of the effective dose of the target tissues. Not only is it implausible, according to Canada, that the dose-response relationship should be linear at the cellular level (level of mathematical modelling of carcinogenesis), but this is even more implausible at the level of the exposure-effects relationship. In fact, the deposition and clearance of solid particles in the lungs may be non-linear saturable processes, in which case, even if the relationship between the dose in the tissues and the risk of cancer were linear, the relationship between dose and exposure would be infra-linear, perhaps even with a threshold, and the resulting relationship between exposure and risk would also be infra-linear, or with a threshold. And this would also depend on the type of fibres.

3.103 Finally, Canada emphasizes that the available epidemiological studies which make it possible to study the relationship between lung cancer and exposure to asbestos are statistically compatible with the linear model, but when an a priori non-modelized method such as “adjustment by least squares weighted by distance” is applied, the form of
the resulting curves is generally infra-linear, except in the case of workers exposed to 100 per cent amosite. Infra-linearity and even a threshold have been noted by various researchers in the case of chrysotile-cement workers, and miners exposed to vermiculite contaminated by tremolite. For the reasons outlined above, in Canada’s view, the linear model must be presented as a possible model yielding an upper risk limit, and not as yielding the most accurate or most probable risk estimate. Finally, the possibility of a threshold must be considered plausible and even very probable, even if it is difficult to determine this threshold quantitatively.

3.104 The European Communities affirm that the carcinogenic character of asbestos for humans is internationally recognized. Since 1977, the WHO has recognized that all varieties of asbestos, including chrysotile, are carcinogenic, causing cancer of the lung or mesothelioma. In 1986, the ILO advised lawmakers, through Convention No. 162, to have asbestos replaced by less harmful materials or technologies as soon as possible. In 1996, the WHO recommended that asbestos, including chrysotile asbestos, be replaced by harmless substitutes, wherever possible. In 1998, WHO reaffirmed the carcinogenic effect of chrysotile asbestos, particularly with respect to mesothelioma, continued to promote substitution and noted that the risk was very widespread among numerous categories of workers.

3.105 According to the EC, asbestos is at the root of a public health problem and chrysotile is the cause of most asbestos-related diseases. In 1998, the WHO reaffirmed that chrysotile “has been associated with an increased risk of pneumoconiosis, lung cancer and mesothelioma in numerous epidemiological studies of exposed workers”, which confirms the conclusions of the 1996 INSERM report. Since asbestos began to be used for industrial purposes, chrysotile has accounted for about 95 per cent of the asbestos consumed in France. Since 1988, chrysotile has accounted for all the asbestos consumed in France. According to the EC, these figures show that most asbestos-related diseases are caused by chrysotile and not amphiboles, as Canada would have the Panel believe. This is confirmed by recent publications. Thus, according to the study by Stayner et al.:

“Our review of both the toxicologic and epidemiologic literature strongly supports the view that occupational exposure to chrysotile asbestos is associated with an increased risk of both lung cancer and mesothelioma.”

3.106 The EC note that the study coordinated by J. Peto and published in 1999 found that in six European countries (France, Germany, Italy, Netherlands, Switzerland, United Kingdom) nearly 10,000 people died of mesothelioma between 1990 and 1994. Moreover, it estimated that during the period 1995-2029 about 200,000 people would die of mesothelioma. If these figures are extrapolated to all the countries of Western Europe and deaths due to lung cancer are included, then the results show that exposure to asbestos could lead to about 500,000 deaths by cancer between now and 2029. In France, the number of deaths due to mesothelioma is increasing steadily. A recently published study, similar to that made by Peto et al., predicts that the annual number of deaths due to mesothelioma in France will go on increasing up to 2020. It is estimated that in France over the entire period 1996-2020 a total of 20,000 men and 2,900 women will die of mesothelioma.

3.107 The EC state that chrysotile poses a threat to an extremely large section of the population. Several scientific studies show that a huge range of occupations and economic sectors is at risk. For a long time, the asbestos producing and consuming countries considered that the risks of cancer were wholly restricted to “primary users” (asbestos miners and workers in the asbestos-processing industry). In the early 1990s, an international scientific consensus emerged to the effect that the asbestos-related risks for secondary users (users of asbestos-based products, servicing and maintenance workers, “do-it-yourself” enthusiasts, and specialists in asbestos removal and containment) had been historically underestimated. According to the EC, many sectors are exposed to asbestos and they are usually far from fitting the description of the traditional asbestos mining and processing industries. Thus, 4 per cent of construction industry trades and 4 per cent of
jobs in the industrial manufacturing sector are exposed to asbestos. As noted by the WHO in its report 203 on the risks of chrysotile asbestos: “Risks are likely to be greater among workers in construction and possibly other user industries”. The recent study by Y. Iwatsubo et al. shows that the great majority of cancer cases occur among “secondary users”. These workers are often unaware of the risk they are running, since when asbestos is mixed with other materials, particularly cement, its presence is not readily discernible.

3.108 The EC stress that the international scientific community does not recognize the existence of a threshold of harmlessness for chrysotile. It is important to note the distinction between the establishment of occupational exposure limits and the existence of a threshold of harmlessness below which there is no risk to health. The occupational exposure limits take into account various criteria, including the technical feasibility of obtaining sufficiently low exposure levels in the workplace and the technological means of measuring actual levels for monitoring purposes. Thus, the occupational exposure limit does not correspond to a threshold of harmlessness which, in the case of asbestos, has never been found. Since as long ago as 1976, the IARC has recognized that “at present it is not possible to assess whether there is a level of exposure in humans [to asbestos] below which an increased risk of cancer would not occur”. This position received support from the WHO in its 1998 report, which states that “no threshold has been identified for carcinogenic risks”, thus confirming the conclusions of the 1996 INSERM report. According to the EC, all the scientific studies show the existence of a linear relation between the dose of asbestos inhaled and the risk of cancer. In 1998, the WHO noted that “there was a clear dose-response relationship (...)”. The results of more recent studies confirm the impossibility of identifying a threshold below which asbestos would not present any risk for the populations exposed. As pointed out in the Collegium Ramazzini report, “The strictest occupational exposure limits in the world for chrysotile asbestos (0.1 f/cc) are estimated to be associated with lifetime risks of 5/1,000 for lung cancer and 2/1,000 for asbestosis (Stayner et al., 1997).” The EC incorporate here, by reference, the report by the Collegium Ramazzini communicated to the Panel on 7 May 1999.

3.109 The EC state that the number of cases of asbestos-related diseases correlates with the amount of asbestos imported. An analysis of the data for ten Western countries shows a very strong correlation between the incidence of mesothelioma and per capita asbestos consumption measured in terms of imports. In each country, the number of cancer cases increases in proportion to the increase in asbestos imports. Accordingly, in view of the acknowledged carcinogenicity of asbestos, whatever the variety, the number of deaths recorded or predictable over 30 years, the absence of a threshold of harmlessness, the proportion of total asbestos consumption represented by chrysotile and the direct link between the amount of asbestos used and the number of cancer cases, it is essential to stop the risk from spreading by banning all future use of asbestos, whether amphibole or chrysotile.

3.110 The EC state that France has the right to establish its own level of protection against the inhalation of asbestos fibres. The scientific findings available to France when it took its decision to ban asbestos were unambiguous. It has been scientifically established that: (i) chrysotile is a toxic material which has a dramatic effect on health (mesothelioma, cancer of the lung, asbestosis) and is no less toxic than other varieties of asbestos; (ii) there is no threshold of harmlessness as far as the carcinogenicity of chrysotile is concerned and large numbers of people (several millions) are at risk; (iii) controlled use is neither applicable nor effective; (iv) the substitutes for asbestos in asbestos cement (PVA, cellulose) and paraaramids give no cause for concern, asbestos-cement accounting for the vast majority (90 per cent) of the asbestos being used at the time of the ban; (v) there is no evidence of the other substitutes (man-made mineral fibres) being carcinogenic in humans. Among these substitutes, only ceramic fibres, which are very rarely used (to withstand very high temperatures), are carcinogenic in animals and should therefore be employed with due care.

3.111 The EC emphasize that the scientific data on which France based its case were of very high quality. The substitute fibres are more than 10 microns in diameter, which makes it physically impossible for them to penetrate into the pulmonary alveoli. With regard to
the effects of asbestos on health, France based its case on the joint report of INSERM\textsuperscript{175}, one of the most important world biological and medical research bodies. INSERM’s report consisted of a critical and reasoned review of the world scientific literature by a multidisciplinary team of 11 scientific experts. The method and scope of this work (12 chapters of discussion supported by 1,200 bibliographic references) make it fundamentally different from any monograph or fragmentary study. The quality of the work has been acknowledged by the entire scientific community, including the Canadian experts. The United States, in its third-party submission, fully endorses the conclusions of the INSERM report, including those concerning linear extrapolation to low doses. The EC consider, therefore, that Canada cannot criticize France for basing its case on this report. France has been concerned with the question of substitutes for asbestos ever since it began considering a ban. At the same time as it requested a report on asbestos from INSERM, in the autumn of 1995, France requested an initial report on man-made mineral fibres from the \textit{Groupe de surveillance des atmosphères de travail} \textsuperscript{176} (G2SAT) in order to obtain an initial insight into the harmfulness of the most suspect fibres which the proponents of the controlled use of asbestos are always stressing. This group based its conclusions on previous scientific research conducted over a period of many years.\textsuperscript{177} The conclusions of G2SAT, submitted to the Government in June 1996, were confirmed by the results of the INSERM report on man-made fibres begun shortly after the submission of the report on asbestos. The fibres used as a substitute for asbestos in asbestos-cement were found to have given no cause for concern. This was confirmed by the report of the CSTE\textsuperscript{178} of DG XXIV of the Commission of the European Communities, and then by the United Kingdom’s COC.\textsuperscript{179} The scientific data on substitute fibres, on which France based its decision to ban asbestos, are thus broadly supported by other scientific authorities. According to the EC, Canada cannot maintain that France has replaced asbestos with the “unknown” risk of substitutes.

3.112 The EC state that, in the light of this assessment of the risk based on sound and internationally recognized scientific findings, France has adopted a risk management regime. As “safe” use is insufficient to eliminate the risk of an excess of cancers and as France is applying, on the one hand, the principle of replacement by a less dangerous product and, on the other, the principle of reduction of the risk to the lowest level technically feasible, it has decided to impose a ban with exceptions. The EC contend that this is the only solution capable of preventing the spread of the risk associated with materials containing asbestos. The measure is sharply reducing asbestos consumption, which is known to be very closely linked with the incidence of asbestos-related diseases. This solution offers every technical guarantee of safety since provision is made for exceptions when substitution does not ensure equivalent performance. At the same time, France has undertaken to explore the question of substitute products\textsuperscript{180} in greater depth. It is also simultaneously strengthening the measures designed to protect workers exposed to “\textit{in situ} asbestos”, namely, the hundreds of thousands of maintenance workers, as well as the much smaller numbers engaged in more dangerous work such as asbestos removal and containment. France is also reinforcing its building surveillance measures by monitoring false ceilings, as well as flocking and lagging.

3.113 The EC emphasize that the aim of the Decree is consistent with the WHO and ILO recommendations. It is intended to establish an obligation to replace asbestos and products or materials containing asbestos whenever alternative materials or techniques are available, assuming always that they constitute a lower risk. The Decree provides for temporary exceptions to the rule in those very few cases in which there is no lower-risk substitute capable of ensuring equivalent performance. For these residual uses of asbestos, enterprises must request a waiver and undertake to carry out research with a view to abandoning the use of asbestos as quickly as possible. An enterprise which replaces asbestos with another material must rigorously: (i) ensure that, in the light of the knowledge available, the substitute represents a reduced risk to the health of a worker handling the product; and (ii) test the finished product in order to verify that it offers every technical guarantee of safety appropriate to its end use. Some three years after the Decree came into force, the use of asbestos in France is now much reduced and only involves a few precisely targeted and not very widespread industrial applications. The EC estimate that almost 100,000 tonnes of carcinogenic fibres have therefore been eliminated during this period in
the form of materials potentially dangerous for the user and have been kept out of French buildings and factories.  

3.114 According to the EC, Canada’s case is based on incomplete and largely erroneous data. In its arguments, Canada relies heavily on assertions which have no scientific foundation and, more often than not, are based on old or partial reports that are scientifically obsolete or of very debatable value. Canada makes many erroneous statements in support of its case and seeks to obscure well-established facts by creating confusion or omitting important data. The EC contend that many of Canada’s assertions are unfounded or erroneous, for example:

(a) The characteristics of chrysotile:

(i) The failure to make a distinction between the risk of mesothelioma and the risk of cancer of the lung: Canada regularly omits to point out that, whereas the risk of mesothelioma is lower for chrysotile than for amphiboles, this is not true for cancer of the lung;

(ii) The assertion that the risk due to chrysotile is “undetectable”, whereas many scientific studies show that its effects are indisputable.

(b) The French data:

(i) The assertion that amphibole fibres were “widely used” in France is absolutely wrong;

(ii) The assertion that France has favoured French products, whereas the EC has emphasized that most of the products used as substitutes for asbestos are imported into France.

(c) “Safe” use:

(i) The assertion that there is no scientific evidence of risks linked with so-called “modern” use (also called “controlled” or “safe” use) of chrysotile, whereas numerous scientific publications cited by the EC (but not mentioned by Canada) clearly show the opposite to be true;

(ii) The assertion that “there are effective methods for reducing dust creation” and that the safe use of asbestos is “based on proven scientific knowledge”, whereas all the data by the EC show the opposite to be true.

3.115 The EC state that Canada is seriously confusing the issue or making significant omissions.

(a) Confusing the issue:

(i) Confusing the level of chrysotile fibres in the environmental ambient air with that in the workplace by suggesting that the very low levels of fibres encountered in the ambient air of cities and buildings are the only problem potentially linked with asbestos. The level of 0.0005 fibre per ml of air, which Canada always cites to show that the corresponding risk is “undetectable” or zero, is about 100,000 times lower than that created by certain routine operations on asbestos-cement parts;

(ii) Systematic confusion between friable products and amphiboles with non-friable products and chrysotile. In
reality, raw asbestos, whether of the amphibole or chrysotile variety, is always spontaneously friable. It is quite wrong to associate amphiboles exclusively with friable products, such as flocking and lagging, and chrysotile exclusively with non-friable products, such as asbestos-cement. Since amphiboles were banned, all asbestos-containing products, whether friable or non-friable, have been based on chrysotile.

(b) Omissions

(i) Systematic omission of the circumstances of occupational exposure most dangerous to health. Thus, in its description of the circumstances of exposure in the construction industry, Canada mentions construction and demolition activities, but simply disregards the fact that once asbestos has been used in the construction of a building, and until the latter is demolished, any installation containing asbestos is likely, during its life cycle, to require some form of plumbing, heating or electrical work carried out by professionals or amateurs.

(ii) Systematic omission of the high occupational levels of exposure associated with everyday uses. Canada cites only a few figures indicating low levels of exposure, omitting to mention the very high figures encountered daily, although they are well-known and have been widely published;

(iii) Systematic omission of the data on the ill-effects on health associated with the use of chrysotile in Canada itself. Thus, Canada seems to be unaware of its own statistics, which reveal a very sharp and rapid increase in cancers caused by chrysotile. Canada must be aware of the publicly-funded scientific research being carried out in Canada by well-known research teams, the results of which are widely published in the scientific press. Canada never refers to this work, which shows that in Canada the risks due to chrysotile are high and at least as great as in the other industrial countries;

(iv) Systematic omission of the most recent scientific research confirming the risks associated with chrysotile, in particular the reports of international organizations to which Canada itself belongs. Thus, Canada cites two World Health Organization reports, one of which dates from 1986 and the other from 1988. It appears that, for Canada, research and the acquisition of new knowledge came to a halt more than ten years ago. Thus, Canada makes no reference to report 203 of the World Health Organization published in 1998 (although it is clearly identified as being an update of the 1988 report), which recognizes the dangers to health posed by chrysotile asbestos and clearly expresses doubts about the possibility of effectively controlling exposure in the building industry.

3.116 The EC state that Canada is using reports whose scientific value is highly debatable or citing reports too selectively. Canada considers that France’s ban on all forms of asbestos is not scientifically well-founded. This criticism is mainly based on two reports. These reports are, however, of very debatable scientific value or cited only selectively, with
the omission of those elements that conflict with Canada’s point of view. According to the EC, the report of the Académie Nationale de Médecine, signed by E. Fournier, is of debatable value. As pointed out by Prof. Terracini of the University of Turin: “a major limitation of this report is the total lack of bibliographic references. In addition, it is severely unbalanced on several issues”. Drawn up by someone who had never previously published anything in the field of asbestos-related risks, the report can be no more than the expression of a personal opinion. In this respect, Prof. Terracini’s conclusion is unequivocal:

“This report has been obviously written by persons with no knowledge of the epidemiological methods and of the subtleties of causal inference. It provides no original information and can hardly be of any use in an overall evaluation of the case”.

3.117 The EC state that the report of the Royal Society of Canada was prepared in haste at the request of the Canadian Government. It should be noted, first of all, that on its own admission the Royal Society of Canada’s Panel of scientific experts had to work too quickly, as it several times acknowledges (see, for example, page 15 of the English text), was not able to reach a consensus among its members (page 15 of the English text) and worked on an incomplete draft of the INSERM report (page 1 of the English text), which obviously explains certain erroneous interpretations resulting from a misunderstanding of the text. The arguments developed in the Royal Society of Canada report have already been the subject of a detailed analysis which showed that the criticisms of the INSERM report were based on an over-hasty reading of an incomplete document and are not well-founded. The EC note, furthermore, that Canada used the Royal Society of Canada report partially and selectively and, in any event, failed truly to reflect the opinion of the group of scientific experts on the main conclusions of the INSERM report. In this connection, the following extracts from the Royal Society of Canada report should be cited:

“The Panel agreed with INSERM’s findings on the following points: all asbestos fibres are carcinogenic, regardless of their mineralogical nature; the risk of lung cancer is higher for longer and finer fibres for low dose and dose-rate exposures, the linear no-threshold model is used by all regulatory agencies that have performed quantitative risk assessments; research should be done on substitutes for asbestos; the linear no-threshold hypothesis for low exposure levels is not the only possible strategy in risk assessment, but evidence is not available to demonstrate that a different hypothesis is a superior predictor of risks from low exposures.” (pages 4-6 of the English text)

3.118 The EC stress that the points on which the Royal Society of Canada says it agrees with the INSERM report are the essential points on which the French decision to ban asbestos is based.

“Our hope was that we could give consensual answers. We could not always do so. In retrospect, the hope seems to have been foolishly optimistic. Scientists cannot achieve consensus on contentious issues after two weeks of reading and two days of face-to-face discussion.” (pages 14 and 15 of the English text)

3.119 In conclusion, the EC emphasize that Canada’s assertion that the experts who analysed the INSERM report “sharply criticized” the methods employed and “very severely criticized” its conclusions is, at the very least, tainted with partiality and does not truly reflect the contents of the Royal Society of Canada report. Moreover, it is inconsistent with almost all the scientific data from the relevant international institutions.

5. “Controlled” or “safe” use of chrysotile asbestos

3.120 Canada emphasizes that the public’s perception of the harmful health effects of using different types of asbestos in various applications has exerted a greater influence on risk management decisions than risk evaluation has. This phenomenon is the result of a
conceptual confusion between the terms “hazard” and “risk”. Information on “hazards,” erroneously presented as information on “risks”, has provoked unjustified fears with regard to the undesirable effects of asbestos, such as cancer, in the population. In Regulation No.1488/94 of the European Commission, hazard is described as being “the adverse effects which a substance has an inherent capacity to cause”. Risk, on the other hand, is defined as “the likelihood that an adverse effect will be caused under the known or the reasonably foreseeable conditions of use”. Thus, it is appropriate to distinguish between the “hazards” associated with chrysotile fibres and the “risks” associated with the modern uses of chrysotile fibres. According to Canada, the characterization of the hazardous properties of a substance is not the same as an evaluation of the real risk of all its uses. The evaluation of the associated hazards is an important element but not sufficient in itself to evaluate the risk, which also encompasses evaluating the exact nature of the substance, data on long-term exposure and estimating the probable risk under current conditions of use. Thus, it is essential that the issue of the risks associated with a product be evaluated in the light of its various uses.

3.121 Canada points out that, today, chrysotile represents more than 97 per cent of the asbestos used. Its use is limited to a certain number of high-density applications, above all construction products based on chrysotile-cement, friction materials based on chrysotile and a number of plastic, resin and bitumen materials containing chrysotile, all of which are safe applications. The distinction between these uses and past uses is vitally important when evaluating the real risks posed by chrysotile. In Canada’s view, an evaluation of the effects of asbestos in the workplace or in the environment must take into account the latest risk analyses, the current occupational exposure limits and modern applications using chrysotile exclusively in high-density materials. The risk can be managed through numerous forms of government action. In the case at issue, two forms of government action are examined for the purpose of determining the appropriateness of the French measure: controlled use; and the complete ban on asbestos, without distinction among the types of asbestos fibres and their multiple uses.

3.122 According to Canada, the appropriate manner of regulating asbestos must be seen in a broad context. In a world in which global trade is becoming increasingly extensive, it is all the more important to adopt a uniform approach to the formulation of policies and regulations. The most effective and objective way to deal with these matters is to seek an international consensus whenever possible. With regard to asbestos, in Canada’s view, the scientific data provide a solid basis for reaching sensible decisions regarding the regulation not only of asbestos, but also of the growing number of potentially hazardous fibres in general, some of which are being used in France as a substitute for asbestos. The lessons to be learned from the uncontrolled use of asbestos point to the importance of establishing appropriate regulations. The most logical and consistent approach to regulation is to reach an international consensus based on the best scientific data. Thus, controlled use means establishing and applying appropriate regulations to ensure the rigorous control of exposure to chrysotile and products containing chrysotile, with the aim of reducing to an undetectable level the risks associated with the extraction, crushing, manufacturing, shipping and handling of this product. Chrysotile is expressly mentioned because the use of amphiboles is banned virtually everywhere at the present time. The principle of controlled use also means that certain uses for which exposure cannot be controlled to an acceptable degree should be banned.

3.123 Canada considers that one of the most in-depth reports on the effects of asbestos and asbestos-containing products is the report published in 1984 by the Royal Commission on Matters of Health and Safety arising from the Use of Asbestos in Ontario. This report, still considered one of the most exhaustive studies of asbestos, recommends that the use of chrysotile be continued under conditions of controlled use. In 1988, under the aegis of the International Programme on Chemical Safety of the World Health Organization (WHO), a working group was convened to evaluate the environmental contamination caused by asbestos. In its recommendations, the working group drew a distinction between the risks posed by brittle, low-density products and the risks posed by high-density products such
as chrysotile-cement. The working group recommended that the use of low-density products be discouraged all over the world. It also concluded that the normal use of products containing high-density chrysotile, including chrysotile-cement, does not pose a risk to the general public. In 1989, a group of experts met at Oxford under the aegis of the WHO and formulated recommendations for exposure to asbestos limits in the workplace. In their final report, the participants recommended a maximum exposure of 2 f/ml (weighted average over eight hours), an objective that can be reached everywhere at low cost, and expressed their desire to see efforts made to reduce this rate to 1 f/ml.

3.124 Canada emphasizes that, apart from the regulation of exposure to chrysotile limits in the workplace, the issue of asbestos in buildings, both with regard to occupants and to maintenance and repair crews, has been addressed on several occasions. A study by the HEI published in 1991 concluded that there is not sufficient evidence of health risks to the occupants of buildings where sprayed asbestos is used to justify the removal of intact materials containing asbestos. According to Canada, the conclusions of the HEI are in agreement with those of the experts from around the world who attended a symposium held at Harvard University in 1989 on the health aspects of exposure to asbestos in buildings. The report of this symposium states that the dust diffusion values in buildings where materials containing asbestos have been used are extremely low. The report also states that the risk posed by installed asbestos is very low, both in absolute and relative terms, and that this risk is much less acute than most other environmental hazards, including cigarette smoke and radon, for instance. Nonetheless, the report continues, increased protection should be afforded to public works and maintenance crews exposed to brittle asbestos that can be disturbed.

3.125 Canada points out that all international standards on asbestos establish parameters for its controlled use. Chief among these is Convention 162 concerning Safety in the Use of Asbestos and the accompanying Recommendation 172. These standards were adopted in 1986 by representatives of governments, industry and labour unions meeting under the aegis of the International Labour Organization (ILO). Convention 162 recommends strict regulation of chrysotile, but does not recommend a total ban, except for crocidolite and sprayed asbestos. Recommendation 172 sets forth the minimum labour standards to be observed in order to prevent and control the risks associated with occupational exposure to chrysotile. Countries which ratify Convention 162 concerning Safety in the Use of Asbestos make a commitment to implement laws and regulations prescribing the measures to be taken to prevent and control the health risks associated with occupational exposure to asbestos and to protect workers against these risks. In addition, there exists a general reference code, the collection of the International Labour Office entitled Code of Practice on Safety in the Use of Asbestos. The guidelines contained in this Code are aimed at preventing, eliminating or reducing as much as possible the exposure to asbestos dust and safeguarding the health of workers engaged in operations that could give rise to such exposure. The first part of the Code deals with workplace regulations, methods of prevention, packaging, shipping and intermediate storage of asbestos, the elimination of debris and the protection of workers’ health. The second part deals with prevention in certain activities: mining extraction, the manufacturing and utilization of asbestos products (asbestos-cement, friction products, etc.), insulation work, construction work and demolition work. Finally, the ISO has issued a number of standards applicable, inter alia, to the diverse applications making use of asbestos. These standards cover the use of a broad range of modern products containing chrysotile-cement, including tubes, corrugated, ribbed and smooth panels, shingles and pipes. In particular, they contain guidelines for construction sites where asbestos-cement products are in use.

3.126 Canada notes that, in the United States, the attempt by the EPA to ban asbestos failed. In 1989, the EPA adopted a regulation that provided for the prohibition and progressive elimination of 94 per cent of all commercial uses of asbestos over a period of seven years. In so doing, the EPA had wholly disregarded all the evidence submitted to it indicating that the past, outdated uses of asbestos did not provide any justification for a complete ban and that chrysotile asbestos can be used in a safe manner. The EPA did not consider either the uncertainty surrounding the health risks of using substitute products or the ad-
ditional costs of using such substitute products. The EPA regulation banning asbestos was appealed in court. The Court of Appeals concluded that the risks presented by the EPA were not supported by the scientific facts. The Court ruled that the EPA would have to prove that the products covered by the ban posed an unacceptable risk. Thus, the ban was annulled and the EPA was ordered to consider all the evidence that it had initially disregarded. In 1997, practically all the uses that had originally been banned were re-admitted by the EPA.

3.127 Canada points out that, prior to adoption of the Decree, France maintained a policy of controlled use. Indeed, it appears that the legislative provisions on asbestos provided adequate protection to workers and consumers. A review of French legislation shows that the hazardous uses of asbestos were already banned before the Decree. Sprayed asbestos was banned for all buildings in 1978. Amphiboles were banned in 1994, including marketing, use and importation of these products. The levels of exposure to asbestos dust were strictly regulated. Successive reductions in the exposure limits to airborne dust in workplaces had been adopted. The regulation currently in effect in France prescribes a workplace exposure limit of 0.1 f/ml. A large number of products that had traditionally contained asbestos had already been banned by successive Decrees. Finally, measures had been taken to protect the population against passive exposure to asbestos in buildings. A Decree imposes the obligation on all building owners, with the exception of owners of single-family dwellings, to search for the presence of sprayed asbestos or heat insulation containing asbestos and brittle materials likely spontaneously to release asbestos fibres into the air and to verify the good condition of these materials by means of an evaluation grid or dust measurements, so as to determine whether work would be necessary. In light of all these actions, it may be concluded that the French Government had already regulated the risks associated with the use of asbestos, including those deriving from the use of chrysotile fibre, prior to the adoption of the ban. According to Canada, there have been no new scientific findings that would justify the change in the asbestos regulations adopted by France.

3.128 The European Communities indicate that Canada implies that chrysotile previously accounted for a smaller proportion of the asbestos consumed. This statement is wrong, as can be seen in the figures for global consumption of chrysotile and amphibole asbestos. In France, for example, since 1945, around 97 per cent of the asbestos used has been chrysotile-cement. Furthermore, the EC highlight below the inadequacies of safe use for protecting the health of workers and the population. The EC welcome Canada’s recognition that “The principle of controlled use also means that certain uses for which exposure cannot be controlled to an acceptable degree should be banned” (Canada does not, however, define what these uses are nor their frequency). Like the WHO, the EC consider that all servicing and maintenance operations cannot be controlled because of their frequency and their ubiquity. Moreover, according to the EC, international scientific data show that controlled use does not suffice to prevent the risk of cancer in asbestos-processing industries (manufacture of asbestos-containing products) nor in removal activities. Under such circumstances, the EC consider that it is justified to ban all uses of asbestos.

3.129 The EC point out that Canada cites the reports of the Ontario Commission and the WHO: one of which dates from 1984 and the other from 1988. It appears that, for Canada, research and the acquisition of new knowledge came to a halt more than ten years ago. Thus, Canada makes no reference to report 203 of the World Health Organization published in 1998 (although it is clearly identified as being an update of the 1988 report), which recognizes the dangers to health posed by chrysotile asbestos and clearly expresses doubts about the possibility of effectively controlling exposure in the building industry. The EC stress that this document confirms the conclusions of the INSERM report. The EC note that, according to Canada, “a group of experts met … under the aegis of the WHO … [and] … recommended a maximum exposure of 2 f/ml … [and] … expressed their wish that efforts be made to reduce this rate to 1 f/ml”. Here again, the document cited by Canada dates from 1989. The EC note that, since then, many findings have highlighted the risks at much lower levels: some of these findings were analysed in the INSERM report.
3.130 The EC point out that the INSERM experts were aware of the HEI report, cited by Canada, and analysed it in detail in 1996 – observing that:

- It only deals with risks associated with the presence of persons inside buildings containing asbestos, excluding other exposures, occupational exposure in particular; INSERM’s mandate related to all exposures; concerning exposure in buildings, INSERM’s findings are the same as those of the HEI, but INSERM focused above all on what is considered by all experts to be the major risk of asbestos, namely, occupational exposure.

- The model used by the HEI to establish the very low risks linked to very low levels of exposure usually to be found in well-maintained buildings is exactly the same as that used by INSERM (linear model without a threshold) and so disparaged by Canada. The information concerning this model can be supplemented by citing the report of the Royal Society of Canada, to which Canada often refers: “but the Panel agrees [with INSERM] that regulatory agencies use the model in this form”. Lastly, the EC consider that it is tendentious to state, as Canada does, that in France “the hazardous uses of asbestos were already banned before the Decree”. This statement takes it as a fact that uses such as work on asbestos-containing materials, which were certainly not banned in France, are not hazardous. The EC reject this statement.

3.131 The EC point out that the available scientific data show that a high mortality rate persists despite the so-called “safe” use of chrysotile asbestos. Surveys carried out more than thirty years after the introduction of a “controlled use” policy in the United Kingdom indicate a significant excess of deaths from lung cancer and mesothelioma, not just among workers in plants where “safe” use was enforced, but above all in the population working outside such plants. According to the manual of the Canadian Asbestos Institute, the “safe” use policy is based on an “at-source” control system which consists of an “agreement” among producers, chrysotile exporters and manufacturers of asbestos-containing products, whereby the latter undertake to comply with rules designed to protect their employees and, in particular, to ban the on-site cutting of asbestos-cement. This is possible because Canadian production methods enable “made-to-measure” units to be manufactured at the plant, thus obviating the need for on-site cutting. However, as the manual admits, voluntary participation by manufacturers remains the cornerstone of this policy. The manual also provides for the supply of the products concerned to be stopped, after consultation with the national authorities, in the event of any failure to comply with this “agreement”. According to the EC, this procedure is limited in a number of ways: (i) it covers manufacturing companies only; and (ii) it does not protect the huge numbers of occupational users engaged in various activities because manufacturers have no power of supervision over these individuals. The question also arises as to how this “agreement” is implemented when chrysotile is exported. Rigorous implementation should have led to a halt in Canadian exports of chrysotile asbestos to France because the French manufacturers had not introduced “made-to-measure” production. In practice, there is clearly no way of ensuring that the Canadian method is actually applied in the importing countries. The EC state that the available scientific data show that despite the “safe” use of chrysotile asbestos, exposure entails fatal disease in a large number of cases. The data from the United Kingdom indicate a significant excess of deaths from lung cancer and mesothelioma, not just among persons employed outside the plants, but also among persons who started their working lives in plants operating a “safe” use policy after that policy was introduced in the United Kingdom in 1969.

3.132 The EC indicate that there are risks associated with “safe” use during the manufacture of asbestos-based products. Natural asbestos fibre, which is a highly resistant material and has exceptional insulation capacities, was used in France for a wide variety of industrial purposes from 1945 onwards. The Canadian concept of the “modern use of asbestos-cement” does not reflect historical reality. For the past 40 years, the production of...
asbestos-cement has consisted of “encapsulating” asbestos in cement, but in the EC’s view this process does not guarantee the safety of the product. When asbestos-cement is used for occupational, para-occupational or domestic purposes, it is usually sanded, crushed or sawn, thus releasing its carcinogenic fibres into the environment in the form of dust. In addition to its use in the asbestos-processing industry (mainly to produce asbestos-cement), asbestos has been used in a wide range of areas, such as construction (flocking and thermal insulation of piping), production of a wide range of consumer goods (floor tiles, brake linings, textiles and boards) and heavy industry (shipbuilding, metallurgy). The EC contend that the available scientific studies demonstrate the limits of “safe” use. The 1996 HSE study carried out in the United Kingdom on the cancer risks incurred by asbestos workers after 1969, when the “safe” use of asbestos was adopted in that country, should be examined in detail. The EC note that a detailed analysis of the study is in fact attached to the 1996 report by the Panel of the Royal Society of Canada, which has been quoted extensively by Canada without acknowledging its source. However, the study indicates that, in spite of strictly “controlled” use – because it focuses solely on workers in the processing industry - there is a significant net excess of mesothelioma in workers who operated solely under conditions of “controlled use”, i.e. after 1969 (when the rules entered into force in the United Kingdom). It follows that “controlled” use does not enable deaths from mesothelioma to be avoided, even in specific manufacturing branches with a limited workforce, which in principle are easy to demarcate and control.

According to the EC, “safe” use is not applicable to servicing and maintenance activities, but mesothelioma fatalities occur mainly among “secondary users” of asbestos. The aforementioned 1995 study by J. Peto shows that 95 per cent of mesothelioma cases in England and Wales occurred in occupations not covered by “controlled use” policies. Also of relevance is the study by M. Siemiatycki, which is never referred to by Canada, even though it was carried out by one of that country’s leading research teams (the one which published the study on women living near asbestos mines). Siemiatycki’s study in Montreal concentrates essentially on workers exposed when dealing with asbestos-containing materials. The study points to a net excess of lung cancer cases and a high risk of mesothelioma associated with chrysotile exposure (risk multiplied by a factor of between 4.4 to 14.6). The worst affected occupations are in the building servicing and maintenance sectors: plumbers, gas fitters, carpenters and electricians are some of the workers most exposed to asbestos. These workers are subjected to exposure peaks that are sometimes very substantially above the current limit values for dust. For example, a roofing worker using a grinder outside to repair corrugated roof sheeting made of asbestos-cement is subjected to a maximum exposure level of 41 f/ml, 410 times in excess of the limit.

The EC contend that, in a legal framework which provides for a total ban on asbestos, the ISO limit continues to be a useful instrument for protecting the health of workers coming into contact with asbestos-containing materials. Such contact is unavoidable, given that millions of tonnes of asbestos were used in France and are still present in many structures. However, merely applying the ISO limit does not contain the risk, and hence “safe” use is not sufficient. The ISO 7337 standard referred to by Canada recommends working methods and tools for cutting asbestos-cement products on-site in such a way as to keep dust emission levels as low as possible. The EC note that, when ISO 7337 was published in 1984, it represented a major step forward in relation to the rules applied prior to that date but it does not guarantee a sufficient level of protection in terms of the health and safety target (maximum limit of 0.1 fibre/ml) adopted by the vast majority of countries. For instance, in the case of a worker using a tool such as a handsaw, application of the ISO standard shows that he is exposed to a level 30 times in excess of the maximum limit of 0.1 f/ml authorized in France and the United States. Cutting an asbestos-cement pipe with a jig saw fitted with a dust collection system (as referred to in this standard) entails exposure of between 7 and 12 times in excess of the maximum limit, whereas manual tools (hammer, chisel) without dust collection entail exposure 20 times in excess of that limit. Although the equipment and rules set out in the ISO standard - which is currently being revised - form the basis of prevention for specialists coming into contact with asbestos-cement materials on-site, they are inappropriate for do-it-yourself enthusiasts or
non-specialists (para-occupational or domestic use). The EC state that there is a high level of mesothelioma among servicing and maintenance workers. The aforementioned study by Y. Iwatsubo\textsuperscript{212}, which was conducted in France on individual cases in the general population (and which covered 405 mesothelioma patients and 389 persons not suffering from the illness between 1987 and 1993) showed a clear increase in the mesothelioma risk among servicing and maintenance workers. Even in Canada, the study\textsuperscript{213} carried out by the Quebec Health and Safety Commission (CSST) showed that the risk of mesothelioma has increased steadily in Canada since 1967, essentially among servicing and maintenance workers. The study points to a particularly rapid growth in the incidence of this illness in the maintenance sector. In cases where exposure had been short-term, servicing and maintenance workers were more numerous than any other category. The study also shows that the occurrence of mesothelioma is caused by chrysotile.

3.135 The EC emphasize that “safe” use in the general population at risk is impossible. Apart from hundreds of thousands of construction and servicing and maintenance workers, other persons run the risk of inhaling asbestos dust. Do-it-yourself enthusiasts are a prime example of a section of the general population at risk. Exposure is “unwitting” in so far as many of the individuals concerned are unaware that their activities may expose them to a risk of breathing in asbestos fibres with fatal consequences. As Canada points out, materials containing asbestos-cement have the same innocuous appearance as most of the materials that building workers and do-it-yourself enthusiasts are used to handling. The EC point out that Canada describes para-occupational and domestic exposure situations in the following terms “Thus, in the eyes of the consumer chrysotile-cement products and fibro-cement products are like products in all aspects, unless a data sheet is available showing which fibres makes up their composition”.\textsuperscript{214} Even when they are aware of the risks associated with asbestos, there is no simple test enabling para-occupational and domestic users to check whether materials not described as dangerous do in fact contain asbestos fibre. Under the current rules, manufacturers are merely required to mark or label the outer packaging of the materials sold.

3.136 The EC conclude that all these data show that it is extremely difficult to pinpoint the risks associated with breathing in asbestos and the use of asbestos products cannot be efficiently controlled. As the HSC notes in its proposed amendments to the rules introduced in the United Kingdom in 1992, efficient control of the use of chrysotile asbestos cannot be guaranteed, even at the stage which in principle is easiest to control, namely production. “Absolute control of the manufacture, and particularly the use of chrysotile asbestos products can never be guaranteed – some people may continue to be exposed, unknowingly, to relatively high levels of fibres during the installation, maintenance or removal/disposal of products containing chrysotile asbestos.”\textsuperscript{215} Controlled use procedures cannot be effectively applied to safeguard the hundreds of thousands of persons who are exposed on a daily basis in industries where health and safety arrangements are minimal, such as the building industry, which accounts for at least 25 per cent of mesotheliomas. The 1998 WHO Environmental Health Criteria 203\textsuperscript{216} state that “it is proven that the risk is probably higher among workers in the building industry and perhaps in other industries where asbestos is used”. Given the extremely large numbers of persons concerned, the difficulty of assessing the risk, the complexity of individual and collective protection systems and their negative effect on dexterity, the need to use special equipment and the overall cost engendered by the requisite arrangements mean that asbestos-containing materials cannot viably be used in a manner that will protect workers’ health effectively. The EC accordingly consider that the principle of “controlled use” cannot be applied to the indeterminate risks incurred over the very wide spectrum of occupations where workers come into contact with asbestos in many ways, in particular in the servicing and maintenance operations for which “safe” use is not a practicable option. Without actually specifying them, Canada acknowledges that “certain uses for which exposure cannot be controlled to an acceptable degree should be banned”.\textsuperscript{217} The EC regret that Canada does not specify these uses nor indicate how frequently they occur. In addition, given the population groups exposed unwittingly, it is unacceptable that materials involving such a high risk should continue to be used in France. The various international organizations dealing with the protection of workers against asbestos are well aware of these facts.
Canada notes that the EC classify exposures to asbestos into three broad categories, namely: (i) occupational exposures of workers; (ii) para-occupational and household exposures; and (iii) environmental exposures. They group occupational exposures of workers into two sub-categories, namely “primary” users (e.g. extraction, manufacturing) and “secondary” users (e.g. construction and maintenance). According to the EC, para-occupational and household exposures mainly involve do-it-yourself enthusiasts. Finally, passive or environmental exposures result from asbestos dust emitted by a natural source of geologic origin, an industrial point source or by asbestos in place in buildings and various facilities.

Canada contends that current uses of chrysotile asbestos do not endanger human health because they are now strictly controlled. These uses include the range of non-friable products where the chrysotile variety alone is used, to the exclusion of the amphibole varieties (crocidolite and amosite), and in which the fibres are firmly bound physically and chemically to the matrix (cement, bitumen, resins, plastics, etc.) of the composite (chrysotile-cement, friction materials, etc.). The French Decree is therefore not necessary to protect health. According to Canada, the EC do not judge the concept of controlled use as proposed by Canada. They judge only one aspect of controlled use, namely responsible use. The EC confuse “controlled use” and “responsible use”. The responsible use of chrysotile, to which they refer, is a voluntary self-regulation initiative of Canadian producers and exporters of chrysotile. Its objective is to sell chrysotile only to businesses which abide by the national regulations required to meet the international standards governing the use of chrysotile. Responsible use is only one component among others of controlled use. Canada emphasizes that, contrary to what the EC endeavours to demonstrate, ironically but quite unconvincingly, controlled use does not require that millions of persons exposed occasionally and unknowingly to emissions of asbestos dust be equipped with elaborate “space suits.” Controlled use is a regulatory approach based on international standards which relies on scientific data applicable to a wide range of hazardous materials. According to the principles on which this approach is based, only products and materials that can be controlled in such a way that the health risks are eliminated throughout their life cycle are permitted. In the case of chrysotile asbestos, controlled use means the application of appropriate regulations aimed at limiting exposures to asbestos by banning certain types of asbestos and certain uses, and prescribing maximum exposure levels as well as work practices and standards.

Canada points out that the controlled use approach is not a Canadian invention. It is based on the principles of controlled use outlined in ILO Convention 162. This provides for protective and preventive measures against health risks due to occupational exposure to asbestos. These measures include: (i) making work in which exposure to asbestos may occur subject to regulations prescribing adequate engineering controls and work practices, including workplace hygiene; (ii) prescribing special rules and procedures, including the authorization of a competent authority in the field, for the use of asbestos or of certain types of asbestos or products containing asbestos or for certain work processes; (iii) where necessary to protect the health of workers and technically practicable, replacement of asbestos by other materials or products evaluated as harmless or less harmful; and (iv) total or partial prohibition of the use of asbestos or of certain types of asbestos or products containing asbestos in certain work processes.

According to Canada, Convention 162 clearly lays emphasis on controlled use and not on banning products. Indeed, the Convention only provides for two specific prohibitions, namely the use of crocidolite and products containing it and the spraying of all forms of asbestos. The general responsibilities of governments, employers’ and workers’ organizations concerned, as well as a national regulatory framework governing safety in the use of asbestos, are also set out in Convention 162. National regulations should: (i) establish procedures for the notification by the employer of certain types of work involving exposure to asbestos; prescribe adequate engineering controls and work practices to prevent or control exposure to asbestos; (ii) enforce laws and regulations through an adequate and appropriate system of inspection, including appropriate penalties; (iii) prescribe limits for the exposure of workers to asbestos and make employers reduce exposure
to as low a level as is reasonably practicable; (iv) measure the concentrations of airborne asbestos dust in workplaces and monitor the exposure of workers to asbestos at intervals; take appropriate measures to prevent pollution of the environment; (v) ensure that employers have established polices and procedures on measures for the education and periodic training of workers on asbestos hazards and methods of prevention and control; (vi) establish standards for respiratory protective equipment and special protective clothing for workers; (vii) recognize contractors qualified to carry out the demolition of plants or structures containing friable asbestos insulation materials, and removal of asbestos from buildings or structures; (viii) ensure that workers who are or have been exposed to asbestos are provided with free medical examinations to supervise their health in relation to the occupational hazard; and (ix) prescribe adequate labelling of containers, including material safety data sheets indicating the asbestos content, the health risks and the appropriate protection measures concerning the materials or the product. Convention 162 is supplemented by the Recommendation concerning Safety in the Use of Asbestos (Recommendation 172), which, in substance, repeats the content of Convention 162 and outlines a set of minimum work standards.223

3.141 Canada notes that the ILO’s Code of Practice on Safety in the Use of Asbestos is intended for all those who are responsible, in the public or private sector, for ensuring the safety of workers when asbestos is used. It is not intended to replace national legislative or regulatory provisions nor applicable standards, but to serve as a guide for the formulation of similar provisions, in particular for authorities and government agencies, specialized agencies as well as businesses and health and safety committees. This Code of Practice aims to: (i) prevent the risk of exposure to asbestos dust at work; (ii) prevent the harmful effects of exposure to asbestos dust on the health of workers; and (iii) indicate reasonable and practically feasible methods and techniques to reduce occupational exposure to asbestos dust to a minimum.224

3.142 Canada indicates that Standard ISO-7337 expressly discourages the use of certain high-speed cutting tools that are not vacuum-equipped and produce excessive quantities of suspended fine dust that can also contain respirable asbestos fibres. France’s member committee approved this Standard.225

3.143 Canada points out that the use of pre-machined parts and fittings, despite the EC’s opinion, is not a far-fetched or utopian solution. According to OSHA: “pre-cut, pre-tapped pipe has received tremendous marketplace acceptance and represents a large majority of sales. [...] This is significant because the use of pre-cut, pre-tapped pipe may reduce or eliminate some types of field fabrication activities.”226 As a result of pre-fabrication, pre-machining, the use of fittings and compliance with work standards, the amount of handling and working of asbestos-cement products is very limited. The ISO has also formulated Standard ISO-14001 - Environmental Management Systems – Specifications and Guidelines for Use.227 The ISO accreditation of a business confirms that it complies with national environmental standards. For instance, if the environmental standards specify a maximum rate of emission of fibres in the air, Standard ISO 14001 is a guarantee of compliance with these standards by the accredited business. In Canada’s view, this is a concrete way of guaranteeing that national standards are observed.

3.144 Canada emphasizes that the practice of controlled use to regulate asbestos, including chrysotile asbestos, is therefore based on well-established international principles. First of all, the permitted uses are based on the varieties of fibre. A clear distinction is made between chrysotile and amphiboles, with certain amphiboles being banned. Secondly, the permitted uses are based on the fixation of the fibres in a binder in such a way that chrysotile asbestos dust cannot be released as a result of normal wear on the product. Almost everywhere, controlled use includes measures such as a ban on processes or products that release or are likely to release dust, for example friable low-density insulation products. Thirdly, the permitted uses are based on control of the average concentrations of asbestos fibres in the workplace. An acceptable limit of 1.0 f/ml or less was recommended by a group of experts that met in 1989 under the auspices of the WHO. The use of dust control equipment combined with the adoption of specific work methods can easily make
it possible to meet this limit in mines, processing plants, as well as at the stage of installation of the product, its repair or removal and disposal of the waste.

3.145 Furthermore, Canada states that controlled use involves the implementation of certain practices and methods of working with materials containing asbestos. To be complete, any regulations based on the principles of controlled use must include appropriate measures and guidelines for the marking and labelling, transport, monitoring of the air, medical monitoring, monitoring of construction activities and waste disposal. The WHO recognizes that controlled use is effective. In its 1998 study entitled *Chrysotile Asbestos*, with respect to the controlled use of chrysotile, the WHO concludes, in the following terms:

“Control measures, including engineering controls and work practices, should be used in circumstances where occupational exposure to chrysotile can occur. Data from countries where control technologies have been applied have demonstrated the feasibility of controlling exposure to levels generally below 0.5 fibres/ml. Personal protection equipment can further reduce individual exposure where engineering controls and work practices prove insufficient.”

3.146 Canada considers that the exclusive use of chrysotile asbestos and the adoption of effective methods to reduce dust are the best guarantees of protection of the health of workers. Canada advocates the controlled use of chrysotile asbestos, relying on scientific research conducted by recognized experts and international organizations, including the WHO. In Canada, the hazardous product regulations apply throughout the life cycle of products. More specifically, all activities involving asbestos – extraction, transport, processing, installation and repair, removal and waste disposal – are closely scrutinized in order to protect not only the health of the public and of workers, but also the environment. In Canada, controlled use has resulted in a ban on friable products containing any type of asbestos in order to protect the public, and in the adoption of measures to ensure the health and safety of workers exposed to asbestos. The latter measures generally fall under provincial jurisdiction and, in Quebec, have been instituted mainly through the Regulations on the quality of the work environment and the Safety Code for the Construction Industry, which are administered by the Commission de la Santé et de la Sécurité au Travail (CSST) (Occupational Safety and Health Board). These two regulatory texts govern working conditions in chrysotile mines, asbestos product manufacturing plants and businesses specializing in the maintenance and removal of sprayed asbestos.

3.147 Canada indicates that, in the early 1990s, Quebec lowered the permitted exposure limit for chrysotile asbestos to 1.0 f/ml, while this limit was set at 0.2 f/ml for crocidolite and amosite. The province has also banned the use of friable insulating materials, as well as crocidolite and amosite. At the same time, Quebec amended its Safety Code for the Construction Industry in order to ensure that workers who have to handle asbestos intermittently on construction sites are adequately protected. Subsection 3.23 applies to all construction sites where work that is likely to release asbestos dust is carried out. The Code establishes a three-tier classification of work: (i) type I jobs - or low-risk activities, i.e. all work with non-friable or high-density products, where the use of hand tools or vacuum-equipped power tools can reduce exposures to well below the exposure limit. Measures must be taken to minimize exposures, prevent the spread of dust and dispose of asbestos waste; (ii) type II jobs - or medium-risk activities - involve minor disturbances or removal of small amounts of friable materials containing asbestos. For example, when removing pipe insulation, workers must wear gloves and special protective clothing, as well as a half-face filter respirator; (iii) type III jobs - or high-risk activities - include large-scale removal of friable asbestos-containing materials, spray application of sealants to asbestos-containing materials, etc., and require very stringent workplace practices. For inside jobs, the ventilation system must be turned off, the work area totally enclosed and maintained under negative pressure, a decontamination centre constructed, and appropriate personal protective equipment issued to workers.

3.148 Canada explains that, before commencing work likely to release asbestos dust, the employer must determine the types of asbestos present in the materials and must also
train and inform workers about the risks, prevention methods and safe work methods. Regulations have also been introduced in Canada to address the disposal of asbestos waste. Provision is made for safe transportation (covered vehicles, sealed packaging, labelling) and disposal of the waste at approved sites. For instance, solid waste containing asbestos is shipped to solid waste disposal or storage sites. In the case of a sanitary landfill site, the solid waste is compacted in uniform layers and covered over. These regulations apply to friable asbestos waste and not to high-density products. At the federal level, again, the Hazardous Products Act prohibits the use of asbestos in the following products or applications: untreated low-quality textile products which can release fibres under normal use; various consumer products, such as toys, modelling compounds and low-density jointing compounds; the sale of loose or raw asbestos to consumers; all asbestos products destined for application by spraying; and finally, the use of crocidolite or crocidolite-containing products.

Regarding labelling, the Workplace Hazardous Materials Information System (WHMIS) was introduced under the Federal Hazardous Products Act. It requires suppliers of all hazardous materials to provide labels with specific pictograms and warning phrases, along with Material Safety Data Sheets, as a condition of sale and importation. WHMIS also requires that workers receive appropriate information and training if called upon to work with or handle such products. It is crucial to measure the emissions of fibres in the air in workplaces, to assess the effectiveness of the control methods and to show that the business complies with the regulations. The measurement of airborne fibres in the workplace is carried out regularly.

3.149 Canada states that, under the Canadian Environmental Protection Act, regulations have been established governing stack emissions from asbestos mines and mills. An emission limit of 2.0 f/ml has been established, in addition to appropriate monitoring methods and administrative controls. Quebec legislative and regulatory provisions prescribe standards for atmospheric emissions of asbestos fibres at all stages of mining operations, including crushing, drying, drilling, storage and processing, as well as conveyors and transfer points, the loading and unloading of asbestos concentrates, not to mention the processing and handling of asbestos waste. In Canada, emphasis is on compliance with requirements. Inspection programmes are in place and strict compliance is monitored so that any necessary legal proceedings can be brought and the maximum penalties imposed.

3.150 According to Canada, the EC rely heavily on a study by the Health and Safety Executive (HSE) entitled “Asbestos-related diseases” to which they refer to show that controlled use is not effective in preventing mesotheliomas and lung cancer among chrysotile-processing workers. This study reveals that workers exposed after 1969 have been affected by exposure to asbestos. In 1969, the standard in England was 2 f/ml, but with an upper exposure limit of 12 f/ml at any time. These levels are much higher than today, in particular the absolute upper limit of 12 f/ml. Secondly, the time required to modernize the obsolete equipment of the period would have to be taken into account. It was only in 1983 that the standard in England was lowered to 1 f/ml in all circumstances. In Canada’s view, the EC wrongly stress the risks of asbestos-related diseases after 10 years of latency in the group, whose exposure to asbestos would have begun after 1969. In fact, in this group, an excess risk of lung cancer, mesothelioma and asbestosis was statistically significant even before the latency period of 10 years. This means that the cohort supposedly exposed to asbestos beginning in 1970 was in fact composed of workers exposed to asbestos prior to 1970 (the explanation advanced by the authors themselves, p.144). Therefore, this is not a genuine cohort of workers exposed for the first time from 1970 onwards. Furthermore, other risk factors (e.g.: smoking rate of 54 per cent in this group versus 42 per cent in the reference population) and other factors not comparable between the cohort and the reference population were already having an impact before the beginning of the exposures to asbestos reported from 1970 onwards. Furthermore, notwithstanding the anomalies and bias cited above, the study cannot constitute a credible test of the Canadian practice of controlled use. In 1969, the new standards constituted only one step toward controlled use; these insufficient improvements were far from equivalent to controlled use as proposed 10 years later. In addition, although England reportedly decided in 1969 to stop importing crocidolite, it did not decrease its imports of amphiboles until 1975 and these imports did not cease until around 1979. Despite all this, there was nonetheless a statisti-
cally significant decrease in the risk of mesothelioma and of other asbestos-related diseases because of a decrease in exposures and perhaps the cessation (barring exceptions) of crocidolite imports.

3.151 Finally, Canada wonders why the EC do not cite the INSERM data and analyses to illustrate the trends in France:

“[Tr.] As a result of the reduction in exposure levels to asbestos in recent decades, practically no further deaths from asbestosis are observed (in the mortality statistics of INSERM in 1990) (INSERM, 1993); 26 deaths from this cause were recorded in France: 24 men and 2 women.”254

If these are the results of the reductions in exposure brought about by only a partial application of the controlled use advocated by Canada, a control similar to that practised in Canada would undoubtedly yield even more positive results in terms of reduction of disease. The EC rely on the study by Siemiatycki to argue that there is a very strong association between mesothelioma and work in the construction sector.235 According to Canada, the EC misinterpret the results of this study. The cancers studied occurred in the early 1980s; they were therefore induced 20-30 years earlier, i.e. in the 1950s. There is absolutely no logical link between these mesotheliomas reported between 1979 and 1985 and the safe use of chrysotile introduced at the beginning of this short period of observation.

3.152 Canada points out that the EC can in no way infer from these results that controlled use does not work. Canada notes that the EC start from the assumption that construction work is by nature uncontrollable. When a study shows an association between mesothelioma and construction or building sub-trades work, they conclude that this represents an association with an uncontrollable exposure. The EC cite a study by Y. Iwatsubo.236 According to Canada, this study does not support a relationship between chrysotile asbestos alone and health effects such as a clear increase in cases of mesothelioma. This study merely reveals what can happen in the absence of a policy of controlled use, two of the main components of which are a ban on amphiboles and control of the average concentrations of chrysotile fibres in the workplace. This study merely reveals the health effects arising from an exposure to mixtures of asbestos fibres that is undefined in terms of concentration and duration.

3.153 Canada considers that the same comments apply to the study by J. Peto on which the EC rely to claim that the bulk of mesotheliomas occur in occupations which fall outside the policies of controlled use.237 Canada cannot agree with the fallacious claim underlying the EC’s argument that controlled use is not feasible with respect to servicing and maintenance workers. A letter from researchers in the Netherlands criticizing the article by J. Peto points out that the higher incidence of mesothelioma in their country has nonetheless stopped rising, even though the controls did not come into effect until 1977 in the Netherlands, i.e. eight years later than in the United Kingdom. These authors propose a hypothesis consistent with Canada’s position:

“[…] the exposure of crocidolite asbestos in construction work may have been more extensive in the UK than the Netherlands. In our country, mesothelioma incidence is clustered in areas with shipbuilding and other heavy industry, where crocidolite was used for insulation […] Although we question the projections of Peto and co-workers, we support their statement that too many people are still being exposed to (crocidolite) asbestos without knowing of it, especially those employed in maintenance and demolition.”238

3.154 Canada notes that the cohort studied by Bégin et al.239, on which the EC base their claim that the risk of mesothelioma has been increasing regularly, extends from 1967 to 1990, and the authors indicate 26 years as the average duration of exposure. Therefore, the cases reported began their exposure between 1941 and 1964. This was obviously not a time when controlled use (low exposure to chrysotile alone) was in effect. According to Canada, the EC’s inference that exposures to chrysotile are responsible for the mesothelio-
mas reported is unjustified and false. In the database of the study by Bégin et al., exposures are usually recorded approximately by categories of use; care must therefore be taken not to make a too specific and consequently abusive interpretation of these data. On the contrary, this study suggests that mesothelioma strikes workers in the secondary and tertiary industrial sectors more than workers in the chrysotile mines and mills. However, several processing businesses in the Montreal area have used amphiboles and particularly crocidolite in the past and construction, sub-trades and insulation workers have been exposed to friable products which very frequently contained amphibole asbestos. Furthermore, the data for Quebec are only reliable since 1984; before this date, the incidence of mesothelioma was considerably under-estimated, resulting in an upward bias in the incidence of mesothelioma during the 1970s and 1980s. Canada considers that the EC’s interpretation of the Bégin study is incorrect. The data from this study corroborate the existence of a risk of mesothelioma in occupations in the secondary and tertiary sectors more exposed to amphiboles, but they cannot in any way specifically “incriminate” chrysotile and its current uses.240

3.155 Canada contends that controlled use in the extraction and processing industries is feasible. The EC have admitted that the control of dust emissions in extraction and processing plants, as well as in plants producing chrysotile-based materials, is effective in controlling the risk of contracting a disease attributable to exposure to asbestos.241 Canada points out that the EC have backed away from their position in this dispute, but this position is quite simply unfounded. In most countries, the processing of chrysotile fibre takes place in highly automated plants, in which very strict control and health measures are applied. These measures include proper work and maintenance methods, appropriate ventilation and dust control systems, as well as information and training for workers. According to the WHO, there is no risk in the extraction and processing industry. “The overall relative risks for lung cancer are generally not elevated in the studies of workers in asbestos-cement production and in some cohorts of asbestos-cement production workers.”242 With respect to emissions from crushing in mills, it adds: “In well-controlled mills, this is largely confined in the mill building, and presents low emissions because the mill air is collected and ducted through control devices.”243

3.156 Canada notes that statistics exist indicating that controlled use is effective. The data collected and compiled by the Asbestos International Association in 1995 cover 28 countries where approximately 25,000 workers are employed. In these countries, 87 to 99 per cent of the workers are exposed to levels of less than 1.0 f/ml.244 In Quebec, the exposure values at raw material extraction sites and in plants which separate chrysotile asbestos fibres from the raw material have declined by an average of 16 f/ml to less than 1 f/ml following the application of control measures, such as technical processes, since 1973.245 Canada contends that controlled use is also effective in the processing industry. Technical processes can be adopted in chrysotile-cement plants. These processes involve, inter alia, dust emission control and confinement. In chrysotile-cement plants, several operations, such as the internal machining of fittings for example, can be performed in such negative pressure enclosures.

3.157 According to Canada, it is essential to stress the latency period in order to assess the effects attributed to exposure to asbestos. The effects reported today reflect the circumstances of uncontrolled exposure which prevailed more than 20 or 30 years ago, and absolutely cannot be used as evidence that effective control (low exposure) of current uses of chrysotile asbestos (excluding amphiboles) is impracticable and solves nothing. G. Berry and M.L. Newhouse conducted a mortality study (1942-1980) in a plant producing friction materials where chrysotile asbestos was used almost exclusively. Compared to national mortality rates, the authors did not observe any excess mortality due to lung cancer, cancer of the digestive tract or any other cancer. The exposure rates were low, and only 5 per cent of the workers had accumulated 100 fibres-years/ml. The authors stated that the experience accumulated at this plant over a 40-year period showed that chrysotile was not associated with any apparent excess mortality.246 The extension of the study over a period of seven years enabled the authors to confirm that there was no excess mortality due to lung cancer or other tumours associated with asbestos or to chronic pulmonary diseases. After
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1950, air quality control measures were gradually improved in this plant, and since 1970, the concentrations of asbestos have not exceeded 1.0 f/ml. The authors conclude as follows: with proper control measures of the ambient air, it is possible to use chrysotile in the manufacturing industry without causing excess mortality.247

3.158 Canada also refers to a study by J. Peto: “Only a few (5 per cent) of British mesothelioma deaths were among workers in regulated occupations.”248 When the latency period is considered, these deaths are due to the regulations in the 1960s. The authors of another study followed 1,970 workers at an asbestos-cement plant where chrysotile was used exclusively and examined their mortality data. They did not note any significant elevation of the standardized mortality ratio for the causes of death examined, i.e. all causes, all cancers, lung cancer and pleural cancer and cancers of the digestive tract. The authors conclude that the general findings of this mortality study suggest that the workers at the chrysotile asbestos-cement plant studied did not face an increased risk with respect to total mortality, mortality due to all forms of cancer, lung cancer and bronchial cancer or to cancers of the digestive tract.249 In another study of 5,645 workers at an asbestos-cement plant, in which the authors did not find any increase in mortality resulting from exposure to chrysotile asbestos over a 20-year period at exposure levels equal to or less than 100 mppcf x years (which corresponds to approximately 15 f/ml x years). The authors state that the demonstration that cumulative low and short-duration exposures did not result in a detectable excessive risk of respiratory cancer could serve as a basis for the development of regulations, because these data make it possible to state that there are low degrees of exposure which are not associated with a demonstrable increase in risk.250

3.159 Canada mentions a cohort study of 1,176 workers at a Swedish plant producing asbestos-cement containing chrysotile, which did not show any excess mortality associated with exposures of approximately 10 to 20 fibres/ml x years.251 In another cohort study, this time of a group of 2,167 workers between 1941 and 1983, the authors did not observe any excess incidence of lung cancer or any other excess mortality associated with asbestos at average fibre concentrations of less than 1 f/ml, although the concentrations were probably higher in certain areas of the asbestos-cement plant.252 Finally, the McDonald study examined what is undoubtedly the largest cohort of asbestos workers ever studied and followed over the longest period, namely the cohort of miners and plant workers at chrysotile mines in Quebec. This study was initiated in 1966 and includes 11,000 workers born between 1891 and 1920 who have been followed since its inception. The researchers made optimal use of all the available measurements of dust concentrations to assess the exposure of each member of the cohort in terms of duration, intensity and period during which this exposure occurred. The mortality data have been published on five occasions, and there is a recent report updating the results of the mortality analysis for the period 1976-1988 period inclusive. One of the main findings of this update is that for several narrow exposure categories of up to 300 mppcf x years, the standardized mortality ratio for lung cancer was around one, with no perceptible trend, and increased significantly above this level of exposure.253 More recently, the same authors again updated their results, this time on a group of 9,780 men followed until 1992. The results for exposures of less than 300 mppcf x years, which is equivalent to about 900 f/ml x years – or 45 f/ml for 20 years – prompt the authors to conclude that, from the standpoint of mortality, exposure in this industry to less than 300 mppcf x years has been essentially harmless.254 With respect to the current exposure levels to chrysotile permitted or recommended and the hesitations one might have to convert mppcf to f/ml, even by applying a prudent conversion factor of 1 mppcf = 3 f/ml, the studies cited previously, including the recent updates, constitute a strong argument in favour of the recommendation by the WHO group of experts of an exposure limit value of 1 f/ml for chrysotile asbestos.

3.160 Canada claims that controlled use is effective on construction sites. However, this in no way alters the fact that there are large quantities of amphiboles and friable materials on construction sites. Certain precautions must be taken during the installation of slabs on construction sites and the burying of chrysotile-cement pipes. When, despite pre-machining and the use of fittings, it is necessary to handle and work with asbestos-cement materials, precautions must be taken. On the other hand, studies have shown that expo-
sure levels can be maintained well below 1 f/ml if proper work methods are followed. Similarly, according to Canada, the application of simple work methods that do not generate dust will protect employees during the removal of chrysotile-cement. According to the WHO, it is friable materials that pose a problem: “Some asbestos-containing products pose particular concerns and chrysotile use in these circumstances is not recommended. These uses include friable products with high exposure potential”. However, Canada notes, the controlled use which the EC claim is too difficult is associated with the use of friable materials, not with the current uses of chrysotile.

3.161 Canada states that the typical exposure levels for various operations performed with low-speed hand tools during the installation of chrysotile-cement products are essentially below 0.1 f/ml and differ significantly from those the EC mention with respect to the application of Standard ISO-337, which they severely criticize. They consider this standard inadequate to guarantee a sufficient level of protection in light of the health objective (limit of 0.1 f/ml) adopted by France. Firstly, based on the rates of dust emission mentioned above, the exposures in many operations remain below the limit of 0.1 f/ml. Secondly, the limit adopted by some countries is lower than that proposed by the WHO. While it is the prerogative of each country to adopt the values it chooses, it does not follow that the higher rates of asbestos dust produced by using certain tools create circumstances of exposure in which workers do not benefit from adequate protection. Also, in cases where exposure rates are higher than the limit of 0.1 f/ml, these rates nonetheless remain well below the value of 1 f/ml suggested by the WHO to guarantee a sufficient level of health protection of workers. In conclusion, Canada points out that, after a presentation of the exposure data concerning the application of Standard ISO-7337, which gives rise to some confusion, the EC recognize that the equipment and the rules outlined in Standard ISO-7337 “form the basis of prevention for specialist workers coming into contact with asbestos-cement materials on-site […].”

3.162Canada does not agree with the EC’s claim that France cannot protect servicing and maintenance workers from exposure to asbestos. This category includes building carpenters, plumbers and electricians. This claim ignores the fact that it is past uses, particularly sprayed-on applications of asbestos (notably amphiboles), that are the cause of the health problems attributable to asbestos in France. That is the conclusion of several epidemiologists, including Dr. Camus, in whose view:

“In the industrialized countries, friable asbestos products inherited from the past represent a potential for exposure and risk today and in the decades to come. In comparison, the potential for exposure to new chrysotile products composed of rigid matrices (mainly cement) seems to be very low and much easier to control. Prevention efforts must therefore focus on the actual risk of asbestos products in place.”256 (WTO translation)

3.163According to the WHO in 1998:

“Data from industries where control technologies have been applied demonstrated the feasibility of controlling exposure levels generally below 0.5 f/ml. Personal protective equipment can further reduce individual exposure where engineering controls and work practices prove insufficient.”257

3.164Canada shares the view that construction workers who work with the parts of buildings containing sprayed-on asbestos are likely to be exposed to asbestos. However, the ban on chrysotile asbestos and the Decree do not alter this situation in any way. The exposure of these workers to asbestos is the consequence of past products and uses, which are prohibited today. The circumstances of workers who work with the parts of buildings containing sprayed-on asbestos cannot be compared in any way with the conditions of servicing and maintenance workers who handle asbestos-cement products. Canada considers that the EC and France cannot use studies or findings concerning the former to justify measures intended to protect the latter.
With regard to controlled use and do-it-yourself enthusiasts, Canada considers France’s concerns to be unfounded for four reasons. First of all, Canada wishes to point out that the EC have not established that intermittent exposures to chrysotile fibres, even at high concentrations, would create a detectable health risk. Secondly, this concern is based on a false perception, that the cutting of high-density non-friable materials containing chrysotile releases large quantities of chrysotile fibres. In fact, even using tools that are not recommended by the guides on safety in the use of asbestos, such as high-speed power tools, the dust emitted contains only a tiny quantity of respirable chrysotile fibres. Thirdly, most of the chrysotile fibres that are released following the use of high-speed saws have been chemically altered. They have a different form in terms of structure and chemical composition. For this reason, they are less hazardous. The same is true of the dust released following the abrasion of chrysotile-based resin or plastic products: this dust contains tiny quantities of chrysotile fibres. Fourthly, cutting any cement material using inappropriate tools such as a high-speed saw releases silica and quartz. Silica is a group 1 type carcinogen according to the IARC classification. The risk incurred by a person who cuts a cement pipe depends on the tool used (high-speed power tool for example) and not on the presence of chrysotile fibres in the cement. Whether or not chrysotile is present, the person should wear an appropriate mask or use the appropriate tools such as hand tools or low-speed power tools producing coarse dust or chips.

Canada considers that the codes of practice intended for all those who are responsible for ensuring the safety of workers when chrysotile asbestos is used and recommended by the ILO or the ISO also support the position that the handling of products in which chrysotile fibres are solidly set in a binder does not expose workers or do-it-yourself enthusiasts to asbestos dust which presents a detectable risk to their health. Unlike professionals, private individuals are likely to work with chrysotile-cement products only very sporadically. According to Canada, the exposures received by these weekend do-it-yourself enthusiasts will only be a fraction of those received by professionals. Consequently, if professionals who work with chrysotile-cement products on a daily basis are not subject to any detectable risk, then logically the risk for private individuals will be even less. Private individuals will generally not carry out heavy operations such as sawing, sanding or demolition of materials. Do-it-yourself enthusiasts will rather perform light work occasionally, for example, drilling a hole to run a cable. Finally, if he works with cement products, whether containing chrysotile or not, a do-it-yourself enthusiast will have to follow simple protection methods, if only because of the presence of other carcinogenic materials such as crystalline silica.

Canada disagrees with the EC’s assertion that the overall costs of the measures necessary to ensure effective control of the uses of asbestos are such that the use of asbestos-based materials is not viable. Most of the measures necessary to implement effective control of applications of asbestos do not entail exorbitant costs. Even if the measures required to control the applications of asbestos should in practice prove too costly, Canada maintains that the economic agents should then be allowed to make the decision to choose the controlled use of chrysotile or the use of substitute fibres or products.

In conclusion, Canada states that effective protection measures for extraction and processing workers have existed since the 1970s. The methods used for this purpose involve relatively simple technologies, such as increased ventilation of work areas, more effective filtration of air containing dust, grinding and processing operations under negative pressure to prevent dust from escaping, exhaust hoods in the workstations directly exposed to fibres, wet-process manufacturing techniques and the mechanization of processes. Workers in the construction, servicing and maintenance sectors are occasionally exposed to friable asbestos products (of all varieties) in the course of their occupational activities. It should be recalled, however, that friable products are not the subject of the present debate. Workers exposed to peaks have a lifetime risk of approximately 20 to 300 per million, which is an “undetectable” risk, i.e. a risk which cannot be demonstrated or measured empirically. Canada maintains that the controlled use of chrysotile asbestos and of current products is possible for all occupations, even those where workers may
occasionally be exposed. It is a matter of enforcing an appropriate framework of use, in the same way as for numerous other hazardous substances used in the workplace.

3.169 The European Communities assert that the so-called “safe” use is ineffective and inapplicable. Attempts to make “safe” use of asbestos concern only a small number of jobs. Canada’s attempt to establish a “safe use” policy is based on “at-source” controls consisting, in particular, of an agreement between the producer-exporters of chrysotile asbestos and the manufacturers of products or materials containing chrysotile. This agreement provides for the transmission of information on the risk and the means of protection to be used by producer-exporters of raw asbestos. It also provides for an undertaking on the part of the manufacturers to inform their employees of the risks associated with inhaling asbestos and to provide the indispensable collective and individual protection equipment. This “understanding” covers only the relatively small asbestos manufacturing sector and presupposes the voluntary participation of the industry. Subsequently, the asbestos-containing products and materials are widely distributed among a very extensive range of users for installation in buildings, industrial plants and vehicles. The installers are professionals assumed to be aware of the risks or occasional professional or non-professional users who may be totally unaware of the risks. Any warnings about the presence of asbestos and the risks associated with using the product (labelling, notices, etc.) have disappeared by this stage, in which the products and materials have already been unpacked and installed (for example, a painted sheet of asbestos-cement in a building or a gasket in an engine).

3.170 According to the EC, “safe use” cannot be applied in the servicing and maintenance sectors. During the entire life cycle of an asbestos-containing product or material in a building, factory or vehicle, servicing and maintenance will be required and the workers are liable to be seriously exposed, throughout their working life, to risks of which they are unaware. Building and industrial maintenance workers constitute a group with high exposure to the risk of mesothelioma and lung cancer. In France, there are several hundred thousand of these workers spread over a wide variety of sectors (metallurgy, construction, vehicle repair, etc.), whereas before asbestos was banned the workers employed by enterprises engaged in the primary processing of raw asbestos into materials - those supposed to be sufficiently well informed to be able to control the risk - numbered less than 1,500. Servicing and maintenance workers may be frequently and unwittingly exposed to peaks of exposure several hundred times higher than the exposure limit established in France, as in many other countries: dry-cutting asbestos-cement generates 41 fibres/ml, or 410 times the limit value.

3.171 The EC point out that, as Canada indicates, the ISO 7337 standard proposes preventive measures (moistening of the material or dust extraction) and recommends the use of hand tools or low-speed tools with a view to limiting fibre emission. These practices are also recommended by the public health authorities in many other countries to limit the exposure of those required to work on-site with materials known to contain asbestos. The large volume of metrological data recorded at work stations at which such equipment is used, including on asbestos-cement, shows that the recommendations of ISO 7337 are inadequate and that the exposure threshold, which itself does not provide absolute protection, is almost always exceeded, the levels being 30 times the limit value for a handsaw and five to 12 times the limit value for a jig saw with dust extraction. Numerous data documenting this finding can be found in the international scientific literature. In most servicing and maintenance situations, because of time and ergonomic constraints, the specialized tools recommended by ISO 7337, which are often heavy and take a long time to set up, are not used (for example, when a hole is to be cut in an asbestos-cement roof, the tool might have to be connected to a cumbersome dust extraction system or a remote water supply network). Thus, the rules of “safe use” cannot be effectively applied on the scale of the hundreds of thousands of people daily exposed to asbestos in sectors as difficult to regulate from the standpoint of occupational health as the construction sector, which generates at least 25 per cent of the cases of mesothelioma.

3.172 The EC point out that all the scientific data available show that the “safe” use of asbestos cannot prevent large numbers of deaths from mesothelioma or lung cancer, nei-
ther in the confined sector of asbestos-containing material production nor in the highly dispersed sectors of construction, servicing and maintenance, and demolition. In particular, the Health and Safety Executive (HSE) study, carried out in the United Kingdom on a population of asbestos-processing workers rigorously subjected to the rules of “safe use” since 1969, shows that the application of these rules cannot prevent a significant excess of mesotheliomas, even in a sector seemingly easy to manage and supervise (involving fixed work stations in a factory environment).

The studies already mentioned by J. Peto\textsuperscript{262}, Y. Iwatsubo\textsuperscript{263} and the CSST\textsuperscript{265} (Commission de la Sécurité et de la Santé au Travail) in Quebec stress the seriousness of the risk for servicing and maintenance workers and indicate an abnormally high level of mesotheliomas in these occupations. The Canadian study shows that in Canada the risk of mesothelioma has been increasing steadily since 1967 and that the increase in the incidence of this always fatal disease has been particularly rapid in the servicing and maintenance sector. Thus, the limitations of all the various attempts to make “safe” use of asbestos have now been revealed and the serious public health failures to which they have led can only be checked by completely halting the spread of the risk.

6. Substitute fibres for chrysotile

\textbf{Canada} points out that, due to the versatility of chrysotile, there exist more than 150 substitute fibres. These substitutes reproduce, for certain specific uses, the thermal resistance, reinforcement capability, chemical resistance and even the acoustical and thermal insulating properties of chrysotile. The most common substitute fibres are aramid fibres, polyvinyl alcohol (PVA) fibres, cellulose fibres, glass fibres, ceramic fibres, rock wool and wollastonite.\textsuperscript{266} The prohibition of chrysotile and the conversion to substitute fibres and fibro-cements have caused an increase in the use of substitute fibres. The conversion took place, however, without sufficient risk analysis. The still undetectable risk associated with modern uses of chrysotile has been replaced by the unknown risks associated with the use of substitute products. Moreover, the use of substitute fibres contributes to an increase in risks associated with products containing substitute fibres that are of lower quality in terms of their physical, chemical and mechanical resistance.\textsuperscript{267} The decision to ban chrysotile was announced the day after the publication of the summary of the INSERM report. In this report, however, the authors admitted that they did not investigate the risks posed by substitute fibres and concluded:

“Although, because of the time limit given to it, the group did not wish to take up the question of substitute fibres, the lack of epidemiological data concerning their harmlessness in the long term cannot hide the findings reached in experimental systems, which show the capacity to cause pathological modifications. Appropriate research should be carried out and developed as a matter of urgency before use of substitute fibres becomes general.”\textsuperscript{268} (WTO translation)

\textbf{Canada} claims that there has been a total conversion to substitute fibres, despite INSERM’s formal warning. It was not until June 1998, one and a half years after the adoption of the Decree, that the Executive Summary – and not the full report – of another INSERM study, on substitute fibres, was published. In the Executive Summary, it is acknowledged that:

“Fibres on which very few toxicological data exist are today being used … on a large scale as a replacement for asbestos; the novelty of their use for such applications is accompanied by a lack of data on their potential effects on human health.”\textsuperscript{269} (WTO translation)

From the beginning, the Executive Summary admits that great vigilance is in order with respect to the possible effects of using substitute fibres:

“Any new fibre proposed as a substitute for asbestos or for any other use should \textit{a priori} be suspected of being pathogenic because of its structure, although this does not prevent an analysis of the possible effects of its physico-chemical characteristics.”\textsuperscript{270} (WTO translation)
3.176 INSERM also stresses that its study is limited to the risks of substitute fibres in the respiratory system, even though the health risks on the whole cannot be confined to respiratory ailments:

“This approach is restrictive in the sense that it focuses in the first place on respiratory diseases. It can be seen that dermatosis caused by fibres affects a large number of workers in contact with such materials … It might be suggested that it would be preferable not to confine studies to the respiratory system, particularly because of the development of vitrous fibres that are soluble in a biological environment. Solubilized products may, sometimes, reach other organs.” (WTO translation)

3.177 Canada points out that the Executive Summary also highlights the differences in the experimental exposure levels used to evaluate the risks of substitute fibres compared to those for chrysotile fibres, even though the modern uses of substitute fibres are similar to those of chrysotile fibres.

“It has been noted that, in general, the animals have been exposed to a number of substitute fibres that is much lower than the level used during experimental exposures to asbestos. It is likely that similar concentrations of asbestos fibre would have yielded results that were of little or no significance in studies of carcinogenicity.” (WTO translation)

3.178 Canada indicates that Directorate General XXIV (consumer protection policies) of the Commission of the EC stated that “for substitute materials (for chrysotile), there is no significant epidemiology base to judge the human health risks (...) hence the conclusion that specific substitute materials pose a substantially lower risk to human health, particularly public health, than the current use of chrysotile, is not well founded.” Dr. J. M. G. Davis’ paper, The Biological Effects of Fibres Proposed as Substitutes for Chrysotile Asbestos: Current State of Knowledge in 1998, conducts a review of the scientific literature on substitute fibres for asbestos and concludes that:

“Replacement is premature in the present state of our knowledge.” [...] The need for full toxicological testing of new fibre products is recommended before these products are marketed. [...] It is of much concern to find that of the three fibre types suggested as chrysotile substitutes, polyvinyl alcohol, para-aramid and cellulose, only one type of aramid (Kevlar) has been tested in a way at all close to what is required.”

3.179 Canada considers that France has replaced the undetectable risk associated with modern uses of chrysotile with the unknown, and perhaps greater, risk associated with the use of substitute fibres for asbestos.

3.180 The European Communities note that the majority of substitute products for asbestos, particularly chrysotile asbestos, are non-fibrous products, for example, PVC. Canada never refers to these products, which competed with asbestos-cement long before the ban. In France, for example, the decision to stop production of asbestos-cement pipes was taken by industrialists before the announcement of the ban because the competition from PVC and ductile iron made this sector non-viable from the economic perspective. The main purpose of the policy adopted in France in 1996 is to replace materials containing asbestos by other harmless materials, particularly non-fibrous materials, for example, by replacing asbestos-cement roofing by galvanized steel or aluminium sheets of an equivalent size or larger, replacing asbestos-cement pipes by ductile iron or PVC (plastic) pipes. According to the EC, fibres to replace asbestos can only be used in certain materials if their harmless-ness or their lower level of risk in comparison with asbestos has been irreputably proved by the international scientific community. The risk assessments proposed by INSERM conclude that, for the various fibres analysed, the risk of cancer is sometimes “plausible” and sometimes “cannot be excluded”, sometimes “there is no convincing element indicating an excess risk” (WTO translation), this is far from the certainties concerning chrysotile,
a proven carcinogen according to all the international authorities. It can also be emphasized that none of the substitute products for asbestos is classed in group 1 of the IARC (proven carcinogen for humans).

3.181 The EC indicate that aramid fibres, polyvinyl alcohol (PVA) fibres, cellulose fibres, glass fibres, ceramic fibres, rock wool and wollastonite are only used as an asbestos substitute in very limited cases where it is not possible to find a non-fibrous substitute. For asbestos-cement materials, which accounted for more than 90 per cent of asbestos use in France, industrialists now use cellulose fibres (the same as those always used in the paper industry), polyvinyl alcohol and aramid fibres, for which the available toxicological data are quite reassuring. According to the EC, it is wrong to state that "The conversion [from asbestos to substitute fibres] took place without sufficient risk analysis". None of the substitute products for chrysotile has been classified as carcinogenic for humans at the international level. Moreover, the replacement of asbestos by less harmful substitute fibres has been done in stages and exercising the utmost caution concerning their harmfulness. The French text provides for the possibility of requesting waivers from the ban on asbestos whenever there is no technically comparable substitute that is less dangerous than asbestos.

3.182 The EC point out that the task entrusted to INSERM by the French Government in 1995 was first to assess the state of scientific knowledge on the risks and dangers of asbestos. Canada indicates that only the Executive Summary of the INSERM report on substitute fibres for asbestos was published in June 1998. This statement is wrong. The full text, in the form of a provisional working document, was published at the same time as the Executive Summary, as was done for the report on asbestos in July 1996. In order to provide transparency and ensure that the findings - which are necessarily succinct in a summary document - are based on detailed scientific arguments, it is INSERM’s policy to provide access to the full text in provisional form (because printing a report in accordance with the requirements of scientific publications necessarily takes a very long time).

3.183 The EC point out that replacement is a necessity acknowledged internationally. In 1986, ILO Convention 162 on asbestos recommended that national legislators should make provision, whenever possible, for the "replacement of asbestos or of certain types of asbestos or products containing asbestos by other materials or products or the use of alternative technology, scientifically evaluated by the competent authority as harmless or less harmful …" (Article 10(a)). The WHO press release of 26 July 1996 also states that it is necessary to effect such replacement, including chrysotile. The WHO states that "consideration should be given to replacing chrysotile by harmless substitute materials whenever it is possible to do so". In 1998, the WHO report drawn up as part of the International Programme on Chemical Safety recommended the use of substitute materials where they are available. Since 1990, in its framework Directive on the health and safety of workers, the European Community has advocated the principle that a dangerous agent or procedure should be replaced by a harmless or less harmful agent or procedure in so far as one exists. In the case of cancer risks, the principle is set out in the Directive on carcinogenic agents, which specifically states that a carcinogen should be replaced by a less dangerous agent in so far as this is technically possible. The EC note that the French Decree contested by Canada in the present case complies with the recommendations of the ILO and the WHO. Substitute products can only be used to replace asbestos if they entail a lower risk for workers according to current scientific knowledge. In view of the proven carcinogenicity of chrysotile and the impossibility of controlling the risk, the French Government opted to prevent any proliferation of the risk by applying the principle recommended by the WHO and the ILO and endorsed by the European Union for carcinogenic risks: replacement by a less dangerous product in so far as this is technically possible.

3.184 The EC point out that the wide variety of substitute products reflects the diverse possible uses of asbestos. There is no natural or synthetic product with the same set of properties as asbestos. Accordingly, there is no one substitute for asbestos, but various substitutes which are sometimes combined with each other for certain applications. When referring to "substitute products", Canada appears to be referring solely to mineral fibres,
whereas in fact a very wide range of products and materials is used to replace asbestos. It
goes without saying that the materials obtained after asbestos is replaced have different
physical and mechanical properties from the asbestos-containing material. As a result,
each substitution operation entails careful checks by manufacturers on the properties of
the new material and, sometimes, a complete redefinition of the product’s field of application
(watertight gaskets and braiding, for example).

3.185 The EC explain that these substitute products can be of three types. Firstly, prod-
ucts which do not use fibres: (i) replacement of asbestos-cement pipes by ductile iron or
PVC (plastic) pipes, which were on the market long before France banned asbestos; and
(ii) replacement of asbestos-cement roofing by galvanized steel or aluminium sheets of an
equivalent size or larger. Secondly, products using fibres shown to be harmless or less
harmful: (i) replacement of asbestos in cement-fibre materials by cellulose, PVA or aramid
fibres (asbestos-cement accounted for more than 90 per cent of asbestos use in France);
and (ii) replacement of asbestos in friction surfaces (brakes, clutches) by metal (copper),
aramid fibres or granular materials. Thirdly, products containing less harmful man-made
mineral fibres: (i) replacement of asbestos in watertight braiding (textile-asbestos) by glass
fibres or rock fibres in most cases, or by ceramic fibres, which are essentially used as a
substitute for amphibole asbestos and for very rare uses (above 1,200°C). The cost of using
these fibres is considerably higher than that of using asbestos (50 per cent more).

3.186 The EC point out that none of the substitute products for chrysotile is classified as
carcinogenic for humans. The lower carcinogenicity of the fibres regarded as most danger-
ous was already known in France when the decision was taken to ban asbestos. In the
course of discussions on the eventualitly of a ban, the Ministry of Labour asked the Groupe
sur la surveillance des atmosphères de travail (G2SAT), which meets under the aegis of the
French Senior Council for the Prevention of Occupational Risks, to draw up a report using
existing international data on the comparative carcinogenicity of asbestos and man-made
mineral fibres. The report284, which was presented in June 1996, indicates a clear hierarchy
in the relative toxicity of substitute fibres. The results were confirmed by a 1997 survey on
synthetic mineral fibres carried out by INSERM. None of the man-made mineral fibres
investigated showed proven carcinogenicity for humans. According to this report, glass
wool and slag wool do not cause fibrosis or cancers in animals; rock wool only causes
fibrosis in animals and then only at very high levels of exposure. These fibres are used
only for certain rare applications (watertight braiding as a substitute for textile asbestos);
ceramic fibres cause cancers and fibroses in animals and have been diagnosed as the cause
of pleural plaques in humans. These fibres are used mainly as a substitute for amphibole
asbestos in very restricted and highly specific cases (temperatures in excess of 1,200 de-
grees). The European Community Directive285 relating to the classification of man-made
mineral fibres placed mineral wool in category III (unproven suspicion of carcinogenicity
in animals) and ceramic fibres in category II (proven risk for animals). The EC point out
that ceramic fibres essentially provide a substitute for amphibole asbestos and their use is
strictly regulated (Labour Code, Articles R-231-55 et seq.). None of the substitute products
for chrysotile is classified as a “proven carcinogen for humans” (group 1). Since it is known that
chrysotile asbestos is classified as a “proven carcinogen for humans” (group 1), according
to the EC, there is therefore a clear hierarchy of risk.

3.187 The EC emphasize that there are no data giving rise to concern as to the
carcinogenicity of fibres used as a substitute for asbestos in cement fibres. The findings of
the Scientific Committee on Toxicity, Ecotoxicity and the Environment are unambiguous
in this connection. The SCTEE was set up by Commission Decision 97/579/EC of 23 July
1997 setting up scientific committees in the field of consumer health and food safety (Offi-
cial Journal L 237 of 28 August 1997, p.18). The SCTEE’s brief is to provide the Commis-
sion with high-quality scientific opinions in all cases in which it has to be consulted. It may
also be consulted on other issues of special relevance to consumer health and food safety.
Its composition and work are based on the principles of excellence, independence and
transparency. The SCTEE has examined the comparative carcinogenicity of chrysotile and
the main fibres used as a substitute for asbestos in cement fibres (which accounted for
more than 90 per cent of the asbestos used when the ban came into force in France). The
investigation focused on cellulose fibres, para-aramid fibres and PVA.
3.188 The EC indicate that the SCTEE’s findings are unambiguous as to the hierarchy of risks. Its conclusions are as follows: (i) the SCTEE notes that the three fibres investigated have been used for many years and yet no worrying data have emerged over this long period as to their carcinogenicity; (ii) there is no doubt whatsoever that chrysotile is carcinogenic; (iii) chrysotile also causes asbestosis and chronic illnesses, which is not the case for the fibres investigated; (iv) chrysotile has carcinogenic effects on the lung and causes mesothelioma, whereas there is no evidence to suggest that para-aramids do so (no long-term research is available for the other two fibres, but it should be recalled that they have been in use for many years); (v) predictive studies - i.e. experimental studies on animals - show that chrysotile has far more substantial and lasting inflammatory and proliferative reactions; and (vi) the substitute fibres are considerably larger (in diameter) and have much less fragmentation capacity than chrysotile, which has very high “fibrillation” (separation of fibres lengthways into even finer fibres). The result is that the substitute fibres are far less respirable. Three interdependent parameters were used to assess the risk associated with substitute products as compared with chrysotile asbestos: (i) the dose of fibres in the air: the fibre’s diameter determines how long it remains suspended in the air; accordingly, the smaller the diameter, the higher the carcinogenic risk; (ii) the physical characteristics of fibres, in particular their dimension (diameter and length) and their fragmentation capacity (“fibrillation”); the greater the degree of fibrillation, the higher the risk of the fibres being inhaled, and thus the higher the risk of cancer; and (iii) the biopersistence (or “durability”) of the fibre in the lung tissue and inside the macrophages, given that long fibres are eliminated slowly. It is accepted that a ratio of length to diameter of more than 3:1 entails a higher risk of cancer. The SCTEE’s study thus concludes that chrysotile asbestos is present in high concentrations in the air because of its small diameter and is very easy to inhale because of its high degree of fibrillation. The substitute fibres for asbestos-cement are PVA and cellulose fibres. Para-aramids are used for other applications (brakes, friction). These three types of substitute fibre are the most frequently used.

### Comparative Table of the Characteristics of Fibres Examined by the SCTEE

<table>
<thead>
<tr>
<th></th>
<th>Length (microns)</th>
<th>Diameter (microns)</th>
<th>Fibrillation</th>
</tr>
</thead>
<tbody>
<tr>
<td>Chrysotile</td>
<td>&gt; 5</td>
<td>&lt; 1</td>
<td>+++</td>
</tr>
<tr>
<td>PVA</td>
<td>&gt; 5</td>
<td>10 – 16</td>
<td>+/-</td>
</tr>
<tr>
<td>Para-aramids</td>
<td>&gt; 5</td>
<td>10 – 12</td>
<td>(requires considerable abrasion to produce a large number of fibrils)</td>
</tr>
<tr>
<td>Cellulose</td>
<td>&gt; 5</td>
<td>12 – 40</td>
<td>Exposure data suggest that fibril production is very limited</td>
</tr>
</tbody>
</table>

3.189 The EC note that, from these observations, the SCTEE unanimously concluded that: (i) there is sufficient evidence to show that all forms of asbestos, including chrysotile, are carcinogenic; (ii) on the other hand, there is no evidence of cancer being caused in humans by fibres of any of the three substitute products examined; and (iii) consequently the dangers of chrysotile are probably greater (in the scientific sense) than those associated with the substitute fibres examined. The SCTEE adds that, in its opinion, no study which might have led it to a different conclusion was omitted from the examination.

3.190 The EC also note that the findings of the Committee on the Carcinogenicity of Chemicals in Food, Consumer Products and the Environment (COC) transmitted to the Health and Safety Executive (HSE) in July 1998 confirm the results of the SCTEE study. This Committee also examined cellulose fibres, para-aramid fibres and PVA fibres. In particular, the COC’s work focused on comparisons between the results of carcinogenicity studies and studies on the physical properties of fibres (dimensions, fibrillation). Its conclusions were identical to those of the European Committee. In addition, the COC noted that the levels of exposure to those fibres were considerably lower than those associated with the use of chrysotile. The COC concluded its study as follows:
“The evidence presented to the Committee on fibre dimensions, studies in animals including that of biopersistence in the lung, indicate that the carcinogenic risk posed by PVA fibres, para-aramid fibres or cellulose fibres is likely to be less than posed by chrysotile ... Additional reassurance can be gleaned by noting that these materials are unlikely to form significant amounts of respirable fibres under normal working conditions and that occupational exposures to these materials will be below the control limit for chrysotile.”

3.191 The EC emphasize that, in France, as in many other countries, manufacturers had for more than ten years been seeking asbestos-free substitutes which would meet their technical requirements and would be compatible with their health and safety obligations. In the vast majority of situations, zero-risk substitution is possible: either (i) by using non-fibrous products which are not carcinogenic (such as plaster for fire protection instead of limpet spraying); or (ii) by using fibres on which no worrying data have emerged after decades of use (cellulose or PVA in cement fibres). Moreover, replacement by man-made mineral fibres is strictly regulated: (i) mineral wool (glass or rock fibres), which have not been proven dangerous even to animals in high doses and which are used for a limited number of applications (watertight braiding); these fibres are subject to occupational exposure limits and to the rules designed to protect workers against chemical risks; and (ii) ceramic fibres, for which manufacturers are currently seeking replacements; these fibres are subject to rules which are even stricter than those concerning chemical risks. Under these rules, ceramic fibres must be replaced by a less dangerous agent wherever technically possible, and their sale to the general public is prohibited.

3.192 The EC point out that exceptions to the asbestos ban are possible where using asbestos is a safer option. Manufacturers may qualify for exemption from the asbestos ban where no asbestos-free substitute performing an equivalent function is available which, on the one hand, represents less of a danger to workers and, on the other, offers users all the technical safety guarantees. When replacing asbestos by a substitute material, it is necessary to follow a carefully defined procedure laid down in the specifications which set out the performance requirements that the manufacturer’s substitute material must satisfy under very precise conditions of use. Once selected, the substitute product must not just be tested but must also undergo feasibility studies over a period of time. This may take the form of qualification or approval procedures carried out by domestic or international third parties; these procedures are lengthy and complex in sectors such as the aerospace, nuclear or chemical industries. The EC also note that the Decree contested by Canada does not make the use of substitute products mandatory, be they fibrous or non-fibrous.

3.193 In the EC’s view, in many instances it is not necessary to use fibrous substitute products. Asbestos-cement can be fully replaced by products with no indication of carcinogenicity; these may be either non-fibrous products (ductile iron, plastic) or fibrous products (cellulose, PVA). In the case of fire protection products, for instance, there is no need to use substitute fibres. In France, traditional plaster-based products or hydraulic binders loaded with mineral aggregates such as perlite or vermiculite have been used to protect concrete or steel structures against fire since 1978 when flocking was banned. In practice, manufacturers decide to use fibrous substitute products only if these are the sole products offering the requisite technical performances (previously obtained using chrysotile). Residual use of asbestos is made on an occasional, highly specialized basis. It is most common in the nuclear, chemical, petrochemical and aerospace industries.

3.194 The EC contend that Canada cannot claim that Decree No.96-1133 is designed to "promote" domestic substitute products because France records a substantial trade deficit for PVA and para-aramid fibres. It should also be noted that PVA fibres are produced in only two plants worldwide, one in Japan and one in China. As regards cellulose fibres, the fraction used to produce fibre-cement is extremely small. The bulk of the cellulose raw material (94.5 per cent) is used by the paper industry and the remainder (5.5 per cent) goes mainly to the textile and construction industries. France imports about 40 per cent of the cellulose it uses, mainly from Canada and the United States. In 1998, France imported 380,000 tonnes from Canada, against 371,000 tonnes in 1997 and 366,000 tonnes in 1996."
Canada asserts that the undetectable risk associated with current uses of chrysotile has, as a result of the Decree, been improperly replaced by the unknown risk of substitute fibres. At the most, science indicates that, although certain substitutes can be considered “less harmful” than chrysotile, their harmlessness has not been proved. Furthermore, the ban on chrysotile promotes a false sense of security among the population, which is likely to reduce the vigilance of workers, do-it-yourself enthusiasts and the population in general when chrysotile substitutes are used. According to Canada, this is the key issue, i.e. whether the ban on asbestos effectively protects the French population while restricting international trade as little as possible. The most common chrysotile substitute fibres include aramid fibres, polyvinyl alcohol (PVA) fibres, cellulose fibres, glass fibres, ceramic fibres, rock wool, slag wool, glass wool and wollastonite.

In Canada’s view, nothing is less certain than the “proven” character of the safety or low toxicity of substitute fibres, as the EC assert. First of all, the EC stress that substitute fibres “have been in use for many years”. Although little accustomed to criticizing others for a lack of historical perspective, Canada considers that use of approximately 20 years does not constitute “use for many years” of substitute fibres. It is sufficient here to refer to the CSTEE (DG XXIV) document, which speaks more appropriately of a “relatively short” period of use extending over “approximately 20 years.” Similarly, a working group which reported on substitute fibres to the British Health and Safety Executive (HSE) noted:

“Carcinogenic risks posed by chrysotile-substitutes and chrysotile cannot be based predominantly on an epidemiological assessment since substitute fibres have only been used for periods of up to approximately 20 years.”

Canada notes that INSERM itself warns against the fact that “[Tr.] very few toxicological data exist” on asbestos substitute fibres. Canada also draws attention to the conclusions of the CSTEE (DG XXIV), to which the EC refer extensively, with respect to the safety of substitute fibres:

“For substitute materials [to chrysotile], with the exception of vitreous fibres, there is no significant epidemiology base to judge the human health risks [...] hence the conclusion that the use of specific substitute materials poses a substantially lower risk to human health, particularly public health, than the current use of chrysotile, is not well founded.”

Canada points out that the CSTEE (DG XXIV) also states that: “no epidemiological studies or observations in humans of the long-term effects of p-aramid or PVA have been reported in the scientific literature”. It goes on to add: “overall, acute and subacute toxicity data on the three substitute fibres are very meagre and do not allow for a proper comparison with chrysotile”. The WHO, in a study on substitutes published in 1993, which is similar in scope to the 1998 report on chrysotile, states in conclusion that:

“All fibres that are respirable and biopersistent must undergo testing for toxicity and carcinogenicity. Exposures to these fibres should be controlled to the same degree as that for asbestos.”

Canada notes that the Occupational Safety & Health Administration of the United States Department of Labor underlines the risk of using substitute fibres and products:

“Although the introduction of asbestos substitutes and alternatives enables manufacturers to avoid contact with asbestos, many of these surrogates pose occupational health hazards of varying degrees.”

Canada asserts that, not only is the safety or low toxicity of substitute fibres not “proven”, what is more, the EC and the United States misrepresent the situation of substitute fibres in an effort to compensate for the irrationality of the total ban on asbestos and the blind use of substitute fibres. For example, the United States claims that “none of these [replacement] fibers have been found to cause either malignant or non-malignant respira-
Canada considers that this is not true, as can be seen from the IARC data and its classification of carcinogenic substances. The IARC classifies substances according to the following four categories:

(a) group 1: the agent (mixture) is carcinogenic to humans. The exposure circumstance entails exposures that are carcinogenic to humans;

(b) group 2 (two classifications):  
(i) group 2A: the agent (mixture) is probably carcinogenic to humans. The exposure circumstance entails exposures that are probably carcinogenic to humans;
(ii) group 2B: the agent (mixture) is possibly carcinogenic to humans. The exposure circumstance entails exposures that are possibly carcinogenic;

(c) group 3: the agent (mixture, or exposure circumstance) is unclassifiable as to carcinogenicity in humans;

(d) group 4: the agent (mixture, or exposure circumstance) is probably not carcinogenic to humans.

3.201 Canada explains that these groups are not based on the toxicity of the substances, but rather on the available information. To the EC, which state that chrysotile is a carcinogenic agent in group 1, Canada responds that this category, which covers three and a half pages in the IARC document, includes many products such as silica, tobacco smoke and wood dust; it also includes alcoholic beverages, salted fish and oral contraceptives. The IARC warns against the reasoning of saying that a substance in group 1 is more hazardous than a substance in group 2.

"These categories refer only to the strength of the evidence that an exposure is carcinogenic and not to the extent of its carcinogenic activity (potency) nor to the mechanisms involved. A classification may change as new information becomes available."

For example, a substance may be classified in group 2B if there is sufficient data from animal experiments confirming its carcinogenicity, but the data for humans are still “inadequate”. Rock, glass and slag wools and refractory ceramic fibres are classified in group 2B, i.e. possible carcinogens. Polyvinyl alcohol, aramid and glass fibres are classified in IARC group 3, which is characterized by “inadequate” evidence of carcinogenicity in humans and “inadequate” or “limited” evidence from animal experiments. For Canada, it is not surprising that knowledge is still insufficient with respect to substitute fibres that have been used industrially for only about 20 years. According to the French Institut national de recherche et de sécurité (INRS) (National Research and Safety Institute), “[Tr.] some authors are of the opinion that glass fibres should be considered carcinogenic.” The IARC, for its part, reveals that glass fibre can cause mesotheliomas, at least in animals.

3.202 Canada also indicates that the United States claims that none of the chrysotile substitute fibres have been associated with respiratory diseases similar to those associated with asbestos. However, the United States Occupational Safety and Health Administration (OSHA) points out the hazards of synthetic mineral fibres:

"Several epidemiologic studies have demonstrated statistically significant elevations in the risk of lung cancer and other respiratory system cancers among workers employed in fibrous glass and mineral wool manufacturing facilities. [...] The most recent follow-up of fibrous glass exposed workers in the U.S. study still demonstrate a significant excess of lung cancer. [...] Numerous studies have
demonstrated that fiber glass and mineral wools produce lung cancer, mesotheliomas, and sarcomas in experimental animals.”305

The accumulation of evidence has prompted the United States to classify glass fibre, glass wool and ceramic fibres as “reasonably anticipated to be a carcinogen”.306

3.203 Canada notes that the French INRS states that refractory ceramic fibres “[Tr.] cause mesotheliomas with an experimental incidence comparable to and indeed greater than (at high doses) asbestos (chrysotile)”307 Again according to the INRS: “[Tr.] In light of the results of animal experiments, RCFs [refractory ceramic fibres] are carcinogenic and fibrogenic when inhaled”.308 According to the United States EPA, these fibres are probable carcinogens.309 The EC cite the larger diameter of three of the substitute fibres, namely PVA, para-aramid and cellulose fibres, to argue that they pose a lower health risk. However, this is a “nominal diameter” which does not prevent a significant proportion of fibres from having a diameter which falls well within the range of respirable diameters.310

3.204 According to the WHO in a 1993 study on substitute fibres:

“All fibres that are respirable and biopersistent must undergo testing for toxicity and carcinogenicity. [...] The data available suggests para-aramid fibres fall within this category. Furthermore, other respirable organic fibres should be considered to fall within this category.”311

In light of the IARC classification, the WHO, EPA and INRS data, and the two INSERM reports, Canada considers that one cannot therefore objectively speak, as the EC do, of the “proven” safety or lower toxicity of substitute fibres. In Canada’s view there is “no clear hierarchy in the relative toxicity” of chrysotile fibres and substitute fibres, as the EC claim and would like to believe.

3.205 Canada considers that the use of chrysotile substitute fibres has an impact not only in terms of creating a risk for the French population because of the toxicity of substitute fibres and the false sense of security that the Decree promotes. The use of substitute fibres also has a direct impact on the question of controlled use, one of the key issues in this dispute. Canada notes that, in order to counter Canada’s argument that there is a means less restrictive for international trade than a total ban, the EC cite “the impossibility” of the controlled use of chrysotile. Yet, in the same breath, they minimize the risks of substitute fibres by stating that the strict regulations which govern them eliminate any hazard. For Canada, these strict regulations constitute nothing less than a controlled use of substitutes.

3.206 Canada notes that the EC claim, for example, that refractory ceramic fibres, despite their known toxicity, do not pose a risk to the French population since they are strictly regulated, citing in support of their argument the relevant articles of the Labour Code. They continue to praise controlled use, boasting that “replacement by man-made mineral fibres is strictly regulated”. Concerning glass and rock wools, the EC claim that “These fibres are subject to occupational exposure limits and to the rules designed to protect workers against chemical risks.” As for refractory ceramic fibres, the French population is apparently protected since “These fibres are subject to rules which are even stricter than those concerning chemical risks.” Canada is quite intrigued by this about-face as to the merits of controlled use and wonders why the EC cite the feasibility of controlled use for substitute fibres and summarily reject the feasibility of controlled use in the case of chrysotile fibres?

3.207 According to Canada, the EC are endeavouring to make the replacement of chrysotile by fibres “shown to be harmless or less harmful” a “necessity acknowledged internationally”. There are two major flaws in this reasoning. The first flaw is that replacement must be done only when necessary. The ILO, through Convention 162312 and Recommendation 172313 therefore provides for replacement “where necessary”. As Canada has already pointed out, high-density non-friable chrysotile materials account for 97.5 per cent of the current uses of chrysotile. These are no longer the friable products that the WHO,
like Canada, considers problematic. In Canada’s opinion, it is these friable products whose replacement must be contemplated and that have already been largely replaced (97.5 per cent).

3.208 Canada notes that, as the EC emphasize, the WHO is of the opinion that replacement must be done whenever it is possible to replace chrysotile asbestos with “safe substitute materials”. As Canada has stressed, the chrysotile substitute products used in France are not, as the WHO prescribes, “safe”. Chrysotile asbestos is not replaced by products “shown to be harmless or less harmful”. Nor is it replaced by products which, according to the ILO, should be “harmless or less harmful”. The replacement of chrysotile asbestos by substitute fibres in fibro-cement and friction products is not a “necessity acknowledged internationally”. At best, the Decree on a ban, which imposes replacement, is an unjustified restriction on international trade in violation of the WTO Agreements. At worst, it is a measure which endangers the French population by replacing an undetectable risk by an unknown risk and by fostering a false sense of security.

3.209 The European Communities assert that the principle of replacing one substance by another that is less dangerous or harmful or by a new technique is a general rule of primary prevention applicable to human and environmental protection. It is based on abolishing the risk at the source by totally abandoning the use of the substance shown to be harmful in order to eliminate its effects. Where the use of asbestos fibres is concerned, this basic rule is applied by all the international bodies that draft recommendations designed to protect human health. According to the EC, positions on this rule have been taken repeatedly: (i) by the ILO since 1986; (ii) by the WHO, in particular in its press release dated 26 July 1996 concerning the chrysotile variety of asbestos, according to which “when available, substitute materials evaluated as safer than chrysotile should be considered”; and (iii) by the European Community in its 1990 framework Directive on the health and safety of workers and later in the Directive on carcinogens.

3.210 The EC point out that, when banned in France, asbestos fibre was not being used as such but mainly as a component in the following products or materials: (i) asbestos-cement materials (pipes, ducts, roof tiles, roof and wall cladding, planting tubes, etc.), which accounted for more than 90 per cent of the chrysotile used in France; (ii) friction materials (clutch and brake linings); (iii) insulation (board, felt); (iv) textile products (sealing braid, tape, blankets); (v) seals and gaskets; and (vi) miscellaneous products (mouldings, plastics, glues, coatings, mortars, etc.). Since asbestos has so many uses, there is no single natural or synthetic product which, alone, could serve as a universal replacement for asbestos in all the products and materials that contain it. According to the EC, there is thus no single substitute for asbestos, only replacement solutions which rely on different substitutes depending on the application, sometimes used in combination to provide a material or product with equivalent performance. Thus, the ban on asbestos is being applied by steering the market for asbestos-containing products and materials towards: (i) existing alternative technologies which do not rely on a substitute for asbestos (ductile iron or plastic pipes, aluminium roof and wall cladding, etc.); and (ii) products and materials containing one or more substitutes, some of which are fibrous but less dangerous.

3.211 The EC state that, depending on the application, asbestos has been replaced by: (i) cellulose, polyvinyl alcohol and polypropylene fibres in asbestos-cement materials; (ii) para-aramid fibres and glass fibres (combined with other non-fibrous materials) in brakes and other friction products; (iii) man-made mineral fibres (glass, rock) and often non-fibrous materials (perlite, vermiculite, silicates, etc.) for insulating panels, blocks and coatings; (iv) para-aramid and man-made mineral fibres combined, depending on the technical requirements, with metal thread (steel) or carbon or polytetrafluoroethylene (PTFE) fibres in asbestos-containing textile products; seals and various moulded products; or (v) special glass fibres (calcium and magnesium silicates) or refractory ceramic fibres in products designed to withstand very high temperatures. The price of asbestos fibres varies from F 5 to 9 per kilo depending on the origin and the physical characteristics of the fibres. The various substitute products, whether or not fibrous, are generally more expensive than...
asbestos (with the exception of some which cost more or less the same). The finished products containing asbestos substitutes are all more expensive than the asbestos-containing products serving the same purpose. In general, the price ranges of the main substitutes are as follows: (i) less than or about F 12/kg: cellulose, man-made mineral fibres (glass, rock), perlite, vermiculite; (ii) from F 12 to 60/kg: glass filaments, steel thread, polypropylene and polyvinyl alcohol fibres; (iii) from F 60 to 120/kg: special glass fibres (high-temperature), polytetrafluoroethylene (PTFE) fibres; and (iv) from F 120 to 200/kg: para-aramid and refractory ceramic fibres.

3.212 The EC indicate that these various substitutes are used as replacements for asbestos only on a very small-scale as compared with their widespread use in other industries: (i) between 10 and 20,000 tonnes of cellulose for fibro-cement, as compared with about 5 million tonnes consumed in France in the paper and textile industries; (ii) only a few thousand tonnes of polypropylene or polyvinyl alcohol fibres, which are mostly used in the textile and packaging industries; and (iii) less than 1 per cent of man-made mineral fibres, mainly used for insulating buildings. In general, France imports about 40 per cent of the cellulose it uses, in particular from Canada and the United States. Thus, in 1998, France imported 380,000 tonnes from Canada as compared with 371,000 tonnes in 1997 and 366,000 tonnes in 1996. The EC also note that there are only two factories in the world that produce polyvinyl alcohol fibres, one in Japan and the other in China. Finally, overall, France’s trade balance for these two products is regularly in deficit.

3.213 The EC assert that none of the substitutes for chrysotile is classified as carcinogenic for humans. The main substitutes are cellulose and polyvinyl alcohol fibres used to replace chrysotile in materials and products formerly made of asbestos-cement, together with the para-aramid fibres used in jointing, textiles and friction products. According to the EC, these are substances which give no cause for concern after decades of massive use throughout the world for other purposes. Thus: (i) cellulose fibres have also been used for several centuries for making paper pulp (several million tonnes are consumed in France every year); (ii) polyvinyl alcohol has been widely used since 1936 for making film for wrapping food and in the textile industry; and (iii) since the 1960s, para-aramid fibres have been extensively used for making clothing and twine. Only fibres less than 3 microns in diameter can penetrate deep into the lungs. The fibres which must be taken into account in making a metrological assessment of a working environment have been defined by the WHO in accordance with the following dimensional parameters: (i) length greater than 5 microns; (ii) diameter less than 3 microns; (iii) ratio of length to diameter greater than 3 microns [3:1]. The polyvinyl alcohol and para-aramid fibres used to replace asbestos are 2 to 8 mm (i.e. 2,000 to 8,000 microns) long and 10 to 16 microns in diameter. Cellulose fibres, which are 12 to 40 microns in diameter, can give rise to finer particles (fluff), which are said to irritate the respiratory pathways. These fibres are more than 10 microns in diameter, which means that it is physically impossible for them to penetrate into the pulmonary alveoli.

3.214 The EC consider that the results of the CSTEE and COC studies are unambiguous as regards the carcinogenicity of chrysotile and the fact that the three fibres mentioned above have given no cause for concern over their long period of use. Man-made mineral fibres such as glass or rock fibres are used for gaskets, friction materials and sealing braid required to withstand temperatures up to 1,200°C. These substances have been the subject of studies analysed in two reports which have concluded that they are less harmful for humans than chrysotile asbestos. The EC emphasize that, unlike chrysotile fibres, which have a diameter of from 0.1 to 1 micron and which separate lengthways into even finer crystalline fibrils (0.020 micron), all the synthetic fibres used in France to replace chrysotile, whether of organic (polyvinyl alcohol, para-aramid) or mineral (glass, rock) origin, retain the diameter determined by the manufacturing process throughout the life cycle of the fibre, even when emitted by a material in the process of being machined. Refractory ceramic fibres, classed as group 2 carcinogens (i.e. carcinogenic in animals but not in man), are only used in industry for a few applications where very high temperatures must be withstood. In accordance with the substitution principle, they are progressively being replaced by other, less harmful substances as technology evolves.
7. The INSERM report

3.215 Canada states that, at the request of the Direction des Relations du Travail (Ministry of Labour) and the Direction Générale de la Santé (Ministry of Social Affairs), INSERM mandated a “joint commission of experts” to study the health effects of the main types of exposure to asbestos. This group of 11 researchers was set up in the summer of 1995 and met some ten times between August 1995 and March 1996. It submitted its Executive Summary to the French Government on 21 June 1996. This Executive Summary was made public in July 1996 and the final report in November 1997. The final report has two sections: an analysis section and an Executive Summary. The analysis section contains a review of the scientific literature and summarizes the state of knowledge, allowing for a better understanding of the health risks of various circumstances of exposure to asbestos. In particular, this section covers measurement methods, the physico-chemical properties of asbestos, the main circumstances of exposure and the epidemiology of diseases linked to asbestos. The Executive Summary contains a recapitulation and presents the findings and recommendations on future studies and research to be carried out.

3.216 Canada notes that the main findings in the Executive Summary are as follows: concerning the evaluation of the risks, INSERM estimated the number of deaths in France in 1996 attributable to passive exposure to asbestos at 1,950. Concerning risk management, the group of experts pointed to certain facts: (i) asbestos is carcinogenic and should be replaced with substitute fibres, as soon as technically possible, in accordance with European law; however, the researchers do not have sufficient information on the risks of substitute fibres; (ii) chrysotile entails a level of risk comparable to or lower than the other types of asbestos, depending on the diseases considered; (iii) there is not enough data on the number of persons exposed to asbestos to assess the risks under varying situations of exposure; (iv) the risk assessment was effected on the basis of the regulation legal exposure limits; (v) the quantification of the risks is different from an evaluation of the risks, which would involve ethical, social, economic and political considerations; (vi) the researchers express grave reservations about the systematic removal of sprayed asbestos from buildings; (vii) extreme caution is called for with regard to the control of the conditions of occupational exposure to asbestos; and (viii) the researchers question the low level of compensation afforded to persons suffering from diseases caused by occupational exposure to asbestos.

3.217 Canada notes that the group also issued recommendations on future studies and research to be carried out in the following areas for which the data are still insufficient: (i) knowledge of site contamination; (ii) knowledge of exposure of people; and (iii) monitoring the trend in health risks associated with exposure to asbestos in order to assess the risk of low-level exposure. Finally, the research group makes the following recommendations: (i) further epidemiological research should be conducted into the risks associated with exposure to asbestos, as the researchers had not been able to gain an in-depth understanding of the topic; (ii) given the lack of data on the safety of substitute fibres, research should be conducted into this area without delay; (iii) more extensive research into the control of environmental health risks should be conducted; and (iv) a policy of coordinated studies and research into the environmental health risks should be elaborated. According to Canada, the INSERM report does not offer any new scientific findings. However, it proposes a model for assessing the risks of contracting lung cancer and mesothelioma from low-level exposure, which is based on a simple extrapolation of the risks observed from higher levels of exposure, without distinction between amphiboles and chrysotile. The report indicates, however, that: “[Tr.] This extrapolation does not produce certain scientific knowledge, but only an aid to understanding the implications for risk management.”

3.218 Canada submits to the Panel four studies that are critical of the INSERM report. The studies are by the Royal Society of Canada, Dr. Graham W. Gibbs, Dr. Jacques Dunnigan and the epidemiologist Michel Camus. Furthermore, certain key conclusions of the INSERM report obviously contradict a study by the French Académie nationale de médecine. The critical studies reveal that INSERM studied the health effects of asbes-
tos on the basis of hypothetical exposure situations that bear no relationship whatsoever to actual exposure. INSERM did not have at its disposal any data whatsoever on the number of persons exposed to asbestos in France. It did not sufficiently emphasize the fundamental pathogenic distinction between amphibole fibres and chrysotile fibres. The conclusions of the INSERM report are based on the risks associated with exposure sustained in the context of outdated asbestos applications, i.e., the use of amphiboles or fibre mixtures in brittle materials such as spraying asbestos and heat insulation. Moreover, the methodology of the INSERM report utilizes a linear risk model that assigns, by pure extrapolation, theoretical risks to low-level exposure, even though no epidemiological study to date has detected a higher health risk resulting from low-level exposure. Finally, the INSERM report does not investigate substitute fibres, a problem that the researchers themselves considered to be “inseparable” from the question of whether to ban chrysotile. Canada also notes that the INSERM report was qualified by Claude Allègre, France’s Minister for Education, as “mediocre”. Mr. Allègre also said that “[Tr.] There is nothing exceptional about the report in terms of scientific rigour, courage or scientific initiative […] The report contains no clear findings and can be used to support any and all arguments.”

According to Canada, the INSERM report is based on hypothetical data and does not study the exposure actually sustained by the French population. Instead, it proceeds to make “the most plausible, though uncertain estimates”. The INSERM researchers themselves concluded that “[Tr.] there is currently a great deal of uncertainty about occupational or passive exposure that has existed or now exists within the population”. In Canada’s view, therefore, the INSERM researchers could only have based their conclusions on hypothetical exposure data. These “hypothetical” conclusions concerning the number of deaths attributable to asbestos among an exposed population (for example 10,000) cannot be applied to the French population because the number of persons exposed to asbestos in France, and the rate of exposure, remain unknown.

Canada notes that the Royal Society of Canada (RSC) writes that “… the report fails adequately to address the relevance of the available studies to the key present question: are current exposures associated with increased risk?” This shortcoming undermines the usefulness of the INSERM report, prompting the RSC to express its concern over the fact that “… the risk characterization is less satisfactory because actual exposure data were not used”. The RSC stresses that the hypothetical estimation of exposure inevitably results in an overestimation of the risks actually incurred by the French population.

Canada notes that, when calculating ongoing indoor passive environmental exposure, INSERM applied the high exposure level of 0.025 f/ml. For occupational exposure levels, it applied another very high rate, 0.1 f/ml. Moreover, INSERM applied these exposure levels to the risk models applicable to cohorts for whom the exposure levels were between 2 f/ml and 250 f/ml, i.e., at levels that are 20 to 10,000 times higher. In its conclusions and recommendations, INSERM stresses the fact that, besides having used hypothetical exposure rates, the number of persons actually exposed to asbestos – the collective risk of the French population – is unknown and has not been the subject of any research. Canada points out that the conclusions presented as the risk per 10,000 persons refer to a population of 10,000 persons actually exposed to the hypothetical – and exaggerated – levels applied by INSERM.
“The estimates we have furnished correspond to ‘individual’ risks; we in fact estimated the probability that a person who might find himself in a hypothetical situation of exposure to asbestos would develop lung cancer or a mesothelioma because of this (it is indeed an individual risk, even if it is expressed in terms of ‘per 10,000’.” [italics in the original].”342 (WTO translation)

3.221 Canada notes that, according to INSERM, “These figures alone do not give an overall estimate of the number of deaths from cancer in France at the present moment attributable to exposure to asbestos” (WTO translation).343 The report concludes that “it is essential to obtain information on the number of persons exposed to asbestos […] in order to be able to estimate the number of cases of lung cancer and mesothelioma caused by exposure to asbestos […] in the French population (‘collective’ risk).”344 (WTO translation). INSERM estimates the number of cancer deaths in 1996 linked to exposure to asbestos at 1,200. It arrives at this number by applying to all of France the results of a study of the population of Glasgow, without analysing the methodology of this study or the rates of exposure to asbestos of this population.345 Yet INSERM points out that “[T]r. France began using asbestos later and used less asbestos than other countries, and no doubt proportionately smaller quantities of amphibole-type fibres. Because of these differences, there is no simple way to transpose to France the results of the projections regarding mesotheliomas [and cancers] recently produced for Great Britain”.346 As for the number of mesothelioma deaths, Canada notes that INSERM does not reveal how it arrived at its estimate of 750. Canada also notes that the French Senate and National Assembly347 and the Académie nationale de médecine estimate this number at 600, of which up to 30 per cent are “unrelated to asbestos”.348 Furthermore, according to the Académie nationale de médecine, an exposure of 0.025 f/ml for eight hours a day over 60 years would entail a risk of contracting mesothelioma of “one case per million, a figure equivalent to zero in biological mathematics”.349

3.222 In Canada’s view, INSERM’s risk estimates do not pay sufficient attention to the crucial distinction between the effects of amphiboles and those of chrysotile. Moreover, the risk estimations presented as being linked to chrysotile exposure are often based on parameters for exposure to amphiboles or amphibole-chrysotile mixtures. According to Canada, it is well known that the pathogenic potential of amphiboles is much greater than that of chrysotile350 and, even if it did not pay sufficient attention to this fact, the INSERM report nonetheless recognizes the much higher mesothelioma risk associated with exposure to amphiboles, compared to chrysotile exposure.351 In this regard, Dr. Gibbs remarks that “when they [INSERM] estimate the risk of mesothelioma, they fail to take into account the differences in risk associated with these various fibre types.”352 He continues: “The INSERM report grossly overestimates the mesothelioma risks due to exposure to ‘commercial chrysotile’ only. The hypothetical risks in the report would be more accurately characterized as those associated with amphibole or mixed commercial amphibole-chrysotile exposures.”353 The comments of the RSC are similar:

“The Panel believes that the risk of mesothelioma from chrysotile exposure is likely overestimated, since it is based on a single study that involved a small amount of crocidolite in addition to chrysotile; and ignored several studies of pure chrysotile exposure, all of which indicate a smaller mesothelioma risk than calculated by INSERM.” [italics in the original]

3.223 Canada indicates that, in addition to mentioning the use of risk estimation parameters associated with amphiboles or mixed fibres, Dr. Gibbs points out that INSERM applied the parameters deriving from the exceptionally high risk levels of the textile industry sector, as opposed to the risks posed by the chrysotile-cement sector, which accounted for more than 90 per cent of the modern uses of chrysotile in France prior to the adoption of the Decree:

“The INSERM authors have chosen to use as their factor for estimating lung cancer risk, a value reflecting risks in the textile industry sector or mixed asbestos fibre industries. The use of such a factor may be reasonable if INSERM re-
ports that the factor reflects mixed exposure or is based on the odd man out – the high textile risk. The INSERM report is misleading in inferring that these are the risks associated with commercial chrysotile. They do not reflect the values which are relevant to the chrysotile friction industry, mining and milling industries or chrysotile asbestos-cement industry.355

3.224 Canada states that INSERM examined outdated exposure, but did not consider any risk study based on current chrysotile exposure. Under the misleading sub-title “3.1.1. Estimation of the risks associated with exposure to asbestos” (WTO translation), the report claims that 1,950 deaths were linked to exposure to asbestos in 1996. According to Canada, these deaths are in no way attributable to the current circumstances of exposure to chrysotile. Considering the latency period of 15 to 50 years in the case of mesothelioma and 15 to 30 years in the case of cancers356, the 1,950 deaths are the result of exposure caused by the outdated practices of the 1950s and 1960s. Moreover, Canada considers that it is very likely that these exposures were to amphibole-chrysotile mixtures and pure amphiboles. Important reservations are in order with regard to this conclusion of INSERM. The RSC states that:

“the INSERM estimates of 750 deaths in France from mesothelioma and 1,200 from lung cancer in 1996 refer to deaths in 1996 but from occupational exposures at a much earlier date: they are not deaths due to exposures in 1996. Although INSERM is well aware of this, their report is not explicit enough.” [underlined in the original]357

3.225 Canada notes that, according to INSERM, “the immense majority of these deaths can indisputably be explained by circumstances of occupational or para-occupational origin”358, which is to say, exposure that is neither environmental exposure nor exposure to asbestos indoors359, and which, according to INSERM itself, poses no risk whatsoever according to the epidemiological data.360 Canada notes that, two months prior to the publication of the INSERM report, the French Académie nationale de médecine published a study urging calm in the face of the widespread fears incited by exaggerated media coverage of cancer and mesothelioma cases. The Académie resolutely stated that the extremely low current levels of exposure are not a source of health risks.361 Canada is surprised that INSERM paid no heed to this observation, which was well-known in French scientific circles at the time.

3.226 Canada also points out that INSERM did not investigate the case of intermittent exposure to modern chrysotile applications. According to INSERM:

“It is clearly established that the highest risks of mesothelioma are now in occupations where conditions of exposure are intermittent … In the 1960s, the main professions affected were the production and use of asbestos […] In contrast, in the 1980s and 1990s, the highest risks concerned work that involved handling materials [i.e. friable materials] containing asbestos.”362 (WTO translation)

3.227 Canada points out that the INSERM report does not look into the question of exposure to chrysotile fibres during intermittent operations on structures made of chrysotile-cement, such as sanding, sawing and drilling. INSERM could only have observed, in accordance with existing studies, that the emission levels in such cases are extremely low or even non-existent.363 Moreover, structures made of chrysotile-cement, by their nature, require very little maintenance, and when they do, the fibre emissions can be minimized by observing simple procedures and safety precautions. Dr. Gibbs points out that the risks identified by INSERM do not correspond to the values applicable to the friction materials industry, nor to the mining, machining or asbestos-cement industries.364 Canada also refers to the conclusions of the Académie nationale de médecine, which determined that: “[Tr.] no illness due to asbestos has yet been formally proven in France outside an occupational type of exposure”.365 These conclusions run counter to those of INSERM and underscore the growing scepticism with regard to the scientific value of the arguments presented as justifications for the banning of chrysotile. According to Canada, these doubts, in fact, originated in the French scientific community.366
3.228 Canada states that, in its risk assessment, INSERM used a linear model that overestimated the risks of low exposure. This model offers a method of extrapolating the risk of low exposure to chrysotile from the risk of high exposure. Canada points out that the epidemiological data, however, indicate no detectable risk whatsoever deriving from low exposure, as INSERM itself acknowledged in the case of environmental exposure and indoors exposure to asbestos. INSERM admits that to date, it does not possess "direct and unequivocal scientific expertise on the value of the risk of lung cancer and mesothelioma that may exist among humans exposed to 1 f/ml or lower levels". The estimates that INSERM arrives at "have to be considered as orders of magnitude given that there is no definite knowledge regarding the risk of cancer at resulting from exposure to levels below 1 f/ml [italics in the original]. INSERM saw its role as being "to provide scientifically verified elements of knowledge concerning the health risks linked to exposure to asbestos, but risk management is not its responsibility". Nonetheless, INSERM performs extrapolations for low exposure even though "This extrapolation does not produce certain scientific knowledge, but only an aid to understanding the implications for risk management." Canada considers that the INSERM report, however, as a scientific evaluation of the risks, should be more than "an aid to understanding" based on "plausible uncertain estimates" when it leads to the adoption of a Decree instituting a total ban such as that adopted by France.

3.229 Canada notes that Dr. Dunnigan remarks that "the authors of the report themselves acknowledge that the methodological basis used to generate risk data at low levels of exposure to chrysotile asbestos cannot provide 'certain scientific knowledge'." He considers that:

"the INSERM report alone cannot serve as a sufficiently credible and comprehensive basis on which to conclude that the only way of protecting the health of workers and of the general public is purely and simply to ban all varieties of asbestos and all its uses." (WTO translation)

3.230 Canada adds that, citing studies in support of his view, Dr. Gibbs also notes the existence of exposure levels below which the health risks are undetectable. In line with the other criticism, the RSC notes the error in the conclusions drawn from the extrapolations of low-exposure risks from high-exposure risk data: "the transfer of risk coefficients calculated from high exposure settings […] involves possible errors". Canada notes that the INSERM conclusions on the risks associated with low exposure are in sharp contradiction to the most recent scientific publications.

3.231 Canada indicates that one of the consequences of the Decree adopted on the heels of the INSERM report has been the replacement of chrysotile fibre by substitute fibre. The ban on chrysotile as the way of managing the risks cannot therefore be viewed in isolation. In this respect, INSERM reveals that the Decree banning asbestos could have unsuspected consequences:

"It must be noted that the group of experts was unable or unwilling to consider some key questions relating to the problem of risks associated with human exposure to fibres. Because of the short time frame, the health risks associated with exposure to substitute fibres were not addressed […] The problem [the ban on asbestos] is, however, inextricably linked to the choice of substitute fibres […] The group of experts considers that it does not have sufficient information to decide on the possibility of replacing asbestos by a substitute product that is completely without risk in all the situations in question." (WTO translation)

Canada indicates that, after having reviewed the INSERM report, all the experts cited above concluded that the report is not supported by real, current data relative to the exposure of the French population to the modern uses of chrysotile. This report cannot be seen as a credible, scientifically grounded basis for justifying a measure as excessive as the total ban on chrysotile and all its uses.
3.232 Canada points out that, following INSERM’s recommendation on further studies on risks associated with substitute fibres, INSERM was commissioned to prepare another report, this time on the health risks of substitute fibres. The Executive Summary of this report was made public in June 1998, a surprising 18 months after the ban on asbestos took effect. As Canada notes, the Executive Summary points to the scarcity of data on the substitute fibres and the novelty of their use, stating that little is known about their effects on human health. It also states that, had experiments on asbestos been conducted with exposure levels similar to the ones used for testing substitute fibres, it is likely that the experiments on asbestos would have shown that asbestos poses little, if any, significant risk. The full INSERM report on substitute fibres has still not been published, one year after the release of its Executive Summary. Canada states that, when the WTO consultations were held in Geneva on 8 July 1998, the EC assured the Canadian delegation that the publication of the full report was imminent. In Canada’s view, it increasingly appears that France is more concerned with banning chrysotile than informing the French population of the danger of substitute fibres, the use of which is effectively imposed because of the ban on chrysotile. Canada considers that the telling criticisms of the INSERM report by France’s Minister for Education, Mr. Claude Allègre, by the French Académie nationale de médecine, the Royal Society of Canada, and internationally renowned scientific experts, cast considerable doubt on whether the INSERM report constitutes a credible basis for so extreme a measure as a ban on a hitherto widely used product.

3.233 The European Communities emphasize that INSERM ranks among the most important biological and medical research institutes in the world. It comprises some 200 research laboratories employing around 10,000 researchers and covers all fields of biomedical research and all relevant scientific disciplines. It is a government research body, financed by the French State and placed under the dual responsibility of the Ministry of Research and the Ministry of Health, and it is totally independent of economic or other interests linked to the asbestos and substitute fibres industries. In the EC’s view, INSERM thus had all the competence and independence required to give a scientific opinion on the threat posed by asbestos to human health. INSERM’s report was drawn up by 11 scientific experts from various disciplines, who also gave hearings to several other experts. As its name suggests, the report was commissioned by the authorities to provide information on the risks connected with the main types of exposure to asbestos. INSERM’s mandate was not to make recommendations on how to combat the harmful effects of asbestos, but to make available the scientific knowledge necessary for any decision to be taken. The INSERM report was based on an analysis of numerous international scientific research projects on the effects of asbestos. The EC emphasize that most of its findings, including all those on which France based its decisions, have been approved by the international scientific community, including the Panel of the Royal Society of Canada, which the Canadian Government commissioned to analyse the contents of the report.

3.234 The EC point out that the INSERM report’s principal conclusions, especially those on the carcinogenic nature of chrysotile, the absence of a harmlessness threshold, the importance of intermittent exposure among end-users of asbestos and the very large number of workers affected by this type of exposure, have since been confirmed by recent studies and by international bodies such as the WHO. The first of its kind in France on the subject of asbestos, the INSERM report comprises a critical and well-argued review of the international scientific literature. The methodology and scope of this work (12 chapters of discussion, supported by 1,200 bibliographical references and followed by a summary of some 60 pages) radically distinguish it from any monograph or fragmentary study advocating one or other of the theories which have contributed to the debate. The report itemizes the main effects which exposure to asbestos is known to have on human health: asbestosis, benign pleural damage, bronchial cancer and mesothelioma. Even though 166 new cases were recognized as occupational diseases in France in 1995, asbestosis is a disease whose most serious forms have become rare in industrialized countries as a result of the reduction in levels of exposure. For this reason, the Executive Summary focuses on the risk of bronchial cancer and mesothelioma.
3.235 The EC note that the authors of the report drew the following main conclusions:

(a) Most cases of pleural mesothelioma that have emerged to date in industrialized countries are of occupational or para-occupational origin. Among the occupations affected, the cases which now occur involve the handling of asbestos-containing materials installed in buildings or equipment. The diverse occupations which carry a high risk of mesothelioma include welders, dockers, laboratory technicians, painters and decorators, jewellers, metal workers, car mechanics, railway workers, workers in thermal power stations, etc. If the number of deaths caused by mesothelioma is compared with the number of people in each occupation, it is found that the jobs which are proportionally most affected are sheet-metal workers (including shipyard workers) and industrial coachbuilders; then come plumbers, carpenters and electricians. The building trades alone currently account for one quarter of all deaths from mesothelioma. Since the 1950s and 1960s, the number of cases of mesothelioma has been growing by 5-10 per cent each year, depending on the country. In France the growth rate has been 25 per cent every three years; the rate of seven cases per million inhabitants per year between 1979 and 1981 reached 16 cases per million inhabitants per year between 1991 and 1993. The number of deaths caused by asbestos in France in 1996 is estimated by INSERM at some 1,950 (750 cases of mesothelioma and 1,200 cases of lung cancer). The risks associated with exposure, which may be occupational or para-occupational (domestic cleaning of work-clothes worn by workers handling asbestos) or environmental (geological or industrial in origin), have been incontestably proved. As regards the effects on health of “passive” exposure in buildings, a risk cannot be ruled out even though the epidemiological data are not sufficient to demonstrate what that risk is. As to whether lung cancer can be causally attributed to occupational exposure to asbestos, the group of experts concluded, on the basis of an analysis of available data, that such cancer can develop where no prior fibrosis is present. As regards the role of tobacco, the group of experts pointed out that the proportion of cases of lung cancer attributable to asbestos is identical in smokers and non-smokers. In other words, the fact that a worker who has developed lung cancer smokes does not mean that the cancer is not due to asbestos. It follows that, where an individual has been exposed to asbestos and has developed lung cancer, the fact that the person concerned smokes gives no indication as to the probability of asbestos having had a role in causing the cancer. This information, fundamental to the recognition of the occupational origin of diseases, provided justification for the arrangements implemented under the December 1995 plan.

(b) The group of experts considers that the carcinogenic effects of asbestos fibres cannot be differentiated according to whether the fibres are of the chrysotile or amphibole type. It found that the carcinogenic effects of asbestos fibres marketed under the designation “chrysotile” were just as serious as the effects of the fibres of the amphibole type, as far as lung cancer was concerned. As regards the risk of mesothelioma, the effect of chrysotile fibres is incontestable, even though less serious than that of certain amphiboles. The morphological characteristics rather than the geological origin of the fibres seem to be the decisive factors, with the longest fibres being the most carcinogenic. In order to estimate the risk of low levels of exposure (<0.1 fibre/cm³), INSERM’s experts extrapolated to low levels, on a linear basis and without a threshold, the risks observed at high levels of exposure. This hypothesis (i.e. that there is a linear relationship between the risk and
the level of exposure and that there is no threshold) is among the theories currently accepted on the subject of carcinogens and is, for example, included in the ILO encyclopaedia. It is therefore the most plausible hypothesis. The estimate is said to be “uncertain” because it is not based on observations which can validate or invalidate it beyond any doubt. INSERM’s expert group estimated, on this basis and in line with knowledge to date, what additional risk of cancer would arise over a lifetime from continuous exposure to asbestos in various reference situations, for example: (i) for 10,000 men aged between 20 and 65 who are subject to continuous occupational exposure (1,920 hours) corresponding to 0.1 f/ml or cm³, the risk is of 30 additional deaths; (ii) for 10,000 people (half of them men, half women) who are subject to continuous passive exposure during their school years (between the ages of 5 and 20) and during their working lives (1,920 hours between the ages of 20 and 65) corresponding to 0.025 f/ml (or 25 fibres per litre), the risk is of nine additional deaths.

3.236 The EC point out that any doubts with regard to the carcinogenicity of chrysotile asbestos have been removed by the joint report: chrysotile is carcinogenic, just as much as amphiboles when it comes to lung cancer but less than amphiboles when it comes to mesothelioma (ratio of 1 to 3 but same order of magnitude). As regards mesothelioma, this information has been largely confirmed by the study published by Camus et al. on cancer mortality among women living in close proximity to chrysotile asbestos mines in Quebec. This study shows a net excess of mesothelioma cases (the risk is at least seven times higher than among women living elsewhere in Quebec). Moreover, low doses do have an effect. The limit of 0.1 f/ml or cm³ therefore corresponds to a not negligible risk of 30 extra cases of cancer per 10,000 people exposed. The values above which work must be carried out to protect populations (25 fibres per litre) carry a theoretical residual risk of nine extra cases of cancer per 10,000 people exposed. The EC indicate that this figure should be compared with the level of risk which is usually recognized as applying to the population in general, namely one extra case of cancer per 1 million people. The risk population is large. For 1996, the report gave an estimate of 1,950 deaths due to asbestos (750 cases of mesothelioma and 1,200 cases of lung cancer). The vast majority of mesothelioma cases are of occupational or para-occupational origin. However, a new aspect which emerged during the period 1980-1990 was that it was no longer those working in the asbestos industry (i.e. asbestos production and processing), who were most at risk. The highest risk is now observed among people handling asbestos-containing materials (e.g. sheet-metal workers, industrial coachbuilders, plumbers, carpenters, electricians, heating engineers and cable specialists). The EC emphasize that the population concerned comprises all finishing trades in the construction industry, as well as a good number of other occupations, in other words, several million people, of whom (despite a low individual risk) a large number will contract mesothelioma. The French study recently published by INSERM shows that around 25 per cent of men born between 1930 and 1939 have been exposed to asbestos during their working life.

3.237 The EC consider that procedures for controlled use cannot be implemented effectively when hundreds of thousands of people are exposed daily in sectors with little supervision as regards health and safety, such as the building industry, which accounts for at least 25 per cent of mesothelioma cases. The study by Y. Iwatsubo confirms that in France most cases of mesothelioma occur among servicing and maintenance workers. The EC therefore consider that the principle of “controlled use” cannot be applied to the diffuse risks affecting a range of very varied trades practised in extremely diverse situations and involving in particular servicing and maintenance work, where it is not possible to implement rules on “safe use”. The EC point out that Canada itself acknowledges the limits of controlled use since it recognizes that certain applications have to be prohibited where exposure cannot be controlled.

3.238 The EC note that “safe” use is not a guarantee even when it is practised. Safe use is contested in two quite official publications:
(a) A study conducted by the Quebec Workers’ Health and Safety Commission (CSTT). This study shows that the risk of mesothelioma increased steadily in Canada between 1967 and 1990, principally among servicing and maintenance workers. Of the 120 cases of mesothelioma identified in the study, 49 were miners and mill workers, 50 were workers in the asbestos industry and 21 were servicing and maintenance workers. Of the 25 per cent of these cases which involved short exposure, the most affected group were the servicing and maintenance workers. The study shows that the occurrence of mesothelioma is indeed due to chrysotile asbestos and that the incidence of the disease is growing more rapidly in the servicing and maintenance sector. This shows that, even in Canada, which specializes in safe use, the efficacy of the method remains to be proved. The EC note that, despite the existence of this CSTT study, the Canadian party declared during the consultation meeting held on 8 July 1998 that it did not have any study at its disposal which covered the servicing and maintenance sector.385

(b) an HSE study386, mentioned in the annex to the report produced for Canada’s Royal Academy in 1996 following the publication of the INSERM report. It shows that despite strictly controlled use - it focuses exclusively on production workers -, there was still a 1.28 per cent excess of mesothelioma cases, as opposed to 4.61 per cent among people who had worked prior to 1969, when legislation was introduced in Great Britain. From this it can be concluded that controlled use does not prevent deaths from mesothelioma, even in specific sectors of industry employing limited numbers which are easy to regulate and control.

3.239 The EC point out that the confirmed carcinogenicity of chrysotile and the impossibility of controlling the risk in all situations prompted the French Government to stop the risk spreading by applying the principle laid down at European Union level for carcinogenic risks and also recommended by the WHO387 and the ILO388: replacement by a less dangerous product where technically possible. The European Union, in its general Directive on the health and safety of workers, lays down the principle that a dangerous agent or process should be replaced by a safe or a less dangerous agent or process where one exists. As regards cancer risks, the Community Directive on carcinogens389 contains the same principle in that it expressly provides for the replacement of a carcinogen by a less dangerous substance where this is technically possible. The WHO press release of 26 July 1996 states that safe substitutes should be used to replace chrysotile whenever this is possible. Moreover, in a “criterion document” dating from September 1998, the WHO states that the risk is probably higher among construction workers and perhaps also among other user groups. The ILO Encyclopaedia of occupational health and safety states that the most effective means of prevention is to avoid the use at the workplace of substances which are recognized as causing cancer in humans.

3.240 The EC assert that the INSERM report thus confirmed the necessity of the measures taken in 1996 in accordance with the guidelines adopted in July 1995. It also provided additional information, on the basis of which the French Government took a coherent set of further measures. The manufacture, import, export and sale of asbestos-containing products, including asbestos-cement, were prohibited with effect from 1 January 1997. This decision was enacted in the form of Decree No.96-1133 of 24 December 1996 laying down a general prohibition on asbestos to apply as of 1 January 1997. Asbestos-cement products were banned totally. A few very limited and strictly regulated exceptions are allowed for industrial uses where no substitute product would ensure the safety of workers and users. The purpose of the ban is to stop the spread of the risk posed by the manufacture and marketing of asbestos-containing products, in other words to prevent the “inflow”. The EC emphasize that, contrary to the impression given by Canada, there was never any intention of promoting the systematic removal of asbestos already present. The prohibition measure is based on reasoning similar to that followed by Canada when it banned
amphiboles\textsuperscript{390}: the aim is to replace a dangerous agent by agents considered less dangerous on the basis of available scientific knowledge. The measure was taken at a time when the Groupe scientifique pour la surveillance des atmosphères de travail (G2SAT), working under the Senior Council for the Prevention of Occupational Risks, had already produced a report setting out the hierarchy of the risks from the most suspect replacement fibres (mineral wools and ceramic fibres). Following the prohibition of asbestos, a redundancy programme was needed to assist companies forced to reduce their staff as a result of ceasing to manufacture asbestos-based products. Three firms were affected: of a total of 1,370 employees, 490 found themselves surplus to requirements but 400 of them qualified for early retirement. One establishment employing 126 people was closed.

3.241 The EC indicate that the authorized exposure threshold, applicable to the manufacture of products containing chrysotile, was brought down to 100 fibres per litre (as opposed to 300, the limit which was originally to apply until 1 January 1998). Since 1994, the limit for occupational exposure in the United States has been set by the Occupational Safety and Health Administration at 0.1 fibre/cm\(^3\) (or ml). France adopted the same value in 1996 since it was the lowest limit value obtained by phase-contrast optical microscopy. A lower value would have required the replacement of all existing measuring equipment.\textsuperscript{391} This provision was designed to reduce further exposure to the inhalation of asbestos dust in manufacturing or processing plants which qualified for an exemption.

3.242 The EC explain that a certification procedure by an accredited organization was introduced at the beginning of 1997 for firms called upon to remove or to contain asbestos. This procedure makes it possible to verify the quality of the work carried out and the quality of the measures taken to protect the health of the workers. The certificate is issued following a study carried out within the company to investigate working methods, the training provided for managers and workers, and the equipment available; the study is followed by an on-site audit to assess the quality of actual practice. Since the summer of 1997, companies have only been allowed to carry out the removal of “friable” asbestos (such as flocked asbestos or lagging) if they possess a certificate qualifying them to do so. The aim of this measure is to tighten up the management of the serious risks involved in asbestos removal (especially removal of flocked asbestos). The surveillance measures applicable to flocked asbestos and lagging have been extended to false ceilings. The EC emphasize that at no time was it planned, as in the United States, to call for the systematic removal of all asbestos. On the contrary, information was widely circulated stating that a higher risk could be created if asbestos were removed too quickly or in the wrong circumstances.

3.243 The EC point out that technological and scientific monitoring was stepped up. Two measures were adopted: (i) epidemiological monitoring of mesothelioma cases was introduced, with the assistance of INSERM, in order to ascertain the conditions in which tumours appear and to measure the actual risks of exposure; and (ii) INSERM was asked to produce a joint experts’ report on the risks linked to the fibres used as substitutes for asbestos. The EC note that, in any event, the use of replacement products classed as dangerous is subject to legislation on chemical hazards (obligation to conduct risk assessment and protect workers, recommended limit values, etc.). If the replacement products are classed as carcinogens, the rules are even stricter and the principle of replacement by a less dangerous product where technically possible applies.

C. LEGAL ARGUMENTS

3.244 The European Communities claim that, according to the general procedural rule applicable to the settlement of disputes, first spelled out by the Appellate Body in its report on United States – Measure affecting Imports of Woven Wool Shirts and Blouses from India\textsuperscript{392}, and since restated on several occasions\textsuperscript{393}, it is for the complaining party to establish inconsistency with a provision of a WTO Agreement before the burden of proving consistency with that provision shifts to the defending party. In this instance, Canada has not produced any convincing evidence to show that the contested measure infringes any of the provisions invoked. In particular, Canada has not produced any evidence that so-called “safe” use offers a feasible and reliable way of protecting the health of all those affected.
1. Agreement on Technical Barriers to Trade ("TBT" Agreement)

(a) Applicability of the TBT Agreement

3.245 Canada contends that the main objective of the TBT Agreement is to prevent technical regulations and standards from creating unnecessary barriers to international trade. Canada also recognizes the right of Members to take the necessary measures to ensure the protection of people’s health and lives. According to Canada, the Decree is subject to the disciplines of the TBT Agreement and is incompatible with its Article 2.2, 2.4, 2.8 and 2.1. More specifically, the Decree is an unnecessary barrier to trade. It is neither based on existing international standards nor on the performance of asbestos fibres and products containing such fibres. Finally, the Decree prevents these products being imported into the French market and is discriminatory in the domestic market with respect to these same imported products.

3.246 Canada contends that the TBT Agreement applies to the Decree because it is a "technical regulation" within the meaning of Annex 1 to the TBT Agreement. The Decree is a technical regulation because it considers all asbestos fibres and all materials, products or devices that contain such fibres, with the exception of four, as products that are supposed to pose risks for people’s health and safety. The Decree prohibits all chrysotile products for which there is a substitute that represents "a lesser occupational health risk than chrysotile fibre" and which, for the purposes of the end-use, gives "technical guarantees of safety" equivalent to the guarantees for chrysotile fibre. More specifically, the Decree is a technical regulation because it contains several aspects of the above definition. It is a document that sets forth a characteristic of a product, a process and a production method for a product. The document also deals with labelling requirements. Moreover, compliance with the contents of the document is mandatory.

3.247 According to Canada, the ordinary meaning of the word document is "anything … written … relied upon to record or prove something". The Decree contested is something written, i.e. a general administrative instrument issued by the Executive and designed to make the public aware of the Government’s decision to consider unacceptable any risk of exposure to all varieties of asbestos. The Decree was issued by the French Prime Minister. It was published in the Official Journal of the French Republic. It is therefore a document, in accordance with the definition. The ordinary meaning of the word "characteristic" is "that which constitutes a recognizable distinctive feature". The Decree describes a recognizable distinctive feature of the products. In fact, in accordance with its purpose, which is to ban asbestos totally, the main provisions of the Decree are designed to prohibit asbestos fibre, regardless of whether it is amphibole fibre or chrysotile fibre, in the manufacturing and processing of materials, products and devices that are placed on the French market. The characteristic of these materials, products and devices, as laid down in the Decree, is the absence of asbestos fibres. The Decree refers in particular to the processing of all varieties of asbestos fibres, whether or not included in materials or products. By so doing, it imposes restrictions on the processes and production methods related to asbestos fibres, including chrysotile fibres. On an exceptional and temporary basis, chrysotile fibre is permitted in the manufacturing and processing of four products, if there is no supposedly less harmful substitute fibre. Moreover, the Decree stipulates that these manufacturing and processing activities are subject to the standards governing exposure to asbestos dust in places of business. With respect to the aspects in the second part of the definition, Canada points out that the Decree includes administrative provisions applicable to products which, exceptionally, contain chrysotile fibre. In Article 3, the Decree provides for a sophisticated declaration mechanism for the purpose of obtaining an exemption from the ban on asbestos. The Decree also deals with the labelling requirements for a product. It sets out rules for labelling chrysotile-containing products that are covered by a temporary exemption from the ban. Finally, with respect to the mandatory aspect of the document, it is prescribed that all the above provisions of the Decree are binding. Compliance with the characteristics of the products and their related processes and production methods, as well as with the applicable administrative formalities, is mandatory. Article 5 provides for the penalties applicable in the event of a breach.
3.248 Canada claims that the fact that the Decree qualifies as a technical regulation under the TBT Agreement is confirmed in France’s notification to the Committee on Technical Barriers to Trade. France gave notice of the measure it took pursuant to Article 10.6 of the TBT Agreement on 21 February 1997. It also recognized the applicability of the TBT Agreement in the notification document. According to Canada, the EC also recognized the applicability of the TBT Agreement in the justification given pursuant to Article 2.2 thereof. They maintained that the Decree was not more trade-restrictive than was necessary in order to fulfill legitimate objectives, i.e. the protection of human health and safety, taking account of the risks non-fulfilment would create. Finally, at the consultations on 8 July 1998, in his opening statement, the representative of the EC conceded the applicability of the TBT Agreement to the French measure. The relevant passage is as follows:

“The only relevant provisions therefore seem to be those of the TBT Agreement. It is for that reason, indeed that the EC notified the French measures as technical regulations under the TBT Agreement.”

3.249 Canada concludes that, by describing the characteristics of the products, by specifying the related processes and production methods, by setting up a declaration mechanism for the purpose of obtaining exemptions from the ban on asbestos, by the labelling provisions for chrysotile-containing products covered by an exemption, as well as by its mandatory nature, the Decree banning asbestos falls within the meaning of “technical regulation” under Article 1 of Annex 1 to the TBT Agreement. According to Canada, the EC themselves recognized the applicability of the TBT Agreement in official documents to the WTO and during the consultations which preceded the establishment of the Panel.

3.250 The European Communities maintain that the Agreement on Technical Barriers to Trade (TBT) does not apply to the Decree. Even if the Panel were to consider that the TBT Agreement is applicable, the EC maintain that the Decree is not incompatible with the provisions of the said Agreement.

3.251 The EC claim that, contrary to Canada’s argument, the Decree cannot be construed as a “technical regulation” within the meaning of the TBT Agreement, which does not cover general prohibitions on the use of a product for reasons to do with the protection of human health. In order to interpret the provisions of this Agreement correctly, it should be remembered that the “customary rules of interpretation of public international law”, in particular those arising out of the 1969 Vienna Convention on the Law of Treaties, must be used to interpret the provisions of the General Agreement and of the WTO Agreement. Those rules call for an examination of the ordinary meaning of the terms of a treaty, read in their context and in the light of the object and purpose of the treaty considered. The Appellate Body has indicated in this regard that “A treaty interpreter must begin with, and focus upon, the text of the particular provision to be interpreted. It is in the words constituting that provision, read in their context, that the object and purpose of the states parties to the treaty must first be sought.” Therefore, an interpreter is not free to adopt a reading that would result in reducing whole clauses or paragraphs of a treaty to redundancy or inutility.

3.252 The EC contend that the TBT Agreement does not cover general prohibitions. It follows from the preamble (notably the third and fifth paragraphs), from the background to the TBT Agreement and from the actual wording of several of its provisions that the fundamental objective of this Agreement is to monitor the adoption and application of the “standards” and “technical regulations” that cover the detailed characteristics of products or their methods of production. The TBT Agreement could, for example, be applied to monitor a “technical regulation,” that set a minimum resistance level for seat belts. Another example would be a “technical regulation” that laid down percentages for ingredients in chocolate and prescribed methods of production. On the other hand, it is not the object and purpose of the TBT Agreement to deal with general prohibitions such as that applied by the French Decree to asbestos and asbestos-containing products. According to the EC, this specific object and purpose of the TBT Agreement can be seen clearly in Annex 1 thereto, which gives the definition of what should be understood by “technical regulation”: “Document which lays down product characteristics or their related processes and production
methods ...”. In this specific case, the definition means that the TBT Agreement cannot apply to the Decree. It follows from the above definition that a technical regulation is a document which lays down the characteristics or processes and production methods with which a specific/identified product must comply, in particular if it is to be released for free circulation on a given market.

3.253 The EC claim that the definition of technical regulation should not therefore apply to prohibition measures that cover all products in general. These products should continue to be covered by the General Agreement alone. To adopt any other approach would be equivalent to nullifying the effect of certain provisions of the GATT, in particular Articles I and III, which are applicable in cases of general prohibitions. The TBT Agreement must be considered as the specific application of the principles of the GATT 1994 to technical regulations. In the EC’s view, it is inconceivable that the negotiators of the TBT Agreement wished it to apply in every case to Members’ regulatory measures affecting products, and in particular to general prohibition measures. The position of the EC is fully supported by the position taken by other Members and other Panels. The Panel report on United States - Gasoline noted that:

“The United States argued that the TBT Agreement had been designed to elaborate on the disciplines of Article III of the General Agreement for a very specific subset of measures (technical regulations, standards and conformity assessment procedures). The fact that a measure was in writing, mandatory and applied to products did not make it a technical regulation. Excise taxes, for instance, met all these criteria but were not ‘technical regulations’. Similarly, the term ‘technical regulation’ was not so broad as to cover all governmental regulations affecting products. For example, government regulations requiring factory smokestacks to have devices to reduce emissions were not technical regulations, though they were in writing, mandatory and specified ‘characteristics’. [...] The United States concluded that the complainants were interpreting the term ‘technical regulation’ out of context and such an interpretation, if accepted, would introduce into the TBT Agreement many measures which were in fact not intended to be covered.”

3.254 The EC claim that the French Decree is not a “technical regulation” within the meaning of the TBT Agreement. The Decree, taken as a whole, is a general ban on asbestos fibres and asbestos products. The prohibition applies to production, processing, importation, exportation, disposal, supply, sale, and marketing. The Decree specifies neither the characteristics nor the production processes and methods for asbestos fibres, asbestos-containing products nor the products exempted from the prohibition measure.

3.255 The EC contend that the Decree does not lay down “product characteristics”. The EC fully agree with Canada’s conclusion that the Decree constitutes a document, but they are surprised by Canada’s choice of a definition for the word “characteristic”. According to the very Robert dictionary on which Canada’s argument is based, the word “characteristic” may be used either as an adjective or a noun. The first definition given by Canada (“which constitutes a recognizable distinctive feature”) is the definition corresponding to the adjective. On the other hand, the second definition (that which serves to characterize) given by Robert, and strangely forgotten by Canada, specifically corresponds to the noun. It is on the latter definition that the Panel should base its analysis because, in the above-mentioned definition in Annex 1, the word “characteristic” is used as a noun and not as an adjective. In the light of this definition, the French Decree cannot be regarded as “laying down product characteristics”. The EC note that Canada maintains that the characteristic is “the absence of asbestos fibres” and that the products covered are “materials, products and devices that are placed on the French market”. However, the EC do not consider that in the French measure the words “the absence of asbestos fibres” serve to characterize products placed on the French market because “the absence of asbestos fibres” does not constitute a characteristic, much less the characteristic of products placed on the French market. One of the critical weaknesses of Canada’s argument lies in the fact that it has eliminated the existence of a possessive phrase, as denoted by the word “of”, between “the
characteristics” and “a product”. For the Decree to be able to lay down the “characteristics” of a product, it would have in one way or another to designate the product(s) to which the said “characteristics” relate. The Decree does not designate any product but lays down the principle of prohibition. It follows that the contested measure cannot be examined on the basis of the TBT Agreement as it has general scope and does not concern any specific product whose characteristics are spelled out in the French Decree. In the light of the foregoing, for the EC it is clear that the Decree does not lay down the “characteristics” of asbestos fibres but simply prohibits their use on French territory. It is equally clear that the Decree does not lay down the “characteristics” of asbestos-containing products. Nor does the Decree define the technical characteristics of the products, which may enjoy an exemption from the general prohibition laid down. Such products may or may not contain asbestos.

3.256 The EC assert that the Decree does not lay down “production processes and methods relating to a product”. The term “process” may be defined as a “means, a practical method of doing something, of obtaining a result”.415 The term “method” may be defined as a “logically ordered set of principles, rules or stages making it possible to reach a result”.416 The Decree does not lay down any means or ordered set of rules governing the production (extraction and processing) of asbestos fibres. The prohibition applies to asbestos fibres, so it is not possible to say how they should be produced because they may no longer be produced. The same is true of asbestos-containing products. In terms of the definitions given for production “process” and “method”, the Decree does not specify any means nor lay down any ordered set of rules governing the manufacture of asbestos-containing products. It merely bans any type of asbestos-containing product. In other words, the prohibition applies to asbestos-containing products, and nothing is said about how they should be produced. Nor does the Decree define the production processes and methods for the products that may be exempted from the general ban. Such products may or may not contain asbestos. The EC therefore conclude that the Decree is not a technical regulation within the meaning of the TBT Agreement, neither for asbestos fibres nor for asbestos-containing products, nor even for products enjoying temporary exemptions. This conclusion follows logically from the object and purpose of the TBT Agreement, which is not to cover general prohibition measures.

3.257 According to the EC, the fact that France has notified the Decree to the Committee on Technical Barriers to Trade in no way prejudges the applicability of the Agreement. The Decree was notified, in good faith, for the sake of transparency vis-à-vis all the Members of the WTO and in response to repeated requests by Canada. Any other interpretation would create additional obligations for WTO Members and would induce them to discontinue, or at least reduce, notifications of their general legislation to WTO Committees. According to the EC, Members must therefore continue to notify their legislation, without notification as such having any legal consequences for the status of the measure concerned or entailing any obligations to which Members are not normally held.

3.258 Canada reiterates that the TBT Agreement applies to the Decree and claims that the EC’s interpretation, based on Article 31 of the Vienna Convention on the Law of Treaties, is incorrect. In this regard, Canada points out that the three third parties to the dispute unanimously recognize the applicability of the TBT Agreement in this case. The EC are therefore alone in disputing its applicability. Canada maintains that the EC’s interpretation is not based on Article 31 of the Vienna Convention and the Decree falls specifically within the scope of the TBT Agreement because it represents the type of measure covered by the definition of “technical regulation” in Annex 1 to the Agreement, including general prohibitions within the scope of the TBT Agreement is not contrary to either its object or its purpose, which are to prevent technical regulations and standards from creating unnecessary obstacles to international trade.

3.259 Canada claims that the EC depart from the ordinary meaning of the terms of the definition of “technical regulation” and refer to the preamble and history of the TBT Agreement to assert that its object and purpose are not to deal with general product prohibitions, but to lay down the characteristics or processes or production methods which a specific
product must meet. This distinction made by the EC between prohibitions that apply to all products without distinction and measures aimed particularly at a specific product is not supported by the TBT Agreement. There is no provision which stipulates that general prohibitions do not fall within the scope of the TBT Agreement. The EC’s interpretation of Annex 1 to the Agreement is contrary to the principle of effectiveness.407 According to this principle, a treaty must be given the interpretation which renders it meaningful and effective, rather than that which would render it ineffective. However, in Canada’s view, the consequence of the EC’s interpretation would be that it would suffice to give a measure intended to hinder trade the form of a general prohibition in order to allow that measure to evade the disciplines of the TBT Agreement. Thus, as the United States notes, by manipulating the form of a measure, it would be possible to render the TBT Agreement ineffective. Contrary to what the EC claim, their interpretation in such circumstances is not supported by the precedents. The excerpt from the Panel Report in United States – Gasoline cited by the EC does not relate to a position taken by the Panel but rather to an aspect of the American arguments in this case. The Panel that examined this case never sanctioned this American claim and did not discuss it. Furthermore, this excerpt, cited out of context, is contradicted by the position expressed by the United States in this dispute. In interpreting the TBT Agreement as applying only to a description of characteristics or of processes and production methods specific to a particular product and not to general prohibitions that apply to any product without distinction, the EC take into account only one part of the provisions of the Decree, in this case those concerning the ban on asbestos, and fail to mention the transitional provisions. However, in Canada’s view, in order to determine whether the Decree satisfies the criteria of the definition of “technical regulation”, all its provisions must be examined, both those concerning the ban on asbestos and those dealing with exceptions to this ban.

3.260 Canada maintains that the Decree is a “technical regulation” as this expression is defined in Annex 1 to the TBT Agreement. Five of the elements of the definition of a “technical regulation” are found in the Decree. These are: (i) a description of the characteristics of a product, including by the negative; (ii) a description of the related processes and production methods; (iii) a description of the administrative provisions which apply to a product; (iv) the marking or labelling requirements which apply to a product; and (v) mandatory compliance.

3.261 Firstly, Canada points out that the term “characteristic” means “a distinguishing trait, quality or property”.408 In this sense, the Decree describes a distinctive characteristic of a product. Indeed, in accordance with its objective, which is to prohibit asbestos totally, the main provisions of the Decree are intended to prohibit asbestos fibres, regardless of whether they are of the amphibole or chrysotile variety, in the manufacture and processing of materials, products and devices marketed in France. The characteristic of these materials, products or devices, in whose manufacture and processing asbestos fibre may not be incorporated, is thus laid down in the Decree. This characteristic is that no product shall contain asbestos. Secondly, the Decree also covers the processing of all types of asbestos fibres, whether or not they are incorporated in materials or products. In so doing, it imposes restrictions on processes and production methods involving asbestos fibres, including chrysotile fibres. The principle of a total ban is accompanied by a limited number of exceptions. Chrysotile fibres are permitted in the manufacture and processing of products when no substitute fibre deemed less harmful exists. The Decree stipulates that the manufacture of chrysotile-based products is subject to the standards of exposure to asbestos dust in buildings.409 Hence, the Decree lays down a production process for products containing chrysotile as defined by the EC. Thirdly, by instituting a reporting mechanism to obtain or maintain an exemption from the ban on asbestos, the Decree specifies the administrative provisions applicable to products. Canada considers that the EC confirm this point when they describe in detail the registration procedure under the annual reporting process in the context of exemptions from the ban on asbestos. Fourthly, the Decree also deals with requirements regarding marking or labelling of a product. It sets out labelling standards for chrysotile-based products that are the subject of a provisional exemption from the ban.410 Fifthly, compliance with the main provisions of the Decree is mandatory. Products which are marketed in, offered for sale in or imported into
France must not contain asbestos. Compliance with the ban on asbestos is mandatory, unless the public authorities agree to grant a waiver, in which case compliance with the terms and conditions of the exception are mandatory. Fines are applicable in case of non-compliance.

3.262 Canada claims that, by notifying the Committee on Technical Barriers to Trade, the EC recognized that the Decree is a technical regulation and consequently the TBT Agreement applies. Furthermore, the EC provided the Committee on Technical Barriers to Trade with a justification of the Decree based on the disciplines of Article 2.2 of the TBT Agreement. The EC certainly considered this Committee the competent forum to receive the notification and the justification. The EC’s most recent official statement concerning the applicability of the TBT Agreement was made during the consultations of 8 July 1998, when the representative of the EC stated that the provisions of the TBT Agreement were the only provisions invoked by Canada that seemed relevant to contesting the Decree. Canada concludes that, in the light of the foregoing, it is clear that the Decree is a “technical regulation” and, therefore, subject to the TBT Agreement.

3.263 The European Communities contend that the TBT Agreement does not cover general prohibitions. A measure cannot fall within the TBT Agreement unless it satisfies, in particular, the definition of “technical regulation” contained in Annex 1 to the Agreement. The fact that the definition of “technical regulation” is narrow is not a matter of chance, but signifies that the authors intended to limit the scope of the Agreement. This is equally apparent from the text of Article 1.2 of the TBT Agreement itself. The object and purpose of the TBT Agreement, like that of the TBT Agreement of 1979, is to “further the objectives of the GATT 1994” (second preambular paragraph) in the fields of standardization and conformity assessment in order to ensure that technical regulations and standards do not create unnecessary obstacles to international trade (third and fifth preambular paragraphs). According to the EC, it follows from the above that the purpose of the TBT Agreement is to deal with technical regulations and standards. It is not to solve market access problems linked with general prohibitions. However, this does not result in a legal vacuum for measures of this type inasmuch as they continue to be covered by other legal provisions, in particular Article III of the GATT.

3.264 The EC maintain that the Decree does not fall within the scope of the TBT Agreement. As far as asbestos fibres are concerned, it is not the purpose of the Decree to lay down the characteristics of asbestos or of processes and methods of asbestos production. The Decree purely and simply bans asbestos as a product or a raw material. As far as other products are concerned, the Decree means that they may not contain any asbestos. This, however, would not be enough for the Decree to be equated with a “technical regulation” within the meaning of the TBT Agreement. In fact, where these products are concerned, the EC emphasize that the Decree does not identify the products that must not contain asbestos. All products, without more precise identification, are subject to the ban. This applies both to cement and to any other product (for example, tennis rackets). Furthermore, the EC repeat that the Decree is not concerned with the characteristics of other products (whether or not containing asbestos). In fact, the horizontal prohibition on incorporating asbestos in products has the effect of prohibiting, on French territory, products that contain asbestos. The general prohibition eliminates these products from the French market. A technical regulation, on the other hand, presupposes that the product concerned can always be supplied on the market. This is comparable, for instance, to a situation where a domestic law prohibits in general and without discrimination the production, importation and use of drugs, alcohol, etc. on grounds of public health. The EC note that even Canada does not dispute the fact that the declared objective of the Decree is to protect human health. Canada only disputes the existence of a risk in this case, not the intention of the French authorities to protect human health. Thus, in the EC’s view, it cannot be maintained that the Decree lays down the characteristics of a product which no longer exists. The same applies to the production processes and methods for the product, which are linked to its characteristics. The Decree does not lay down production processes and methods for a product which no one any longer has the right to manufacture.
3.265 The EC therefore reaffirm their conclusion that the TBT Agreement is not applica-
table to a measure imposing a general ban on a product, in this case asbestos and asbestos-
containing products. As regards the provisions of the Decree concerning exemptions from
the ban, the EC reiterate that these provisions do not fall within the scope of the TBT Agree-
ment. The EC have already pointed out that the Decree does not define either the technical
characteristics of the products that may be exempted from the general ban adopted. These
products may or may not contain asbestos. In addition, this very limited number of prod-
ucts will anyhow be phased out as soon as substitute products that can ensure a lower
level of risk and guarantee the same security for users become technically available. So,
even in the case of limited and transitional exceptions, the sole preoccupation of the French
authorities has been to protect human health because, for the time being, there are no alter-
native products that can guarantee a lower level of risk to human health (for example, the
risk of accidents if the use of asbestos were to be banned immediately in situations of
exposure to very high temperatures, etc.) The EC add that the Decree does not define
either the production processes and methods for the products that may be exempted from
the general ban adopted. These products may or may not contain asbestos. It follows from
these two observations that the provisions of the Decree concerning exemptions from the
general ban do not fall within the scope of the TBT Agreement either. In fact, it is neither
the purpose nor the effect of the French measure to lay down the characteristics of these
products or to impose production processes or methods for their manufacture.

(b) Article 2.1 of the TBT Agreement

3.266 Canada claims that the Decree is incompatible with Article 2.1 of the TBT Agree-
ment because it subjects chrysotile fibre and chrysotile-cement products imported from
Canada and from any other country to less favourable treatment than like PVA, cellulose
and glass fibres, and like fibro-cement products, of French or foreign origin. Article 2.1 of
the TBT Agreement restates the principles of non-discrimination set forth in Articles I:1
and III:4 of the GATT. In order to determine incompatibility with Article 2.1 of the TBT
Agreement, the Panel must determine: (i) that the measure in question is a technical regu-
lation; (ii) that the products involved are like products; (iii) that these like products are of
French origin or originate in any other country; and (iv) that, by virtue of the technical
regulation, imported chrysotile fibre and chrysotile-cement products are subject to less
favourable treatment than like products. Given that the allegations of incompatibility with
Article 2.1 of the TBT Agreement are being submitted to examination by a Panel for the
first time, Canada maintains that examination of the matter should take into account the
precedents relating to Articles I:1 and III:4 of the GATT 1947 and the GATT 1994.415

3.267 Canada asserts in the first place that, as it has shown in the paragraphs regard-
ing the applicability of the TBT Agreement, the Decree banning asbestos is a “technical
regulation” and is subject to the application of Article 2. Secondly, as Canada has pointed
out in relation to Article III:4416, Canadian chrysotile fibre is like PVA, cellulose and glass
fibre. Similarly, chrysotile-cement products are like fibro-cement products. Thirdly, Canada
contends that there are substitute fibres of French origin that are “like products” to chrysotile
fibre. There are also fibro-cement products of French origin that are “like products” to
chrysotile-cement products of Canadian origin. PVA fibres from China, Korea and Japan
intended for the manufacture of fibro-cement products are also imported into France. Lastly,
in 1997, France imported over 157,000 tonnes of fibro-cement products originating in “any
other country”.417

3.268 Canada contends that chrysotile fibres and the products containing them are
subject to less favourable treatment than substitute products of French or foreign origin.
Following the criteria applied to Article III:4 of the GATT, a technical regulation is incom-
patible with Article 2.1 of the TBT Agreement if this regulation subjects imported chrysotile
fibre and chrysotile-cement products to treatment less favourable than that accorded to
like PVA, cellulose and glass fibres, and to like fibro-cement products of national origin or
originating in any other country.418 The terms of Article 2.1, providing that “treatment no
less favourable” be accorded, are identical to the terms of Article III:4 of the GATT: “shall
be accorded treatment no less favourable”. In Canada’s view, the inclusion in the TBT
Agreement of terms similar to those of the GATT reveals the intention of the Members to have them interpreted in the same manner. Article 2.1 thus includes the same obligation with respect to national treatment as that set forth in Article III:4 of the GATT. Article 2.1 also requires the Members to treat like products respecting the principle of most-favoured-nation treatment. It provides for treatment no less favourable than that accorded to like products originating in any other country to be accorded. The wording of Article 2.1 also repeats in full the concept set forth in the disciplines of Article I of the GATT, namely, that any advantage granted to any product originating in any other country shall be accorded to the like product originating in the territory of all other contracting parties.

3.269 Canada points out that the fact that the expression “no less favourable” was repeated throughout the GATT Agreements was highlighted by the Panel in the case United States – Section 337 of the Tariff Act of 1930. The relevant passage is as follows:

“These words are to be found throughout the General Agreement and later agreements negotiated in the GATT framework as an expression of the underlying principle of equality of treatment of imported products as compared to the treatment given either to other foreign products, under the most favoured nation standard, or to domestic products, under the national treatment standard of Article III.”

The use of the same wording indicates that the disciplines of Article 2.1 of the TBT Agreement are the same as those of Article III:4 of the GATT and that they are applied in the same manner. In order to address the question of whether less favourable treatment is accorded to products of Canadian origin than that accorded to like products originating in any other country, Canada considers that the Panel should examine the criteria of “the actual equality of the opportunities offered”. As Canada has already stressed, the Decree affects areas that are governed by a technical regulation as defined in the TBT Agreement. There is no similar measure on PVA, cellulose or glass fibres nor on fibro-cement products. In fact, it is undeniable that the prohibitions ordered in the French technical regulation with regard to chrysotile fibre and products containing it constitute a denial of the actual equality of the opportunities for competition offered to chrysotile fibre and chrysotile-ce ment products on the French market. The bans do not apply to PVA, cellulose or glass fibres nor to fibro-cement products imported into France. Canada concludes that the Decree banning asbestos is incompatible with the provisions of Article 2.1 of the TBT Agreement because it discriminates against chrysotile fibre and chrysotile products, as opposed to PVA, cellulose and glass fibre and fibro-cement products.

3.270 The European Communities draw attention to the following points presented as an alternative, should the Panel consider that the TBT Agreement is applicable to the French measure, which they nevertheless contest.

3.271 The EC point out that Article 2.1 of the TBT Agreement may be considered as a specific application to technical regulations of Articles I and III of the GATT 1994. The sole argument put forward by Canada is the claim that a measure’s compatibility with Article 2.1 of the TBT Agreement should be examined taking “into account the precedents relating to Articles I:1 and III:4 of the GATT 1947 and the GATT 1994”. As the EC show in the section relating to the application of Article III:4 of the GATT, the Decree does not discriminate between imported products and like national products.

3.272 Canada emphasizes that Article 2.1 of the TBT Agreement can be considered as a specific application to technical regulations of Articles I and III of the GATT 1994 and that the EC are in agreement on this point. Canada refers back to its previous arguments on the issue of the applicability of the TBT Agreement, on the one hand, and on the issue of likeness on the other. These arguments are the same as those made with respect to Article III:4 of the GATT. Canada notes that, unlike the case of Article III:4, the origin of like products is of no consequence for Article 2.1 of the TBT Agreement as this Article covers national treatment and most-favoured nation treatment. Finally, on the issue of the less favourable
treatment of imported products compared to like products, Canada refers to the arguments it puts forward in relation to Article III:4 of the GATT.

(c) Article 2.2 of the TBT Agreement

3.273 Canada emphasizes that Article 2.2 of the TBT Agreement requires the Members of the WTO to ensure that technical regulations are not adopted with a view to or with the effect of creating unnecessary obstacles to international trade. A technical regulation creates an unnecessary obstacle to international trade if its objective is not legitimate or if the regulation is more trade-restrictive than necessary to fulfil this objective, taking account of the risks that non-fulfilment would create. In Canada’s view, in order to determine whether a technical regulation is compatible with Article 2.2, the Panel must first determine if the objective that the regulation is supposed to fulfil is part of the range of legitimate objectives listed in Article 2.2. If the objective is not on this list, the regulation is incompatible with Article 2.2. If it is determined that the objective is part of the range of objectives stipulated in Article 2.2, the Panel must then determine whether the technical regulation is more trade-restrictive than is necessary.

3.274 The European Communities point out that a reading of Article 2.2 reveals two basic criteria enabling a WTO Member to adopt a restrictive technical regulation: (i) there must first be a legitimate objective, such as the protection of human health; and (ii) then the Member’s technical regulation must not be more trade-restrictive than is necessary to fulfil this legitimate objective, taking account of the risks that non-fulfilment would create. According to the EC, the Decree meets those two conditions. The EC refer the Panel to the arguments they put forward in Section III.B and in relation to Article XX(b) of the GATT. A legal analysis relating to Article XX(b) of the GATT also applies in the context of Article 2.2 of the TBT Agreement as far as the test of necessity is concerned.

(i) Legitimate objective

3.275 Canada considers that, in this case, the objective claimed by France, namely, the protection of the health of workers and consumers, corresponds to the objectives identified in Article 2.2.

3.276 The European Communities claim that, as they emphasize in connection with Article XX(b) of the GATT, the aim of the French measure is to halt the spread of the risks associated with the use of asbestos fibres and asbestos-containing products and thus reduce the number of deaths among the French population. The Decree is therefore perfectly in keeping with policies designed to protect human health. The EC note that Canada does not dispute this.

3.277 Canada states that it has acknowledged that the objective sought by France when adopting the Decree, namely, the protection of human health, is one of the objectives included in Article 2.2 of the TBT Agreement. However, contrary to what the EC would like one to believe, Canada did not acquiesce in the argument that the objective is to halt the spread of the risks associated with the use of asbestos fibres and asbestos-containing products. The examples in the list of objectives set out in Article 2.2 of the TBT Agreement include protection of human health and safety. Halting the spread of the risks associated with asbestos can be a way of protecting human health. This is not, however, a legitimate objective within the legal framework of the TBT Agreement. Inasmuch as Canada does not dispute the legitimacy of the objective, understood as being the protection of human health, the question which is submitted to the Panel for examination in this case under Article 2.2 is whether the Decree is more restrictive than is necessary to fulfil this objective, taking account of the risks non-fulfilment would create.

3.278 The European Communities point out that, contrary to Canada’s claims, they have neither changed nor modified the objective of the Decree: the objective in banning asbestos was to stop the spread of the risk of asbestos-related diseases for the purpose of protecting human health so the objective is the same.
“For this purpose, technical regulations shall not be more trade-restrictive than necessary to fulfil a legitimate objective, taking account of the risks non-fulfilment would create ...” (Article 2.2, second sentence)

Canada maintains that the applicable test, for the purposes of determining if the technical regulation is more trade-restrictive than necessary to fulfil the objective claimed by France, consists of two different criteria. The Panel must first determine if the Decree makes it possible for the objective claimed by France to be fulfilled, in other words, if there is a rational link between the Decree and the objective. Then, to the extent that this first criterion is met, the Panel must ask itself if the trade effects of the technical regulation are necessary, taking account of the risks that the absence of this technical regulation would create. For this purpose, it is essential to consider the existence of alternative regulatory approaches, less trade-restrictive, that would make it just as possible to achieve the objective being pursued as the technical regulation being challenged. Canada maintains that the Decree banning asbestos creates an unnecessary obstacle to the international trade in chrysotile fibre and products containing it. First of all, the Decree has no rational link with the objective of protecting the health of workers and consumers sought by France. Secondly, its detrimental effects on trade are not necessary if the actual risks that would exist in the absence of a total ban are taken into account. Controlled use is an alternative likely to be less restrictive to the competition opportunities in the French market. Although a violation of Article 2.2 of the TBT Agreement exists if only one of the two criteria is not met, Canada contends that the Decree imposing the total ban on asbestos does not meet either of the two criteria. Canada adds that the Decree is not in compliance with the applicable international standards, so the EC do not benefit from the presumption in Article 2.5 of the TBT Agreement.

Canada claims that the first question before the Panel is whether there is a rational link between the Decree and the objective of protecting the health of workers and consumers being pursued by France. In other words, does a total ban make it possible to fulfil the objective of protecting people’s health? This stage of the analysis is essential in order to prevent technical regulations that only have tenuous links – or even no links at all – with the objective being pursued from finding any justification. Canada maintains that the Decree does not make it possible to fulfil the objective of protecting the health of workers and consumers, that is to say that there is no rational link between the Decree and the objective, for the following three reasons: firstly, the risks against which the Decree is supposed to protect workers and consumers are linked to uses of asbestos fibres that were already prohibited when the Decree was adopted; secondly, chrysotile fibre and modern products containing it do not present any detectable risk for health; and thirdly, the Decree replaces the undetectable risk of modern chrysotile-based products with the unknown – and perhaps greater – risk of substitute products for asbestos.

Canada states that the first difficulty in terms of the rational link is the fact that asbestos problems are derived from old uses of asbestos fibres such as the use of sprayed asbestos, which was prohibited in France in 1978. The current risks to human health are linked to exposure to asbestos dust released from brittle materials found in asbestos-treated buildings. The Decree does not achieve the protection of the population against the risks of asbestos already installed. Canada considers that there is no rational link between the ban on hard chrysotile products and the protection of the health of workers and consumers because these products do not present any detectable risk for health. Nothing in the INSERM report links a total ban approach to any risk that might be derived from these chrysotile products. This is an essential element in determining the existence of a rational link because France has invoked this report as the only justification for its measure. INSERM has only evaluated hypothetical risks linked to past uses of asbestos, covering all varieties of fibres. The concentrations of asbestos measured in the atmosphere today and on which INSERM was supposed to base its risk projections almost exclusively contain fibres released by brittle materials installed over 20 years ago. Canada maintains that the INSERM report, on which the ban is based, is therefore not relevant to the decision to adopt a policy on a ban. The only epidemiological data recorded in the report concern workers whose illnesses appeared several decades after exposure to asbestos fibres as the health effects of
asbestos take 20 to 35 years on average to emerge. Moreover, the INSERM report explicitly states that almost all deaths attributable to asbestos nowadays are the result of past exposure caused by uncontrolled processing or spraying.\textsuperscript{421} The decision to ban modern chrysotile products does not lead to reduced risks resulting from past uses nor does it even mitigate the damaging effects of brittle materials that are still to be found in certain buildings.

3.282 Canada contends that the second difficulty in terms of the rational link is that INSERM has not evaluated whether the production or use of modern products containing chrysotile poses a risk for the health of workers and consumers. Such a study would undoubtedly have led INSERM to conclude that modern uses of chrysotile do not pose an actual health risk given the negligible emission of chrysotile-cement products. As Canada emphasized in its factual arguments (see Section III.B above), the health risks associated with modern chrysotile products are undetectable. Moreover, Canada notes that the EC have admitted that the practice of the controlled use of chrysotile is effective and appropriate in asbestos extraction and processing industries.\textsuperscript{422} INSERM does not draw any conclusions on the harmless nature of the production and use of dense and non-brittle chrysotile products.\textsuperscript{423} Nowhere is there any discussion on the specific risks related to the banned chrysotile-cement products. However, that is the necessary implication of the French statement according to which “while the principle of controlled use in the asbestos industry can be admitted, on the other hand, it is not possible to control the risk incurred in occupations related to the construction industry and in other occupations concerned”.\textsuperscript{424} (WTO translation) This statement implies, wrongly, that exposure to chrysotile-cement products is hazardous for construction industry workers. In Canada’s view, the context that gave rise to the announcement and the adoption of the Decree proves that, in order to appease public opinion, the French Government used INSERM’s report, which takes into account the consequences of past and prohibited uses of asbestos, in order to assign risks incorrectly to modern uses of chrysotile fibre and products containing it. The Decree does not make any distinction between hard chrysotile products, for which health risks are at undetectable values, and asbestos-based brittle products, used as insulating material in the 1970s. The use of free fibres in insulating products, or products where the fibre is not encapsulated, was prohibited in view of the difficulty of controlling the dust caused by these applications. According to Canada, it is essential to note that in the modern uses of chrysotile fibre, for instance in chrysotile-cement products and chrysotile-reinforced plastics, the fibre is sealed in a matrix and cannot be released into the environment.

3.283 Canada asserts that the third difficulty in terms of the rational link between the order and the protection of health is the blind faith placed in substitute fibres for asbestos. The human health effects of most of the fibres used to replace chrysotile fibre are not known. According to Canada, the data prove that some substitutes that have been the subject of careful studies are often as, or even more, harmful than chrysotile. The total ban on chrysotile fibre and products containing it, and their replacement with substitute products, implies that substitute products are safe. However, INSERM has not been able to determine if these fibres are safe in view of the lack of epidemiological data currently available on this subject. The ban substitutes the undetectable risk of hard chrysotile products for the unknown – and perhaps greater – risk of substitute fibres. A consistent approach to risk management would have required the French Government, in the case of exposure to fibres and substitute products, to try to ensure a degree of health protection equivalent to what it intended through the total ban on chrysotile. In other words, a consistent approach would have required exposure to substitute fibres to be controlled in the same way as chrysotile. Based on the foregoing, Canada maintains that there is no rational link between the Decree and the objective of protecting public health being pursued by France. Firstly, the Decree does not address the true problem of asbestos in France: work on the sprayed asbestos products in place for at least 20 years. Secondly, the Decree ignores the scientific fact that products containing chrysotile imprisoned in a matrix are harmless. Thirdly, the Decree imposes substitute products whose risks are unknown, which in itself puts the objective being pursued by France in jeopardy.

3.284 Canada concludes that, inasmuch as the conditions of the first criterion (namely a rational link between the Decree and the objective claimed by France) have not been met,
the Decree is incompatible with Article 2.2 of the TBT Agreement. Nevertheless, should the Panel conclude that there is a rational link between the Decree and the objective claimed by France, the Decree would still be incompatible with Article 2.2 of the TBT Agreement because, as Canada shows below, its effects are more trade-restrictive than necessary, taking account of the risks non-fulfilment of the objective would create. Canada asserts that an alternative solution less prejudicial to competition opportunities in the French market and with similar effects on the health of workers and consumers was available, i.e. control-led use.

3.285 Canada contends that the detrimental effects of the Decree on competition opportunities are not necessary, in view of the risks that the absence of technical regulations would create. The words “technical regulations shall not be more trade-restrictive than necessary to fulfil a legitimate objective, taking account of the risks non-fulfilment would create” in Article 2.2 of the TBT Agreement imply a necessity criterion. Canada notes that the two factors to be considered when examining necessity are, on the one hand, the risks that the absence of technical regulations would create and, on the other, the existence of a less trade-restrictive alternative measure that would make it possible to attain the objective being pursued just as effectively. As regards whether there are risks involved in not fulfilling the objective being pursued which could result from the absence of the technical regulations in force, the third sentence of Article 2.2 contains a non-exhaustive list of relevant factors that must be taken into consideration. In the context of this dispute, the scientific and technical information available, as well as the intended end-uses of products, in other words, chrysotile fibre and the products containing it, must be considered. By applying this criterion to the facts of this dispute, Canada maintains that the total ban on asbestos, without distinction as to fibres, completely eliminates the French market for chrysotile fibre and products containing it, whereas the practice of controlled use of asbestos fibres, in terms of the types of fibre and their uses, allow fulfilment of the French objective of protecting the health of workers and consumers while authorizing certain safe uses of chrysotile fibre and products containing it. Accordingly, Canada claims that controlled use is an alternative that is less trade-restrictive and makes it possible to fulfil the objective claimed by France in the same way as the Decree. Consequently, if the actual risks of the modern uses of chrysotile are considered in light of the relevant criteria, specifically the scientific information and the intended end-uses, in Canada’s view it becomes clear that the total ban on asbestos is not necessary, inasmuch as there is a less trade-restrictive measure that makes it possible to fulfil the objective claimed by France.

3.286 Canada states that, in the first place, there is no credible scientific information supporting a total ban on asbestos. The scientific information on which the Decree is based are included in INSERM’s report, but this does not constitute a credible basis for justifying a total ban on all varieties and all uses of asbestos. INSERM’s report attempts to use circumstances of high levels of exposure to a mixture of amphiboles and chrysotile in an effort to extrapolate the risks for a population that no longer experiences such exposure. Although this technique can in itself be deemed dubious, INSERM’s report also raises a number of other questions when it is subjected to the criticism of peers in the scientific community. In Canada’s view, INSERM’s report contains conclusions that do not stand up to criticism. First, the report does not study the data on exposure actually sustained by the French population. The INSERM researchers relied on hypothetical data. Moreover, INSERM’s risk assessments do not adequately make the distinction between the effects of amphiboles and the effects of chrysotile. The risk assessments presented as being linked to exposure to chrysotile are often based on parameters of exposure to amphiboles or to amphibole-chrysotile mixtures. The lack of a study of the risks posed by current exposure to chrysotile appears clearly in light of the alarmist conclusions of the report on cancer cases in France attributable to asbestos. These deaths are linked to past occupational exposure and not to current circumstances of exposure to dense products containing chrysotile. The INSERM researchers did not look at exposure to asbestos dust during incidental and/or intermittent work on structures where chrysotile-cement was used. INSERM used a linear model in its evaluation of the risk, extrapolating from cases of low exposure to chrysotile the risk involved in cases of high exposure to amphiboles or mixtures of fibres. However, as recognized by INSERM, the epidemiological data in cases of environmental
exposure and exposure indoors do not indicate any detectable risk for low exposures. Finally, one of the effects of the Decree is to force the use of substitute fibres even though health risks associated with exposure to such fibres have not been addressed by INSERM.

3.287 Secondly, Canada asserts that the intended end-uses of modern products containing chrysotile fibre have not been taken into consideration in the evaluation of the human health risks. However, the intended end-uses of chrysotile, chrysotile-cement products for instance, do not present any detectable health risk. INSERM did not identify any risk linked to the modern uses of chrysotile fibre and products containing it. In assessing the risks, France has not taken into consideration the intended end-uses of modern products containing chrysotile. The known assessments of the risks linked to the modern uses of chrysotile indicate that the risks faced by an individual are so small as to be undetectable. According to Canada, the ILO confirms this as follows: “The handling of products containing asbestos in which asbestos fibres are solidly fixed in a bonding material so that dust cannot form does not present a health hazard.” The assessments of the risks faced by workers who come into contact with chrysotile-cement products once the material is installed indicate that the risks are undetectable.

3.288 Thirdly, Canada contends that the total ban on asbestos is the most restrictive measure from an international trade standpoint. It leads to a complete closure of the domestic market for these products. In France, in the case of asbestos, the ban has caused the entire French market for chrysotile fibre and products containing it to disappear. As the statistical tables in the section on French imports of chrysotile fibre from Canada show, the French market for Canadian exports has disappeared. Canada maintains that there is an alternative solution that makes it possible to fulfil the objective of protecting human health while being significantly less trade-restrictive. This alternative to the ban, i.e. controlled use, does not result in the effects previously described. Canada indicates that the practice of the controlled use of asbestos is based on recognized scientific principles and on an international consensus. Controlled use was available to France as a means of fulfilling the objective of protecting against risks associated with the uses of asbestos, without creating an unnecessary obstacle to international trade. In fact, that was the approach adopted by France until the adoption of the ban. Controlled use is an internationally endorsed regulatory approach based on scientific data applicable to an entire range of risk-related materials. The principles underlying this approach are that only products and materials that can be controlled in such a way that the associated risks are maintained at an acceptable level throughout their entire life cycle are authorized. In the case of asbestos, controlled use involves, in particular, maximum reduction of the quantity of dust released in a work environment, the adoption of specific work methods, the prohibition of amphiboles (crocidolite and amosite) and low-density brittle insulation products and the prohibition of processes or products that release dust. On the other hand, this form of regulation would allow the use of chrysotile fibre and high-density products such as chrysotile-cement. Controlled use is considerably less trade-restrictive than the ban on asbestos. The drop in Canadian chrysotile asbestos exports in 1997, the year immediately following the adoption of the Decree, clearly shows that the controlled use practices that had been implemented in France prior to the ban on asbestos did not create the same obstacles to trade as those created by the total ban on asbestos as soon as it was adopted.

3.289 Canada maintains that the total ban on asbestos, without distinction as to the types of fibre, completely eliminates the French market for chrysotile fibre, even though the practice of the controlled use of asbestos fibres, in terms of the types of fibre and their uses, allows the manufacturing of safe chrysotile products, while having the same effects on human health as the ban. The detrimental trade effects of the ban are not necessary in order to fulfil France’s objectives just as effectively as the Decree. The measures based on controlled use that existed in France at the time of the announcement of the ban, would make it possible to fulfil the objective of protecting human health without creating unnecessary obstacles to trade. Canada considers that the excessive effects of the ban can only be explained by the political desire of the French Government to respond in a spectacular way to the pressure of public opinion. Canada claims that the Decree banning asbestos is incompatible with Article 2.2 of the TBT Agreement. Firstly, the Decree does not make it
possible to fulfil the objective being pursued by France, that is the protection of the health of workers and consumers. Secondly, the detrimental effects of the total ban are not necessary in view of the actual risks resulting from the modern uses of chrysotile. The ban has done nothing more to protect health than the controlled use already in place. The only uses allowed at the time of the Decree’s adoption were the use of chrysotile fibre in dense materials, which, according to the available scientific information and considering the end-uses, did not pose – and still does not pose – detectable risks for people’s health. Canada therefore concludes that the Decree is an excessive measure because controlled use is a less trade-restrictive alternative that makes it possible to fulfil the objective being pursued by France.

3.290 The European Communities respond that the distinction made by Canada between, on the one hand, the need to determine whether the Decree permits the fulfilment of the objective cited by France (“rational link” in Canada’s own words) and, on the other, whether the effects of the technical regulation are necessary, taking account of the risks that non-fulfilment would create, is artificial and at variance with the actual text of Article 2.2. of the TBT Agreement. The wording of Article 2.2 shows that the test laid down therein is in line with that used in connection with Article XX of the GATT and developed by Panel practice. In following the terms of Article 2.2 of the TBT Agreement, according to the EC, a dual examination has to be carried out: (i) determine whether the measure is the only one that allows the objective set by the Member to be attained, or whether there is a less restrictive measure whereby this objective can also be achieved; (ii) assess the risks which a failure to take the measure concerned would create, taking account, in particular, of available scientific and technical information or end-uses of products. The EC claim that, applied to the Decree, these two criteria show that the measure is compatible with Article 2.2.

3.291 The EC note that Canada repeats throughout its submission that a less restrictive measure (namely the “safe” use of asbestos) would enable the objective chosen by the French authorities to be achieved. However, the factual arguments of the EC, like their comments relating to Article XX(b) of the GATT, emphasize that “safe” use is: (i) insufficient to halt the spread of the risks linked to exposure to asbestos in the production and processing industries, even though the number of workers in those industries is limited and they are therefore, in principle, easy to supervise and monitor (this means an excess of mesotheliomas even with “safe” use); (ii) ineffective in halting the spread of the risks linked to occasional, and often unwitting, exposure to asbestos. The principle of “safe” use cannot apply where the risks affect a range of very varied occupations operating in a wide variety of situations (this means that it is genuinely impossible to ensure “safe” use among do-it-yourself enthusiasts and those exposed to para-occupational risks). For further details, the EC refer the Panel to the comments made in the factual section and those relating to Article XX(b) in their legal arguments. On that basis, the EC consider that the Panel should also conclude that the prohibition of asbestos and asbestos-containing products is the sole measure that will enable the objective set by the French authorities to be achieved.

3.292 The EC claim that failure to ban asbestos and asbestos-containing products would create risks for human health. The EC assert that, even though the burden of proof lies on Canada, the latter has done nothing to show that the replacement of the horizontal prohibition by “safe” use would not create risks to human health. In the EC’s view, the risks that would be created by a failure to impose a ban are evident in the light of the available scientific information, the available technical information and the end-uses of asbestos and asbestos-containing products.

3.293 The EC maintain that the consequence of using asbestos and asbestos-containing products is to expose certain sections of the population to asbestos to a degree which scientific studies recognize as dangerous to human health. The spread of the risks linked to such exposure cannot be halted by “safe” use. In the mining and processing industries, in principle the easiest to monitor, the limits to the “safe” use of asbestos are already apparent. For example, the 1996 study of the British HSE, mentioned in the factual section, finds a significant excess of deaths due to mesothelioma among workers who began working in asbestos mines after the introduction of “safe” use. The EC note that, throughout its sub-
mission, Canada considers that asbestos inhalation is a threat only to production and processing workers and affects several thousand persons at the most. In the para-occupational and domestic context, however, hundreds of thousands of persons are exposed to asbestos (very often unwittingly) and may even be subject to exposure levels greatly in excess of the current limit values for asbestos dust. For example, as the EC indicated in Section III.B, a roofing worker using a grinder in the open air to repair corrugated roof sheeting made of asbestos-cement is exposed to a peak level of 41 f/ml, 410 times in excess of the limit value. The EC point out that a 1992 study by the Quebec CSST shows that the risk of mesothelioma has been rising steadily in Canada since 1967, chiefly among repair and maintenance workers. This finding is even more relevant to those persons exposed to asbestos inhalation in a non-occupational context. The publication by Camus et al. shows a net excess of mesotheliomas in women living near chrysotile asbestos mines in Quebec, seven times higher than the level found among other women in Quebec. Canada never quotes this study.

3.294 The EC point out that, as they indicated in the factual section (see Section III.B) and in their comments on Article XX of the GATT, all these findings were taken into account in the INSERM Report. These findings are also in line with the most recent scientific studies. In the EC’s view, Canada cannot discharge the burden of proof by trying to justify the “safe” use of asbestos by reference to texts which are over 15 years old and do not guarantee an adequate level of protection, given the health objectives adopted by the vast majority of countries. The EC emphasize in this connection that recent texts, not quoted by Canada, confirm the ineffectiveness of such “safe” use. This can be clearly seen in the 1998 WHO report, which states:

“Some asbestos containing products pose particular concern and chrysotile use in those circumstances is not recommended. These uses include friable products with high exposure potential. Construction materials are of particular concern for several reasons. The construction industry work force is large and measures to control asbestos are difficult to institute. In-place building materials may also pose risk control to those carrying out alterations, maintenance and demolition. Minerals in place have the potential to deteriorate and create exposures.”

3.295 The EC state that the technical information indicates that “safe” use cannot halt the spread of the risks linked to exposure to asbestos, in particular where persons are unwittingly exposed (repeatedly or occasionally). The fact that “safe” use cannot halt the spread of the risk follows logically from the fact that it would be completely unrealistic to apply to the general public the extremely stringent rules applicable to a small “targeted” population (primary users). In fact, as the EC have already stated, it is impossible to implement “safe” use effectively when hundreds of thousands of persons are exposed every day in areas of activity with little supervision in terms of health, such as the construction industry, which accounts for at least 25 per cent of the cases of mesothelioma occurring in France, not to mention the millions of do-it-yourself enthusiasts unwittingly exposed to asbestos inhalation when performing everyday tasks such as cutting. As regards the specific means of ensuring the effectiveness of “safe” use, the EC point out that Canada, even though it bears the burden of proof, considers that the risk of asbestos fibres escaping during cutting or sawing of asbestos-containing products would not exist if items were supplied “pre-cut”.

3.296 The EC note that Canada also refers to an ISO standard which is over 15 years old. Canada states that “… the cutting of plates or tiles for roof covering is not a source of emission if the simple techniques of standard ISO-7337 are followed”. The EC note that these “simple” techniques include “the use of chains that break the pipes by the effect of pressure, low-speed saws and saws equipped with a vacuum extractor, as well as the moistening of the materials prior to any action”. Use of these techniques also presupposes that do-it-yourself enthusiasts will put on a “diver’s suit” before starting any operation that will bring them into contact with asbestos. In reality, these techniques are not sufficient to halt the spread of the risks. The EC emphasize that, in the real world, not all items are pre-cut. In the real world, the thousands of people who are unwittingly exposed to asbes-
tos inhalation will not put on protective clothing to perform everyday tasks. In the EC’s view, the fact that Canada is unable to demonstrate any specific or realistic methods of halting the spread of the risk linked to asbestos exposure, without a total ban on asbestos and asbestos-containing products, is due to the absence of any scientific basis. Once products are placed on the market, there is no longer any realistic means of monitoring the use of asbestos, and in particular the everyday operations (cutting, sawing, etc.) in which many persons may be engaged. The EC maintain that the “safe” use of asbestos advocated by Canada is therefore inapplicable and does not enable the legitimate objective set by France to be achieved. Furthermore, the EC note that Canada states that the French Decree means “blind recourse” to substitute fibres. According to the EC, this statement reflects a misunderstanding of the French legislation. In practice, manufacturers carry out technical tests with a view to replacing asbestos products by substitute products. If these tests are inconclusive, and if the manufacturers can demonstrate that there are no substitute products with the same technical characteristics as asbestos, they can submit an application for a waiver so that they can continue to use asbestos. The Decree provides for such waivers, which are granted after a detailed assessment.

3.297 Concerning the intended end-uses of the products, the EC note that Canada maintains that the “modern” uses of chrysotile fibre are such that the fibre is bound in a matrix and cannot be released into the environment. The EC point out in this connection that Canada is claiming that this is an innovation when it is not. As the EC have already indicated, for the past forty years the production of asbestos-cement has consisted of “encapsulating” asbestos in cement (10 percent of asbestos fibres in 90 percent of cement). The EC emphasize that, in talking of “modern” use or of “modern” products, Canada is misleading the Panel by trying to promote a “clean” image of chrysotile asbestos; although “chrysotile asbestos” sounds better than “amphibole asbestos”, chrysotile is also classified by the WHO as a “category I” product known to cause cancer in man. The Panel should know that the “encapsulation” in question by no means guarantees the harmlessness of asbestos-cement, for example. Once the use of asbestos-cement is authorized, it is no longer possible to monitor such use. A variety of operations such as cutting, sanding, crushing or sawing are bound to be applied to asbestos-cement, in the occupational, para-occupational and domestic contexts. During these operations, large numbers of carcinogenic fibres will be released in the form of dust. Consequently, the EC conclude that the “modern” use of chrysotile asbestos, to which Canada refers in its submission, is illusory and can only conceal the grave risks linked to the use of asbestos-cement, in particular by persons who are subject to unwitting, repeated or occasional exposure.

3.298 Canada emphasizes that, once it is acknowledged that an objective is legitimate, it must be determined whether the technical regulation constitutes a rational and necessary measure. A measure is said to be rational if it is carefully designed, based on the legitimate objective. It must be neither arbitrary nor based on irrational considerations. A measure is said to be necessary if it is not more trade-restrictive than is necessary to fulfil the legitimate objective, taking account of the risks non-fulfilment would create. The measure must therefore have a minimal impact on trade. The risks that the absence of a measure would entail must also be assessed. The foregoing is based directly on the text of Article 2.2, particularly the second sentence. In Canada’s view, the application of this provision requires that the following three questions be answered: (i) does the technical regulation make it possible to “fulfil” the “legitimate objective” (rational measure); (ii) is the technical regulation “more trade-restrictive than necessary” to fulfil the legitimate objective (necessary measure); and (iii) have the alleged harmful effects (“the risks non-fulfilment would create”) been assessed in light of the “available scientific and technical information” and the “intended end-uses of products”? Unlike the United States, Canada considers that the obligation under Article 2.2 does not have to be interpreted in a similar way to Article 5.6 of the Agreement on the Application of Sanitary and Phytosanitary Measures (“SPS Agreement”). Article 5.6 of the SPS Agreement refers to a specific test that the United States suggests be transposed to Article 2.2 of the TBT Agreement, even though the terms of each Article are different, calling for separate approaches. To cite only one example, contrary to the footnote to which Article 5.6 of the SPS Agreement refers, Article 2.2 of the TBT Agreement does not require the demonstration of the existence of another applica-
ble measure that is “significantly” less restrictive to trade. Canada notes that the EC subscribe to this viewpoint as they propose to interpret the criterion of necessity in Article 2.2 of the TBT Agreement in a similar way to the criterion of necessity in Article XX of the GATT 1947 and 1994.

3.299 Canada considers that the question that must be answered is whether the technical regulation embodied by the Decree has been carefully designed to fulfil France’s objective? This stage of the analysis is essential in order to prevent the justification of technical regulations that only have tenuous links with the stated objective. Canada maintains that there is no rational link between the Decree and France’s objective and that the total ban on asbestos does not allow this objective to be attained for three reasons: (i) the Decree does not address the real problem of asbestos in France, which is the amount of contact with friable materials containing amphiboles that have been in place for at least 20 years; (ii) the Decree ignores the indisputable scientific evidence concerning the safety of products in which chrysotile fibres are trapped in a matrix; and (iii) the Decree imposes substitute products whose risks are unknown, thereby compromising France’s objective. These three points will be addressed in turn.

3.300 Regarding the first point, Canada contends that the risks associated with asbestos in France are attributable to amphibole asbestos and to sprayed-on applications, both of which are no longer used. When setting out its arguments on controlled use and workers in the servicing and maintenance sector in the factual section (Section III.B above), Canada emphasized that, in the industrialized countries, it is the friable asbestos products inherited from the past which represent by far the major potential for exposure and risk today and in the years to come. That is the situation in France, where the health problems associated with asbestos are essentially attributable to friable materials containing amphiboles (or a mixture of amphiboles and chrysotile), used in the 1950s, 1960s and 1970s in processes such as spraying and insulation. The French Académie nationale de médecine came to this conclusion earlier, in April 1996, in its report on the protection of the populations exposed to the inhalation of asbestos fibres in public and private buildings. The relevant passage reads as follows:

“A number of epidemiologists have found that unprotected contacts with the parts of buildings containing asbestos expose, even today, too many professionals to diseases comparable to those described prior to the current regulations. This is unfortunately a simple statement of the fact that improper working conditions pose the same hazards as the working conditions of the 1950s.”

3.301 According to Canada, inasmuch as the hazard already exists, France, like the other industrialized countries, cannot prevent it. France has already ordered a mandatory survey of apartment buildings containing sprayed-on asbestos. Construction workers will be able to refer to this register and take the appropriate measures. France banned spraying in 1978 and amphiboles in 1994. The Decree therefore has the effect of prohibiting chrysotile-cement and other non-friable chrysotile-based products. In Canada’s view, it is obvious that the Decree does not in any way constitute a solution to France’s public health problem, which is associated with amphiboles and friable products containing asbestos. Canada maintains that there is no rational link between the objective, which is to solve the health problems caused by the large quantities of friable asbestos materials in place in existing buildings, and the Decree, which is obviously not designed to fulfil this objective.

3.302 Regarding the second point, Canada states that chrysotile fibre and high-density non-friable products do not pose any detectable health risk. Nor is there a rational link between the ban on high-density products and the protection of health because these products do not pose any detectable health risk. Today, 97 per cent of chrysotile fibres are used in high-density non-friable materials. Asbestos-cement products account for 90 per cent of the world market. Brake linings represent about 7 per cent of the market. Various products, such as plastics, seals, etc. account for the remaining 3 per cent market share. These products are non-friable and only chrysotile fibres are used. In these products, the fibres
are firmly bound physically and chemically to the matrix of the composite and cannot easily be released in biologically significant concentrations. Canada contends that the health risks associated with these non-friable high-density chrysotile products are undetectable. The word “undetectable” here means “below the limit of detection”. This expression means that, by using the most recent methods and techniques and the most rigorous statistical analyses, the risk associated with conditions of exposure is so low, if there is even any risk at all, that it is “below the limit of detection”. The source of the asbestos risk which the French want to stop from spreading does not come from these high-density chrysotile-based products. In Canada’s view, there is no rational link between the objective of protecting human health and the Decree, whose objective is to prohibit the marketing, sale and importation of non-friable high-density chrysotile products because their health risks, if any, are undetectable.

3.303 Thirdly, Canada claims that the Decree replaced the “known” and undetectable risk of non-friable chrysotile-based products with the “unknown” risk of substitute fibres, thereby increasing the risk to human health. The absence of a rational link is even more obvious if it is considered that the Decree encourages the replacement of chrysotile by substitute fibres. As a result of the Decree, the undetectable risk associated with high-density chrysotile-based products has been replaced by the unknown risk of substitute fibres. The ban on chrysotile asbestos thus creates a false sense of security in the population, lowering the vigilance of workers and of the public who are the most likely to come into contact with the substitute products. Canada notes that the EC claim that the harmlessness or less harmful nature of substitute fibres is proven. As it emphasized in its arguments concerning substitute fibres, Canada maintains that not only does this claim not stand up to analysis, but the description given by the EC is incorrect. The EC are endeavouring in this manner to justify the irrationality of the total ban on chrysotile fibres and their ill-considered replacement by substitute fibres. According to Canada, the WHO is of the opinion that replacement must be done whenever it is possible to replace chrysotile fibres with “safe substitute materials”. Canada recalls that on several occasions it has emphasized that the chrysotile substitutes used in France are not “safe” products or products whose “harmlessness” or “less harmful” nature are proven. Nor is chrysotile asbestos replaced by products which, according to the ILO, should be “harmless or less harmful”. The fact that substitute products do not contain asbestos is not sufficient grounds to believe that these products are safe or less hazardous. In many cases, fibrous substitute materials are assumed to be safe by users and safety measures such as dust control and removal are not observed. Canada considers that banning a material whose risks are known, and even over-estimated, and replacing it indiscriminately with substitutes whose effects are unknown seems to be an odd choice given France’s objective of protecting public health. Canada maintains that there is no rational link between the objective of protecting human health and the Decree, whose effect is to replace a product - chrysotile - whose risks are known and undetectable with substitutes - fibres or products - whose risks are unknown. The Decree does not result in increased protection of public health; it contributes rather to fostering in the population a false sense of security and potentially a greater risk to human health.

3.304 Canada concludes that, inasmuch as it has no rational link with France’s objective, the Decree is inconsistent with Article 2.2 of the TBT Agreement. However, in the event that the Panel should find that there is a rational link between the Decree and France’s objective, the Decree would nonetheless be inconsistent with Article 2.2 of the TBT Agreement because, as demonstrated below, its effects on trade are more restrictive than is necessary, taking account of the risks that non-fulfilment of France’s objective would create. An alternative solution that is less trade-restrictive and fulfils the same objective of protecting human health was available, namely controlled use.

3.305 Canada maintains that the prejudicial effects of the Decree on trade are not necessary, taking account of the risks that the absence of a technical regulation would create. The terms of Article 2.2 of the TBT Agreement - “technical regulations shall not be more trade-restrictive than necessary to fulfil a legitimate objective, taking account of the risks
non-fulfilment would create” - suggest the application of a criterion of necessity. The two factors to be considered in examining necessity are, on the one hand, the risks that the absence of a technical regulation would create and, on the other, the existence of a less trade-restrictive alternative measure that would make it possible to fulfil the stated objective just as effectively. According to Canada, the Panel must therefore first determine what risks the absence of a Decree would create. To assess these risks, the relevant factors to take into consideration include the available scientific and technical information and the intended end-uses of the products. In this respect, Canada is of the opinion that the protection of human health would be just as effectively ensured if the ban on chrysotile prescribed by the Decree were replaced by controlled use measures, considering the undetectable risks of the high-density chrysotile-based products which account for virtually the entire market. To determine whether the effects of the technical regulation on trade are necessary, taking account of the risks that its absence would create, Canada considers that it is essential to consider the existence of less trade-restrictive alternatives that would make it possible to fulfill the objective just as effectively as the disputed measure. In examining whether the prejudicial effects of the technical regulation are appropriate given the stated objective, the Panel must consider the existence of a less trade-restrictive alternative solution that would make it possible to fulfill the desired objective just as effectively. In Canada’s view, the controlled use of asbestos is a less trade-restrictive alternative which, nonetheless, makes it possible to fulfill the objective of protecting human health.

3.306 Canada contends that the Decree is not based on scientific information. It notes that there are two types of “asbestos” risk in France: (i) the risk of diseases that might occur among those exposed while working with modern high-density products containing chrysotile; (ii) the risk of diseases that might occur among persons working in buildings containing friable asbestos materials. The risks and diseases associated with friable materials that were put in place in France using a process – spraying – that is now banned are not affected by the Decree and are therefore not the subject of this debate. Indeed, France banned spraying, amphiboles and friable products well before the date of the Decree. The only risks and diseases concerned by the Decree are those associated with contact with modern high-density products containing chrysotile. None of the “available scientific and technical information” supports the total ban on asbestos (in fact, chrysotile asbestos) adopted by France. Canada maintains that the risk assessment used by France as a basis for the ban on chrysotile – the INSERM report – does not take into consideration the scientific information concerning products in which chrysotile fibres are solidly set in a binder in such a way that no dust can be released. Furthermore, using models derived from exposures to high doses of asbestos dust containing amphiboles, INSERM claims to make an assessment of the risks of modern applications of chrysotile. It assumed that the risk is proportional to the exposure for levels considerably lower than those that have been the subject of epidemiological study. Canada considers that such extrapolations are scientifically unfounded. According to the INSERM group of experts, this is the “most plausible uncertain” risk assessment.

3.307 Canada states that the figures obtained by INSERM based on these models are average values subject to large variations. INSERM did not consider the risks of the intermittent or discontinuous exposures that occur while working with high-density products in which chrysotile fibres are incorporated, such as asbestos-cement, the composite in which more than 90 per cent of chrysotile fibres imported into France were incorporated before the adoption of the Decree. In this application, chrysotile fibres are solidly bound to the cement in such a way that no asbestos dust is released into the ambient air. Most manufacturers now offer products pre-cut and pre-drilled in the plant. Special techniques such as pre-cutting are used to avoid any cutting operations. When cutting or drilling cannot be avoided, work methods exist which make it possible to limit the dust emissions to a level significantly lower than the safe exposure values. This is also true for the other modern chrysotile-based products in which the fibres are incorporated or encapsulated in a matrix. According to Canada, the EC have not demonstrated that chrysotile fibres encapsulated or incorporated in a cement matrix or a plastic composite pose a health risk. Canada states that it has refuted all the studies on which the EC rely in order to argue that controlled use
is not effective in keeping risks to human health to an undetectable level during all stages of the life cycle of modern chrysotile-based products (extraction of chrysotile, processing, incorporation in manufactured products, distribution, sale, use and disposal). On the other hand, credible scientific information indicates that emissions of asbestos dust from non-friable high-density chrysotile-based products are minimal and represent, for those exposed, an undetectable level of risk. Canada contends that INSERM did not correctly assess the risk to human health from the manufacture or use of high-density chrysotile-based products. Such a study would undoubtedly have prompted INSERM to conclude that modern uses of chrysotile do not pose a genuine health risk.

3.308 Canada maintains that high-density chrysotile-based products do not pose a risk to human health in light of their intended end-uses. Canada also claims that France dismissed controlled use, a practice less trade-restrictive than a total ban on chrysotile, even though controlled use is equally effective in protecting human health. Canada emphasizes that the array of measures aimed at controlling the use of asbestos in place in France when the total ban on asbestos was adopted enabled the objective of protecting human health to be attained without creating an unnecessary obstacle to trade. The French legislative provisions on controlled use were quite elaborate. Several measures had been taken by the public authorities, including: (i) a ban on sprayed-on applications of asbestos for all buildings; (ii) a ban on the marketing, use and importation of all varieties of amphiboles; (iii) rigorous control of the average concentrations of asbestos fibres in the workplace (exposure limit value of 0.1 f/ml); (iv) a ban on an entire range of products containing asbestos; and (v) a mandatory survey of apartment buildings containing sprayed-on asbestos and insulation. This array of measures offered optimal protection of human health.

3.309 Canada claims that France did not have to ban asbestos completely. It merely had to continue to apply the measures already in place. In fact, the use of relatively simple methods and rules is sufficient to ensure safe use of chrysotile at each stage of its life cycle, as can be seen in the case of chrysotile-cement, which accounts for virtually all of the world market for products in which chrysotile is incorporated. Canada notes that controlled use makes it possible, to borrow the expression of the EC, to halt the spread of the risk associated with exposure to chrysotile in the production and processing sectors. The following control measures are effective in maintaining emissions of chrysotile dust below the permitted limit values: dust control equipment, regular verification of chrysotile concentrations, wet process, confinement, policy on work clothing, showers and medical examination. Similarly, during the installation and maintenance of chrysotile-cement products, controlled use affords workers a level of protection corresponding to an undetectable risk. If the methods prescribed by Standard ISO-7337 are followed, the exposure levels generated during cutting, sawing, drilling or sizing operations on chrysotile-cement products can be maintained well below values that represent a detectable risk. Canada claims that the total ban on asbestos constitutes the most restrictive possible measure from the standpoint of international trade. It has resulted in the total closure of the French market to chrysotile asbestos. In the case of high-density chrysotile-based products, controlled use is entirely practicable, effectively protects human health and has a less drastic effect on international trade. That is why controlled use is a preferable alternative to a ban in the context of Article 2.2 of the TBT Agreement. Canada states that when, in accordance with Article 2.2 of the TBT Agreement, the risks of modern uses of chrysotile are taken into consideration in light of the scientific information and the intended end-uses of the products, the total ban on asbestos is not justifiable in terms of this Article because controlled use is less trade-restrictive and makes it possible to fulfil the objective of protecting human health.

3.310 Canada maintains that the preamble to the TBT Agreement cannot be invoked to justify non-compliance of a technical regulation with Article 2.2. The preamble to the TBT Agreement cannot be used to justify the Decree. The preamble outlines the goals and rationale of a treaty. It does not confer any rights and does not impose any obligations. In the case of the preamble to the TBT Agreement, the statement that the Members can choose the measures necessary for the protection of health is explicitly limited by the obligations contained in the Agreement. Paragraph 6 of the preamble to the TBT Agreement stipulates
that the principles outlined therein (notably that no country should be prevented from
taking "...measures necessary for the protection of human ... health") are subject to two
requirements. First of all, the measures in question must not constitute a means of arbi-
trary or unjustifiable discrimination between countries where the same conditions prevail
nor a disguised restriction on international trade (in this connection, see also Canada’s
arguments in the section dealing with Article XX(b) of the GATT). Secondly, the measures
adopted must comply with the obligations contained in the TBT Agreement, notably Arti-
cle 2.2. In Canada’s view, it is clear that the preamble cannot be invoked to justify a techni-
cal regulation which, like the Decree, is inconsistent with Article 2.2.

3.311 With regard to the precautionary principle, to which the EC refer, Canada notes that it
was recently mentioned by the Appellate Body in Japan – Measures Affecting Agricultural
Products that the preamble and Articles 3.3 and 5.7 of the SPS Agreement refer to the pre-
cautionary principle. However, again according to the Appellate Body, the precautionary
principle cannot, in itself, justify a violation of any of the obligations of the SPS Agreement.
If this is true in the case of the SPS Agreement, it is even more true in the case of the TBT
Agreement, which cites this principle much more vaguely. Nor can the precautionary
principle be invoked to justify the attainment of a zero risk.431

3.312 Canada concludes that the Decree is inconsistent with Article 2.2 of the TBT
Agreement. The Decree does not make it possible to fulfil France’s objective, namely, bet-
ter protection of human health. The prejudicial effects on trade of the total ban prescribed
by the Decree are neither necessary nor justifiable. The ban has done nothing more to
protect health than the controlled already in place. The only uses allowed at the time of the
Decree’s adoption were those in which chrysotile fibres are incorporated in high-density
materials. Yet these materials, according to the available scientific information and consid-
ering their end-uses, at the time of the adoption of the Decree did not pose - and still do not
pose - a detectable health risk. To summarize, the Decree is an excessive measure in view
of the fact that controlled use is a less trade-restrictive alternative that enables the French
objective of protecting human health to be attained. Neither the preamble to the TBT Agree-
ment nor the precautionary principle can justify the measure taken by the French Govern-
ment in breach of the obligations contained in the TBT Agreement.

3.313 The European Communities respond that it is important to be aware that a
Member of the WTO can establish the level of health protection it deems appropriate in its
territory. In this connection, the EC point out that, for example, the sixth paragraph of the
preamble to the TBT Agreement clearly states that Members are free to choose the level of
health protection they deem appropriate. This principle was noted by the Appellate Body,
which pointed out that:

"WTO Members have a large measure of autonomy to determine their own poli-
cies on the environment (including its relationship with trade), their environ-
mental objectives and the environmental legislation they enact and implement.
So far as concerns the WTO, that autonomy is circumscribed only by the need to
respect the requirements of the General Agreement and the other covered agree-
ments."432 [italics in the original]

3.314 The EC claim that the Members of the WTO benefit from a similar "large meas-
ure of autonomy" in the field of human health protection. The question might arise whether
this appropriate level is limited by the word “necessary” or by the fact that the measure
must not be applied in a manner that would constitute a means of arbitrary or unjustifiable
discrimination between countries where the same conditions prevail or a disguised restric-
tion on international trade. In this respect, the EC consider it important to make a distinc-
tion between the “level” deemed appropriate by the Member and the “measure” taken by
that Member to achieve the chosen level. The EC note that, in the context of Article XX(b)
of the GATT, all the panels which have examined the concept of necessity have concluded
that it was not the necessity of the objective pursued by the measure concerned that should
be examined but whether or not it was necessary to submit the imported products to the
measure contested.431 In this connection, the Panel on United States - Restrictions on Imports
of Tuna stated that:
“... Article XX(b) allows each contracting party to set its own human, animal or plant life or health standards. The conditions set out in Article XX(b) which limit resort to this exception, namely that the measure taken must be ‘necessary’ and not ‘constitute a means of arbitrary or unjustifiable discrimination or a disguised restriction on international trade’, refer to the trade measure requiring justification under Article XX(b), not however to the life or health standard chosen by the contracting party.”

3.315 According to the EC, it follows from the above that whereas the trade measure that makes it possible to achieve the desired objective must satisfy certain conditions, there is no restriction on the level of protection chosen by the Member. Accordingly, the EC consider that France was free to choose the level of protection it deemed appropriate in the present case, i.e. to halt the spread of the risk linked with the use of asbestos fibres and products containing such fibres.

3.316 The EC maintain that the test of “necessary” in Article 2.2 of the TBT Agreement is a formalization of the previous practice relating, in particular, to Article XX(b) of the GATT. The EC would, however, like to point out that, even though the test of necessity within the context of Article 2.2 of the TBT Agreement corresponds, in particular, to the test in Article XX(b) of the GATT, it nonetheless remains true that the burden of proof, within the context of the TBT Agreement, lies with the party which invokes a specific provision of the latter. As the EC pointed out above, the Panel on United States - Gasoline introduced into the GATT case law concerning the test of necessity established in the GATT 1947. In particular, the Panel noted that:

“... the term ‘necessary’ had been interpreted in the context of Article XX(d) by the panel in the Section 337 case which had stated that: a contracting party cannot justify a measure inconsistent with another GATT provision as ‘necessary’ in terms of Article XX(d) if an alternative measure which it could reasonably be expected to employ and which is not inconsistent with other GATT provisions is available to it. By the same token, in cases where a measure consistent with other GATT provisions is not reasonably available, a contracting party is bound to use, among the measures reasonably available to it, that which entails the least degree of inconsistency with other GATT provisions.”

3.317 The EC note that the Panel on Thailand – Cigarettes adopted the same reasoning when examining a measure under Article XX(b). That Panel saw no reason not to adopt the same interpretation of “necessary” under Article XX(b) as under Article XX(d), stating that:

“The import restrictions imposed by Thailand could be considered to be ‘necessary’ in terms of Article XX(b) only if there were no alternative measures consistent with the General Agreement, or less inconsistent with it, which Thailand could reasonably be expected to employ to achieve its health policy objectives.”

3.318 The EC consider that the concept of necessity under Article XX(b) of the GATT is similar to that in Article 2.2 of the TBT Agreement. The criterion of necessity under Article 2.2 of the TBT Agreement is also based on whether or not the measure adopted is more restrictive than necessary to fulfil a legitimate objective. In this sense, Article 2.2 echoes the test of necessity in Article XX(b) of the GATT which involves, inter alia, examining whether a measure consistent or less inconsistent with the GATT (and hence less restrictive) is available and could be employed to fulfil the Member’s objective.

3.319 As for the second sentence of Article 2.2 of the TBT Agreement, which reads “… taking account of the risks non-fulfilment would create”, the EC consider that, here again, this is an integral part of the implementation of the test of necessity under Article XX(b) of the GATT. In fact, according to the EC, a restrictive measure is “necessary” only if there are risks associated with the non-adoption of the measure in question. The purpose of this
sentence is to deny Members the possibility of adopting measures, under cover of protecting human health, without having taken into account the risks associated with the use of the product banned. Such risks should be assessed, in particular, on the basis of the available scientific and technical information. Clearly, then, the degree of necessity depends on the nature of the objective, as well as on the risks its non-fulfilment would create. As the EC claimed, under the so-called “safe” use policy there continues to be a significant excess of deaths in the asbestos production and processing sector. Moreover, this policy does not provide protection for the secondary populations exposed to asbestos (carpenters, electricians, do-it-yourself enthusiasts, etc.), although the number of deaths caused by asbestos in these groups has steadily increased and now accounts for more than a quarter of deaths due to asbestos.

The EC note that France, through the INSERM report, analysed the risks associated with the use of asbestos when it adopted its Decree. Furthermore, the scientific evidence predicted an increase in the annual number of deaths by mesothelioma in France up to the year 2020. It is estimated that in France the total number of deaths by mesothelioma over the whole of the period 1996-2020 will be 20,000 for men and 2,900 for women. Furthermore, the study by Peto et al., which analyses mesothelioma mortality in England and Wales during the period from 1979 to 1990, shows that about 95 per cent of all the deaths during that period involved workers belonging to the “secondary” user group. It follows that, in the present case, the risks created by non-fulfilment are mortal risks. Accordingly, as so-called “safe” use is inapplicable and ineffective, the EC maintain that banning asbestos and asbestos-containing products was the only measure that could be employed to achieve the level of protection deemed appropriate by France, which is to halt the spread of the risk associated with the use of this product. The EC therefore conclude that the test of necessity is the same under Article XX(b) of the GATT 1994 as under Article 2.2 of the TBT Agreement.

The EC point out, however, that the fact that the test of necessity should, in substance, be applied in the same way in each of the above-mentioned provisions does not mean that the distribution of the burden of proof is the same under each of these provisions. Within the context of Article 2.2 of the TBT Agreement, as distinct from Article XX(b) of the GATT, the burden is on the complaining party to first establish a violation. Article 2.2 of the TBT Agreement cannot be understood as an exception to another provision of the TBT Agreement. Considering the structure and context of the TBT Agreement, the Report of the Appellate Body in the Hormones case is particularly relevant. In fact, the complaining Member must first demonstrate the availability of a consistent or less inconsistent alternative measure that can be employed to achieve the level of protection deemed appropriate by the defending Member. The EC consider that Canada has not shown that the French measure was not necessary, within the meaning of Article 2.2 of the TBT Agreement, to protect human health in accordance with the level of protection deemed appropriate by France.

Canada contends that, contrary to the requirements of Article 2.2 of the TBT Agreement, the Decree has no rational link with France’s stated objective: it does not protect the health of French workers or of the public in general. The requirement for such a link flows naturally from the text of Article 2.2, which states that regulations shall not create “unnecessary obstacles to international trade”. As the preamble to the TBT Agreement prescribes, the rational link element ensures that measures are not “applied in a manner which would constitute a means of arbitrary or unjustifiable discrimination”. Three facts supported by science demonstrate the lack of a rational link: (i) the risks posed by the high-density chrysotile products at issue in this case are undetectable; (ii) amphiboles are far more dangerous than chrysotile; and (iii) substitute fibres have not been shown to be harmless. Canada addresses each of these factual issues in turn.

Canada notes, first of all, that high-density chrysotile products do not threaten the health of French workers, but are nonetheless wrongly targeted. The Decree banning such products therefore has no rational connection with the protection of human health. According to Dr. Henderson, chrysotile-cement products pose a “negligible risk to health”
because of their normal weathering, erosion or general degradation. "... there is little or no dispute among experts on this issue." At the meeting with the parties on 17 January 2000, he added that the risk of cancer and mesothelioma is consistently higher with friable products than with high-density products and that the risk associated with manufacturing high-density chrysotile-cement products was "extremely low." Canada further states that interventions on the high-density products at issue, namely chrysotile-cement exterior construction panels, roofing tiles and pipes, are rarely, if ever, required. Moreover, with the adoption of proper working procedures and the use of proper tools as recommended in international standards, such interventions pose no detectable risk to human health. According to Canada, the WHO specifically targets past uses as being problematic: "There is potential for widespread exposure of maintenance personnel to mixed asbestos fibre types due to large quantities of friable asbestos materials still in place." One of the five conclusions of the WHO 1998 Report on asbestos is: "Some asbestos-containing products pose particular concerns [...] these uses include friable products with high exposure potential." Secondly, Canada claims that the Decree has no rational connection with the objective because it targets chrysotile, whereas amphiboles are the real danger. It does nothing either to address the serious problem posed by the enormous quantity of friable products containing amphiboles still in place. The ban merely relieves public pressure on the French Government to find a solution to the real problem. According to Canada, three of the four experts agree that a clear distinction should be drawn between the toxicity of amphiboles and chrysotile. Amphiboles may be up to 100 times more dangerous than chrysotile for the induction of mesotheliomas. At the meeting of 17 January, Dr. Henderson repeated that "most if not all these mesotheliomas are a consequence of exposure to asbestos-containing materials that included a mixture of asbestos types" [sic]. Canada notes that a 1997 report of the French Ministry of Labour (Ministère du Travail, Comité G2SAT), submitted to the Panel by the EC, recognizes that, as a result of the chemical dissolution process that takes place in the lungs, chrysotile's carcinogenic activity is practically nil:

"It has been shown that chrysotile is much more easily eliminated from the human lung than other forms (amphiboles). Moreover, it has practically no carcinogenic activity." Secondly, Canada contends that the Decree has no rational connection with the protection of human health because it imposes use of substitute fibres that are not proven to be safe. Canada notes that, at the meeting with the experts, Drs. Henderson, de Klerk and Infante agreed that, perhaps with the exception of glass fibres, which should be presumed to be carcinogens, the information on substitutes is meagre. This is confirmed in the recently published INSERM Report on substitute fibres, which "underlines some grave concerns that should rapidly be dissipated." At the second substantive meeting with the parties, Canada drew to the Panel's attention the key conclusions of the INSERM report on substitute fibres. Canada notes that, in this report, INSERM admits that very little is known about their potential impact on human health:

"Fibres on which very few toxicological data exist are today being used ... on a large scale as a replacement for asbestos; the novelty of their use for such applications is accompanied by a lack of data on their potential effects on human health." Canada points out that INSERM concluded that, in epidemiology, no significant excess risk of cancer has ever been detected from exposures to asbestos at the same exposure levels used to evaluate the carcinogenicity of substitutes: "No significant increase in the risk of cancer due to exposures to asbestos at levels comparable to those estimated for substitute fibres has ever been detected." Canada notes that, at the meeting, the experts did not deny this statement. INSERM also concluded that it was unable to demonstrate that substitute fibres are not carcinogenic:
"Globally, it was not possible to reach a firm conclusion for any of the types of fibre or any type of cancer [...]"

It must, however, be strongly emphasized that the data available do not allow the existence of a risk of cancer caused by exposure to substitute fibres to be eliminated [...]"

On the basis of the epidemiological data currently available, therefore, no conclusion can thus be drawn concerning the carcinogenicity of various types of substitute fibre.453 (WTO translation)

3.327 Canada points out that, based on these findings, INSERM recommends that all proposed substitutes be suspected of inducing pathologies: “Any new fibre proposed as a substitute for asbestos or for any other use should a priori be suspected of being pathogenic.”454 (WTO translation) Canada adds that a host of other studies also show that many substitute fibres present substantial health risks. According to the United States Occupational Safety and Health Administration (OSHA), glass fibres “are reasonably anticipated to be a carcinogen”.455 The United States Environmental Protection Agency (EPA) has further concluded that refractory ceramic fibres (RCF) present a significant risk of serious harm to human beings from cancer.456 This is consistent with the findings of the WHO’s International Agency for Research on Cancer (IARC), which has classified glass wool, rock wool, slag wool, and refractory ceramic fibres as “possibly carcinogenic to humans”.457 Finally, Dr. Infante has concluded that: “on a fiber-per-fiber basis, glass fibers may be as potent or even more potent than asbestos”.458 Canada notes that, at the meeting with the experts, Dr. Infante reiterated that glass fibres should be presumed to be carcinogenic. Canada submits that there is a convergence of opinions within the scientific community that there is no solid scientifically validated evidence to support the allegation that the use of substitutes is safe. Thus, the Decree has replaced the undetectable risk of chrysotile with the unknown risks of substitute fibres. The French ban is not rationally connected with its objective of protecting human health because the risk of substitutes has not been addressed.

3.328 Canada contends that the ban is not necessary because a less trade-restrictive measure is available. Why is it so “necessary” for France to protect its citizens against the undetectable risk posed by chrysotile high-density products, but not against the risks of substitutes? Canada notes that several variations of the word “necessary” have been put forward to justify France’s ban on chrysotile. In Canada’s view, however, necessity, in the WTO context, is not and should not be understood as political necessity “to be seen to be doing something” in order to assuage public opinion. The French National Assembly and Senate clearly acknowledged these facts in a joint report on asbestos published in 1997:

“It the climate in society changes as a result of pressure from shaken public opinion [page 19] … [The Decree is] a decision which concerns public opinion and whose aim is to provide reassurance [page 57].”459 (WTO translation)

3.329 Canada contends that necessity, in the WTO context – and more specifically in the context of the TBT Agreement– is a narrower concept than France and the EC suggest. A contested measure, to be in compliance with Article 2.2 of the TBT Agreement, must be “necessary to fulfil a legitimate objective”, i.e. to protect human health, not to meet the French population’s concerns about health. The TBT Agreement was not negotiated to allow governments to legislate to appease public opinion. It was specially negotiated to counter such recourse to technical barriers to trade in the absence of scientific evidence. Canada considers that a decision in favour of the EC would eviscerate the TBT Agreement. Under the necessity test of Article 2.2 of the Agreement, a measure will be found inconsistent if a less trade-restrictive means is available to reach the same policy objective. As a GATT Panel formulated the rule in respect of Article XX of the GATT, a measure will be deemed necessary only “if there were no alternative measure consistent with the General Agreement, or less inconsistent with it, which [the Party] could reasonably be expected to employ to achieve its health policy objectives”. In other words, even assuming that the chosen end is legitimate; the measure must not be an excessive or over-reaching means to achieve a legitimate end. 460
3.330 Canada claims that a ban is the most extreme and restrictive trade measure available. The Decree, as it stands, is excessive and does not meet the necessity criterion because a less trade-restrictive alternative could have properly achieved the protection of human health. France could have reached the same objective by setting up a regulatory regime under which prohibitions and authorizations of asbestos products were decided not on the basis of the existence or non-existence of substitutes, as the Decree does, but rather in the light of two guiding principles: (i) an assessment of risk made on a product-by-product and use-by-use basis; and (ii) an analysis of the non-feasibility and ineffectiveness of controlled use for each product. Canada claims that France could and should have used these two guiding principles to determine which chrysotile products are to be authorized in France, instead of the criterion of the non-availability of substitutes. If it had, the range of chrysotile products and their uses authorized in France would have been broader and international trade less restricted.

3.331 Canada contends that, first of all, the Decree is not based on a credible risk assessment and, secondly, that decisions to ban or exceptionally to authorize asbestos products under the Decree do not stem from an analysis of the feasibility and effectiveness of controlled use.

3.332 Canada considers that, if France had based its chrysotile policy on a credible product-by-product and use-by-use risk assessment, it would not have banned high-density chrysotile products. At the very least, such a risk assessment would have led the French authorities to authorize the use of a certain number of high-density chrysotile products. Canada claims that the INSERM report certainly cannot be considered a credible and sufficient risk assessment of high-density chrysotile products. The French Government used the INSERM report as the scientific basis to justify its ban on high-density chrysotile products, even though INSERM did not recommend banning them. Irrespective of the fact that the text of the report does not support the ban on high-density products, its serious deficiencies, the timing and circumstances of its adoption relative to the announcement of the French ban, and the strong criticism to which it has been subjected by members of the French and international scientific communities cast very real doubt on its credibility and its sufficiency as a scientific basis for the French ban. Canada notes that critics have identified six major problems with the INSERM report:

- INSERM’s risk assessment is based on hypothetical data and therefore has no real factual relation to the situation actually prevailing in France today.
- INSERM’s risk assessment is often misleadingly based on data from exposure to amphiboles or mixed fibres, instead of to chrysotile fibres alone.
- The report does not make clear that, because of the long latency period for asbestos-related diseases, the estimated deaths in the 1990s and after are almost exclusively the result of occupational and para-occupational exposures in the 1950s and 1960s.
- The report did not even examine what is surely the key issue relating to the ban, namely, exposures to current chrysotile products.
- INSERM’s extrapolations from data based on high exposure levels and exposures to friable products greatly exaggerates the risk from low exposure levels to products where chrysotile is encapsulated in a hard matrix, specifically chrysotile-cement and friction products.
- Finally, the INSERM report acknowledges that, “This problem is, however, indissociable from the choice of substitute fibre” (WTO translation) for which little information is available. It was only in November 1999, almost three years after the ban on chrysotile took effect, that INSERM’s report on substitutes was published, concluding that there was a lack of information on substitutes. It should also be recalled that INSERM did not recommend banning high-density chrysotile products.
3.333 Canada emphasizes that all available scientific evidence indicates that there is no detectable risk of mesothelioma or lung cancer as a result of exposure to chrysotile in the manufacture or use of friction products or in the manufacture of high-density chrysotile-cement products. The same is true for the use of high-density cement products. One of the most challenging issues in international trade law is striking the right balance between keeping markets open and allowing States to regulate in order to accomplish other legitimate goals, such as the protection of human health. In Canada’s view, one manner of resolving this issue – which has been used by WTO panels – is to determine whether the formal scientific analysis on which the decision to adopt trade-restrictive measures is based constitutes a credible and sufficient justification for the measure in question. It is often unrealistic to expect a panel of trade experts to resolve a complex scientific controversy. In fact, it is not its role to do so. In this case, there is so little scientific evidence for a ban that the Panel can only reasonably conclude that France has not conducted a credible and adequate scientific evaluation to justify this extremely trade-restrictive measure. Consequently, in assessing the WTO consistency of the French ban, Canada asks that the Panel disregard the INSERM report.

3.334 Canada also points out that, at the meeting on 17 January 2000, the experts mentioned two seemingly anomalous studies. Neither overcame the great weight of contrary evidence. First, Dr. Henderson in particular relied on the Australian Mesothelioma Register (Leigh et al., 1999). However, this is not a controlled study but simply a set of observations. It might be expected that any large enough subset of the population (such as garage workers or schoolteachers or lumberjacks) would experience some instances of mesothelioma or lung cancer. The question that must be asked is whether the number is more than expected? The only way to know for sure is by observing controls. According to Canada, other more carefully conducted studies suggest that Leigh’s data do not show excess risk. An examination by electron-microscopy of lung tissue in 221 definite or probable cases of mesothelioma in the Register and of 339 age-and-sex-matched controls (Rogers et al. 1991) suggest that the mesothelioma cases in the Register cannot be attributed to chrysotile exposure. Dr. Henderson puts a lot of emphasis on the Australian Mesothelioma Register to justify the non-feasibility of controlled use at all stages of the lifecycle of asbestos-cement products. But, in addition to criticism by Dr. McDonald regarding the use of such data, Canada stresses that the Australian experience is unique and that it is not prudent to draw general conclusions applicable elsewhere. Crocidolite amphiboles were used massively until the end of the 1970s in Australia, while amosite was used until the mid-1980s. Many houses were built with asbestos-cement products containing mainly amphiboles. Leigh et al. write that:

"From about 1940 to the late 1960s all three types of asbestos were used in [the asbestos-cement manufacturing industry], crocidolite then being phased out. Amosite was used until mid 1980s. Much of this industry output remains in service today in the form of ‘fibro’ houses and water and sewage piping. By 1954 Australia was number four in the Western world in gross consumption of asbestos-cement products, after USA, UK and France, and clearly first on a per capita basis. After World War II to 1954, 70,000 asbestos-cement houses were built in the State of New South Wales alone (52% of all houses built). In Australia as a whole, until the 1960’s, 25% of all new housing was in asbestos-cement."

3.335 Secondly, Canada maintains that the apparently high risk of lung cancer in the Charleston cohort, where exposure was predominantly to chrysotile, has only been observed in two other textile plants where substantial amounts of amphibole fibres were used. The “textiles mystery”, as it has been called by Dr. McDonald (1998), is anomalous and has yet to be explained. However, it is evident that it is limited to the textile process. Perhaps the numerous very long and fine fibrils produced in carding or the use of carcinogenic mineral oil spray for dust control and facilitation of weaving are responsible. These factors are peculiar to the textile industry and have no relevance whatsoever to friction or high-density cement products. In Canada’s view, it is evident that the French ban is not supported by credible and sufficient scientific evidence.
3.336 Canada contends that a balanced assessment of high-density chrysotile products would have forced the French authorities to conclude that this type of product is not dangerous for the public. Canada notes that, according to the experts appointed by the Panel, the risk to human health associated with the various uses of chrysotile throughout its life cycle is "overwhelmingly a workplace issue". The INSERM report on asbestos also supports the view that the asbestos issue concerns occupational and para-occupational exposures. Chrysotile-cement products do not pose a health risk from normal weathering, erosion or general degradation, and "there is no dispute among experts on this issue" [sic]. The removal of chrysotile-cement products does not pose any danger either, except in Dr. Infante’s view. Nor do these products endanger the general public via environmental exposure. As for the French do-it-yourself enthusiast, he will rarely, if ever, come into contact with a chrysotile-cement pipe, an exterior construction panel on a commercial building or brake linings, let alone saw or puncture one of these. Canada notes that, as Dr. Henderson confirmed, a do-it-yourself enthusiast therefore runs very little risk. The experts agree that one must turn to cumulative exposure to assess the risk. It can readily be understood that the cumulative exposure of do-it-yourself enthusiasts is inconsequential because, for the few who may be exposed, this is probably nothing more than a once-in-a-lifetime exposure to chrysotile from high-density products. Canada concludes that banning high-density chrysotile products allegedly because they subject the general public to risk is not supported by a balanced, product-by-product, use-by-use risk assessment.

3.337 Canada points out that a balanced risk assessment of high-density chrysotile products would have led the French authorities to conclude that these products are not dangerous to workers for two reasons. Firstly, most workers are not in contact with high-density chrysotile products and, when they are, their exposure is intermittent and therefore the cumulative exposure is very low. Secondly, controlled use is both feasible and effective. The term “worker” encompasses quite a diverse group of people. The Panel, in its questions to the experts, grouped workers in the following categories (see Question 1): (i) miners and millers; (ii) workers in the manufacturing industry (friction materials and chrysotile-cement products); (iii) employees of the textile industry; (iv) workers in the construction industry; (v) workers in the renovation, maintenance and heat insulation industry; and finally (vi) workers in the asbestos removal industry. Canada points out that it is defending high-density chrysotile products, not asbestos textiles. Therefore whether or not these workers are at risk is irrelevant to this proceeding. Regarding miners and millers, there has been no mining or milling activity in France since 1965. Canada also contends that these workers are not at risk. Workers manufacturing high-density chrysotile products likewise are not at risk. Article 2.2 of the TBT Agreement specifically points to the analysis of the risks that “related processing technology” entails. Canada concurs with the EC and the experts that, with controlled use, processing technologies used to manufacture high-density products pose no threat to human health.

3.338 According to Canada, no epidemiological studies show that workers in the high-density chrysotile products manufacturing industry incur excess risks of lung cancer or mesothelioma. As Dr. Thomas concluded for an asbestos-cement factory, “the population of the chrysotile-cement factory studied are not at any excess risk in terms of total mortality”. Studies of the friction products manufacturing industry show no chrysotile-related increase in lung cancer risk for persons exposed to the equivalent of up to 9 f/ml for 40 years. Workers in the maintenance, renovation, asbestos removal and heat insulation sectors, as well as electricians, are likely to be in contact with in-place friable asbestos and amphiboles, not primarily high-density chrysotile products. In Canada’s view, banning high-density chrysotile products is not necessary to protect these workers against the risks of amphiboles and friable products. Authorizing high-density chrysotile products in France does not increase the level of risk for these workers. The WHO specifically identifies the threat of friable asbestos and amphiboles for maintenance personnel. Dr. Henderson is of the view that, for these workers, mesotheliomas are almost invariably associated with amphiboles:

"... for data on mesotheliomas among electricians, carpenters, plumbers, insulation workers and so forth (it is acknowledged that most if not all these mesothe-
liomas are a consequence of exposure to asbestos-containing materials that included a mixture of asbestos types, including chrysotile and one or more of the amphiboles) ...

3.339 Canada maintains that the ban on high-density chrysotile products is not necessary to protect workers in the maintenance, renovation, asbestos removal and heat insulation sectors, and electricians. Banning high-density chrysotile products on the grounds that such products increase the risks for this category of workers is inconsistent with a balanced product-by-product, use-by-use risk assessment. Some workers and do-it-yourself enthusiasts may encounter high-density chrysotile products intermittently, for example, some construction workers who might occasionally intervene, depending on the nature of their work, on cement pipes, cement external wall panels or cement roof tiles; or perhaps mechanics who handle brake linings; or a do-it-yourself enthusiast who is occasionally exposed when working on his home. Canada considers that, in most cases, infrequent exposures lead to such low cumulative exposures that, even without the use of proper tools and procedures, the experts have agreed that the health of workers is not in danger. In the other cases, application of controlled use, including procedures and the use of tools prescribed by international standards, reduces the risk to an undetectable level.

3.340 Canada asserts that the French ban is not based on evidence that controlled use is unfeasible and ineffective. The regulatory system established by the Decree does not determine bans and authorizations for asbestos products in the light of a product-by-product examination of the feasibility and efficacy of controlled use. Canada notes that, if this were the case, France would have authorized high-density chrysotile products for which it has been established that controlled use is applicable and effective, for example, asbestos-cement pipes and brake pads and linings. Before commenting on controlled use as applied to work on asbestos-cement pipes and brake linings, Canada wishes to return to the meeting of 17 January 2000 with the experts and then describe briefly the concept of controlled use.

3.341 Canada claims that controlled use is an approach that is not specific to the asbestos industry. It is a general method of risk management for all products or technologies which, in the absence of controls or regulations, might involve risks. Moreover, the experts confirmed this when they specifically recommended the controlled use of substitute fibres. The experts also recommended applying the principles of controlled use as a way of dealing with the sensitive issue of amphiboles and friable products already in place. Canada notes that Dr. Musk referred to the principles of controlled use as a way of dealing with the problem of asbestos in place, in particular friable products and products containing amphiboles. The implementation of controls, the issue of permits and best work practices have already been advocated by Dr. Musk. Canada also notes that Dr. Infante refers to the principles of controlled use in the context of use of substitute fibres. He suggested that best work practices (use of low-speed cutting tools and wearing a mask) should be used to reduce the levels of exposure. This also forms part of controlled use as applied to the chrysotile industry. He also referred to United States standards for the control of nuisance dusts such as glass fibre. Dr. Infante also declares at the same time, paradoxically, that controlled use is unfeasible for chrysotile products because workers are not sufficiently informed or do not use the recommended equipment. If controlled use is applicable and effective for substitutes, amphiboles and friable material already in place, Canada wonders why it would not be applicable and effective for high-density chrysotile products.

3.342 Canada notes that, during the meeting with the experts, the question of whether it could be expected that regulations on controlled use would be respected was raised. In Canada’s view, official decisions should definitely not be taken on the basis of a hypothesis that the law will not be observed. Dr. Infante nevertheless commented that controlled use was not feasible in the United States due to the high number of breaches of the two United States standards on exposure to asbestos established by the OSHA. Canada wishes to point out in this respect that, of the 3,349 breaches of the two standards recorded during the period 1998-1999, only 16 concerned exceeding the United States exposure standard of 0.1 f/ml. Even more significantly, these 16 breaches were exclusively related to exposure to
3.343 Canada maintains that controlled use can be summarized by the three major recommendations in ILO Convention 162. Firstly, exposure should be brought down to levels that do not involve any health risk. Low levels of exposure allow asbestos-related diseases to be eliminated. Secondly, use of crocidolite should be banned. In general, scientific studies show that exposure to chrysotile is much less dangerous than exposure to amphiboles. Thirdly, friable asbestos materials such as sprayed and insulating materials should be banned. Asbestos-related diseases appear in the secondary production industries, when installing insulation containing asbestos and in building work involving friable asbestos materials. The fixing of an exposure limit in the workplace is only one aspect of protecting the health of workers. The means used to ensure observance of the exposure limit are of course an important part of the effort. These means include dust collection, monitoring, employing best work practices and, if necessary, using protective breathing equipment. Proper monitoring of exposure limits in workplaces is an essential criterion for correct application. Lastly, other aspects of protection in the workplace specifically concern secondary users, for example, building workers, servicing and maintenance workers, who work with asbestos-cement, or mechanics who handle brake linings. Controlled use mainly involves reducing the number of situations in which fibres are released (for example, by pre-cutting of asbestos-cement pipes) and eliminating unprotected work on high-density products.

3.344 Canada points out that the means employed for controlled use effectively reduce exposures, which then fall within the exposure limits, as recognized by the WHO:

"Exposure is dependent upon such factors as the extent of control, the nature of the material being manipulated and work practices [...]"

Levels dropped considerably between the 1930’s and the late 1970’s and have continued to decline substantially to the present day, owing to the introduction of controls.  

3.345 Canada notes that the WHO goes further and states that even partial application of controlled use methods, techniques or procedures is effective because it brings exposure levels down below 0.5 fs/ml. If personal protective equipment is used, the exposure levels drop even lower:

"Data from industries where control technologies have been applied demonstrated the feasibility of controlling exposure levels generally below 0.5 f/ml. Personal protective equipment can further reduce individual exposure where engineering controls and work practices prove insufficient."

3.346 Canada notes that the WHO also indicates that controls eliminate fibre suspended in the air when it becomes necessary to work on high-density chrysotile products, as may occasionally be the case for certain workers in the construction industry.

3.347 Canada adds that, according to the OSHA as well, controlled use is effective:

"There is substantial information available concerning both health effects and current exposure levels, and the potential for risk reduction is great using control methods already in use in some workplaces where mineral fibers or asbestos are manufactured or used."

3.348 Canada draws attention to the study by J. Peto, which shows the effectiveness of regulations, one aspect of controlled use. The 1995 study shows that only 5 per cent of the cases of mesothelioma recorded were in jobs subject to regulation. Taking into account the latency period, the British regulation of the asbestos industry in the 1960s, even though it allowed high levels of exposure and the use of friable materials and amphiboles, already
constituted an effective way of limiting the risks of using asbestos. J. Peto underlines the
greater effectiveness of the regulations in the 1980s by highlighting the marked decrease in
exposure levels. For example, the 1995 regulations in France, i.e. before the adoption of the
Decree, sufficed to protect the population against risks related to high-density chrysotile
products. Canada points out that it stressed, and the experts agreed, that for primary
users, namely, workers involved in the mining and processing of chrysotile products such
as asbestos-cement and friction materials, occupational exposure levels are low because of
controlled use and, consequently, the risk that these workers will contract an asbestos-
related disease is undetectable. Canada acknowledges that it is aware of the EC’s objective
and the experts’ concerns regarding controlled use. It is not so much the risk during min-
ing of chrysotile and manufacturing that are the cause of concern, but the risks linked to
occupational exposure of secondary users. Canada declares that, in order to show that the
EC’s objection to controlled use among secondary users is excessive, it will comment on
both products, namely, asbestos-cement pipes and friction materials.

3.349 Canada asserts that no-one can seriously contest that controlled use is as effec-
tive as a ban for the protection of human health against risks caused by working with
asbestos-cement water pipes buried in the ground. Asbestos-cement pipes are almost never
cut during their installation. They are delivered on-site ready to use and it is not necessary
to carry out any work likely to produce dust. Prefabrication of elements subsequently
assembled on the spot thus makes it possible to eliminate exposure of the workers to
chrysotile fibres. If the pipes have to be cut, the use of hand tools or tools with vacuum
dust extractors allows the risks to be eliminated. Workers use hand tools or low-speed
mechanical tools, which create large dust particles or chips, rather than machines that work
through abrasion. High-speed mechanical tools are equipped with an effective and spe-
cially designed vacuum dust extractor. According to the OSHA, the application of control
measures leads to average levels of exposure of 0.00253 f/ml for those working with asbes-
tos-cement pipes. Canada adds that the risk of asbestos-related disease is based on the
hypothesis of persistent exposure. There are no data on cumulative “lifetime” levels for
persons exposed intermittently. Workers using asbestos-cement pipes are, however, ex-
posed intermittently. In Canada’s view, the EC will readily acknowledge that few handy-
men have to work with asbestos-cement pipes. Workers usually install asbestos-cement
pipes by heavy machinery and not manually. Buried in the ground, asbestos-cement pipes
are quite safe. They can also be taken out of the ground mechanically. Canada asserts that
controlled use is also feasible and effective for mechanics working on brake linings. They
are in contact with the brake linings intermittently so their cumulative exposure to chrysotile
is very low. In addition, simple methods such as moistening and vacuum dust extractors
make it possible to reduce exposure to levels at which the risk is undetectable, as already
indicated in studies cited by Canada and discussed with the experts.

3.350 Regarding the next objection that, according to some theoretical models not
empirically verified, workers in general are subject to risks even if cumulative exposure is
within the limits laid down under controlled use policies, Canada responds that the lower
the exposure, the lower the risk. The question is to what extent levels of exposure to a
substance can be lowered in order to achieve a significant and measurable reduction in the
risk for workers? Canada notes that the most recent data published, as well as the latest
retrospective reviews on the relationship between exposure and the effect of chrysotile
asbestos are revealing: when levels of exposure to around 40 f/ml over 20 years (or 20 f/
ml over 40 years) are reached, the excess risk of lung cancer related to chrysotile asbestos,
although it still exists, has become undetectable. If a safety factor is introduced by impos-
ing a level that is ten times lower, for example, 2 f/ml over 40 years, it is obvious that the
risk, although it still exists, remains undetectable. Canada points out that Drs. de Klerk
and Musk agree that current epidemiological data does not show an excess health risk at
low levels of exposure to chrysotile, lower than the exposure limits under controlled use.
Dr. Henderson acknowledges that “no increase in risk of mesothelioma has been identified
at very low-level exposures”. A review of eight studies of cohorts exposed to chrysotile by
Browne and Gibbs led the authors to conclude that “there exist levels of exposure below
which risks are for practical purposes zero”.

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Canada maintains that, if France had based its asbestos regulations on a serious examination of the feasibility and effectiveness of controlled use product-by-product, it would – at the very least – have authorized asbestos-cement pipes and friction materials. But this is not the case: the French Decree bans asbestos-cement pipes and friction materials, two products which handymen and non-specialized workers never or only exceptionally use and for which controlled use is feasible and effective. The Decree is therefore not based on any serious examination of the feasibility and effectiveness of controlled use product-by-product, which means that it does not meet the necessity criterion in Article 2.2. According to Canada, France could have regulated the application and use of asbestos through non-discriminatory technical regulations dealing \textit{inter alia} with monitoring the average concentrations of asbestos fibres in the workplace, which France in fact does for the exemptions allowed. In addition, France could have regulated the manufacture of high-density materials by eliminating work likely to release dust, just as the construction industry gives workers proper training on how to handle these materials. The effect or the purpose of such measures would not have been to create unnecessary obstacles to international trade and at the same time would have enabled the objective of protecting health to be attained.

To conclude on Article 2.2 of the TBT Agreement, Canada points out that its examination shows that the French ban on chrysotile asbestos does not enhance the protection of human health. The objective sought by France is not achieved by a ban. There is no rational link between the French ban on chrysotile and France’s objective. Moreover, within the meaning of Article 2.2 of the TBT Agreement, the French measure is not necessary in regard to the objective sought. The ban does not address the real health problem associated with asbestos: the friable products and the amphiboles already in place. It is a bold step for the ban to replace chrysotile by the risk of substitute fibres. Furthermore, the ban, by covering high-density chrysotile products, does nothing to raise the level of protection of human health. The French Decree is an excessive measure “more trade-restrictive than necessary” to achieve France’s objective because the objective could be achieved by a measure that is less trade-restrictive and which Canada has presented: a regulatory system that is, first of all, really based on a rigorous examination of the risks product-by-product and use-by-use and then by an examination of the feasibility and effectiveness of controlled use product-by-product.

The European Communities wish to recall that any Member of the WTO has the right to establish the level of health protection it deems appropriate in its territory. The EC note that, within the context of Article XX(b) of the GATT, all the Panels which have examined the concept of necessity have concluded that it was not the necessity of the objective pursued by the measure concerned that should be examined but whether or not it was necessary to submit the imported products to the measure contested in order to achieve the chosen level of protection (e.g. United States - Section 337, Thailand - Cigarettes and United States - Gasoline Panel reports). It follows that, whereas the trade measures that make it possible to achieve the desired objective must satisfy certain conditions, there is no restriction on the level of protection chosen by the Member. Accordingly, the EC consider that France was free to choose the level of protection it deemed appropriate in the present case, i.e. to halt the spread of the risk linked with the use of asbestos fibres and products containing such fibres. The EC again point out that, even though the test of necessity within the context of Article 2.2 of the TBT Agreement corresponds, in particular, to the test in Article XX(b) of the GATT, it nonetheless remains true that the burden of proof within the context of the TBT Agreement lies with the party which invokes a specific provision of the Agreement to establish the inconsistency. The EC indicate that the arguments it develops on the necessity criterion are also applicable in the context of Article XX(b) of the GATT.

The EC maintain that the necessity criterion under Article 2.2 of the TBT Agreement involves, \textit{inter alia}, examining whether a less restrictive measure is available to a Member and could be employed to fulfill that Member’s legitimate objective. The EC note in this respect that, in their written and oral replies, the scientific experts all agreed that banning the use of all types of asbestos, including high-density asbestos-containing cement products, to which Canada attempts to limit the scope of this dispute, was in fact the
only real option available to France to achieve its legitimate objective of protecting human health. This is because all types of asbestos are classified as proven human carcinogens and because the so-called "controlled use" advocated by Canada is in practice unfeasible and unrealistic. The EC note that Dr. Henderson gave a number of reasons why it is unrealistic and impossible to apply such a use in practice and cited examples of concrete cases in which he personally observed the absence of any kind of "controlled use". Dr. Infante also explained that in the United States several hundreds of violations continue to arise, whether in the manufacturing of asbestos-containing cement products or in subsequent downstream situations, despite the controls exercised and the fines applied by the United States authorities. Dr. de Klerk, when discussing this issue, said that "at best it is imprudent" to wish to continue exporting asbestos, for example to least developed or developing countries, because it is impossible for them to apply such a controlled use.

3.355 The EC also recall that, since the beginning of this dispute, Canada has been avoiding a precise definition of what it means by the term "controlled use". As a last attempt to confuse the legal debate, in its comments on the replies of the experts, Canada finally defined the way it understands the concept of "controlled use". But that description of "controlled use" has never been explained in such detail until now, none of the experts chosen by the Panel have ever heard of it before, and they have explicitly stated that they know of no international standard or recommendation suggesting that such a use will substantially reduce or eliminate the risks of asbestos-related diseases. In addition, all the experts found that the conditions of use mentioned by Canada in its comments are impossible to apply in practice in a way that will enable the level of protection chosen by France to be attained. According to the EC, Dr. Henderson and the other experts agreed with the EC's argument that the problem France faces because of the existing asbestos in place and the risks associated with its removal do not in any way justify the continued use of asbestos and the perpetuation of the serious risks to human health through the further introduction of asbestos into the environment. France, like other countries, has been trying to deal in the best way possible with the problems for human health posed by the asbestos in place and it took measures for this purpose a long time ago. But experience and science have shown that these measures cannot eliminate the risks to human health. In the EC's view, there is no provision in the TBT Agreement nor in the WTO Agreements that could oblige France to continue using chrysotile asbestos, due to the difficulties it (like other countries) faces in eliminating the risks from past mistakes when the use of chrysotile asbestos was allowed.

3.356 As for the second sentence of Article 2.2 of the TBT Agreement ("taking account of the risks non-fulfilment would create"), the EC consider that, here again, this is an integral part of the implementation of the test of necessity under Article XX(b) of the GATT. In fact, a restrictive measure is "necessary" only if there are risks associated with the non-adoption of the measure in question. The purpose of this sentence is to deny Members the possibility of adopting measures, under cover of protecting human health, without having taken into account the risks associated with the use of the product that is banned. Such risks should be assessed, in particular, on the basis of the available scientific and technical information. The EC comment that they have shown, and all the experts have agreed, that under the so-called "safe use" policy there continues to be a significant excess of deaths in the asbestos production and processing sector. Moreover, this policy does not offer any meaningful and realistic protection, in particular for the secondary or downstream users exposed to asbestos (carpenters, electricians, do-it-yourself enthusiasts, etc.) because they cannot be aware of the risks in all situations and the risks frequently become apparent many years after exposure. The EC point out that they have also asserted (and all the experts have agreed) that there exist several safer alternative products that can replace nearly all the uses of asbestos. Indeed, all the scientists agreed with Dr. Infante and Dr. Henderson that the available scientific evidence shows that none of the substitute non-fibrous or fibrous products is dangerous to human health or as dangerous as asbestos, including chrysotile asbestos. Dr. Infante also stated that it is unreasonable to continue using chrysotile asbestos simply because there exist some doubts about some fibrous products.
3.357 The EC also wish to respond to Canada’s claim, completely unjustified in their view, that the French ban on asbestos was taken for reasons of “political necessity”, not out of real concern to protect human health. This is plainly wrong and contradicts all the scientific evidence coming from so many countries that already banned asbestos before France and from so many international institutions such as the WHO, the IARC, the ILO, etc., which all recommend that asbestos be banned and replaced. The EC ask whether Canada can really argue that all these countries and all these international institutions have done so for reasons of political necessity? Certainly not in the EC’s view.

(d) Article 2.4 of the TBT Agreement

3.358 Canada maintains that Article 2.4 lays down the principle that, where technical regulations are required and relevant international standards exist, Members shall use them, or the relevant parts of them, as a basis for their technical regulations if they are effective and appropriate for the fulfillment of the objective being pursued. Under Article 2.4, it is up to the Panel to determine: (i) whether a technical regulation on chrysotile is required; (ii) whether there are international standards concerning chrysotile; (iii) whether the international standards are an effective and appropriate means for the fulfillment of the objective pursued; and (iv) whether the Decree is based on international standards. Canada recognizes that it is important for governments to take action to manage the risks associated with the use of asbestos fibre. Therefore, Canada does not dispute the fact that action is required to manage the risks associated with asbestos fibres and their uses. However, in this case, it is the excessive nature of the French action that is being challenged. Such action must be based on existing international standards that recognize the controlled use of asbestos, which, in practice, eliminates any health risk, i.e. the objective pursued by France. Canada submits that, in the case of chrysotile fibre, a regulation and not a ban is required pursuant to Article 2.4. Canada points out that it in fact implements and encourages the regulation of the “safe” or “controlled” use of chrysotile fibre, the required and approved form of action. Prior to the adoption of the Decree, France practised controlled use. Canada has already emphasized that a set of measures existed prior to the adoption of the Decree and guaranteed the protection of people’s health in France.

3.359 Canada claims that international standards, within the meaning of Article 2 of Annex 1 to the TBT Agreement, exist for the controlled use of chrysotile fibre. Convention 162 and Recommendation 172 of the International Labour Organization concerning safety in the use of asbestos by workers constitute such international standards.480 They set forth rules and guidelines for the use of asbestos, in particular chrysotile fibre, as well as processes and production methods for products containing it. The ILO Code of Practice on Safety in the Use of Asbestos – endorsed by Convention 162 and Recommendation 172 - constitutes another international standard.481 The Code contains standards and practical instructions aimed at preventing the risks faced by workers using chrysotile. These standards govern all the stages of exposure to asbestos by proposing safe procedures and methods; from the extraction of the fibre to the transportation of products containing it, from the manufacturing of products containing fibre to their maintenance. Standards for the use of chrysotile-cement materials are included in the Code for chrysotile-cement products – Guidelines for On-Site Work of the International Organization for Standardization (International Standard ISO-7337).482

3.360 Canada contends that existing international standards are effective and appropriate for fulfilling the objectives of protecting human health. International standards such as those in Convention 162, Recommendation 172, the ILO Code of Practice and standard ISO-7337 provide for controlled and safe use of asbestos. The wording of these standards indicates very clearly that asbestos fibres should only be replaced if it is established that this is necessary to protect the health of workers and if it is technically feasible. However, replacement of chrysotile asbestos fibre in modern materials or products, where it is sealed in a matrix and cannot be released into the environment, is not necessary to ensure the protection of workers’ health because these products do not pose any detectable risks against which workers would need to be protected. Canada states that it has already clearly shown the absence of a detectable health risk linked to the controlled use of chrysotile fibre, in-
cluding its incorporation in safe modern materials. As the measures set forth in the international standards described above are, with respect to controlled use, both effective and appropriate to protect people’s health from the risks posed by exposure to chrysotile fibre, the total ban on the modern uses of chrysotile was not necessary.

3.361 Canada asserts that the Decree is not in compliance with international standards because it imposes a total ban on chrysotile fibre rather than providing for the controlled and safe use of this fibre and of its uses. Canada contends that a ban is not necessary to protect people’s health and, consequently, is not in compliance with the relevant international standards. International standards impose an approach that is quite different from the complete ban without distinction as to the type of fibre or its use. International standards impose an approach whereby regulation of asbestos must take into account the type of fibre, the products in which a type of fibre is incorporated and the intended uses of each product. Convention 162 and Recommendation 172 provide for the banning of crocidolite and materials containing brittle sprayed asbestos, as well as the elimination of certain other uses (work processes), if the national authority deems these measures to be necessary for the protection of workers, but only if the substitute products have undergone a thorough scientific evaluation of their effects on health. According to Canada, the EC have recognized that controlled use would ensure adequate protection of the health of workers in the chrysotile industry (extraction, manufacturing of chrysotile-cement, for example). Canada considers that, given this recognition of the effectiveness of the principles of controlled use upon which international standards are based, the EC must prove that the international standards are ineffective and inappropriate in order to comply with Article 2.4. Canada submits that the burden of proof rests with the EC. Canada also maintains that the Decree is not necessary pursuant to Convention 162 or Recommendation 172 because, according to the EC themselves, there are only certain specific categories of workers, on projects involving brittle materials containing asbestos, that are at risk. The Decree does not resolve the problem that these workers face. Canada also indicates that the EC cannot claim that Convention 162 or Recommendation 172 are ineffective or inappropriate when the Decree does not offer the same objective safety warranty with respect to substitute products. In fact, the Decree takes for granted that any product replacing asbestos is safer than chrysotile, whereas Convention 162 and Recommendation 172 require that the replacement be carried out only if it has been “scientifically” and “thoroughly” evaluated that each substitute product proposed is safer. In providing that no asbestos shall be used in the manufacture and processing of materials, products and devices, without distinction as to the variety of fibre or its use, the Decree is the most restrictive type of technical regulation that can exist. The international community has nevertheless developed standards that are less trade-restrictive. These standards enable people’s health to be protected in an effective and appropriate manner. France has chosen to ignore these international standards and opted for a total ban. In view of the foregoing, the Decree banning asbestos violates the provisions of Article 2.4 of the TBT Agreement.

3.362 The European Communities contend that, for the purposes of the TBT Agreement, the term “standards” has the precise meaning it is given in Annex 1 to the Agreement. In this case, however, the international texts referred to by Canada (the ILO, WHO, ISO texts) do not meet the definition concerned. In any event, the EC consider that the French authorities used the texts referred to by Canada in its submission as a basis for their Decree, within the meaning of Article 2.4 of the TBT Agreement.

3.363 The EC point out that, as already stated, the object and purpose of the TBT Agreement, as derived from its preamble, its background and the actual wording of several of its provisions, are to monitor the adoption and application of “standards” and “technical regulations” which cover the detailed characteristics of products or their methods of production. The object and specific purpose of the Agreement are bound to have an impact on the meaning to be given to the term “standards” mentioned in Article 2.4 of the TBT Agreement. This impact is moreover recognized by the Agreement itself, whose Article 1.1 and 1.2 are particularly explicit. As Annex 1 contains a definition of “standard”, according to the EC, the drafters of the TBT Agreement must have wished to use a specific definition of “standard” for the purposes of the Agreement’s application. This specific definition ap-
pears in Annex 1 and it follows from this definition that the TBT Agreement encourages the use of international standards, but solely those which can provide rules, guidelines or characteristics for products or related processes and production methods. But in this case, the EC note that the international texts referred to by Canada do not satisfy the definition contained in Annex 1 to the TBT Agreement and cannot therefore be used “as a basis” for technical regulations. Neither the ILO and WHO documents nor the ISO standards can be considered as laying down the characteristics of asbestos fibres or an ordered set of rules for the manufacture of this product. Even less can they be considered as laying down the characteristics of asbestos-containing products or an orderly set of rules for the manufacture of those products. These are texts that certainly do not satisfy the definition of “standard” within the meaning of the TBT Agreement.

3.364 In the alternative, the EC maintain that, even if the Panel were to consider that the texts cited by Canada in its submission are “standards” within the meaning of the TBT Agreement, it has to be recognized that these texts were used “as a basis” for the adoption of the Decree. The phrase “as a basis for” could be compared with the term “on the basis of” used in the SPS Agreement, a term for which the Appellate Body (Hormones) indicated: “…we disagree with the Panel’s interpretation that ‘based on’ means the same thing as ‘conform to’.”488 “A thing is commonly said to be ‘based on’ another thing when the former ‘stands’ or is ‘founded’ or ‘built’ upon or ‘is supported by’ the latter.”489 If this definition is followed, the EC conclude that the international texts quoted, or sometimes not quoted, by Canada serve “as a basis” for the Decree.

3.365 The EC indicate that, as early as 1986, ILO Convention 162 stated:

“Where necessary to protect the health of workers and technically practicable, national laws or regulations shall provide for one or more of the following measures:

(a) replacement of asbestos or of certain types of asbestos or products containing asbestos by other materials or products or the use of alternative technology, scientifically evaluated by the competent authority as harmless or less harmful, whenever this is possible.

(b) total or partial prohibition of the use of asbestos or of certain types of asbestos or products containing asbestos in certain work processes.”490

3.366 Similarly, ILO Recommendation 172 also indicated in 1986 that:

“… asbestos should be used only when its risks can be prevented or controlled; otherwise, it should be replaced, when technically feasible, by other materials or the use of alternative technologies, scientifically evaluated as harmless or less harmful.”491

3.367 The EC note that, more recently, a WHO report specifically dealing with chrysotile was even more categorical. The summary of its conclusions and recommendations states:

“Exposure to chrysotile asbestos poses increased risks for asbestosis, lung cancer and mesothelioma in a dose dependent manner. No threshold has been identified for carcinogenic risks. […] Where safer substitute materials are available for chrysotile, they should be considered for use. […] Some asbestos containing products pose particular concern and chrysotile use in those circumstances is not recommended. These uses include friable products with high exposure potential. Construction materials are of particular concern for several reasons. The construction industry work force is large and measures to control asbestos are difficult to institute. In-place building materials may also pose risk control to those doing alterations, maintenance and demolition. Minerals in place have the potential to deteriorate and create exposures.”492
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3.368 The EC conclude from these texts that: (i) the banning or replacement of asbestos fibres or asbestos-containing products may be decided in cases where this is necessary to protect the health of workers and is technically feasible; (ii) where substitute materials are considered safer, they must be used to replace asbestos; (iii) control of the use of asbestos, including chrysotile, in the construction industry is difficult to introduce. According to the EC, this conclusion by the WHO contradicts Canada's statements that the "safe" or "modern" use of asbestos would do away with any risk connected with its use. The Decree is completely in line with these conclusions. The EC add that Article 2.4 of the TBT Agreement also provides that international "standards" must be ignored when they are "ineffective or inappropriate". Such is clearly the case and the ISO standard provides a perfect illustration of this point. At the time it was published in 1984, this standard represented a major step forward in relation to the arrangements prior to that date, but it does not guarantee sufficient levels of protection in the light of the health objective adopted by the vast majority of countries, and by France in particular. In the light of the foregoing, the EC therefore maintain that the Decree is compatible with Article 2.4 of the TBT Agreement.

3.369 Canada asserts that Article 2.4 of the TBT Agreement sets out the principle that, where technical regulations are required and relevant international standards exist, Members shall use them, or the relevant parts of them, as a basis for their technical regulations when such international standards are an effective and appropriate means for the fulfilment of the objective pursued. This means that, in this case, the Panel must determine: (i) whether a technical regulation concerning chrysotile is required; (ii) whether there are relevant international standards concerning chrysotile; if so, (iii) whether the international standards are effective and appropriate for the fulfilment of the objective pursued; and (iv) whether the Decree is based on these international standards.

3.370 Canada acknowledges that it is important for governments to take action to manage the risks posed by asbestos. However, in the case of chrysotile fibre, this action cannot legally take the form of a ban. A regulation concerning the "controlled" use of chrysotile fibre and products containing it is the required and appropriate form of action. Canada contends that there are relevant international standards concerning controlled use of chrysotile fibre. These are to be found in Convention 162 and Recommendation 172 of the ILO, the ILO Code of Practice on Safety in the Use of Asbestos, and Standard ISO-7337 – chrysotile-cement products – Guidelines for on-site work practices. Canada rejects the EC's claim that these are not standards within the meaning of Annex 1 to the TBT Agreement. The ILO and ISO documents fully meet the criteria of the definition in Annex 1 to the TBT Agreement. They are documents approved by recognized bodies that provide "process and production methods" for chrysotile-based products. Canada notes that the EC have not, however, disputed the relevance of these international standards. The ordinary meaning of the word "relevant" is: "bearing on or pertinent to the matter in hand". The standards cited deal with the same product: asbestos. Furthermore, they have an identical objective, namely, the safe use of asbestos. These international standards are relevant because they relate to the use of chrysotile asbestos in a safe and controlled manner.

3.371 Canada contends that international standards such as those contained in Convention 162, Recommendation 172, the ILO Code of Practice and Standard ISO-7337 prescribe a safe and controlled use of asbestos. According to Convention 162, the use of substitute products is encouraged only "Where necessary to protect the health of workers and technically practicable [...] by other materials or products or the use of alternative technology, scientifically evaluated by the competent authority as harmless or less harmful ..." According to this standard, chrysotile fibre should not be replaced in high-density products because it is trapped in a matrix from which it cannot escape. Furthermore, Canada has also stressed that controlled use of high-density chrysotile-based products presents, at most, an undetectable risk to human health. As the measures prescribed by these standards are both effective and appropriate to protect human health against the risks posed by exposure to chrysotile, a total ban on chrysotile, and consequently on modern uses of chrysotile, is not necessary. According to Canada, the EC have not provided evidence that it is necessary, in this case, to set aside these international standards on the grounds that they are ineffective or inappropriate. In order to avoid rendering Article 2.4 ineffective,
interpretation consistent with the precepts of the international law of treaties requires that, in order to set aside a relevant international standard, a Member must provide evidence of a “fundamental” problem and not a simple allegation that the relevant standards are not appropriate.

3.372 Canada points out that Article 2.4 of the TBT Agreement requires that Members use relevant international standards or their relevant parts as a basis for their technical regulations. According to the ordinary meaning of the words, this means that a technical regulation must be based on international standards or relevant parts thereof. In other words, a technical regulation must take what is set out in the international standards as its basic principle or starting point. The technical regulation adopted by a Member does not have to be identical to the international standards, but when they are relevant, the technical regulation must use them as a foundational or logical starting point. Canada contends that the international standards concerning asbestos are relevant. The international standards do not call for a complete ban without distinction as to the type of fibre or its use. They propose rather an approach whereby the asbestos regulations must take into account the type of fibre, the products in which a type of fibre is incorporated, and the intended uses of each product. The international standards thus propose a less trade-restrictive approach while at the same time stressing the protection of human health in an effective and appropriate manner. France has chosen to ignore these international standards, preferring instead a total ban. What is more, the French technical regulation deviates so much from the international standards that their basic principles are no longer recognizable. Canada points out that the EC have offered no explanation of the reasons that might have justified France’s total rejection of these standards as the logical starting point for its regulations on asbestos. Given the foregoing, the Decree banning asbestos is inconsistent with the provisions of Article 2.4 of the TBT Agreement.

3.373 The European Communities contend that the texts cited by Canada do not fall within the scope of the TBT Agreement and are therefore not relevant under Article 2.4 thereof because they do not correspond to the definition of “standard” in Annex 1 to the Agreement. In fact, the terms of the ILO, ISO and WHO texts, in their particular context, do not lay down guidelines or characteristics for products or related production processes and methods, nor is that their object and purpose. At best, they can be treated as assessments of the risks created by asbestos and asbestos-containing products rather than as establishing international technical standards or conformity assessment procedures. The EC point out, for example, that the Constitution of the ILO (preamble) specifically provides for “the protection of the worker against sickness, disease and injury arising out of his employment”. As for the Philadelphia Declaration (Article III-g) concerning the aims and objectives of the ILO, it provides for “adequate protection for the life and health of workers in all occupations”. The ISO texts are conceived in the same spirit. Similarly, Article 3 of ILO Convention 162 of 1986 states that “National laws or regulations shall prescribe the measures to be taken for the prevention and control of, and protection of workers against, health hazards due to occupational exposure to asbestos.” According to point 1(i) of the scope and definitions of ILO Recommendation 172 of 1986, “The provisions of the Asbestos Convention, 1986, and of this Recommendation should be applied to all activities involving a risk of exposure of workers to asbestos in the course of work.” Thus, the EC consider that the ILO and ISO texts are not relevant standards within the meaning of Article 2.4 of the TBT Agreement.

3.374 The EC, after responding to the question of whether or not a standard was relevant, emphasizes that Article 2.4 raises the question of whether the standard is effective or appropriate. This Article prescribes in particular that “…Members shall use [these international standards] … except when such international standards or relevant parts would be an ineffective or inappropriate means for the fulfillment of the legitimate objectives pursued, for instance because of fundamental climatic or geographical factors or fundamental technological problems”. Before considering the case at issue, the EC would like to spell out the circumstances in which a standard might be considered ineffective or inappropriate within the meaning of this provision. The EC consider that the level of protection
deemed appropriate by the Member could be a factor in making international standards ineffective or inappropriate. Within the context of the TBT Agreement, a Member is free to choose the level of protection it deems appropriate. An international standard is only effective or appropriate if it enables the Member to achieve the legitimate objective it has set itself. Advances in scientific knowledge can also lead to the application of obsolete standards being ruled out. A systematic comparison of the WTO Agreement (in this specific case the TBT Agreement) with the NAFTA Agreement - a GATT Article XXIV agreement negotiated at the same time as the WTO Agreements – reveals that, on the subject of technical barriers to trade, the NAFTA Agreement provides as follows:

"1. Each Party shall use, as a basis for its standards-related measures, relevant international standards or international standards whose completion is imminent, except where such standards would be an ineffective or inappropriate means to fulfill its legitimate objectives, for example because of fundamental climatic, geographical, technological or infrastructural factors, scientific justification or the level of protection that the Party considers appropriate.

2. A Party's standards-related measure that conforms to an international standard shall be presumed to be consistent with Article 904(3) and (4).

3. Paragraph 1 shall not be construed to prevent a Party, in pursuing its legitimate objectives, from adopting, maintaining or applying any standards-related measure that results in a higher level of protection than would be achieved if such measure were based on an international standard."

The EC therefore consider that scientific reasons or the level of protection deemed appropriate by the Member are factors that could render an international standard ineffective or inappropriate within the meaning of Article 2.4 of the TBT Agreement. In the present case, the EC consider that the standards cited by Canada do not make it possible to achieve the level of protection deemed appropriate by France insofar as: (i) there can be no doubt that chrysotile asbestos is a proven carcinogen; (ii) there is no exposure limit (threshold) for chrysotile asbestos and asbestos-containing products; (iii) so-called “safe” use is not applicable in all circumstances nor to every type of person who may come in contact with asbestos or asbestos-containing products and, moreover, does not eliminate every risk; and (iv) there are substitute products that are safe, or safer, than chrysotile asbestos.

The EC therefore consider that scientific reasons or the level of protection deemed appropriate by the Member are factors that could render an international standard ineffective or inappropriate within the meaning of Article 2.4 of the TBT Agreement. In the present case, the EC consider that the standards cited by Canada do not make it possible to achieve the level of protection deemed appropriate by France insofar as: (i) there can be no doubt that chrysotile asbestos is a proven carcinogen; (ii) there is no exposure limit (threshold) for chrysotile asbestos and asbestos-containing products; (iii) so-called “safe” use is not applicable in all circumstances nor to every type of person who may come in contact with asbestos or asbestos-containing products and, moreover, does not eliminate every risk; and (iv) there are substitute products that are safe, or safer, than chrysotile asbestos.

The EC therefore consider that scientific reasons or the level of protection deemed appropriate by the Member are factors that could render an international standard ineffective or inappropriate within the meaning of Article 2.4 of the TBT Agreement. In the present case, the EC consider that the standards cited by Canada do not make it possible to achieve the level of protection deemed appropriate by France insofar as: (i) there can be no doubt that chrysotile asbestos is a proven carcinogen; (ii) there is no exposure limit (threshold) for chrysotile asbestos and asbestos-containing products; (iii) so-called “safe” use is not applicable in all circumstances nor to every type of person who may come in contact with asbestos or asbestos-containing products and, moreover, does not eliminate every risk; and (iv) there are substitute products that are safe, or safer, than chrysotile asbestos.
The texts cited by Canada as international standards come from the ILO and the ISO. Canada points out that it did not mention, for example, Environmental Health Criteria for Chrysotile Asbestos (203) of the WHO as a standard within the meaning of Article 2.4 of the TBT Agreement because it does not see how this text could be considered a standard when the WHO itself presents it as a risk assessment. Furthermore, contrary to what the EC suggests, the WHO recommendations are not to cease using chrysotile asbestos and to replace it with substitute products. Rather, the WHO suggests that consideration be given to replacing chrysotile by harmless substitute products where such materials are available. Canada refers the Panel to the arguments already put forward concerning Article 2.4 and adds some comments relating to the two issues: on the one hand, the effectiveness and appropriateness of the international standards and, on the other, the question of whether these standards were used as a basis for the Decree.

3.378 Canada contends that the international standards are effective and appropriate. Article 3.1 of Convention 162 of the ILO provides that “National laws or regulations shall prescribe the measures to be taken for the prevention and control of and protection of workers against health hazards due to occupational exposure to asbestos”. The purpose of the ILO Code of Practice is similar. The object of international standard ISO 7337 is to provide “guiding principles for the tools and methods to be used on site so as to keep dust emissions as low as possible [...]”. In Canada’s view, the international standards must be appropriate because their purpose and the objective of protecting human health put forward by France are one and the same. Canada notes that the EC give two reasons to justify the inappropriate and ineffectiveness of the standards, namely: (i) scientific grounds; and (ii) the level of protection deemed appropriate by France. The EC obfuscate on this point. They reject the standards of the ILO and the International Labour Office on the pretext that they recommend controlled use, a means of protection that is unacceptable to France, which now only cites the WHO. Canada points out that, in their first written submission, the EC did not reject the ILO standards but claimed that they recommended a ban on asbestos and its replacement by less harmful substitute fibres. Canada can only explain this sudden change in position by implicit recognition that the standards mentioned by Canada do not recommend a ban and the replacement of all types of asbestos and all their uses. Regarding the reasons for ineffectiveness and inappropriateness, namely, the scientific grounds, it is unthinkable to claim that progress in scientific knowledge leads to disregard for international standards. These standards recommend taking effective and appropriate measures to protect human health against the risk of exposure to chrysotile. To disregard an international standard, it is necessary to provide evidence of a fundamental situation that makes the standard ineffective or inappropriate. According to Canada, the EC have not shown that recent scientific discoveries have made the international standards cited by Canada obsolete, standards which, moreover, reflect the WHO’s recommendations in its recent assessment of the risks of chrysotile asbestos.

3.379 As far as the second reason for ineffectiveness and inappropriateness is concerned, namely, the non-fulfilment of the level of protection deemed appropriate by France, Canada does not contest a Member’s freedom, within the framework of the TBT Agreement, to choose the level of protection it deems appropriate, but nevertheless considers that the international standards are effective and appropriate because they enable France to achieve its objective, namely, to protect human health. Canada notes that the EC hold the contrary position for the following four reasons: (i) chrysotile is a proven carcinogen; (ii) there is no exposure limit (threshold) for chrysotile; (iii) safe use is not feasible in all circumstances and for all persons and does not eliminate all the risks; and (iv) there are substitute products which are safe, or safer, than chrysotile. Under such circumstances, the EC point out that the only texts that are effective and appropriate are those which advocate ceasing to use asbestos, banning it and replacing it with substitute products. Canada, for its part, considers that the international standards make it possible to achieve the level of protection of human health deemed appropriate by France for four reasons: (i) chrysotile is less harmful than amphiboles; (ii) the understanding of certain mechanisms by which the fibres cause certain diseases makes plausible the existence of a threshold of exposure below which no cancer can develop; (iii) controlled use is possible; and (iv) the risks associated with substitute fibres are not known.
3.380 Canada maintains that there is a broad consensus among the scientific community that chrysotile asbestos is less harmful than amphibole asbestos. The WHO draws a distinction between chrysotile and amphiboles as far as the risks to human health are concerned. Three of the four experts consulted by the Panel recognize that chrysotile asbestos is less dangerous. The international standards, as well as the national regulations in a number of countries, take into account the difference in the harmfulness of different types of asbestos by laying down higher exposure limits for chrysotile than for amphiboles. Canada recalls that it has already explained that the difference in harmfulness can be explained by the different chemical compositions, physical properties and biopersistence of amphiboles and chrysotile. The question of controlled use has already been examined. Canada asserts that controlled use is feasible and effective and gives workers and do-it-yourself enthusiasts adequate protection. The control measures have proved their effectiveness in the mining and processing sectors. According to Canada, studies show that it is possible to produce chrysotile fibre and manufacture asbestos-cement or friction materials in places where exposure levels are strictly controlled and kept below safe levels, where workers are not exposed to any increased risk of disease due to exposure to chrysotile.

3.381 With regard to secondary users, in Canada’s view this is a false problem. Threats to the health of secondary users are due to friable materials, in particular sprayed asbestos already in situ. As Canada has already pointed out, this debate does not concern friable materials, but high-density products (water pipes, roof tiles, cladding, guttering, friction linings), in other words, products in which the fibres are encapsulated and cannot be released into the ambient air. The recommended methods of installation eliminate the need to cut or drill these products on building sites because they are delivered in a number of pre-cut and pre-drilled sizes according to the buyer’s specifications. If such operations are necessary, the risks can be reduced to undetectable levels by using proper hand tools or tools equipped with vacuum dust extractors. By using appropriate tools and practices (moistening, for example), the emission of fibres during intermittent work can be kept at levels below those deemed safe. In Canada’s view, the effect of intermittent exposure on the health of workers naturally depends on protective measures, but also on the intensity and duration of the exposure. It is in fact cumulative exposure that determines the risk. For those working with chrysotile intermittently, cumulative exposure levels are usually low, precisely because the work is intermittent. This is even more true for do-it-yourself enthusiasts, who work even less with high-density products than construction workers. Canada points out that these considerations come from Dr. Henderson’s reply to question 5(e) by the Panel: “… the risks from occasional or infrequent interventions on chrysotile-only products (e.g. by home ‘handymen’) – although not quantifiable because of absence of data – must be very small for lung cancer and mesothelioma, and non-existent for asbestosis”.

3.382 Canada rejects the EC’s claim that the substitutes most commonly used are “substances which have given no cause for concern after decades of massive use [...]”. The most recent data available on the dimensions of substitute fibres, their biopersistence and the reactions of the human organism to exposure to them do not enable any definitive comparison with chrysotile fibre to be made. This is also the opinion of the European Commission’s Scientific Committee on Toxicity, Ecotoxicity and Environment: “… the conclusion that specific substitute materials pose a substantially lower risk to human health, particularly public health, than the current use of chrysotile, is not well founded …” In Canada’s view, the Panel should be circumspect concerning the opinions of the experts on substitute fibres. Those who have adopted a position on this subject based themselves on a very small number of scientific data. According to INSERM’s report on substitute fibres:

“Taking into account the current uncertainties regarding the effects on humans of exposure to substitute fibres for asbestos, it is important to ensure that the levels of exposure among users of products containing substitute fibres for asbestos are as low as possible.”

3.383 For the same reasons, Canada believes that the international standards enable France to give the population the protection it considers appropriate in the health sphere.
3.384 Canada also considers that the effective and appropriate international standards have not been used “as a basis” for the adoption of the French Decree. Canada refers the Panel to the arguments in Section III.B above and the replies to the questions by the Panel (see Annex II to this report) concerning the meaning of the obligation to use the international standards or their relevant parts “as a basis” for a technical regulation. Neither the ILO nor the International Labour Office recommends replacing all types of asbestos for all applications. Convention 162 only advocates the use of substitute fibres “Where necessary to protect the health of workers and technically practicable […] by other materials or products, or the use of alternative technology […] scientifically assessed by the competent authority as harmless or less harmful […].” Article 5 of the ILO Code of Practice on Safety in the Use of Asbestos is in the same vein. Canada asserts that France did not proceed in the manner called for in the standards. It did not assess the risks for each product, each use and each application. If it had done so, it would have realized that chrysotile fibre does not have to be replaced in asbestos-cement products, which alone account for almost all applications of chrysotile. Dr. de Klerk points out that: “Asbestos-cement products containing only chrysotile pose no measurable threat to health.”

3.385 Canada notes that in the Environmental Health Criteria for Chrysotile Asbestos (203), considered by the EC to be a standard within the meaning of the TBT Agreement, the WHO recommends that chrysotile asbestos should only be replaced by safer substitute materials where this is possible. In France, however, the products replacing chrysotile are not safer. As the INSERM report on substitute fibres states:

“On the basis of the epidemiological data currently available, no conclusions can be drawn concerning the carcinogenicity of the various types of substitute fibre.”

3.386 Canada contends that, to date, no comparative scientific study has established beyond any doubt that substitute products are harmless or less harmful than chrysotile asbestos. Some recent studies even show that certain fibrous substitutes (for example, PVA fibres and cellulose fibres) have a higher degree of biopersistence than chrysotile. Canada also maintains that many scientific studies, which it mentioned during the meeting with the experts on 17 January 2000, show that, at the low levels of exposure currently prevalent in the chrysotile products industry, it is not possible to measure an increased risk for human health. International standards on asbestos do not call for a total ban without drawing distinctions that take into account the type of fibre or the use. The standards propose an approach under which asbestos regulations must take account of the type of fibre, the products in which the fibre is incorporated, and the uses of the product. The international standards thus propose a less trade-restrictive approach while at the same time laying emphasis on the effective and appropriate protection of human health. France has ignored these international standards and has banned asbestos. Moreover, the technical regulation it has adopted is so far from the international standards that their bases can no longer be perceived. According to Canada, the EC has not provided any valid explanation to justify rejecting the standards as the starting point for French regulations on asbestos. For the foregoing reasons, Canada asks the Panel to find that the Decree is inconsistent with the provisions of Article 2.4 of the TBT Agreement.

3.387 The European Communities contend that the texts cited by Canada to suggest that there exist international standards that could have been used as a basis by France are not relevant under Article 2.4 of the TBT Agreement because these texts do not correspond to the definition of a “standard” in Annex 1 to the TBT Agreement. In fact, the terms of the ILO and ISO texts mentioned by Canada do not lay down any guidelines or characteristics for products or related process and production methods nor is that their object and purpose. At best, they can be seen as assessments of the risks caused by asbestos and asbestos-containing products rather than as establishing international technical standards or conformity assessment procedures. They also aim at providing a minimum level of protection for workers, not the substantial reduction or elimination of the risks, which is the level of protection chosen by France. Thus, the EC consider that the ILO and ISO texts are not relevant standards within the meaning of Article 2.4 of the TBT Agreement and therefore
are not applicable in this case. Accordingly, in the light of the foregoing, the EC consider that the Decree is compatible with Article 2.4 of the TBT Agreement.

(e) Article 2.8 of the TBT Agreement

3.388 Canada maintains that the Decree is based on provisions relating to products in terms of their design or their descriptive characteristics, whereas it would have been appropriate to base it on their performance. Based on this fact, the Decree is incompatible with Article 2.8 of the TBT Agreement, which provides that a Member shall regulate products “Wherever appropriate ... in terms of performance...”. Regulating chrysotile fibre “in terms of performance” requires an analysis of the intended uses of this fibre. Canada maintains that the appropriateness criterion in Article 2.8 has been met to a large extent in this case inasmuch as the potential risks of chrysotile fibre make regulating it “in terms of performance” necessary, even essential. International standards address the uses of chrysotile in terms of the risks that they present. These international standards provide for technical regulation of chrysotile based on its performance. Until the adoption of the Decree, France regulated asbestos in terms of the risks that each of its uses entailed. For a long time, the technical regulation of chrysotile, in France, was therefore based on the performance of this product. According to Canada, France has not offered any justification to show that, henceforth, it would no longer be “appropriate” to follow this approach. Examination of the “wherever appropriate” criterion for regulating a product “in terms of performance” rather than “descriptive characteristics” must be done on a case-by-case basis. In this case, chrysotile fibres have no usefulness or commercial value as such. They are only used as inputs in a limited range of finished products. Chrysotile fibres only present an actual risk in certain conditions of past use, whereas hard modern products that contain these fibres do not present any detectable risk. It therefore appears quite appropriate to regulate chrysotile in terms of its performance rather than its descriptive characteristics.

3.389 Canada maintains that it is utterly inappropriate and contrary to the objectives of the TBT Agreement to consider that a product must be banned because it is potentially hazardous per se because many products would fall into this category, at least in part. In the particular case of products containing fibres, the potential risks depend on the type of fibre used, on the way in which it is incorporated in a product and the precautionary measures that are taken at the time of manufacturing, handling and using the product. If the purpose of Article 2.8 is to have technical regulations based on “performance”, it is because the potential risk of a product is generally in fact closely linked to its use. Bagged chrysotile exported by Canada is an inert product as long as it is not handled or used for particular purposes. It is these uses that can be the source of some risk. Modern products where chrysotile fibre is imprisoned do not pose any detectable risk for health. Article 2.8 requires, wherever appropriate, that a Member comply with this fact when drawing up its technical regulations. Canada maintains that, with respect to chrysotile fibre, only a regulatory approach based on an analysis of the risks, use-by-use, product-by-product, is in keeping with Article 2.8 of the TBT Agreement or, more generally, with the objectives of this Agreement. Until the adoption of the Decree, France recommended this approach, and rightly so, as it regulated uses (e.g. spraying) or specific products containing asbestos (e.g.: toys), taking account of the hazards inherent in some of these applications. Based on the foregoing, Canada concludes that, inasmuch as the Decree was not based on performance even though it was utterly appropriate to do so, France has violated Article 2.8 of the TBT Agreement.

3.390 The European Communities contend that it is important to note that Article 2.8 applies to a sub-category of “technical regulations” i.e. “technical regulations based on product requirements”. In this instance, it is clear that the Decree is not “based on” requirements relating to asbestos or asbestos-containing products. Article 2.8 is therefore inapplicable in this instance. According to the EC, the purpose of the provision is to ensure that technical rules that aim to ensure a given quality or minimum performance are, as far as possible, technically neutral and do not therefore prescribe a particular process or technology but simply set objectives to be achieved. This is perfectly clear from the English text of Article 2.8 of the TBT Agreement.
3.391 The EC contend that the ban on asbestos, even if it were considered to be a technical rule, is not a technical rule that falls within the scope of this provision. In any event, Article 2.8 of the TBT Agreement means that, wherever appropriate, the technical regulation shall be based on the performance of the product in question (i.e. based on requirements connected with the performance of the product, for example, “the product must be safe, watertight and non-flammable”), and not based on the design or descriptive characteristics of the product (i.e. not specify in detail how these requirements of safety, watertightness and non-flammability are to be attained). However, asbestos and asbestos-containing products present a major risk to human health, and in particular to the health of persons who are exposed repeatedly, occasionally or unwittingly. With the exception of the waivers granted when there is no substitute product less harmful to health or safer, there is no possible “use” of the product. Only a prohibition is capable of halting the spread of the risk. The EC consider that, under such conditions, a technical regulation that aims to prohibit the “use” of a product cannot set out the circumstances or conditions in which asbestos or asbestos-containing products are to be used. Consequently, the Decree is compatible with Article 2.8 of the TBT Agreement.

3.392 Canada maintains that the Decree is inconsistent with Article 2.8 of the TBT Agreement because the Decree’s requirements relating to chrysotile asbestos and products containing it were based on the design or descriptive characteristics of these products, whereas it would have been appropriate to base them on performance. Indeed, the Decree prohibits a product because it contains chrysotile asbestos. This ban is a product requirement. The scope of the ban (whether it affects a product or not) depends on its design or a descriptive characteristic (the product does or does not contain chrysotile asbestos). According to Canada, the EC interpret Article 2.8 incorrectly. This Article does not apply only to a sub-category of technical regulations that are “technical regulations based on product requirements”. Nor does it have the limited goal of ensuring that technical regulations aimed at ensuring minimum product performance or quality are neutral. Contrary to what the EC maintain, Canada contends that there are possible uses of chrysotile asbestos that do not involve a detectable risk to human health and can be subject to controlled-use measures. The EC wrongly claim that only a ban is capable of halting the spread of the asbestos risk.

3.393 Canada contends that the EC, in their reply to a question from Brazil, acknowledged the relevance of a regulatory approach based on product performance and, implicitly, their obligation, pursuant to Article 2.8 of the TBT Agreement, to regulate chrysotile asbestos and asbestos-containing products based on their performance. Canada notes that the EC write that the assessment of the risk must be specific to each product and that the national risk management decisions that result differ depending on each product in question. With respect to chrysotile fibre, only a regulatory approach based on an assessment of the risk on a use-by-use, product-by-product basis is compatible with Article 2.8 of the TBT Agreement and, more generally, with the objectives of the TBT Agreement. Moreover, prior to the adoption of the Decree, France advocated this approach because it regulated chrysotile asbestos based on uses (for example, spraying) or based on the hazards inherent in applications of specific products containing it (for example, toys). Canada claims that, inasmuch as it did not base the Decree on the performance of chrysotile asbestos and products containing it when it would have been possible to do so, France violated Article 2.8 of the TBT Agreement.

2. General Agreement on Tariffs and Trade (GATT)

(a) Applicability of Article III and/or Article XI of the GATT

3.394 Canada maintains that the Decree is incompatible with Articles XI:1 and III:4 of the GATT. The Decree should be examined in the light of these two Articles because it comprises two distinct aspects: on the one hand, it prohibits imports and, on the other, it contains discriminatory internal regulations. The wording of the Decree clearly shows these two aspects. With regard to the first aspect, the Decree prohibits the import of all varieties of asbestos fibres and any product containing them. According to Canada, this aspect is
subject to Article XI:1 of the GATT. With regard to the second aspect, the Decree affects domestic market use of all varieties of asbestos fibres and all products containing them. Canada considers that this point is subject to Article III:4 of the GATT. Just as two specific aspects of the same measure can be examined in the light of two different agreements, similarly, two specific aspects of the same measure can be examined in the light of two different articles of the same agreement; in this case, Article XI:1 and Article III:4 of the GATT. Specifically, Canada maintains that the first aspect of the Decree, which aims to prohibit the import of all varieties of asbestos fibres and any product containing them, is incompatible with Article XI:1. The second aspect of the Decree, which aims to prohibit, in particular, the domestic marketing and transfer on any basis of all varieties of asbestos fibres and any product containing them, is, according to Canada, incompatible with Article III:4 of the GATT.

3.395 The European Communities claim that the Decree has to be seen as an “internal regulation” providing wholly identical treatment for “like” domestic and imported products within the meaning of Article III:4 of the GATT, and must therefore be considered fully compatible with that Article. The EC note that Canada maintains that the measure contested is contrary to both Articles III:4 and XI:1 of the GATT. However, either the measure is an “internal regulation”, in which case it falls under Article III:4, or it concerns only the importation of products, in which case it must be judged in the light of Article XI:1. In the present case, the Decree only establishes a single measure that applies in a non-discriminatory manner to asbestos and covers both domestic and imported products.

3.396 The EC contend that Canada wrongly maintains that the Decree comprises two separate measures: one relating to the use of asbestos and asbestos-containing products on the domestic market (falling under Article III:4) and the other relating to imports of asbestos and asbestos-containing products (falling under Article XI:1). Such a distinction fails to take account of the relationship between Article III:4 and Article XI:1 and, according to well-established GATT practice, a single measure that applies both to domestic and to imported products is necessarily covered as a whole by Article III:4 of the GATT if it is imposed on an imported product at the time or place of importation.

3.397 The EC point out that the Decree prohibits the manufacture, processing, possession for sale, offer, importation and exportation, domestic marketing, or transfer under any title of all varieties of asbestos fibres or any product containing them. The Decree also provides for an exemption from the ban on asbestos and asbestos-containing products when there are no substitute products capable of performing a function equivalent to that of asbestos and of guaranteeing both the technical safety of use and a lower risk to workers’ health. This is a measure prohibiting the use (in any form whatsoever) of asbestos and asbestos-containing products, “which applies” both to domestic asbestos and asbestos-containing products and to imported asbestos and asbestos-containing products at the time of importation. In other words, the import ban is merely the logical corollary of the general prohibition on the use of asbestos and asbestos-containing products. The fact that the Decree applies at the border (by banning imports) so as to enforce the French measure prohibiting the use of asbestos does not render Article III:4 inapplicable.

3.398 The EC note that, in a different case, a Panel held that: “The fact that Section 337 is used as a means for the enforcement of United States patent law at the border does not provide an escape from the applicability of Article III:4”. Similarly, the Panel Report on Canada - Import, Distribution and Sale of Certain Alcoholic Drinks by Provincial Marketing Agencies raised the question of whether the measure contested (in that instance, minimum prices) fell under Article X:1 or Article III:4. It noted that, according to the Interpretative Note Ad Article III, a regulation is subject to the provisions of Article III if it “applies to an imported product and to the like domestic product” even if it is “enforced, in the case of the imported product, at the time or point of importation”. Even if the word “import” were deleted from the Decree, this would not alter the position as regards imported asbestos and asbestos-containing products in any way. Their use, in whatever form, would still be prohibited on French territory. The import ban is simply designed to make the prohibition of their use more effective in terms of inspection. For the same reason, in the case of do-
mestic products, the Decree prohibits not only their sale, but also their manufacture. The aim, then, is to halt the spread of asbestos as far upstream as possible.

3.399 The EC contend that Article III:4 must be assessed in the light of the Interpretative Note relating to it. The Panel Report on *United States - Restrictions on Imports of Tuna* is especially instructive for the purposes of the present case. In this case, the United States Marine Mammals Protection Act involved two measures. The first was a regulation governing the fishing practices of national tuna fishermen. Secondly, it also imposed an import ban on tuna or tuna-based products where the commercial fishing techniques used had the incidental effect of killing or severely injuring a greater number of marine mammals than the norms set by the United States. The Panel Report stated in general terms that:

“This suggests that Article III covers only measures affecting products as such. Furthermore, the text of the Note Ad Article III refers to a measure ‘which applies to an imported product and the like domestic product and is collected or enforced, in the case of the imported product, at the time or point of importation’. This suggests that this Note covers only measures applied to imported products that are of the same nature as those applied to the domestic products, such as a prohibition on importation of a product which enforces at the border an internal sales prohibition applied to both imported and like domestic products.”

3.400 The EC indicate that, in this case, the Panel was unable to make use of the Note Ad Article III because the internal measure governing the fishing practices of United States tuna fishermen did not cover a particular product, unlike the import measure. The situation regarding the Decree is radically different and consequently warrants examination solely in the light of Article III:4 of the GATT. The thesis that Articles III and XI cannot both apply to any one measure is apparently confirmed by the rulings of Panels which, when examining a measure, always pose the question whether it falls under Article III or Article XI. The EC note that Canada appears to share this view too, as it argued in the *Hormones* case that: "... in the alternative, the EC import prohibition infringed GATT Article XI, but noted that this claim should only be considered by the Panel if it decided that Article III of GATT did not apply in this case". Besides, it would be totally illogical if one and the same measure were permitted under Article III and prohibited under Article XI, in other words if a measure prohibiting the use of a product on national territory could not be enforced at the border.

3.401 Canada maintains that the Decree is inconsistent both with Articles XI:1 and III:4 of the GATT because it involves two separate aspects. As Canada has asserted, the terms of the text make it clear that, on the one hand, the Decree prohibits imports of asbestos or of asbestos-containing products and, on the other hand, regulates the domestic market. Canada notes that the EC do not recognize the dual aspect of the French measure, but claim that the import ban is intended simply to render more effective, in terms of control, the ban on use. According to Canada, the EC’s reasoning is tendentious in that it is incomplete. Canada notes that, in an effort to demonstrate that the measure is solely an internal measure, the EC claim that removing the “importation” aspect from the text of the measure would not in any way affect the scope of the internal provisions. Canada fully agrees with this reasoning because it is precisely what Canada asserts. However, this is only half of the Canadian reasoning.

3.402 Canada asserts that the corollary of this reasoning highlights the dual aspect of the Decree. Using the same logic as that applied by the EC, Canada can in return claim that if the “internal measure” aspect is removed from the reading of the text of the Decree, this does not change in any way the scope and effectiveness of the ban on the import of asbestos into France. That is why the measure must be examined under both Article III:4 and Article XI:1 of the GATT. Canada notes the argument of the EC and of the United States relating to the Interpretative Note Ad Article III (measures imposed “at the time or point of importation”). However, the Interpretative Note is not applicable because the import ban is not an internal measure imposed, for administrative reasons, at the border. Also, the
Interpretative Note Ad Article III only applies, as indeed the EC maintain, if the measure applies "to an imported product and to the like domestic product". Yet, it is clear to Canada that the aspect of the measure in question, namely the explicit import ban, is not applicable to the domestic product because the domestic product is obviously not "imported". The import ban is therefore aimed specifically at something that is not covered by the internal measure. Canada contends that the Decree has a dual aspect involving Articles III:4 and XI:1 of the GATT.

3.403 The European Communities assert that the Decree must be examined as an internal measure falling exclusively under Article III:4 of the GATT. When a domestic measure applies to both domestic and imported products, Article III must apply. Therefore, the contested measure may be examined as a single measure with identical implications for both domestic products and imports. Article III then rules out the application of Article XI, inasmuch as these two Articles cannot be applied cumulatively to one and the same measure. This interpretation follows both from a textual analysis of the Interpretative Note Ad Article III and from previous practice under the GATT. It also follows from an analysis, by analogy, of other legal systems.

3.404 The EC recall that, as already indicated, the Interpretative Note Ad Article III has been interpreted by the Panel in United States - Restrictions on Imports of Tuna as follows:

"This suggests that this Note [Ad Article III] covers only measures applied to imported products that are of the same nature as those applied to the domestic products, such as a prohibition on importation of a product which enforces at the border an internal sales prohibition applied to both imported and like domestic products."510

In so doing, the Panel recognized that, even if a measure is applied to imported products, the fact that that same measure is also applied to domestic products makes Article III applicable. This in itself rules out any possibility of cumulative application together with Article XI of the GATT.

3.405 The EC indicate that the Panel in United States - Measures Affecting Alcoholic and Malt Beverages noted that:

"...the issue is not whether the practices in the various states affect the right of importation as such, in that they clearly apply to both domestic (out-of-state) and imported wines; rather, the issue is whether the listing and delisting practices accord less favourable treatment – in terms of competitive opportunities – to imported wine than that accorded to the like domestic product. Consequently, the Panel decided to analyze the state listing and delisting practices as internal measures under Article III:4."511

This latter Panel shows that a domestic measure can affect the right to import as such without, for all that, falling under Article XI.

3.406 The EC claim that some legal systems, for example, that of the EC, take the same approach. Thus, with regard to the fiscal measures applicable between Member States, the Court of Justice of the EC makes a distinction between, on the one hand, measures that apply to a product only when imported, which fall under Article 12 of the EC Treaty (Elimination of customs duties or charges having equivalent effect), and, on the other, measures that apply to both imported and domestic products, which fall under Article 95 of the EC Treaty (Internal taxation). The text of Article 95 of the EC Treaty could, as a Panel has already indicated512, be compared to the text of Article III of the GATT. It follows from the above that a measure which applies to both domestic products and imported products must be examined in relation to Article III, insofar as such measure is identical in nature for both domestic products and imports. From a systematic interpretation it also follows that such a measure cannot be examined simultaneously from the standpoint of both Article III and Article XI of the GATT. According to the EC, the students of legal theory have come to the same conclusion.
“This line [between Articles XI and III] may sometimes be hard to draw. The notes to Article III provide that measures applied to domestic products, which are applied to imported products at the time of importation, are to be analyzed under Article III. Thus, if an import were barred because it failed to meet a national product standard, the permissibility of the action would be examined under Article III, not as an import ban under Article XI.”

3.407 The EC assert that, in the case of the French measure on asbestos, the domestic measure and the border measure cover the same products (asbestos and asbestos-containing products) and are of the same nature. The Decree prohibits the importation of asbestos and asbestos-containing products from third countries and, at the domestic level, prohibits the production, processing, possession for sale, offer, exportation and transfer of these products. The practical application of the Decree leads to the same measure - that is, the general ban on asbestos and asbestos-containing products - being applied to all products, irrespective of their origin. As one and the same measure is applied to domestic products and imported products - in the case of the latter at the border - the Communities conclude that only Article III:4 of the GATT is applicable in this case, which rules out the application of Article XI. With regard to the Canadian argument that France does not manufacture asbestos or manufactures it only in extremely limited quantities, the EC wish to point out that asbestos was produced in France in the past. The potential for production remains, insofar as asbestos in its natural form is present on French territory.

3.408 Canada asserts that the Decree should be considered as falling under both Article XI and Article III of the GATT. The Decree’s provisions banning imports of asbestos are incompatible with Article XI:1 and the Decree’s provisions banning the sale of asbestos and other transactions on the French market are incompatible with Article III:4. In this connection, Canada points out that the report in United States – Restrictions on Imports of Tuna, cited by the EC several times to underpin its thesis of systematic interpretation, was not adopted by the Contracting Parties. Canada indicates that, if the Panel considers that the Decree cannot be examined under the two Articles, it should be considered as a measure affecting imports and incompatible with Article XI:1. If the Panel considers that the Decree cannot be examined under the two Articles and cannot be considered as a measure affecting imports, it should be considered as a measure affecting sales and other transactions on the French market and thus incompatible with Article III:4.

3.409 The European Communities contend that the Decree constitutes a single measure that applies to both domestic and imported products alike. As one and the same measure is applied to both domestic products and imported products – in the case of imports at the border - the EC conclude that only the arguments of Canada relating to Article III:4 of the GATT are relevant in this case. According to the EC, this interpretation rules out the cumulative application of Articles III and XI of the GATT, as the Note Ad Article III provides and previous GATT Panel reports have clearly established (see, for example, the first Panel report in the Tuna/Dolphin case). Indeed, the Decree prohibits the importation of all types of asbestos and products containing asbestos from all third countries and, at the domestic level, prohibits the production, processing, possession for sale, offer, exportation and transfer of these products. The practical application of the Decree thus leads to the same result, namely, a general ban on asbestos and products containing asbestos applicable to all products irrespective of their origin. The EC claim that, in order to establish a violation of Article III:4 of the GATT, the complaining party has the burden of showing that there is de jure or de facto discrimination in the treatment of imported compared to like domestic products.

(b) Article III:4 of the GATT

(i) Application of Article III:4 of the GATT

3.410 Canada maintains that, in adopting the Decree banning asbestos, France violates the national treatment disciplines in Article III:4 of the GATT 1994. The effect of the French measure is to favour the French fibre industry and that of substitute products such
as chrysotile fibre and chrysotile-cement products, which is prohibited under Article III:4 of the GATT 1994. According to case law under the GATT 1947 and the GATT 1994, examination of the applicability of Article III:4 to a measure taken by a Member and, if need be, of its compatibility with the disciplines of Article III:4, has two separate facets. The first is to determine whether the measure constitutes a law, a regulation or a requirement affecting the sale, offering for sale, purchase, transportation, distribution or use of an imported product on the domestic market. Canada asserts that the Decree banning asbestos is a regulation affecting these activities. The second facet is to determine whether the products from the territory of a Member imported into the territory of another Member are subject to treatment less favourable than that accorded to like products of national origin in respect of all laws, regulations or requirements affecting, *inter alia*, their supply, offering for sale, or sale. With respect to the second facet, Canada draws attention to the following three points: (i) products like Canadian chrysotile fibre and chrysotile-cement exist; (ii) these like products are of French origin; and (iii) they benefit from treatment more favourable than that accorded to imported Canadian chrysotile fibre and chrysotile-cement products. Canada notes that, in this case, France recognized in specific terms in the Decree banning asbestos that chrysotile fibre and the products containing it are like the substitute fibres and materials, products or devices containing the latter.

3.411 Canada maintains that the likeness of these products is confirmed by application of the criteria established in the case law, i.e. end-use of the product, consumers’ tastes and habits, the physical properties, the nature and quality of the product, as well as tariff classification. According to these criteria, the substitute fibres are like chrysotile fibre and fibro-cement products are like chrysotile-cement products. Furthermore, a substitute fibre manufacturing industry and a sizeable industry for the manufacture of fibro-cement products exist in France. The prohibition of a number of operations - including domestic marketing, sale and transfer on any basis - relating to chrysotile fibre and products containing it unequivocally constitutes treatment less favourable than that accorded to like French substitute fibres and products that are not subject to any prohibition regulations of the same type.

3.412 Canada contends that the Decree constitutes a regulation affecting the sale, placing on the market or purchase of chrysotile fibre and chrysotile-cement products in France. According to Canada, Article III:4 applies to “all laws, regulations and requirements”. Article III:4 applies to the Decree, which is a regulation of the French Government. Pursuant to the terms of Article III:4, a measure must affect “the internal sale, offering for sale, purchase, transportation, distribution or use ...” on the domestic market. The Decree specifically aims to prohibit the manufacture, processing, sale, domestic marketing, possession for sale, offer or transfer on any basis of all varieties of asbestos fibres and products containing them. In Canada’s view, the Decree undoubtedly constitutes a regulation directly affecting the sale, release for sale or purchase of chrysotile fibre and products containing it. The Decree applies to France’s domestic market pursuant to the terms of Article III:4. In prohibiting the sale, release for sale or purchase, the Decree changes the conditions of competition, in the domestic market, between substitute fibres and products containing them of French origin and chrysotile fibre and products containing it originating in Canada. Canada concludes that the conditions regarding the applicability of Article III:4 of the GATT have been met.

3.413 The *European Communities* assert that the Decree has to be examined as an “internal regulation” that provides perfectly identical treatment for “like” domestic and imported products within the meaning of Article III:4 of the GATT and it must therefore be considered fully compatible with that Article.

(ii) *The concept of “like products”*

3.414 Canada claims that, in the case of the Decree, the question of determining whether the treatment accorded is less favourable for the imported product than for the like product of national origin presupposes the examination of the following three points: (i) the existence of products like Canadian chrysotile fibre and chrysotile products; (ii) the French
origin of those like products; and (iii) less favourable treatment accorded to chrysotile-cement products and to imported chrysotile fibre than that accorded to like French products. According to the report by the Working Party on Border Tax Adjustments, likeness of products shall be assessed on a “case-by-case” basis considering, in particular, end-use of the product, consumers’ tastes and habits and the properties, nature and quality of the product. The Appellate Body describes the Working Party report on Border Tax Adjustments as setting out “the basic approach for interpreting ‘like or similar products’, generally, in the various provisions of the GATT 1947.” Tariff classification was added as a supplementary element to the above-mentioned criteria. Analysis of the likeness of products is therefore based on criteria such as (i) end-use of the product; (ii) consumers’ tastes and habits; (iii) the properties, nature and quality of the product; and (iv) the tariff classification. Precedents under the GATT 1947 and the GATT 1994 do not require that all the criteria apply for the purpose of analysing the likeness of products. For example, in the report Japan – Customs Duties, Taxes and Labelling Practices on Imported Wines and Alcoholic Beverages, it was determined that various types of alcohol were like products because they constituted a well-defined and single product at the end-use stage.

3.415 Canada maintains that the term “like” does not mean identical. It is rather a matter of showing that fibres and products that are substitutes for chrysotile fibre and chrysotile-cement products are similar in many ways and, according to the Decree they are not covered by the national treatment rule under Article III:4. More than 150 substitute fibres for chrysotile fibre exist. In the products in which they are incorporated, they attempt to replicate the properties of chrysotile fibre. The most common are the aramid fibres, PVA fibres, cellulose fibres, glass fibres, ceramic fibres, rock wool, and wollastonite. Virtually all the chrysotile fibre of Canadian origin imported into France before the Decree took effect was intended for the manufacture of chrysotile-cement products. Canada uses the example of chrysotile fibre and chrysotile-cement in order to show the similarity of chrysotile fibre and substitute fibres and of products containing chrysotile and those containing substitute fibres. Chrysotile fibre and PVA, cellulose and glass fibres are used interchangeably, the first in the manufacture of chrysotile-cement, the three others in manufacturing fibro-cement. Chrysotile-cement and fibro-cement are used in the manufacture of products such as pipes, pipe fittings (casings, bends, sealing joints), corrugated sheets, insulation and soundproofing board, exterior siding and roofing shingles, floor tiles, slabs, gutters, chimney cowls and sinks.

3.416 Canada asserts that the guiding principle underlying the examination of the question of similarity is to proceed on a case-by-case basis, i.e. taking into account the particular circumstances of each situation. However, in this instance, France undertook to ban chrysotile asbestos for all possible uses and to replace it with like substitute fibres capable of performing an equivalent function. The Decree indicates that products like chrysotile fibre, chrysotile-cement products, as well as all other products using chrysotile fibre, exist. This observation ensues from the terms of the Decree, which require substitute products to perform an “equivalent function” and offer the same “technical guarantees” as chrysotile-based products. Under Article 2 of the Decree, the ban does not apply to certain products that contain chrysotile fibre when, to ensure an equivalent function, “no substitute for that fibre is available which, on the one hand, in the present state of scientific knowledge, poses a lesser occupational health risk than chrysotile fibre to workers handling those materials, products or devices; [and] on the other, provides all technical guarantees of safety corresponding to the ultimate purpose of the use thereof.” In France, chrysotile fibre has been replaced in all products in which it was used, with the exception of four uses. According to Canada, the replacement of chrysotile fibre in all cases in which there exists a substitute fibre capable of performing an equivalent function presupposes that the chrysotile fibre and the substitute fibre are nearly perfectly alike. For example, the replacement of chrysotile fibre in fibro-cement products implies an admission that the substitute fibres have like characteristics. Finally, to satisfy the requirement under the Decree that a substitute fibre must provide all technical guarantees of safety appropriate to the end-use, it is essential that the end-uses of the products manufactured using chrysotile fibre be the same. According to the precedents, products are like products if their end-uses are the same.
3.417 Canada claims that the application of the criteria from case law confirms that PVA, cellulose and glass fibres and chrysotile fibre, on the one hand, and fibro-cement products and chrysotile-cement products, on the other, are like products.

3.418 As far as end-use is concerned, given the nature of chrysotile fibre - a raw mineral resource - Canada maintains that, in accordance with the case-by-case approach, special importance must be attached to the end-use criterion for the product when examining whether chrysotile fibre is like the substitute fibres. Chrysotile fibre has no use in its raw form; it serves as an input in the production of chrysotile materials. The most widespread end-use of chrysotile fibre is chrysotile-cement. Around 90 per cent of Canadian chrysotile fibre previously imported into France was used for this purpose. Like chrysotile fibre, PVA fibre, cellulose fibre and glass fibre are used in the manufacture of fibro-cement. After they have been incorporated in the cement, chrysotile fibre, on the one hand, and PVA, cellulose and glass fibres, on the other, are used for the manufacture of chrysotile-cement products and fibro-cement products respectively. These products constitute, “at the end-use stage”, “a well-defined and single product” intended for the same purposes: whether a chrysotile-cement or a fibro-cement product. The Decree points to the existence of substitute fibres for chrysotile fibre that perform an “equivalent” end function and provide “all technical guarantees of safety to the ultimate purpose of the use thereof”. Canada maintains that the conclusion that chrysotile fibre and PVA, cellulose and glass fibres are “like products” under Article III:4 of the GATT is derived from applying the sole criterion of end-use of the product.

3.419 Canada claims that the fact that chrysotile-cement products and fibro-cement products have the same end-uses is proof that they are “like products”. Chrysotile-cement and fibro-cement boards are used for insulation and soundproofing. Sheeting, shingles and tiles are used for cladding exteriors, roofs and floors. Pipes and pipe fixtures are used for pipeline systems or, for industrial purposes, in the transportation of liquids. Chrysotile-cement and fibro-cement products are like products because they are manufactured using either chrysotile-cement or fibro-cement without distinction, because they are finished products the end-use of which is identical and because the Decree recognizes that the technical guarantees with regard to “end-use” are identical. In Canada’s view, the Decree concludes that there are substitute products for chrysotile-cement products that perform an “equivalent” function and provide “all technical guarantees of safety appropriate to the use” of chrysotile-cement products.

3.420 With regard to the properties, quality and nature of the product, Canada contends that the nature of chrysotile fibre (natural mineral fibre), PVA fibre (synthetic organic fibre), cellulose fibre (natural organic fibre) and glass fibre (man-made mineral fibre) is the same inasmuch as fibres are involved, be they organic or mineral, man-made or natural. Substitute fibres for chrysotile fibre are used to replicate the qualities of chrysotile fibre in view of the end-use. Chrysotile-cement or fibro-cement manufacturers use them for identical purposes, which points to the likeness of the properties, nature and quality of chrysotile, PVA, cellulose and glass fibres. In addition, PVA, cellulose and glass fibres, just like chrysotile fibres, give composites both strength and resistance. Their chemical resistance and their quality as a binder and reinforcing agent are useful in the manufacture of chrysotile-cement and fibro-cement. In Canada’s view, the likeness of the manufacturing processes for chrysotile-cement and fibro-cement shows the likeness between the properties and the nature of the fibres in question.

3.421 Canada states that chrysotile-cement products, like fibro-cement products, are known for their greater durability than cement products not containing fibre. They are also known for their chemical resistance, their insulating properties, both for heat and sound-proofing, and their lightness. Chrysotile-cement and fibro-cement, whether they contain a percentage of chrysotile fibre or other like fibres, are manufactured using the same technical process. Indeed, the Hatschek process and the Mazza process - its derivative for pipe production – are used in the manufacture of chrysotile-cement and fibro-cements. The likeness of the manufacturing processes indicates a convergence as regards the properties, quality and nature of the products. According to Canada, the Decree recognizes that, ex-
cept in four cases, like asbestos-based products exist which offer “the same technical guarantees”. Fibro-cement products have replaced chrysotile-cement products. By offering the same technical guarantees as chrysotile-cement products, fibro-cement products undoubtedly have the same properties, are of the same quality and the same nature. Given that chrysotile-cement products and fibro-cement products have the same intrinsic qualities, that they are manufactured using the same technical process and that the Decree recognizes that they offer the same technical guarantees, for Canada, the obvious conclusion is that chrysotile-cement products and fibro-cement products are “like products”.

3.422 Canada asserts that, according to the consumers’ tastes and habits criterion, chrysotile fibre and PVA, cellulose and glass fibres, which are inputs in chrysotile-cement products and fibro-cement products, are “like products”. Chrysotile fibre, PVA fibre, cellulose fibre and glass fibre are not products of mass consumption. These products are used by a limited number of economic agents, in particular the manufacturers of chrysotile-cement and fibro-cement products, who incorporate these fibres into their products. In so doing, these manufacturers are the consumers of chrysotile fibre for the purposes of the consumers’ tastes and habits criterion. According to Canada, the drop in chrysotile asbestos imports intended for chrysotile-cement products in 1996 and 1997 is a result of the Decree and not of a sudden change in consumers’ tastes and habits. To change the preferences of French consumers, the Decree on the banning of asbestos was necessary. The impact of the Decree does not reflect consumers’ tastes and habits. Businesses manufacturing chrysotile-cement products have converted to substitute fibres or have closed down. The Decree forced these conversions or closures. Canada maintains that, in determining the likeness of products, it is inappropriate to consider the criterion of the habits of manufacturers who were forced to convert to fibro-cements.

3.423 Canada contends that chrysotile-cement products and fibro-cement products are both industrial products and it is almost impossible to distinguish between them based on their external appearance. Thus, in the eyes of the consumer, chrysotile-cement products and fibro-cement products are like products in all aspects, unless a data sheet is available showing which fibre makes up their composition. From the point of view of the tastes and habits of French consumers, chrysotile-cement products and fibro-cement products are interchangeable. The ban on the import of chrysotile-cement products did not cause a fall in the imports of fibro-cement products. Since the Decree took effect, the import of chrysotile-cement products has simply been replaced by the import of fibro-cement products. If products like asbestos-cement products had not been available, imports into France of fibro-cement products, which include asbestos-cement products, would have dropped immediately after the import of chrysotile-cement products was prohibited. According to Canada, the fact that it is impossible to distinguish between chrysotile-cement products and fibro-cement products based on their external appearance and their interchangeability are elements which indicate that consumers perceive these products as “like products”.

3.424 Concerning tariff classification, according to the World Customs Organization, chrysotile-cement products and fibro-cement products are so similar that the 107 six- or eight-digit codes given to chrysotile-cement products in the Harmonized System are identical to the 107 codes given to fibro-cement products. Chrysotile-cement and fibro-cement products are in Heading 68.11 of the Harmonized System, i.e. the category relating to “Articles of asbestos-cement, of cellulose fibro-cement or the like”. The World Customs Organization defines Heading 68.11 as follows:

Chapter 68 – Articles of stone, plaster, cement, asbestos, mica or similar materials

Heading 68.11 – Articles of asbestos-cement, cellulose fibro-cement or the like

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
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<tbody>
<tr>
<td>6811.10</td>
<td>Corrugated sheets</td>
</tr>
<tr>
<td>6811.20</td>
<td>Other sheets, panels, tiles and similar articles</td>
</tr>
<tr>
<td>6811.30</td>
<td>Tubes, pipe and tube or pipe fittings</td>
</tr>
<tr>
<td>6811.90</td>
<td>Other articles</td>
</tr>
</tbody>
</table>
Canada indicates that this category includes fibro-cements made of cellulose, synthetic polymer (i.e. PVA) or glass fibres. As proof that chrysotile-cement and fibro-cement are like products, tiles, for example, made of chrysotile-cement, cellulose-cement or the like are given the same code, i.e. 6811.2011. Chrysotile fibre and PVA, cellulose and glass fibres, on the one hand, and chrysotile-cement products and fibro-cement products, on the other, are “like products” for the purpose of Article III:4 of the GATT. This conclusion derives from the application of criteria such as the end-use of the products, consumers’ tastes and habits, the properties, quality and nature of the products and their tariff classification. Each of the four criteria taken separately leads to the same conclusion.

The European Communities consider that asbestos and asbestos-containing products, on the one hand, and substitute products, on the other, are not “like” products within the meaning of Article III:4 of the GATT. According to established GATT practice, four specific criteria can be used to assess whether products are “like products”: (i) their properties, nature and quality; (ii) their tariff classification; (iii) their end-use; and (iv) consumers’ tastes and habits. The Panel Report on United States - Gasoline held that “those criteria were also applicable to the examination of like products under Article III:4”. On the issue as to what criteria should be applied in considering what are “like products”, the Appellate Body report on Japan - Taxes on Alcoholic Beverages stated that:

“No one approach to exercising judgement will be appropriate for all cases. The criteria in Border Tax Adjustments should be examined, but there can be no one precise and absolute definition of what is ‘like’. The concept of ‘likeness’ is a relative one that evokes the image of an accordion. The accordion of ‘likeness’ stretches and squeezes in different places as different provisions of the WTO Agreement are applied. The width of the accordion in any one of those places must be determined by the particular provision in which the term ‘like’ is encountered as well as by the context and the circumstances that prevail in any given case to which that provision may apply.”

The EC consider that Canada is confusing the concept of “like products” contained in Article III:4 of the GATT with that of “directly competitive and substitutable products” contained in Article III:2 of the GATT, read in conjunction with the relevant Interpretative Note. However, the two concepts are radically different. This can be seen by examining the texts in question. Article III:2 contains a concept that is not explicitly contained in Article III:4. The Panel report on EEC - Measures on Animal Feed Proteins noted that “the General Agreement made a distinction between ‘like products’ and ‘directly competitive and substitutable products’.” Moreover, the Appellate Body has held that:

“If imported and domestic products are not ‘like products’ for the narrow purposes of Article III:2, first sentence, then they are not subject to the strictures of that sentence and there is no inconsistency with the requirements of that sentence. However, depending on their nature, and depending on the competitive conditions in the relevant market, those same products may well be among the broader category of ‘directly competitive or substitutable products’ that fall within the domain of Article III:2, second sentence.”

The EC point out that, in the case at issue, although certain fibrous products (aramids, PVA, cellulose) and non-fibrous products (plastic, ductile iron) are indeed “substitutable” for chrysotile asbestos and products containing it, they are nevertheless not like products. Asbestos has unique physical characteristics and properties that make it difficult to replace for certain industrial purposes, especially in the chemical, petrochemical, aeronautical and nuclear industries. Amphibole asbestos is in fact the only product that is truly a like product to chrysotile asbestos. And it is precisely because asbestos is uniquely suited for certain industrial uses that the Decree envisages some exceptions to the general prohibition.

The EC consider that, in this case, three criteria are pertinent when assessing the similarity of products pursuant to Article III:4. These criteria are the products’ properties,
nature and quality, tariff classification and end-use. As far as the tastes and habits of consumers are concerned, the EC consider that, while this criterion may be relevant in certain cases (everyday consumer goods), it is not relevant in the case of asbestos and asbestos-containing products. The Panel Report on United States - Gasoline also made use of the first three criteria, but ignored the tastes and habits of consumers.

3.430 The EC claim that the properties, nature and qualities of the products are different, both for asbestos fibres and substitute products and products containing asbestos and substitute products. As the EC have explained in their factual arguments, asbestos fibres have a very particular fibrous texture. They consist of bundles of small fibrils, stuck to each other. The fibrils separate very easily lengthways under the effect of machining, shock, vibration, friction (or simply draughts if it is a friable material) to form a cloud of very fine dust, often invisible to the naked eye, which can settle anywhere and penetrate very deep into the lungs. Asbestos fibres are minute in diameter (less than 1 micron) so there may be high concentrations in the air. According to the EC, the properties of asbestos fibres serve to underline their uniqueness. Although they pose a danger to health, their unique characteristics incontestably make them a mineral with exceptional physical and chemical properties. They do not burn, are remarkably resistant to various kinds of chemical attack and show high mechanical resistance to traction. No other product, natural or synthetic, exhibits all of the properties that asbestos fibres possess, for the simple reason that no substitute product has the same characteristics as these fibres. The explanatory notes to the Harmonized System (“HS”) acknowledge the special nature of asbestos fibres, stating that:

“Asbestos is a natural mineral substance produced by the decomposition of certain rocks. It has a very characteristic fibrous texture; it is sometimes silky in appearance and the colour varies greatly, being usually white, but sometimes grey, greenish, blue or dark brown …”537

3.431 The EC indicate that the characteristics of asbestos fibres make them particularly dangerous to health as they increase the risk of cancer. It is the diameter of a fibre that determines how long it will remain in suspension in the air. Consequently, the smaller the diameter, the higher the risk of cancer. In addition, the greater the fibrillation, the higher the risk of inhalation, again increasing the risk of cancer. Since 1977, the WHO has recognized the existence of a link between the characteristics of asbestos fibres and their danger to health by classing them (including chrysotile fibres) in “category I” of proven carcinogens. By contrast, none of the substitute products for chrysotile asbestos is classified as carcinogenic for man. Substitute fibres possess different characteristics from chrysotile: they are much bigger in diameter than asbestos fibres, up to 40 microns thick. Substitute fibres also show a more limited fibrillation capacity. Moreover, many substitute products are not fibrous in texture (for example, plastic, ductile iron or plaster). The EC point out in this connection that Canada mentions only fibrous substitute products (cellulose, p-aramid, PVA), making no reference at all to non-fibrous products. Yet non-fibrous products are very easily and commonly used instead of asbestos-cement, which used to account for 90 per cent of the chrysotile used in France. The EC conclude that the very nature, composition, physical properties and proven effects on human health of chrysotile make it radically different to substitute products, whether fibrous or non-fibrous. For this reason, in 1986 the ILO538 – followed by the WHO in 1996539 and 1998540 – recommended replacing asbestos by less harmful materials or technologies wherever possible.

3.432 Regarding asbestos-containing products and substitute products, the EC point out that asbestos fibres possess such special characteristics that they necessarily have an impact on the properties of the product in which they are incorporated, making that product extremely dangerous to health too. If one compares a PVC pipe with a chrysotile-cement pipe, for instance, it is impossible to say that the products are of the same nature. One is made of cement (with asbestos added), whereas the other is made of plastic. The EC note that if one compares a slab of chrysotile-cement used as a roof covering with a tile or slate, it is evident once again that the products are completely different in kind. They do not have the same properties or qualities either. Here again, in the EC’s view, it is impossible to speak of “similarity” between asbestos-containing products and substitute products.
The EC note that Canada is not wrong when it refers to “convergence” and “resemblance” in terms of the properties, quality and nature of the products concerned. But the fact remains that “convergence” and “resemblance” certainly do not mean “likeness” within the meaning of Article III:4 of the GATT. The EC claims that this lack of “similarity” between asbestos-containing products and substitute products is also confirmed by the fact that, as the materials obtained after replacing asbestos have different physical and mechanical properties from the material which contained asbestos, each substitution operation requires, on the part of manufacturers, careful checks on the properties of the new material and sometimes a complete redefinition of the field of application of the product. This is especially true in the case of gaskets and plaited sealing twine.

3.433 The EC assert that the tariff classifications are different, both for asbestos fibres and substitute products and for asbestos-containing products and substitute products. Canada does not deal with the question of tariff classification for asbestos fibres. It should be recalled that the explanatory notes to the Harmonized System (“HS”) recognize the special nature of asbestos fibres by stating that “Asbestos … has a very characteristic fibrous texture …”541 Given the differences due to the properties, nature and quality of asbestos fibres, the Harmonized System takes the logical step of classifying them under a single tariff heading: HS 25.24. This heading includes chrysotile asbestos fibre. The only other products covered by the same heading are the other varieties of asbestos fibres such as amphibole asbestos fibres. Substitute fibres, on the other hand, all fall under different tariff headings. The EC consider that, in the light of the tariff classification criterion, therefore, there can be no likeness between asbestos fibres and substitute fibres. In the second case – asbestos-containing products and substitute products – the EC point out that many substitute products come under a different tariff heading from asbestos-containing products. For example, rock wool and glass wool fall under heading HS 68.06, which does not cover asbestos-containing products. Some asbestos-containing products, on the other hand, fall under a specific tariff heading such as HS 68.12, which covers in particular clothing, cords, yarn and thread, and gaskets containing asbestos. This heading does not cover any products that do not contain asbestos. As for articles of chrysotile-cement (HS 68.11), the EC must point out that they can be replaced by many other products that fall under different tariff headings. For example: (i) articles of plaster come under tariff heading HS 68.09; (ii) articles of cement are under tariff heading HS 68.10; and (iii) plastic pipes come under tariff heading HS 39.17. By the criterion of tariff classification, the EC conclude that there can be no similarity between asbestos-containing products and products not containing asbestos.

3.434 The EC claim that the end-uses are different. Canada takes the line that the concepts of “like products” and “competitive or substitutable products” are equivalent and that the criterion of end-use is the key to determining what are “like products” within the meaning of Article III:4 of the GATT. However, the two concepts are radically different. Although end-use is the key criterion for determining whether two products are directly competitive or substitutable542, this is not true for “likeness” within the meaning of Article III:4, which is essentially “technical” in nature. The EC note that this was pointed out in the clearest terms in the Panel Report on Japan - Taxes on Alcoholic Beverages.

"In the view of the Panel, the wording of the term ‘directly competitive or substitutable’ does not suggest at all that physical resemblance is required in order to establish whether two products fall under this category. This impression, in the Panel’s view, was further supported by the words ‘where competition exists’ of the Interpretative Note; competition can and does exist among products that do not necessarily share the same physical characteristics. In the Panel’s view, the decisive criterion in order to determine whether two products are directly competitive or substitutable is whether they have common end-uses, inter alia, as shown by elasticity of substitution. The wording of the term ‘like products’ however, suggests that commonality of end-uses is a necessary but not a sufficient criterion to define likeness. In the view of the Panel, the term ‘like products’ suggests that for two products to fall under this category they must share, apart from commonality of end-uses, essentially the same physical characteristics."543
In other words, the end-use is not in itself conclusive in deciding whether products are “like” within the meaning of Article III:4 of the GATT.

3.435 The EC argue further that, even where products may have some end-uses in common, that is not sufficient grounds to class them as “like products”, if each of them also has many other end-uses. As the EC has already asserted above that asbestos and asbestos-containing products are different in terms of their properties, nature, quality and tariff classifications; the criterion of end-use cannot, by itself, invalidate the conclusion that such products are not “like” products within the meaning of Article III:4 of the GATT. In any event, substitute products can be used for many other purposes than the uses to which asbestos fibres or asbestos-containing products can be put, and vice-versa. For the EC, every asbestos-free substitute product necessarily has many uses that asbestos-containing products do not have. In conclusion, the Decree is compatible with Article III:4 of the GATT 1994 inasmuch as: (i) there is no protection of domestic industry; (ii) there is no discriminatory treatment of imported products compared with domestic products de jure or de facto; and (iii) there is no likeness within the meaning of Article III:4 of the GATT between asbestos and asbestos-containing products, on the one hand, and their substitute products on the other.

3.436 Canada contends that case law in the GATT and the WTO indicate that the concept of “like product” in Article III:2 must be interpreted narrowly. This narrow interpretation does not, however, apply to Article III:4, where the concept of likeness must be interpreted more broadly, given the purpose and context of Article III:4. Consequently, Article III:4 encompasses a broader range of like products than Article III:2, first sentence. Canada also refers to its response to question 34 of the Panel (see Annex III).

3.437 Canada indicates that, for the purposes of Article III:4 of the GATT, it does not invoke the argument of likeness with respect to non-fibrous substitutes (e.g. PVC, ductile iron). Nor does Canada extend the argument of likeness to non-fibrous products used as substitutes for chrysotile-cement products. The argument of likeness put forward by Canada is limited to glass fibre, cellulose fibre, and PVA fibre, even though the range of substitute fibres is broader, as well as to fibro-cement products incorporating these types of fibre. Canada does not consider that, in order to demonstrate the violation of Article III:4 of the GATT and Article 2.1 of the TBT Agreement, it is incumbent upon it to cite all the products like to chrysotile or to chrysotile-cement products. For there to be a violation, it simply suffices to demonstrate that, for an imported product or a series of given products, there are like products that enjoy more favourable treatment. According to Article III:4, these like products must be of national origin. Canada notes that the EC go to a great deal of trouble to assert that PVC and ductile iron are not like products to chrysotile fibre and chrysotile-cement products. However, this analysis, although interesting, is completely irrelevant for determining whether glass fibre, cellulose fibre, PVA fibre, as well as fibro-cement products incorporating these types of fibre, are products like to chrysotile fibre and chrysotile-cement products.

3.438 Canada points out that, of the four criteria that can be used to determine the likeness of products, it takes into account end-use and tariff classification, as well as the product’s properties, nature and quality. These criteria show that the products in question are like products. Canada also agrees with the EC that consumer tastes and habits are not relevant in this case.

3.439 Concerning the end-use criterion, Canada emphasizes that chrysotile fibre and chrysotile-cement products are like products to PVA, cellulose and glass fibres, and to fibro-cement products incorporating them. Canada reiterates the importance of an analysis of likeness on a “case-by-case” basis, i.e. based on the circumstances. In this case, because these products are inputs that cannot be used in their raw state, particular importance must be accorded to the end-use criterion in these circumstances. Canada notes that the EC appear to claim that the end-use criterion can be decisive only for determining whether two goods are directly competitive or substitutable according to Article III:2, first sentence. This question might be relevant if Article III:2 was being addressed, but this is
not the case. Canada recognizes that end-use is a “decisive criterion” for determining the “substitutability” or directly competitive nature of two products. In Canada’s view, however, end-use is not the only relevant factor with respect to “substitutability”, as the EC suggest. End-use can be equally important with respect to likeness, as the Panel dealing with Article III:2 in Japan - Taxes on Alcoholic Beverages, asserts: “In the Panel’s view, the wording makes it clear that the appropriate test to define whether two products are ‘like’ or ‘directly competitive or substitutable’ is the marketplace.” The market determines the end-use of a product. Canada notes that the EC maintain that substitutability and likeness are two radically different concepts. Canada takes the opposite view, like the Appellate Body in Korea - Taxes on Alcoholic Beverages, namely, that likeness is only a sub-set of substitutability and that like products are by definition substitutable. In any event, the jurisprudential distinction between substitutability and likeness in Article III:2 is not necessary because in this case it is a question of Article III:4. The concept of substitutability is inevitably implicit in Article III:4 whenever products are deemed to be like products according to the broader criterion of likeness in Article III:4.

3.440 Canada indicates that the tariff classification of chrysotile-cement products is exactly the same for 107 separate products in the Harmonized System (HS). Heading 68.11 of the HS contains “Articles of Asbestos-Cement, of Cellulose Fibro-cement or the Like”. The HS describes the products contained in heading 68.11 as follows: “This heading covers hardened articles consisting essentially of an intimate mixture of fibres (for example, asbestos, cellulose or other vegetable fibres, synthetic polymers [PVA] or glass fibres).” All of the like products cited are found together, under 107 customs codes with six or eight common digits. For example, shingles made of chrysotile-cement, cellulose-cement, PVA-cement and glass-cement fall under code 6811.2011. Canada indicates that, despite these facts, the EC continue to argue, less than convincingly, that “By the criterion of tariff classification, therefore, there is no similarity between asbestos-containing products and products not containing asbestos.” Canada is surprised that the EC simply reject the argument of likeness for the products cited by Canada merely by demonstrating that products which are not cited by Canada (PVC, ductile iron, rock wool, etc.) are not like products for the purposes of Article III:4. Such an approach is unfounded. Canada considers that the fact that glass wool or rock wool products do not fall under the same heading as chrysotile-cement products in no way changes the fact that chrysotile-cement products are found in exactly the same heading as articles containing PVA fibre, glass fibre and cellulose fibre.

3.441 Canada reaffirms its arguments regarding the properties, nature and qualities of the products in question and calls on the Panel not to lend too much weight to the distinctive physical properties of asbestos, put forward by the EC, i.e. that asbestos fibre is silky, and white, grey, green, blue or brown in colour. Canada notes that the EC are also endeavouring to reject the argument of likeness advanced by Canada by stating that the “lower” pathogenicity of substitute fibres makes it impossible to conclude that they are like products. As Canada has already asserted, the inadequate state of scientific knowledge about substitute fibres makes it impossible to conclude that they are less carcinogenic than chrysotile fibres. In any event, the fact that a product has a greater or lesser impact on health is not a criterion that makes it impossible to conclude that they are like products. In this respect, wine and vodka are like products, even if their effects on health are different.

3.442 The European Communities emphasized that it follows clearly from a textual analysis of Article III:4 that the “like products” provision therein does not cover “directly competitive or substitutable” products. In fact, whereas Article III:2, second sentence, read in the light of the corresponding Interpretative Note, provides for an examination of directly competitive or substitutable products, Article III:4 mentions only the consideration of “like domestic products”. This difference in the texts is not a matter of chance. If the authors of Article III:4 had intended to include an analysis of directly competitive or substitutable products, it would have had to appear in the text of Article III:4 or in an Interpretative Note, as was done in Article III:2, second sentence. Any other interpretation would result in giving a provision of the GATT a meaning that the signatories to the Agreement did not intend. The EC recall in this connection that the Appellate Body considered that
"directly competitive or substitutable products" were a “broader” category than “like products”.549

3.443 The EC, having already developed this point, only wish to point out that, at the first meeting of the Panel with the parties, Canada stated that “the Panel in Japan - Alcoholic Beverages stated that all similar products are, by definition, directly substitutable”.550 In the EC’s view, however, the passage from the Panel Report mentioned by Canada needs to be cited in full. In fact, the Panel in Japan - Taxes on Alcoholic Beverages stated that:

“In the view of the Panel, like products should be viewed as a subset of directly competitive or substitutable products. The wording (‘like products’ as opposed to ‘directly competitive or substitutable products’) confirmed this point, in the sense that all like products are, by definition, directly competitive or substitutable products, whereas all directly competitive or substitutable products are not necessarily like products.”551

3.444 The EC also note that a Panel has already concluded that the concept of “like products” in the context of Article III:4 does not cover directly competitive or substitutable products. Thus, in the case of EEC – Measures on Animal Feed Proteins, the Panel observed that:

“Having regard to its own conclusion with respect to ‘like products’, the Panel was satisfied that animal, fish and synthetic proteins could not be considered as ‘like products’ for the purpose of Article III:4. Since the obligations under Article III:4 relate to ‘like products’, the Panel concluded that the non-application of the EEC measures to these products was not inconsistent with the EEC obligations under the Article.”552

The EC point out that, for a better understanding of this conclusion, it should be noted that the same Panel also pointed out that: " … vegetable proteins and skimmed milk powder were technically substitutable in terms of their final use and that the effects of the EEC measures were to make skimmed milk powder competitive with these vegetable proteins”553. The EC maintain that, despite Canada’s unfounded assertions to the contrary, it follows from these two paragraphs that Article III:4 does not cover directly competitive or substitutable products.

3.445 The EC thus confirm the arguments they have already put forward, namely that, even if they can be considered to be competitive or substitutable products, substitutes for asbestos fibres and asbestos-containing products are not “like products” within the meaning of Article III:4 of the GATT 1994. Substitute products, by definition, are only substitutable for other products. In the present case, the substitute products are only substitutable for asbestos fibres and products containing them to a very limited extent. In fact, in view of the multiple uses of asbestos, there is no single natural or synthetic product that alone could replace asbestos generally in all the products or materials that contain it. Thus, there is no substitute for asbestos, only alternative solutions that rely on substitutes which vary depending on the intended application and are sometimes used in combination in order to obtain a material or product that performs an equivalent function.

3.446 The EC claim that the nature/quality/properties criterion is clearly important for assessing likeness within the meaning of Article III:4 of the GATT. This follows from the practice of previous panels, which have always used the physical characteristics for the purpose of determining “likeness” within the meaning of Article III:4, but this is not the case for the other criteria. Thus: (i) the Panel on Measures Affecting Alcoholic and Malt Beverages554 used neither tariff classification, nor end-use, nor consumers’ tastes and habits for determining the “likeness” of the products within the meaning of Article III:4; (ii) the Panel on EEC - Measures on Animal Feed Proteins555 used neither end-use nor consumers’ tastes and habits for determining the “likeness” of the products within the meaning of Article III:4; (iii) the Panel on United States - Gasoline556 did not use consumers’ tastes and
habits for the purpose of determining the “likeness” of the products within the meaning of Article III:4. The EC note, on the other hand, that all these Panels used the nature/quality/property criterion for examining likeness within the meaning of Article III:4 of the GATT, which underlines the importance of this criterion within the context of this provision. The EC add that, among these differences in the nature, properties and quality of the products, the health risk posed by the product must necessarily be taken into account. A dangerous product should be regarded as being different in nature and quality from a harmless or less dangerous product. The EC also emphasize that, even if the criterion of consumers’ tastes and habits might seem to have little relevance to the present case insofar as the products concerned are not consumer goods, it might nevertheless be useful to analyse the consumer’s perception of these products. There can be little doubt that informed users would not choose asbestos or asbestos-containing products after the competent international organizations had decided that asbestos was a proven carcinogen.

3.447 The EC assert that the criteria of likeness, in particular the nature/quality/properties criterion, mean that the substitute products are not “like” products with respect to asbestos and asbestos-containing products. Asbestos fibres are by definition “fibrous” products. It follows logically that, given this marked difference in physical characteristics, no non-fibrous substitute product could be considered a product “like” to asbestos fibres. As for “fibrous” substitutes, they cannot be considered “like” either within the meaning of Article III:4 because the morphology of asbestos fibres is different to that of “fibrous” substitutes. The fibres that must be taken into account in making a metrological assessment of a working environment have been defined by the WHO in accordance with the following dimensional parameters: (i) more than 5 microns in length; (ii) less than 3 microns in diameter; (iii) ratio of length to diameter greater than 3 microns [3:1]. Chrysotile fibres are 0.1 to 1 micron in diameter and separate lengthways into even finer crystalline fibrils (0.020 micron). The EC indicate that substitute fibres, on the other hand, have a different morphology. Thus, the polyvinyl alcohol and para-aramid fibres used to replace asbestos are 2 to 8 mm in length (i.e. 2,000 to 8,000 microns) and from 10 to 16 microns in diameter. Cellulose fibres, which are from 12 to 40 microns in diameter, can give rise to finer particles (fluff), which are said to irritate the respiratory pathways. These fibres are more than 10 microns in diameter, which physically prevents them from penetrating into the pulmonary alveoli. In addition, the EC stresses the analogy with the case United States - Measures Affecting Alcoholic and Malt Beverages. If a beer containing the same substance (alcohol) is not “like” to another beer simply because it contains a different quantity of alcohol, then a fortiori a product containing a different type of fibre cannot be considered to be “like” a product containing asbestos. The EC also mention the case of EEC - Measures on Animal Feed Proteins, in which the Panel noted that “the varying protein contents and the different vegetable, animal and synthetic origins of the protein products before the Panel” were sufficient to conclude that “these various protein products could not be considered as ‘like products’ within the meaning of Articles I and III”. Similarly, the different origin of the fibres present in the substitute products containing them prevents these products from being considered “like” to products containing asbestos.

3.448 The EC note that Canada itself points out, in relation to the quality of fibrous substitutes, that they are often “of lower quality in terms of their physical, chemical and mechanical resistance”. The EC also emphasize that asbestos fibres have certain characteristics that differentiate them from other fibres. Thus, asbestos fibres do not have the same physical characteristics as the substitute fibres, despite Canada’s assertions to the contrary. Furthermore, within the context of this nature/quality/properties criterion, the EC maintain that, being hazardous, asbestos and asbestos-containing products cannot be considered as “like” to other products. In fact, asbestos is a hazardous product classified as such by the competent international bodies (a category I carcinogen) whose use can put the lives of thousands at risk. Its hazardous nature has a marked impact on the consumer’s perception of the product. Thus, in 1989, the Supreme Court of Canada recognized that the risks associated with the use of asbestos were already a matter of common knowledge in 1973. In particular, it observed that:
“It seems clear from the journals, newspapers, magazines and manuals which are discussed in detail in the courts below that asbestos-related health risks had spread well beyond the boundaries of industry knowledge and were in wide public circulation. [...] Quite apart from the many studies and reports on asbestos-related health hazards [...] some of the articles in question were not published in obscure publications. These were prominent articles in the New York Times, the Wall Street Journal, the New Yorker magazine, the Washington Post and others. [...] Did all of this lend a ‘public character’ and ‘notoriety’, within the meaning of Article 2486 C.C., to the asbestos-related health risks that existed in 1970 and 1973? In my respectful opinion, it did.”

3.449 The EC contend that not only North American consumers, but also European consumers, are perfectly well aware of the carcinogenicity of asbestos. Consumers may therefore take a different view of asbestos and asbestos-containing products, on the one hand, and the substitute products, on the other. This difference in perception reinforces the lack of similarity between these products. In these circumstances, it must be concluded that asbestos and asbestos-containing products are not “like” the substitute products within the meaning of Article III:4, in particular because the nature/quality/properties of these products are different from those of asbestos and asbestos-containing products, as is the consumer’s perception of them, due to the health risks associated with the characteristics of asbestos. To conclude, the EC wish to draw attention to the paradox inherent in the observations made by Canada. Thus, Canada developed, at considerable length, arguments according to which amphibole fibres are very different to chrysotile fibres. The Canadian arguments related in particular to their physical characteristics. The EC consider that, resituated within the present context, Canada’s arguments concerning a hypothetical similarity between asbestos and asbestos-containing products, on the one hand, and the substitute products, on the other, are even more surprising.

3.450 Canada emphasizes that, in their second written submission, the EC stated that the tastes and habits of consumers are relevant in order to determine whether products are like products. However, in their first written submission, the EC do not identify the tastes and habits of consumers as a relevant criterion for the purposes of examining the similarity of asbestos products because they are not products in everyday use. In their second written submission, the EC consider that the hazardous nature of asbestos has an effect on the way in which consumers perceive the product. According to the EC, the fact that consumers, rightly or wrongly, perceive chrysotile pipes as different to PVA, cellulose or glass fibre pipes from the risk standpoint supports the hypothesis that chrysotile-cement and fibre-cement are not like products. Canada states that such a hypothesis does not stand up to examination. Consumers’ perception of the health risks of chrysotile and of substitute fibres is not an element that should be taken into account when determining the likeness of chrysotile fibres and substitute fibres. Moreover, it is not appropriate to consider the criterion of the tastes and habits of manufacturers of fibre-cement – the consumers whose tastes and habits count in this case – when determining the likeness of chrysotile fibres and PVA, cellulose or glass fibres. Canada agrees with the practice of the Panel in the three cases cited by the EC in paragraph 3.446 and asks the Panel not to take account of consumers’ tastes and habits in order to determine the likeness of the products in this case.

3.451 Canada claims that, contrary to the EC’s argument, the different origins of substitute fibres (artificial minerals, natural, organic, synthetic organic or natural organic minerals) do not prevent them being considered like products. The higher cost of substitute fibres in comparison with chrysotile fibres, and the fact they have a number of uses in addition to replacing chrysotile fibre, do not run contrary to the likeness of substitute fibres and chrysotile fibres within the meaning of Article III:4 of the GATT. The criterion of the nature, quality and properties of the product imply that chrysotile fibre and chrysotile-cement, on the one hand, and substitute fibres and fibre-cement, on the other, have quali-
ties in common. Canada notes that, in their reply to question 9 of Canada (see Annex II),
the EC recognize the relevance of the chemical composition of the fibres for determining
likeness and relate it to the criterion of the nature, quality and properties of the products.
Asbestos fibres have the same characteristics as substitute fibres. Even if the length, diam-
eter and width/diameter ratio have an effect on pathogenicity (one of the three “Ds”), in
Canada’s view this does not mean that fibres with different dimensions are necessarily not
like fibres. Canada notes that, in paragraph 3.447, the EC state that fibrous substitute prod-
ucts cannot be considered as like products within the meaning of Article III:4 because the
morphology of asbestos fibres is not the same as that of “fibrous” substitute products. The
EC therefore maintain that substitute fibres for asbestos used in fibro-cement are too large
to penetrate the lungs and cannot be taken into account when making a metrological as-
sessment of a work environment according to the dimensional parameters set by the WHO.
In Canada’s view, the EC’s conclusion is contradicted by the Notice concerning chrysotile
asbestos and possible substitute products. This Notice clearly states that the characteristics
of the fibres suggest that substitute fibres (PVA, p-aramids and cellulose) of the critical size
and shape reach human pulmonary alveoli. The dimensional parameters set by the
WHO do not constitute the criterion of the nature, quality and properties according to
which the likeness of fibrous products is determined. They help to identify fibres likely to
reach far into the lungs. That is all. A fibre is a fibre, irrespective of its dimensions. Canada
points out that the potential risks or harmlessness of chrysotile fibres is not an element to
be taken into account in order to determine whether chrysotile fibres can be considered as
“like” products to other fibres. Even if the toxicity of chrysotile is not the same as that of
amphiboles, the chrysotile fibre is a “like” product to amphibole fibres. Likewise, some
substitute fibres are like to chrysotile fibre because of their nature, quality and properties,
irrespective of any differences in their toxicity potential. There is no contradiction between
distinguishing two types of fibres at the scientific level and in relation to their pathogenic-
ity, on the one hand, and applying the criteria utilized in the WTO and the GATT to deter-
mine whether products are like products. An examination of likeness in connection with the
GATT and the WTO is different to an analysis of pathogenicity. The toxicity of a prod-
uct is not recognized as a criterion when examining likeness.

3.452 The European Communities contend that Article III:4 does not cover “directly
competitive or substitutable products”. In fact, whereas Article III:2, second sentence, read
in the light of the corresponding Note, provides for an examination of directly competitive
or substitutable products, Article III:4 mentions only the consideration of “like domestic
products”. This difference in the texts is not a matter of chance. If the authors of Article
III:4 had intended to include an analysis of directly competitive or substitutable products,
it would have had to appear in the text of Article III:4 or in an Interpretative Note, as was
done in Article III:2, second sentence. In the EC’s view, any other interpretation would
result in giving a meaning to a provision of the GATT which the signatories to the Agree-
ment had not intended and would, therefore, not be supported by the generally accepted
principles of treaty interpretation. The GATT case law relies essentially on the physical
characteristics of products, i.e. the nature, quality and properties criterion, when assessing
“likeness” within the meaning of Article III:4 of the GATT. In this case, the substitute
products for asbestos can be non-fibrous or fibrous, although the vast majority is non-
fibrous. The EC note that Canada accepts that non-fibrous substitute products are not
“like” products in this case. In view of the multiple uses of asbestos, there is no single
natural or synthetic product that alone could replace asbestos generally in all of its applica-
tions and for all the products or materials that contain it. Thus, there is no single substitute
for asbestos, only alternative solutions which rely on substitutes that vary depending on the
intended application and are sometimes used in combination in order to obtain a mate-
rial or product that performs an equivalent function. For example, cellulose has been used
for many years in a large number of applications, including as a partial substitute for
chrysotile. The EC conclude that this very large category of substitute products with nu-
merous potential uses, including as a substitute for asbestos, cannot be considered “like”
within the meaning of the case law developed under the GATT.

3.453 The EC point out that, among these differences in the nature, properties and qual-
ity of the products, the health risk posed by the product in question must necessarily be
taken into account. In fact, a dangerous product should be regarded as being different in nature and quality from a harmless or less dangerous product. The EC point out that, at the meeting held on 17 January 2000, the four scientific experts clearly stated that all the substitute products used to replace chrysotile asbestos are safer than chrysotile-containing products. Therefore, the fact that chrysotile is carcinogenic has a special impact on the consumer’s perception of the products containing it, compared to safer substitute products. The EC indicate that, contrary to Canada’s claim, they have not changed their position, but have responded to the developments in the written and oral debates before the Panel. Therefore, even if the criterion of consumers’ tastes and habits might seem to have little relevance to the present case, inasmuch as the products concerned are not to be consumed directly, it is nevertheless very useful to analyse the users’ perception of these products when determining likeness. In these circumstances, in the EC’s opinion, it must be concluded that asbestos and asbestos-containing products are not “like” the substitute products within the meaning of Article III:4 for three reasons: firstly, because the nature, quality and properties of these products are different to those of asbestos and asbestos-containing products; secondly, the consumer’s perception of them is also different because of the health risks associated with the characteristics of asbestos; thirdly, the tariff classification of these products is also different to that of asbestos.

(iii) Imported products … shall not be subject to less favourable treatment

3.454 Canada claims that, by prohibiting sale, offering for sale or purchase, the Decree alters the conditions of competition on the domestic market for substitute fibres and products containing substitute fibres of French origin, on the one hand, and chrysotile fibres and products containing chrysotile fibre from Canada, on the other.

3.455 Canada asserts that France has an important fibro-cement industry. The two major producers of fibro-cement products are Eternit and Saint Gobain. Eternit manufactures PVA fibro-cement products and Saint Gobain glass-cement products. Eternit’s products are manufactured at the former sites of four converted factories: Saint-Grégoire (Ille-et-Vilaine), Terssac (Tarn), Vitry-en-Charolais (Saône-et-Loire) and Thiant (Nord). Two production sites in France are used for the manufacture of glass-cement products: Descartes (Indre-et-Loire), where the factory has been converted, and Dunkerque (Pas-de-Calais), where two production lines have been introduced. PVA fibres, which are industrially synthesized, are of French origin. Glass fibres are also of French origin. Cellulose fibres are produced in France. The products marketed by Eternit are presented as a new generation of cement fibres without asbestos and are intended mainly for exterior siding and coverings. The glass-cement composites of the Saint Gobain group are used for cladding.

3.456 Canada claims that chrysotile fibre and chrysotile-cement products are subject to less favourable treatment than PVA, cellulose and glass fibres and fibro-cement products. The Decree banning asbestos is incompatible with the disciplines of Article III:4 of the GATT because its provisions subject chrysotile fibre and chrysotile-cement products to “treatment less favourable” than that accorded to like substitute fibres and fibro-cement products. According to the Panel’s report in the case United States – Section 337 of the Tariff Act of 1930:

“The words ‘treatment no less favourable’ in paragraph 4 call for effective equality of opportunities for imported products in respect of the application of laws, regulations and requirements affecting the internal sale, offering for sale, purchase, transportation, distribution or use of products.”

The French public authorities, in prohibiting the manufacture, processing, sale, export, release on the domestic market, possession for sale, offer and transfer on any basis of all varieties of asbestos fibres or of any product containing them, have eliminated the opportunities for competition by chrysotile fibre and chrysotile-cement products on the French market. Canada maintains that the incompatibility of the Decree banning asbestos with Article III:4 of the GATT is due to the French Government’s refusal to allow chrysotile fibre
and products containing it imported from Canada the opportunity to compete, a benefit accorded fibres and like products of national origin.

3.457 The European Communities maintain that the contested measure accords with the fundamental purpose of Article III, which is to prevent protectionism. Before looking in detail at the specific provisions of Article III, and especially Article III:4, the interpretation of that article given by the Appellate Body in its Report on Japan - Taxes on Alcoholic Beverages must be borne in mind:

“The broad and fundamental purpose of Article III is to avoid protectionism in the application of internal tax and regulatory measures. More specifically, the purpose of Article III is to ensure that internal measures “not be applied to imported or domestic products so as to afford protection to domestic production”.

Toward this end, Article III obliges Members of the WTO to provide equality of competitive conditions for imported products in relation to domestic products. (…) The broad purpose of Article III of avoiding protectionism must be remembered when considering the relationship between Article III and other provisions of the WTO Agreement.”564 [italics in the original]

3.458 The EC note that another Panel report (United States - Measures Affecting Alcoholic and Malt Beverages) indicated that:

“The purpose of Article III is thus not to prevent contracting parties from using their fiscal and regulatory powers for purposes other than to afford protection to domestic production. Specifically, the purpose of Article III is not to prevent contracting parties from differentiating between different product categories for policy purposes unrelated to the protection of domestic production.”565

In the EC’s view, the measure complained of is not discriminatory, neither de jure nor de facto, inasmuch as it guarantees effective equality of opportunity for domestic and imported products, in accordance with the letter of the requirement spelled out in the Reports cited above and in the Report on United States - Section 337 of the Tariff Act of 1930.566

3.459 The EC maintain that the Decree does not introduce any de jure discrimination. Firstly, the context and circumstances in which the Decree was adopted show that it is in no way intended to discriminate against imported products or to protect domestic products. Its sole purpose is to curb the spread of any risk of death or serious illness linked to exposure to asbestos, in particular among those subject to repeated or occasional – and often unwitting – exposure. Secondly, none of the Decree’s provisions makes a distinction in terms of treatment between French products and “like” imported products. For example: (i) both domestic asbestos fibres and imported asbestos fibres are prohibited; (ii) both domestic products containing asbestos fibres and imported products containing asbestos fibres are prohibited; (iii) both domestic asbestos fibres and imported asbestos fibres may be granted a temporary exemption on the same terms; (iv) both domestic products containing asbestos fibres and imported products containing asbestos fibres may be granted a temporary exemption on the same terms; and (v) both substitute domestic products and substitute imported products are permitted. It is thus clear to the EC that the Decree makes no distinction between imported products and domestic products, and neither its object nor its effect is to protect domestic production, so it is fully compatible with Article III.

3.460 The EC maintain that the Decree does not create any de facto discrimination. In its report on Japan - Taxes on Alcoholic Beverages, the Appellate Body recalled that “… Article III obliges Members of the WTO to provide equality of competitive conditions for imported products in relation to domestic products.”567 In this case, the effective equality of opportunity for domestic and imported products alike is not only reflected in law, but is also clearly evident from the facts. As the EC claimed in their factual arguments, most of the substitute products used in France are imported from various third countries. France has a negative trade balance in these substitute products. The EC note that, in Canada’s
view, the Decree favours the French fibre and substitute products industry. The fact is, however, that France chose to prohibit the use of asbestos on public health grounds. The changeover was a painful process, especially in human and financial terms, because the French measure prohibits exports of asbestos and asbestos-containing products, thus closing off external outlets for its domestic industries. Moreover, far from recommending the use of particular substitute products, the Decree leaves it to businesses to decide whether to replace asbestos by substitute fibre products or traditional products (plaster, ductile iron, etc.) of their choice. The Decree thus satisfies the “neutrality” requirement in terms of the choices made by enterprises, as set down in the Panel Report on United States - Measures Affecting Alcoholic and Malt Beverages:

"The Article III:4 requirement is one addressed to relative competitive opportunities created by the government in the market, not to the actual choices made by enterprises in that market. Producers located in the states in question have the opportunity to choose their preferred method of marketing." 

The EC conclude that the Decree does not produce de facto discrimination between domestic products and imported products.

3.461 Canada maintains that the origin of the fibres claimed by Canada has not been disputed for all the like products, with the exception of PVA fibre. Canada notes that the EC claim that PVA fibres are produced only in China and Japan. Canada therefore wonders how it is possible for France to export PVA fibres if it has no industry manufacturing or processing them. In 1998 alone, France exported F 41 million worth of PVA fibres. These fibres exported by France are undoubtedly of domestic French origin.

3.462 Canada contends that the Decree discriminates by subjecting chrysotile fibre and products containing it to less favourable treatment than that accorded to similar substitute fibres and fibro-cement products containing them. Canada rejects the EC’s arguments that the Decree does not constitute any form of de jure or de facto discrimination.

3.463 Canada maintains that the Decree, by prohibiting the manufacture, processing, sale, export, domestic marketing, possession for sale, offer or transfer on any basis of all varieties of asbestos fibres or of any product containing them, constitutes de jure discrimination in that it does not treat like products of national origin (substitute fibres or products containing them) in the same way. Not only does the ban apply only to asbestos fibre and products containing it, but, what is more, it is applicable only if there are like products to chrysotile fibre or to the products containing it. No exception to the ban will be permitted if there is a like product offering all the technical guarantees. The Decree therefore imposes a less favourable treatment in all cases where like products exist. What is more, the cases where an exception should be permitted, in the absence of a like product, are very rare. It is only on an “exceptional and temporary” basis that asbestos and asbestos-containing products may be granted effective equality of opportunity with like products of national origin. Canada notes that the EC claim that there is no de jure discrimination because asbestos fibres imported and of national origin are subject to the same treatment and because substitute products imported and of national origin are subject to the same treatment. In Canada’s view, the EC are simply not comparing the right products. Article III:4 invoked by Canada provides that imported products “shall be accorded treatment no less favourable than that accorded to like products of national origin”. Canada indicates that the EC fail to compare the treatment given to imported products (chrysotile fibre and fibro-cement containing it) with that reserved for like products (PVA, cellulose or glass fibre and fibro-cement containing them). Canada calls on the Panel to conclude that the text of the Decree treats chrysotile and chrysotile-cement products less favourably than like products, and that the Decree therefore constitutes de jure discrimination.

3.464 Canada also claims that the effective inequality of opportunity available to imported products and to like products of national origin is apparent not only in law in the text of the Decree, but is also reflected in the facts. A total ban on chrysotile fibres and fibro-cement products can only benefit the French substitute fibres manufacturing and
fibro-cement products industries. Canada points out that the EC, in their claim that the Decree does not favour the substitute products industry, cite a trade deficit in the case of PVA (the question of aramid fibres is not relevant to the analysis under Article III:4). In Canada’s view, a trade deficit does not necessarily indicate that an industry is doing better or worse. In reality, the French PVA fibres industry is in better shape than ever; its exports more than doubled between 1994 and 1998. Canada notes that the EC also claim that the national cellulose fibre industry does not benefit from the ban, pointing out that imports from Canada have increased. This claim does not stand up to analysis: it is not because France has imported a marginal additional quantity of Canadian cellulose fibres since the Decree that the French domestic industry has not benefited from the ban to the detriment of imported products. According to Canada, the EC misleadingly use a passage from United States – Measures Affecting Alcoholic and Malt Beverages to claim that the Decree is neutral in its application and does not impose a choice on consumers. On the contrary, the Decree does indeed impose a choice on the French consumer, who is now prevented from using chrysotile fibre or products containing it. Finally, the fact that the asbestos industry in France has or has not suffered is of no consequence in this case. It is a matter of comparing the effects of the Decree on Canadian asbestos interests with its effects on French interests in the substitute products industries.

3.465 The European Communities refer the Panel to the arguments they have already put forward concerning the absence of de jure and de facto discrimination in the Decree’s application (see paragraph 3.457-3.460 above).

3.466 The EC also point out that the Decree does not make any distinction between “like” national and imported products. It prohibits the use of all kinds of asbestos and asbestos-containing products in general, irrespective of origin. The substitute products used to replace asbestos are also treated in exactly the same way, irrespective of their origin. Therefore, the Decree in question does not discriminate in any way whatsoever, neither de jure nor de facto, between all types of asbestos of whatever origin and between all types of asbestos and substitute products for asbestos of whatever origin. Indeed, France imports very large quantities of a wide range of substitute products and treats them in exactly the same way it treats like substitute products of domestic origin, for the purpose of replacing asbestos. The EC also contend that the object and purpose of the Decree in question and the way it was elaborated confirm the view that the intention of the French authorities was not to protect domestic substitute products, but to protect human health from the risks of asbestos-related diseases. It follows that none of the conditions for the application of Article III:4 of the GATT is fulfilled in the present case. The EC maintain that, as Canada has failed to establish a violation of Article III:4 of the GATT, there is clearly no need to examine the applicability of Article XX(b) of the GATT in this case (see, for example, the Panel report on Section 337 in 1989).

(c) Article XI of the GATT

3.467 Canada points out that, in the Japan - Trade in Semi-Conductors case, the Panel noted that “... this wording was comprehensive: it applied to all measures instituted or maintained by a contracting party prohibiting or restricting the importation, exportation or sale for export of products other than measures that take the form of duties, taxes or other charges.” Canada maintains that the Decree is incompatible with Article XI:1 of the GATT, which applies to all measures instituted or maintained by a Member that prohibit or restrict the importation, exportation or sale for export of products, unless the measures are in the form of duties, taxes or other charges. Through the Decree, France is maintaining a ban or restriction on the importation of chrysotile and products containing it, other than by duties, taxes or other charges, contrary to its obligation under Article XI:1 of the GATT. Pursuant to Article 1, paragraphs 1 and 2, of the Decree, in order to protect workers and consumers “the importation ... of all varieties of asbestos fibres, whether or not incorporated into materials, products or devices” is prohibited. Article XI:1 of the GATT applies because one aspect of the Decree deals specifically with prohibiting importation. As a result of the total ban, producers of chrysotile or chrysotile-containing products, from Canada and elsewhere, cannot export their products to the French market. Simi-
larly, French companies cannot import chrysotile fibre or products containing it because, with some limited and temporary exceptions, imports are prohibited and liable to a fine.

3.468 Canada points out that the penal provisions in the Decree prevent French industry from seeking to import chrysotile and, as a result, this product will never reach France’s borders; similarly, a producer of raw chrysotile fibres will refrain from sending his product to France if he knows that its importation is prohibited. Since the Decree took effect on 1 January 1997, the chrysotile-cement industry has ceased operations. As of that date, the former chrysotile-cement industry was forced to convert to the use of chrysotile substitute products. Therefore, rather than speaking of the chrysotile-cement industry, the expression “fibro-cement industry” is now more appropriate. Canada claims that, in adopting the Decree, France instituted, and has since maintained, a prohibition or a restriction on the importation of chrysotile and chrysotile-containing products, originating in Canada and elsewhere, other than duties, taxes or charges, in violation of the provisions of Article XI of the GATT.

3.469 The European Communities claim that, as they indicated above, Article III:4 of the GATT is applicable and rules out application of Article XI.

3.470 Canada states that, in the event that the Panel recognizes the dual aspect of the measure, Canada refers it to the arguments put forward by Canada above. If, however, the Panel decides that the Decree is indivisible and cannot involve two specific aspects that can be examined separately with respect to Article III:4 and Article XI:1 of the GATT, Canada is of the opinion that it must be deemed a measure affecting imports and, consequently, be examined in the light of Article XI:1. Taken as a whole, because of its substance and its true character, the Decree affects imports. It establishes a quantitative restriction on imports of chrysotile asbestos fibres. Among all the types of measure that come within the scope of Article XI:1 of the GATT, the quantitative restriction instituted by the Decree – the ban - is the most severe and the most extreme of all.

3.471 Canada claims that the Decree is a measure restricting or prohibiting imports within the meaning of Article XI of the GATT for a number of reasons. Firstly, the text of the Decree expressly stipulates that the “importation” of asbestos fibres and of asbestos-containing products is banned. According to the terms of Article 1, paragraphs 1 and 2, of the Decree, on the grounds of protecting workers and consumers “the importation ... of all varieties of asbestos fibres, whether or not these substances are incorporated into materials, products or devices” is subject to a ban. Secondly, as France does not produce or extract asbestos fibres on its territory, the ban on manufacturing, processing, selling and domestic marketing is equivalent, in practice, to an import ban on chrysotile asbestos fibres. With respect to chrysotile fibre, the ban on manufacturing, processing, sale and offering for sale are complementary to the import ban. These bans are a means by which France is pursuing a more basic objective, namely a ban on introducing chrysotile asbestos fibres into its territory. According to Canada, the EC acknowledge that the bans on manufacturing, processing, selling and marketing are complementary to the import ban. Indeed, they point out that deleting the word “importation” from the Decree would not change anything in the asbestos situation in France, which is equivalent to saying that the basic goals would in any case be attained: chrysotile asbestos fibres would not be introduced into France. Thirdly, the Decree is a measure prohibiting or restricting imports within the meaning of Article XI by virtue of its goal. Canada notes that the EC claim that the goal indeed is to halt the spread of asbestos as far “upstream” in the production and distribution process as possible. However, on French territory, the importation of chrysotile fibre represents the most “upstream” link in the commercial or industrial chain. The French Government has never concealed the fact that the goal of the ban was to prohibit asbestos on its territory in the future. With respect to asbestos fibres, this stated goal relates more to importation than to use because France does not produce asbestos fibres.

3.472 Canada points out that, even if the Decree were considered an internal regulatory measure, this would not necessarily mean that Article XI:1 would not apply. Article XI:1 can apply to an internal regulation that has the effect of restricting or prohibiting imports.
Excluding any internal regulation from the scope of Article XI:1, simply because it is an internal regulation, would deprive the words “other measures” in the conclusion of Article XI:1 of any useful effect. Such an interpretation would also be contrary to the position adopted by the Panel in Canada - Import, Distribution and Sale of Certain Alcoholic Drinks by Provincial Marketing Agencies. In that case, the Panel accepted the argument of the EC that the requirements of provincial liquor boards with respect to listing and delisting and points of sale, which discriminated against imported alcoholic beverages, were inconsistent with Article XI:1 of the GATT. However, the requirements in question were part of an internal regulatory framework as they concerned the distribution and marketing of alcoholic beverages. This did not prevent the Panel from concluding that these requirements were contrary to the provisions of Article XI:1 of the GATT. The Panel considered “that systematic discriminatory practices of the kind referred to should be considered as restrictions made effective through ‘other measures’ contrary to the provisions of Article XI:1.”

The Panel also considered “that it was not necessary to decide in this particular case whether the practices complained of were contrary to Article III:4 because it had already found that they were inconsistent with Article XI.” Canada claims that, with respect to chrysotile asbestos fibres, the heart of the Decree is an import ban. For the purposes of this debate, which deals with Canadian imports of chrysotile asbestos fibres and their applications, the Decree, even when considered as a whole, must be deemed an import restriction or import ban within the meaning of Article XI:1 of the GATT, and examined in light of this provision.

3.473 The European Communities maintain that the Decree must be examined as an internal measure to which Article III:4 of the GATT alone is applicable, thus ruling out the cumulative application of Article XI in this case.

3.474 The European Communities contend that, even if the Panel were to hold that the Decree is incompatible with Article III:4 of the GATT, it would nevertheless have to find that the measure falls under the exception provided in Article XX(b) of the GATT. Article XX allows Members an exemption from their obligations under the General Agreement subject to certain conditions. According to the precedent set by the Appellate Body in its Report on United States - Shrimps, the analysis of the measure in the light of Article XX must begin with the paragraph of Article XX invoked by the defending party, in this case (b), and continue with consideration of the introductory phrase (“chapeau”) of that Article.

Following this line of approach, the EC will claim that: (i) the Decree is necessary to attain the underlying policy goal, in other words to protect human health and life (test of “necessary” in paragraph XX(b); (ii) the Decree is not applied in such a manner as to constitute a means of arbitrary or unjustifiable discrimination between countries where the same conditions prevail or a disguised restriction on international trade (“chapeau” of Article XX).

3.475 Canada claims that it is clear from the practice of Panels that, in relation to Article XX(b), it is necessary to show that: (i) the policy which prompted the measure for which the provision is invoked falls within the category of policies intended notably to protect human life or health; (ii) the inconsistent measure for which the exemption is invoked is necessary to fulfil the objective of the aforesaid policy; and (iii) the measure is applied in compliance with the requirements of the introductory paragraph or “chapeau” of Article XX. The Appellate Body also noted that “...the ultimate availability of the exception is subject to the compliance by the invoking Member with the requirements of the chapeau.” Article XX permits a “limited and conditional exception from the obligations of the substantive provisions of the GATT” which must, according to the precedents, be interpreted narrowly. Again according to the Appellate Body, “...the measures falling within the particular exceptions must be applied reasonably, with due regard both to the legal duties of the party claiming the exception and the legal rights of the other parties concerned.” Canada notes that the EC assert that they have the right to
establish the level of protection they desire. In any event, this must be done in compliance with their obligations. The Appellate Body has condemned the abuse of rights under Article XX of the GATT. Sir Leon Brittan, for his part, states that the WTO Agreements do not allow a country to invoke zero risk:

“Using the precautionary principle to justify action aimed at reducing risks to zero would clearly be excessive. We should therefore distinguish the precautionary principle from a zero risk approach. To try and adopt the latter across the board could bring us to a scientific standstill since there are risks involved in any new venture.”

3.476 Canada claims that it is up to the EC to demonstrate that the Decree is an exception to Article XX(b). According to Canada, the EC did not discharge the burden of proof that lies with them in the arguments they put before the Panel.

(ii) Policy to protect human health

3.477 The European Communities maintain that, as explained in their factual arguments, because of their unique characteristics asbestos fibres and products containing them are a proven hazard for human health. The risks linked to the use of these fibres and products are recognized, both by the competent international organizations and by scientific studies, in particular the INSEERM study, which was the basis of the Decree. The Decree seeks to halt the spread of this risk and thereby reduce the number of deaths among the French population. The measure taken is the only possible one that enables the spread of the risks due to asbestos exposure to be halted effectively. It therefore falls under the heading of measures for the purposes described in Article XX(b).

(iii) “… necessary to protect human health and life …”

3.478 The European Communities contend that the review in the light of Article XX cannot be allowed to undermine the health protection goal set by the Member concerned. Its sole purpose must be to assess whether the trade measure adopted is actually “necessary” to attain that goal. The Panel report on United States - Gasoline introduced the GATT 1947 rule on the test of necessity into the GATT 1994. The Panel noted specifically:

“… the term ‘necessary’ had been interpreted in the context of Article XX(d) by the panel in the Section 337 case which had stated that: a contracting party cannot justify a measure inconsistent with another GATT provision as ‘necessary’ in terms of Article XX(d) if an alternative measure which it could reasonably be expected to employ and which is not inconsistent with other GATT provisions is available to it. By the same token, in cases where a measure consistent with other GATT provisions is not reasonably available, a contracting party is bound to use, among the measures reasonably available to it, that which entails the least degree of inconsistency with other GATT provisions.”

3.479 The EC note that Panel report on Thailand - Cigarettes followed the same line of reasoning when reviewing a measure in the context of Article XX(b). This Panel saw no reason not to follow the same interpretation of “necessary” for Article XX(b) as for Article XX(d), stating that:

“… the import restrictions imposed by Thailand could be considered to be ‘necessary’ in terms of Article XX(b) only if there were no alternative measure consistent with the General Agreement, or less inconsistent with it, which Thailand could reasonably be expected to employ to achieve its health policy objectives.”

It is this line of reasoning that should be followed in the present case in considering whether the French Decree is “necessary” in order to attain the goal of limiting the number of deaths caused by asbestos. The EC will claim that: (i) the ban is justified by the existence of risks
to the health of the population; and (ii) the ban is the only measure that enables the objec-
tive set by the French authorities (halting the spread of the risk) to be attained.

3.480 The EC maintain that the test of necessity concerns the trade measure and not
the level of protection set by the Member. They recall that the purpose of Article XX(b) is
not to call into question the level of protection that a Member considers appropriate for its
population. The fact that a Member is free to decide what level of protection it considers
appropriate in order to safeguard the health of its population was firmly spelled out in the
Panel Report on United States - Restrictions on Imports of Tuna

591 which stated that:

“The Panel further noted that Article XX(b) allows each contracting party to set
its human [...] health standards. The conditions set out in Article XX(b) which
limit resort to this exception [...] refer to the trade measure requiring justification
under Article XX(b), not however to the life or health standard chosen by the
contracting party. The Panel recalled the finding of a previous panel that this
paragraph of Article XX was intended to allow contracting parties to impose
trade restrictive measures inconsistent with the General Agreement to pursue
overriding public policy goals to the extent that such inconsistencies were una-
voidable.”

592

3.481 The EC point out that this principle was first established under the GATT 1994
in United States - Gasoline, when the Panel stated that:

“... it was not the necessity of the policy goal that was to be examined, but whether
or not it was necessary that imported gasoline be effectively prevented from
benefiting from as favourable sales conditions as were afforded by an individual
baseline tied to the producer of a product. It was the task of the Panel to address
whether these inconsistent measures were necessary to achieve the policy goal
under Article XX(b). It was therefore not the task of the Panel to examine the
necessity of the environmental objectives of the Gasoline Rule...”

3.482 The EC emphasize that there can be no questioning the right of the French authori-
ties to decide the level of protection they wish to offer their population. According
to the EC, this means that in this case the French authorities are free to choose to
halt the spread of the risks due to exposure to asbestos, in particular for people
who are occasionally and often unknowingly exposed to asbestos. The sole pur-
pose of Article XX(b) is to consider whether the Decree is “necessary” as a regula-
tory option. The EC state that the so-called “safe” use advocated by Canada does
not allow the objective of protecting human health, set by France, to be attained.
Canada is thus setting trading interests above the legitimate goal of human health
protection. The Panel Report on Thailand - Cigarettes states that: “this provision
[Article XX(b)] clearly allowed contracting parties to give priority to human health
over trade liberalization”.

3.483 The EC claim that the Decree constitutes the only adequate measure for the in-
tended purpose. As was indicated in their factual arguments, the EC consider that the
Decree pursues the realistic and “reasonable” goal of halting the spread of the risk due to
any kind of exposure to asbestos, especially the risk due to occasional and often unwitting
exposure. The EC maintain that the Decree is justified by the existence of risks to human
health. Canada wrongly argues that asbestos and asbestos-containing products pose “no
detectable risk” to health. In the EC’s view, this claim relies on the contention that the risk
due to levels of exposure in the atmosphere - on which Canada bases its entire submission
- is comparable to the risks from the occasional but very high levels of exposure that
may be faced by a substantial proportion of a country’s population. Canada seems to
suggest that the dangers of inhaling asbestos only concern workers engaged in the extrac-
tion and processing of asbestos, in other words “primary users”, whose number in Quebec
is around 1,700. However, far from affecting only those involved in asbestos extraction
and processing, the danger of inhaling asbestos at levels above the minimum threshold
(0.1 fibre/ml) primarily concerns users of asbestos or asbestos-containing products, in other
words “secondary users”, whether occupational (workers in the textile, building, automobile industries, for example), para-occupational (servicing, maintenance), or domestic (such as do-it-yourself) users. In France, there are millions of such secondary users. Several hundred thousand daily users and several million occasional users thus come into contact with a product which, in 1977, was classed by the WHO as a proven carcinogen for humans (category I) and for which there is, according to the 1998 WHO report, no threshold of harmlessness. The EC note that Canada does not fully discuss the occupational risks and completely ignores the para-occupational and domestic risks, although these account for the vast majority of those who suffer exposure.

3.484 The EC maintain that, even in production and processing, which in principle are easier to monitor, there are limits to the so-called “safe” use of asbestos. The “controlled” use of asbestos does not halt the spread of the risks. The 1996 study by the British HSE found a substantially higher than normal incidence of deaths due to mesothelioma among workers entering the asbestos production and processing industries after 1969, in other words even after the United Kingdom had introduced “safe” use.

3.485 The EC also contend that not only is so-called “safe” use unable to halt the spread of risks from exposure to asbestos in production and processing – where the number of people involved is fairly limited and, in principle, easy to manage and control – but it is completely ineffective in cases of occasional exposure to asbestos. Indeed, the principle of “safe” use cannot be applied to the risks affecting a wide range of jobs involving an enormous variety of situations, especially in servicing and maintenance. These workers may only be exposed to asbestos occasionally, but they are subject to exposure peaks that sometimes far exceed the currently accepted dust thresholds. A worker using a grinder outside on corrugated asbestos-cement roofing is subject to a peak exposure of 41 fibres/ml – 410 times higher than the threshold. The EC note that a 1992 study by the Quebec CSST also shows that the risk of mesothelioma has been steadily rising in Canada since 1967, mainly among servicing and maintenance workers. Not once does Canada quote this study.

3.486 The EC point out that the same finding is all the more true for the general population at risk, in other words those who are exposed to inhalation of asbestos on a non-occupational basis. “do-it-yourself enthusiasts” are a typical example of a large sector of the public subject to exposure who are often unaware that this occasional or regular activity may expose them to a potentially fatal risk. According to the EC, Canada conveys the impression that the INSERM report did not appreciate the risks described above. Yet the INSERM report shows that it came to be realized during the 1980s and 1990s that the highest risk is no longer to workers in the asbestos industry, namely production and processing. The report clearly states that the risk occurs mainly among those who work with materials containing asbestos. With regard to para-occupational and domestic exposure, the INSERM report states:

“These are studies which concern cases of mesothelioma in subjects with no known occupational exposure. Several case-control and cohort studies have shown the existence of cases of mesothelioma (pleural and/or peritoneal) attributed to exposure where the source was usually either soiled work clothing brought home by a person subject to occupational exposure or DIY activities. The levels of exposure to asbestos evaluated in these circumstances can be high, comparable to certain occupational exposures. The existence of an increased risk of mesothelioma among those exposed in para-occupational and domestic circumstances seems clearly established.” 593

3.487 The EC indicate that the INSERM report also states:

“As far as para-occupational exposure is concerned - especially in connection with DIY activities - it is legitimate to consider the exposure peaks as identical to those found in industrial operations of the same nature. The main differences in the level of exposure, in terms of the dose inhaled, are in the length of exposure, as the DIY enthusiast does not generally perform these operations as frequently as the professional.” 595
3.488 The EC maintain that so-called “safe” use will not halt the spread of risk. The EC note that, in Canada’s view, “safe” or “controlled” use of asbestos is possible and that, consequently, the prohibition is not “necessary” within the meaning of Article XX(b) of the GATT. Canada also cites the “negligible emissivity of chrysotile-cement products”. In support of this statement, Canada argues that the “modern” uses of chrysotile fibre mean that the fibre is bound in a matrix and cannot be released into the environment. The EC point out that what Canada describes as an innovation is nothing of the kind. For 40 years, the manufacture of asbestos-cement has involved “encapsulating” asbestos in cement (10 per cent asbestos fibres in 90 per cent cement). The EC indicate that, by referring to what it calls “modern” usage or even “modern” products, Canada is misleading the Panel by seeking to promote a “clean” image of chrysotile asbestos, although the name, while less ominous-sounding than amphibole asbestos, cannot hide the fact that it is still classed by WHO in ‘category I’ along with other products that are proven carcinogens for humans. The Panel should know that such “encapsulation” cannot be guaranteed to make asbestos-cement, for example, harmless. In fact, once the use of asbestos-cement is permitted, its use can no longer be controlled. Inevitably asbestos-cement, whether in an occupational, para-occupational or domestic context, will be worked in various ways – for instance by cutting, sanding, crushing or sawing. These operations will release large numbers of carcinogenic fibres in the form of dust. Consequently, the EC assert that the “modern” use of chrysotile asbestos to which Canada likes to refer is illusory and merely serves to mask the serious risks involved in the use of asbestos-cement.

3.489 The EC maintain that “safe” use is incapable of halting the spread of the risk due to occasional and often unwitting exposure to asbestos, because instituting the kind of tight constraints that are feasible for a small and “targeted” population - primary users - is wholly unrealistic for the general population. How could such use be effectively ensured for the hundreds of thousands of people exposed every day in sectors where there is little control in terms of health, such as the building industry, where at least 25 per cent of mesothelioma cases in France occur? Not to mention the millions of do-it-yourself enthusiasts subject to occasional and often unwitting inhalation of asbestos when performing quite ordinary operations such as cutting, for instance. The EC note that, as regards practical means of ensuring the effectiveness of “safe” use, Canada stated during consultations that there was no risk of asbestos fibres being released through cutting or sawing asbestos-containing products as they were supplied “pre-cut”. Canada also refers to an ISO standard issued more than 15 years ago, in 1984. The EC emphasize that, although the standard may have represented a major step forward at the time as compared with the earlier arrangements, it is not sufficient to guarantee an adequate level of protection, given the health objective of an exposure threshold of 0.1 fibre/ml that is recognized by many countries. The EC note that Canada states that “the cutting of roofing sheets or tiles is not a source of emission where the simple techniques laid down in ISO-7337 are followed”. These “simple” techniques are: “the use of chains to break pipes by means of pressure, the use of low-speed saws or saws equipped with dust extractors, and the wetting of materials before any operation”.

3.490 The EC point out that, in the real world, not all the pieces are pre-cut. And in the real world the thousands of people subject to occasional and unwitting inhalation of asbestos do not put on hermetically sealed protective suits when performing quite everyday tasks. Moreover, the use of a hand saw in accordance with the ISO standard leaves the worker exposed to a level 30 times higher than the permitted limit of 0.1 f/ml. According to the EC, Canada offers no answer to the problem of how to control operations performed further downstream in widely differing circumstances by many different people who are often unaware that they are being exposed to asbestos. The EC note that, once asbestos is on the market, there is no reasonable way of controlling its use and, in particular, of controlling the everyday operations (cutting, sawing, etc.) that many people are likely to perform on asbestos-containing products. In fact, there is clearly no way of ensuring that “safe” use as advocated by Canada can be effectively implemented. According to the EC, recent texts confirm that “safe” use is not feasible and this emerges very clearly from the 1998 WHO report, not cited by Canada, which states:
“Some asbestos containing products pose particular concern and chrysotile use in those circumstances is not recommended. These uses include friable products with high exposure potential. Construction materials are of particular concern for several reasons. The construction industry work force is large and measures to control asbestos are difficult to institute. In-place building materials may also pose risk control to those doing alterations, maintenance and demolition. Minerals in place have the potential to deteriorate and create exposures.”

The EC add that, as the “safe” use of asbestos and asbestos-containing products is unable to halt the spread of risk and anyone affected by current exposure is quite likely to seek redress in the courts at some later stage. However, such claims for damages are notoriously hard to prove and run up against many obstacles, especially on the legal side. There is, then, this additional social justification for recognizing that the ban on asbestos is the only measure truly capable of halting the spread of the risk due to exposure to asbestos. That is the path France has chosen to follow. For the foregoing reasons, it is obvious that “safe” use cannot be implemented effectively and, in any event, significant risks remain for all categories of user.

3.491 The EC note that Canada gives the impression that France has prohibited asbestos and recommended the “indiscriminate use” of substitute products, reinforcing Canada’s view that the measure is not necessary. This claim ignores the purpose of the French rules. Far from recommending the use of specific substitute products, the Decree leaves it to businesses to replace asbestos by whichever products or fibres they choose. The public authorities play no part in the actual choices made by enterprises on the market. In practice, firms will carry out technical tests with a view to replacing asbestos products by substitute products. However, if the tests are inconclusive and if firms can show that there are no safer substitute products, they can apply for an exemption in order to continue using asbestos. Such exemptions are provided for in the Decree and are granted only after careful scientific evaluation. Under the terms of Article 2 of the Decree, the use of asbestos is still permitted in exceptional cases and on a temporary basis, if there is no substitute which is capable of performing an equivalent function and which: (i) on the one hand, in the present state of scientific knowledge, poses a lesser occupational health risk to workers handling those materials, products or devices; and (ii) on the other, provides all technical guarantees of safety corresponding to the use thereof. The EC point out that the replacement of asbestos fibres by substitute fibres is therefore reasonable and justified. In particular, it is grounded in the 1996 and 1998 WHO reports. To conclude on this issue, the EC reject Canada’s claim that the only rationale for the excessive impact of the ban has to be the political will of the French Government to respond in spectacular fashion to the pressure it faced from public opinion. The EC maintain that their arguments set out above show, on the contrary, that the ban was the only appropriate solution capable of curbing the number of deaths due to exposure to asbestos. The foregoing shows that the Decree serves to attain the French policy goal regarding asbestos and is compatible with the test of necessity under Article XX(b).

3.492 Canada asserts that, in order to determine whether a measure falls under Article XX(b), it must first be examined whether the ban imposed by the Decree is “necessary” to protect human life or health. The only ban that the Decree adds, in practice, is the ban on chrysotile asbestos in non-friable products. The only exposures that the Decree could affect are those, if any, to chrysotile encapsulated in high-density products. A measure will be deemed necessary “… only if there were no alternative measure consistent with the General Agreement, or less inconsistent with it, which [the party] could reasonably be expected to employ to achieve its health policy objectives”. The measure that one seeks to justify by invoking Article XX(b) must therefore involve the minimum restriction on international trade. Canada notes that the EC, taking the view that France’s choice cannot be called into question, maintain that the French authorities have the right “to decide the level of protection they wish to offer their population … this means that in this case the French authorities are free to choose to halt the spread of the risks due to exposure to asbestos”. However, in order to determine whether there is an equally effective alternative that is less restrictive for international trade and that is capable of protecting human life or
health just as effectively, Canada considers that the health risk can and must be examined, regardless of what the EC say. The failure to consider whether in fact there is a risk would enable any country to cite a risk - real or not - in support of any prohibition measure.

3.493 Canada is of the opinion that the current uses of chrysotile in high-density non-friable products do not constitute a detectable risk to human health. The risks, which existed in the past and which still persist today in certain cases (amphiboles and friable materials in place), are associated with past uses of asbestos, very often amphiboles, in friable materials. Today, high-density non-friable chrysotile products do not pose a detectable risk. Canada notes that the EC identify the risk as existing both at the levels of extraction and processing and of secondary uses (textiles, construction, maintenance and custodial work, mechanics and do-it-yourself enthusiasts). According to Canada, the EC are clearly trying to mislead the Panel by invoking the risks of the asbestos extraction and processing industry, even though they have already recognized that controlled use is effective here in eliminating the risk. DG III affirmed in April 1997: “the principle of controlled use can be accepted in the asbestos industry”.\footnote{600} INSERM is of the same opinion: “Given the occupational origin of exposures to asbestos, we are beginning to observe, in a few countries which instituted strict measures to protect workers at an early date, a stagnation in the trends of the incidence of mesothelioma.”\footnote{601}

3.494 Canada asserts that the EC also show bad faith in citing the risks of building maintenance workers (electricians, plumbers, sheet metal workers and boiler-makers, etc.) and mechanics. The EC do not explain that these exposures are essentially to friable materials very often containing amphiboles of high pathogenic potential. Canada maintains that the risk associated with current uses of chrysotile, if any, is not detectable. According to the data of the United States Occupational Safety & Health Administration (OSHA), the institution of control measures lowers the average exposures of workers handling asbestos-cement pipes to 0.00253 f/ml and of workers handling asbestos-cement sheets to 0.00727 f/ml. The average exposure of mechanics handling friction products, is 0.00294 f/ml.\footnote{602} Canada indicates that, despite what the EC appear to believe, these hundreds of thousands of professionals have no need to don “space suits” every morning. Canada asserts that controlled use is sufficient. The use of pre-machined parts and fittings is not a far-fetched or utopian solution. According to OSHA: “pre-cut, pre-tapped pipe has received tremendous marketplace acceptance and represents a large majority of sales. [...] This is significant because the use of pre-cut, pre-tapped pipe may reduce or eliminate some types of field fabrication activities.”\footnote{603} As a result of pre-fabrication, pre-machining, the use of fittings and compliance with work standards, workers are not exposed to rates of 3 f/ml.

3.495 Canada claims that the EC is wrong in stating that Canada is comparing the exposures associated with the use of chrysotile to those of ambient air. However, Canada, far from using ambient data, bases its position on data for occupational exposures under controlled use conditions to assert that there is no detectable health risk. As for do-it-yourself enthusiasts, Canada notes that the EC also show bad faith in failing to mention that these exposures are essentially attributable to friable materials containing amphiboles. Furthermore, they fail to mention the conclusions of the Académie nationale de médecine according to which “… no disease due to asbestos has been formally proven in France outside an occupational type of exposure”.\footnote{604} Canada considers that few do-it-yourself enthusiasts, certainly not millions, work with high-density chrysotile products. Inasmuch as the only risk associated with asbestos is that of the past use of amphiboles and the use of friable materials, this risk cannot be eliminated by the Decree. The Decree, which prohibits the contemporary uses of chrysotile, is therefore not necessary - and even less useful - to protect human life or health from the risks associated with past uses of asbestos. Canada therefore asserts that the ban is not “necessary” to protect human life or health because high-density chrysotile products do not pose any detectable risk.

3.496 Canada contends that, in the event that, despite the scientific evidence submitted by Canada, Brazil and Zimbabwe, the Panel finds that contemporary uses of chrysotile are a hazard for human health, an examination of less trade-restrictive alternatives is nec-
In its arguments on controlled use, Canada has underlined the “possibility” and the effectiveness of controlled use. Controlled use indisputably constitutes an alternative to a total ban that is significantly less restrictive for international trade and eliminates the risk just as effectively, if any risk remains today. It follows from this analysis that the Decree banning current uses of chrysotile cannot fall under Article XX(b) because it is not necessary to protect human life or health.

3.497 Canada rejects the EC’s claim that Canada is placing its trade interests ahead of the legitimate objective of protecting human health. Canada is rather of the opinion that the EC are misusing the objective of protecting human health to justify a measure that does not fall under the scope of Article XX(b). The Decree does not fall under paragraph (b) of Article XX and is not consistent with the introductory paragraph of Article XX. The EC therefore cannot take advantage of the exception provided in Article XX(b) to justify a violation of Article XI:1 and Article III:4 of the GATT. Lastly, Canada refers the Panel to the arguments on the concept of necessity in Article 2.2 of the TBT Agreement because, as it emphasized in its reply to question 33 of the Panel (see Annex II), Canada contends that the criterion of necessity in Article 2.2 of the TBT Agreement is in many respects similar to that in Article XX(b) of the GATT.

3.498 The European Communities refer to the arguments put forward in connection with Article 2.2 of the TBT Agreement (see paragraphs 3.353-3.356 above), which also apply in the context of Article XX(b) of the GATT.

(iv) Preamble to Article XX

3.499 The European Communities recall that the Appellate Body stated that the “chapeau” of Article XX applies to: “… the manner in which that measure is applied” and that “If those exceptions [the exceptions under Article XX] are not to be abused or misused, … the particular exceptions must be applied reasonably…” The Appellate Body also noted that:

“The chapeau of Article XX is, in fact, but one expression of the principle of good faith. [… ] One application of this general principle, the application widely known as the doctrine of abus de droit, prohibits the abusive exercise of a state’s rights and enjoins that whenever the assertion of a right ‘impinges on the field covered by [a] treaty obligation, it must be exercised bona fide, that is to say, reasonably.”

Finally, the Appellate Body suggested that Panels should endeavour “to inquire into how the measure at stake was being applied in such a manner as to constitute abuse or misuse of a given kind of exception” [italics in the original] The EC maintain that in the present case there is nothing to support the contention that France acted “in bad faith” or in an “unreasonable”, “improper” or “abusive” manner in exercising its right under Article XX(b) of the GATT.

3.500 The EC maintain that the Decree is “reasonable”. They note that, as has been recognized by the WHO since 1977, asbestos is a product that is a proven carcinogen for humans. It has caused thousands of deaths. So-called “safe” uses of the product cannot prevent deaths from mesothelioma among “primary users” (extraction and processing industries) and cannot feasibly be implemented for all para-occupational and domestic uses, which involve millions of people. Depending on their age, between 18 and 25 per cent of the male population in France have been exposed to asbestos at least once in the course of their working lives. Building industry trades account for a quarter of mesothelioma cases. In view of these figures, France considered that the only measure capable of halting the spread of the risk was the outright ban on the use of asbestos on French territory. The EC contend that, despite Canada’s claims, France did not act on impulse. The French decision is based on a reliable scientific report that involved a critical and careful review of the most recent world scientific literature, proving the risks entailed by the use of asbestos, in particular for those subject to exposure in a para-occupational or domestic context. The report, like the Decree, was adopted after “mature reflection”.

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3.501 The EC claim that the Decree is not applied as a means of imposing arbitrary or unjustifiable discrimination between countries where the same conditions prevail. The prohibition covers products originating in any country (whether domestic or foreign), not only in Canada. The Panel report on United States - Imports of Certain Automotive Spring Assemblies, which dealt with a similar case, stated the following:

"The Panel noted that the exclusion order was directed against imports of certain automotive spring assemblies produced in violation of a valid United States patent from all foreign sources, and not just from Canada. It found, therefore, that the exclusion order was ‘not applied in a manner which would constitute a means of arbitrary or unjustifiable discrimination against countries where the same conditions prevail’."^608

3.502 The Appellate Body Report on United States - Shrimps elaborated on this idea, noting that:

"In order for a measure to be applied in a manner which would constitute ‘arbitrary or unjustifiable discrimination between countries where the same conditions prevail’, three elements must exist. First, the application of the measure must result in discrimination. As we stated in United States – Gasoline, the nature and quality of this discrimination is different from the discrimination in the treatment of products which was already found to be inconsistent with one of the substantive obligations of GATT 1994, such as Articles I, III or XI. Second, the discrimination must be arbitrary or unjustifiable in character. We will examine this element of arbitrariness or unjustifiability in detail below. Third, this discrimination must occur between countries where the same conditions prevail. In United States – Gasoline, we accepted the assumption of the participants in that appeal that such discrimination could occur not only between different exporting Members, but also between exporting Members and the importing Member concerned."^609 [italics in the original]

3.503 The EC point out that, in this case, the application of the French measure does not involve any discrimination^610 between countries, including France, where the same conditions prevail. All countries that export asbestos or asbestos-containing products, including France (which used to have its own asbestos industry), are covered by the prohibition or by the exemptions, with no difference in treatment. As this does not meet the definition of discrimination laid down by the Appellate Body, the EC consider that the Decree is not being applied in such a manner as to constitute a means of arbitrary or unjustifiable discrimination between countries where the same conditions prevail.

3.504 Lastly, the EC maintain that the Decree does not constitute a disguised restriction on international trade. On this question, the Appellate Body noted in its report on United States - Gasoline that:

"...the kinds of considerations pertinent in deciding whether the application of a particular measure amounts to ‘arbitrary or unjustifiable discrimination’ may also be taken into account in determining the presence of a ‘disguised restriction’ on international trade. The fundamental theme is to be found in the purpose and object of avoiding abuse or illegitimate use of the exceptions to substantive rules available in Article XX. [...] The resulting discrimination must have been foreseen, and was not merely inadvertent or unavoidable. In the light of the foregoing, our conclusion is that the baseline establishment rules in the Gasoline Rule, in their application, constitute ‘unjustifiable discrimination’ and a ‘disguised restriction on international trade’."^611

3.505 The EC have already shown that the Decree is not applied in a manner which would constitute a means of arbitrary or unjustifiable discrimination between countries where the same conditions prevail. This is proof enough that the Decree is not being applied in a manner which would constitute a “disguised restriction on international trade".
The EC emphasize in this respect that: (i) the restriction applies to products of any origin (including domestic origin); (ii) the restriction is justified on public health grounds; (iii) the restriction was announced and published; (iv) many other Members of the WTO also apply restrictions to these products; and (v) the restriction takes international standards as a base. The EC therefore hold that the Decree cannot be considered as being applied in such a way as to constitute a “disguised restriction on international trade”. Any other approach would imply that all international legislation on asbestos and asbestos-containing products (such legislation is, in practice, always restrictive) amounts to “a disguised restriction on international trade”. The EC therefore maintain that they did not “abuse” or make “unreasonable” use of their right under Article XX(b) of the GATT.

3.506 Canada claims that, in the event that the Panel finds that the Decree is nonetheless covered by paragraph (b) of Article XX, the EC must justify it under the introductory clause of Article XX. However, “… it does not follow from the fact that a measure falls within the terms of [a paragraph of] Article XX that that measure also will necessarily comply with the requirements of the chapeau”. The introductory clause of Article XX prohibits any arbitrary or unjustifiable discrimination or disguised restriction on international trade. Whether a measure falls under one of these three types is commented on by the Appellate Body: “The fundamental theme is to be found in the purpose and object [of Article XX] of avoiding abuse or illegitimate use of the exceptions to substantive rules available in Article XX.”

3.507 Canada points out that it has already claimed, in respect of Article III:4, that the Decree was discriminatory. The Decree is arbitrary and unjustified first of all because the current uses of chrysotile do not pose any health risk. The ban is also arbitrary and unjustified because it does not have a solid scientific basis. The ban is also arbitrary and unjustified because it takes the INSERM report as a basis for prohibiting current applications of chrysotile, even though this study did not assess the risks associated with modern uses of chrysotile. The ban is also arbitrary and unjustified because it is not motivated by the objective of protecting human life or health but rather by the desire to reassure a panicked population. According to Canada, even the French Senate and National Assembly state that the ban is “… a decision which concerns public opinion”.

Furthermore, the purpose of the Decree is not to protect the population because it imposes, by its very nature, the blind use of substitute fibres whose lower toxicity is far from proven. What is more, the Decree fosters a false sense of security among the “reassured” population because the public is not afraid of the risks of substitute products, even though these risks are unknown and have not been quantified.

3.508 Canada claims that, in addition to constituting arbitrary and unjustified discrimination, the Decree also constitutes a disguised restriction on international trade. In order for a measure to be a “disguised” restriction on trade, it is not necessary that it be “concealed” or “unannounced”. The Appellate Body excludes a narrow reading of the term “disguised restriction”: “It is equally clear that concealed or unannounced restriction or discrimination in international trade does not exhaust the meaning of ‘disguised restriction’.” Hence, the fact that the measure was published does not prevent it from being a disguised restriction on international trade. The ban is a disguised restriction on international trade and contrary to the introductory clause of Article XX in the sense that, under the cover of a public health decision, the Decree favours the French national industry of substitute products for chrysotile and products containing it.

(e) Article XXIII:1(b) of the GATT

3.509 Canada claims that the incompatibility between the Decree and the obligations of France pursuant to the TBT Agreement and the GATT establishes a presumption that a benefit pursuant to Article XXIII:1(a) of the GATT 1994 and of Article 3.8 of the Understanding has been nullified or impaired. However, in the event that the Panel were to conclude that the French measure complies with the TBT Agreement, the application of the latter nevertheless nullifies or impairs the benefits accruing to Canada under these Agreements, pursuant to Article XXIII:1(b) of the GATT. Within the framework of disputes aris-
ing from the GATT 1947, Article XXIII:1(b) was interpreted as meaning that, even if a measure is judged to be in compliance with the provisions of the GATT, such a measure may nevertheless be contested as nullifying or impairing benefits. Canada points out that, traditionally, panels under the GATT 1947 have deemed that three conditions must be met for a case of nullification or reduction of benefits in a non-violation situation. This reasoning was confirmed after adoption of the WTO Agreements in the matter Japan – Measures affecting Consumer Photographic Film and Paper. These conditions are as follows: (i) the negotiation of a tariff concession; (ii) the subsequent adoption of a governmental measure unfavourably disrupts conditions of competition between the product for which the concessions were granted and the imported product that is a like or directly competitive product; and (iii) the adoption of the measure in question could not reasonably have been foreseen at the time of the tariff concession negotiations.

3.510 Canada asserts that these three conditions are present in this case. First of all, asbestos and many asbestos-containing products are subject to tariff concessions granted by the EC during the Uruguay Round Negotiations. At that time, the EC granted initial negotiating rights to Canada for asbestos. Asbestos and asbestos-containing products were the subject of tariff concessions by France starting in 1947; they were taken up again by the European Economic Community after the 1960-1961 Tariff Conference and have been renewed until now. Secondly, the effect of the Decree’s adoption has been to disrupt the competitive relationship in the French market between, on the one hand, chrysotile asbestos fibre and products containing it and, on the other, like and competitive French products. In establishing a total ban, the Decree has destroyed this competitive relationship and has created a monopoly for substitute fibres and products containing them. Thirdly, at the time the tariff concessions were negotiated, Canada could not reasonably have foreseen that France would adopt the Decree. At the time of the negotiations on tariff concessions for asbestos, there was no indication that France was going to abandon its policy of controlled use of asbestos and compromise the value of its commitments by implementing a total ban on chrysotile and any possible use thereof.

3.511 Canada claims that, at the time of the negotiations on the WTO Agreements, it could in no way foresee that France was going to adopt a measure with regard to chrysotile that is obviously incompatible with the treatment it grants to other potentially hazardous products. Other raw materials, such as lead and copper, are potentially hazardous but are not banned. Certain uses of these products are indeed banned, restricted or subject to regulation. Canada was reasonably entitled to expect a similar approach to chrysotile on the part of France. The excessiveness of the French measure could not have been foreseen by Canada. The excess of the measure can be seen in the treatment given to chrysotile-cement products to be withdrawn from the market. As of 1 January 1997, stocks of chrysotile-cement products became “waste” to be handled and stored according to strict orders stipulated in two other measures adopted by France. One of these measures is particularly revealing of the incompatibility and the excessive nature of the overall French regulatory approach to chrysotile. Indeed, a section of the Note relative aux conséquences de l’interdiction de l’amiante et à l’élimination des déchets [Note on the effects of the ban on asbestos and the elimination of waste] states that “for waste containing bonded asbestos […] if the waste is composed of asbestos combined only with inert matter, it may be eliminated in accordance with the circular of 9 January 1997 on the elimination of asbestos-cement waste”. If France recognizes that products containing bonded asbestos are inert, Canada cannot understand why it had to ban chrysotile and dense products containing it. Thus, a benefit accruing to Canada under the WTO Agreement was nullified or impaired. This benefit was seriously nullified or impaired, i.e. more than de minimis. For the foregoing reasons, the Decree has a negative effect on the objective of liberalizing international trade in the WTO Agreement, in violation of Article XXIII:1(b).

3.512 The European Communities indicate that, in order to determine whether a measure nullifies or impairs a benefit enjoyed by a Member by virtue of the General Agreement, in the context of Article XXIII:1(b) of the GATT, it must be shown that: (i) the Member enjoys a benefit accruing to it under the GATT; and (ii) the measure in question nullifies or impairs that benefit. To conclude that a measure does indeed nullify or impair that benefit,
the complainant must establish that: (i) the measure could not reasonably have been anticipated by that Member when the concession was negotiated; and (ii) the measure upsets the competitive relationship between domestic and imported products that prevailed before the measure was adopted. By way of a preliminary remark, the EC stress that the burden of proof for the application of Article XXIII:1(b) is particularly onerous as a result of Article 26:1(a) of the Understanding, which states that: “the complaining party shall present a detailed justification in support of any complaint relating to a measure which does not conflict with the relevant covered agreement”. This provision essentially reflects the practice established under the GATT 1947. A forceful reminder of the weight of the burden of proof was also given under the GATT 1994 in the Panel Report on Japan - Measures affecting Consumer Photographic Film and Paper, which found that, in this context, it was the duty of the complainant to provide “a detailed justification for its claim in order to establish a presumption that what is claimed is true”. In the case at issue, the EC consider that the very brief explanations supplied by Canada are insufficient to discharge the burden of proof incumbent upon it. In any event, the EC claim that Canada’s submission under Article XXIII:1(b) is unjustified for the following reasons: (i) Canada could reasonably have anticipated the French measure when the concession was negotiated under the Uruguay Round; and (ii) the French measure has not upset the competitive relationship between domestic and imported products that prevailed before the measure was adopted.

Before considering the conditions set out above, the EC point to the observations by the Panel on Japan - Measures affecting Consumer Photographic Film and Paper, namely, that the Article in question has been invoked only eight times during the fifty-year existence of the GATT 1947 and that “…most of the cases of non-violation nullification or impairment have dealt with situations where a GATT-consistent domestic subsidy for the producer of a product has been introduced or modified following the grant of a tariff concession on that product”. While acknowledging that Article XXIII:1(b) could be applied in contexts other than subsidies, the Panel stated that it “should be approached with caution and treated as an exceptional concept.”

The EC contend that it is important first of all to note that there can be no “legitimate expectations” in the case of a measure that is taken to protect human health and can therefore be justified, particularly with regard to Article XX(b) of the GATT or Article 2.2 of the TBT Agreement. While a Member may have “legitimate expectations” in connection with a purely commercial measure, no such expectations can be pleaded when it comes to health protection matters. The protection of human health is a fundamental duty of any government and it cannot be compromised or restricted by the concept of non-violation. The EC contend that, by their very nature, science and scientific evidence are constantly evolving and a restriction on the right of Members to protect the health of their people, based on Canada’s arguments, would run counter to the scope, aims and structure of the

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GATT as a whole. For example, a Member might have legitimate expectations of computer markets opening up, but it cannot have legitimate expectations of improved access to the market for a carcinogenic product. On the contrary, if there is any expectation, it is that access to that market may be barred or restricted rather than improved. In the EC’s view, Canada has been unable to prove that the measure in question could not have been anticipated at the time when the concession was negotiated. Canada had good reason to suppose that France would adopt rules prohibiting asbestos.

3.516 The EC point out that, since 1977, asbestos fibres, in particular chrysotile, have been classified as category I carcinogenic products by the WHO. The EC claim that, when the tariff negotiations took place, Canada knew that there was a danger that the product under negotiation could at any time be prohibited by Members of the WTO, particularly if non-hazardous or less hazardous substitutes could be used. In 1986, ILO Convention 162 on asbestos stated that national legislation should provide wherever possible for the “replacement of asbestos or of certain types of asbestos or products containing asbestos by other materials or products or the use of alternative technology, scientifically evaluated by the competent authority as harmless or less harmful.”

3.517 The EC maintain that the scientifically proven carcinogenicity of chrysotile and the impossibility of keeping the risk under control in all cases led the French Government to halt any spread of the risk by applying the principle recommended by the WHO and the ILO - and provided for by the European Union - with regard to carcinogenic products, namely that they be replaced by less hazardous products wherever technically possible. Since 1990, the European Community has provided for the replacement of asbestos under Council Directive 90/394/EEC of 28 June 1990 on the protection of workers from the risks related to exposure to carcinogens at work (Sixth individual Directive within the meaning of Article 16 (1) of Directive 89/391/EEC), Official Journal L 196 of 26 July 1990. Under this Directive, the European Community recommends a framework for the health and safety of workers, including the principle of replacing a dangerous substance or process with a non-dangerous or less dangerous substance or process, where they exist. Regarding carcinogens, the principle is set out in the Directive on carcinogens, which provides explicitly for carcinogens to be replaced by less dangerous substances, preparations or processes where technically possible. In 1983, WTO Members began to ban the use of asbestos, including chrysotile. The EC consider that the ban imposed by France could have been anticipated by Canada. Many WTO Members, including Canada itself, have banned amphibole asbestos for many years. In the EC’s view, this means that Canada considered that it was impossible to envisage a “safe” use of amphibole asbestos. Based on this and the fact that chrysotile asbestos is classified in the same category of products proven to be carcinogenic for humans as amphibole forms of asbestos, Canada should have expected that several Members, including France, would take steps to ban chrysotile asbestos. Moreover, the EC indicate that France is not the first European Union country to have banned chrysotile fibre. Austria banned chrysotile in 1990, followed by Finland and Italy in 1992, and Germany in 1993. As several European Union countries have banned chrysotile asbestos, Canada could easily have anticipated that other European Union countries, including France, would follow suit.

3.518 The EC maintain that Canada’s argument that it could not legitimately have anticipated the ban imposed by the Decree because France did not simultaneously ban other potentially dangerous substances (such as lead and copper) is misleading. There are no provisions in the GATT or the TBT Agreement requiring Members to be so consistent in applying health measures against substances that pose a carcinogenic risk for human health. The EC consider that accepting Canada’s argument would be equivalent to preventing Members entirely from taking measures to protect human health on their territory. In the EC’s view, Canada can no longer claim a legitimate expectation at the time of the concession because such a legitimate expectation would have to concern “improved” market-access opportunities. It is clear that a product which entails risks for human health cannot legitimately offer “improved” market-access opportunities. Moreover, the facts clearly show that in industrialized countries the trend in asbestos fibre imports is going down rather than up. The EC maintain that Canada could not therefore have had a legitimate expecta-
tion that the opportunities for access to the market in chrysotile fibres would “improve”. It also appears that the tariff concession invoked by Canada actually dates from at least as far back as the 1960-1961 negotiating round. The Panel on Japan - Measures affecting Consumer Photographic Film and Paper pointed out that “… the establishment of a case based on expectations from rounds concluded 18 or 30 years ago may be difficult”.

In the case at issue, the EC maintain that Canada must give detailed reasons as to why it could legitimately have expected that France would not adopt measures restricting or eliminating the use of any asbestos product after the Uruguay Round negotiations, given the growing scientific evidence that all types of asbestos and asbestos-containing products were carcinogenic to humans.

3.519 The EC claim that Canada fails to demonstrate how the French measure upsets the competitive relationship between asbestos and fibrous or non-fibrous substitute products. As the EC maintained in the section relating to Article III of the GATT, there is no “similarity” between asbestos and fibrous or non-fibrous substitute products. Because of their different characteristics there can be no distortion of competition between these products. Asbestos and asbestos-containing products (and asbestos-cement in particular) are replaced by many fibrous or non-fibrous products (plaster, ductile iron, etc …). Finally, Canada’s exports were essentially, if not exclusively, of raw chrysotile asbestos, which is like only to amphibole asbestos. The EC note that, on this subject, the Panel on Japan - Measures affecting Consumer Photographic Film and Paper found that:

“… it must be demonstrated that the competitive position of the imported products subject to and benefiting from a relevant market access (tariff) concession is being upset by (nullified or impaired … as the result of) the application of a measure not reasonably anticipated… Thus, in this case, it is up to the United States to prove that the governmental measures that it cites have upset the competitive relationship between domestic and imported photographic film and paper in Japan to the detriment of imports. In other words, the United States must show a clear correlation between the measures and the adverse effect on the relevant competitive relationships.”

3.520 The EC contend that Canada has in no way established such a clear correlation. Furthermore, the conditions of competition on the French market have not been upset. On this point, Canada claims that “the effect of the Decree’s adoption has been to disrupt the competitive relationship in the French market between, on the one hand, chrysotile asbestos fibre and products containing it and, on the other, like and competitive French products”. On this issue, the EC point to the findings of the Panel on Japan - Measures affecting Consumer Photographic Film and Paper: “in an Article XXIII:1(b) case the issue is not whether equality of competitive conditions exists but whether the relative conditions of competition which existed between domestic and foreign products as a consequence of the relevant tariff concessions have been upset”.

3.521 The EC claim that the products for which the competitive conditions must be examined are those covered by the tariff concession. If a tariff concession is granted for asbestos, the competitive conditions to be examined are those concerning Canadian asbestos and French asbestos. It is therefore irrelevant for Canada to try to compare chrysotile with French substitute products because such products cannot be considered in terms of the same relevant tariff concession. In conclusion, the EC consider that Canada has not provided detailed explanations to justify its claim that the Decree nullified a tariff concession which, in its view, it could legitimately have expected under the GATT.

3.522 Canada maintains all its arguments concerning the incompatibility of the French measure with Article XXIII:1(b) of the GATT. It contends that it could not have anticipated, at the time of the most recent tariff negotiations, that France would adopt such a drastic and unreasonable measure as a ban on all forms of asbestos and products containing it. Canada maintains that it could not have anticipated, at the time of the negotiations, that France was going to nullify and impair in this manner its tariff concession concerning chrysotile and products containing it. Canada maintains that, when a complainant proves
that it enjoys a tariff concession and the respondent subsequently adopts a measure that affects the value of this concession, the complainant benefits from the presumption that it could not reasonably anticipate that this concession would be nullified or otherwise impaired by this measure. In such circumstances, it is up to the respondent to prove that the complainant should have anticipated the possibility that such a measure would be adopted.\textsuperscript{634} In Canada’s opinion, the EC have not proved this. Although Canada recognizes France’s right to take action in order to protect human health and the health of workers, Canada could not reasonably have anticipated that France was going to adopt a measure totally prohibiting asbestos, without distinction as to the types of fibre or their use. This measure did not exist at the time of the negotiations and there was nothing to suggest that France was going to adopt such a radical measure as a ban on chrysotile and asbestos-cement. The lack of consistency in the nature of the regulatory intervention, as well as its severity compared to the type of regulatory intervention then in place in France and still in effect now concerning products equally harmful, if not more harmful, than chrysotile, mean that it would have been impossible for Canada reasonably to anticipate that France was going to act in the manner that it did.

3.523 Canada claims that, at the time of the most recent tariff negotiations, conducted by the EC during the Uruguay Round, France maintained a controlled use approach for chrysotile asbestos and there was nothing to suggest at the time that it was going to ban the product suddenly without any scientific reason to justify this maximum increase in the restrictive effects on trade in the products concerned. Similarly, at the time these concessions were negotiated, many hazardous products were, and still remain today subject to a controlled use approach whereby certain uses of these products continue to be permitted. Canada therefore had every reason to believe that the controlled use approach was going to continue to be preferred for these products, including chrysotile asbestos. Canada had every reason to believe that the adoption of such a radical measure as a total ban on asbestos was not going to be adopted. In light of the regulatory approach advocated by France up to that point, the extreme nature of such a measure made it unforeseeable. As Canada has already claimed, there is nothing exceptional in the case of chrysotile and products containing it in terms of risk management, if there is any risk. Other products, which moreover have been proven to be hazardous, are the subject of a controlled use policy. Furthermore, the effect of the French prohibition is to impose the replacement of chrysotile by a range of substitute products of domestic or foreign origin, even though they are suspected of being carcinogenic by the French authority which is presented as being the source of the ban on chrysotile, namely INSERM. In these circumstances, Canada’s reasonable expectations were that the competitive position of imported chrysotile in France and of products containing it would not be affected by a measure such as a total ban, without distinction as to the types of fibre or their use, in favour of substitute fibres of domestic or other origin. Canada’s reasonable expectations were also that this type of measure would not be adopted unless there were exceptional circumstances, which is certainly not the case with respect to the circumstances surrounding the use of chrysotile. There have been no new scientific developments that have changed anything in terms of managing the risks or the effects associated with chrysotile. Finally, Canada’s reasonable expectations were that such a ban would not be adopted in favour of substitute products unless these products had been subjected to a rigorous process of examination to prove that their use satisfies the public health objectives invoked by France. The evidence submitted by Canada in this respect shows very clearly that the substitutes do not meet this requirement.

3.524 Canada maintains that Article XXIII:1(b) of the GATT is applicable to the present case because the Decree radically alters the competitive conditions between chrysotile asbestos exported by Canada and substitute products. As a result of the Decree, the latter enjoy a decisive advantage over Canadian chrysotile asbestos, which can no longer be exported to France. Canada rejects the EC’s claim that an examination of the impact of the Decree’s effect on competitive conditions must be limited to Canadian asbestos and French asbestos, and exclude any examination of the competitive relationship between Canadian chrysotile asbestos and substitute products of French or other origin. Canada points out that this European claim is contradicted by the Panel decisions in \textit{Treatment by Germany of Imports of Sardines}\textsuperscript{635} as well as in \textit{Australian Subsidy on Ammonium Sulphate}\textsuperscript{636}. In the first
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For these reasons, Canada asserts that, without regard to the question of whether the measure is consistent with the provisions of the GATT 1994, it nullifies or impairs the tariff concessions granted to Canada by the EC with respect to access to the French market for chrysotile asbestos and products containing it.

The European Communities point out that the Decree cannot be considered incompatible with the provisions of Article XXIII:1(b) of the GATT. The EC stress that the burden of proof, in the context of Article XXIII:1(b) of the GATT, is particularly heavy and lies with Canada. The EC consider that Canada has not provided sufficient evidence of incompatibility with this Article.

The EC claim that the rules on “non-violation” apply only if the measure in question does not fall under other provisions of the GATT. The EC recall that the Panel on Japan - Measures affecting Consumer Photographic Film and Paper noted that “We reach this conclusion in considering the purpose of Article XXIII:1(b), which is to protect the balance of concessions under GATT by providing a means to redress government actions not otherwise regulated by GATT rules…”637 The EC consider that Article XXIII:1(b) is applicable only if the Panel reaches the conclusion that the Decree is consistent with Article III of the GATT, or possibly with the TBT Agreement if the Panel were to apply that Agreement to the present case. Otherwise, the EC consider that there cannot be “non-violation”. Moreover, in this connection, the EC recall that Article XX of the GATT provides, in particular, that “…nothing in this Agreement shall be construed to prevent the adoption or enforcement by any contracting party of measures: …(b) necessary to protect human … life or health”. Similarly, the preamble to the TBT Agreement states that “…no country should be prevented from taking measures necessary … for the protection of human … life or health …”. The EC therefore conclude that, if the French measure is considered “necessary” for the protection of human health by the Panel and hence if specific rules have been applied in this respect, the provisions of Article XXIII:1(b) of the GATT are inapplicable.

The EC maintain that Canada could not have “legitimate expectations” with respect to a measure taken to protect human health and which can therefore be justified, particularly with regard to Article XX(b) of the GATT or Article 2.2 of the TBT Agreement. While a Member may have “legitimate expectations” in connection with a purely commercial measure, no such expectations can be pleaded when it comes to health protection matters. The EC note, moreover, that the United States shares this point of view.638 The EC indicate that Canada, which was aware of past and present scientific research and of the steps being taken by the relevant international organizations to encourage the rapid replacement of asbestos by substitute products, could not have had “legitimate expectations” arising out of relevant tariff concessions, whatever the date of those concessions.

Canada reiterates its arguments on non-violation set out above. The following comments are limited to rejecting certain claims by the EC in paragraphs 3.526-3.528. Canada
notes that the EC claim that in a situation of non-violation no action can be taken against a measure deemed justified in accordance with Article XX of the GATT. According to Canada, the cases of *Uruguayan Recourse to Article XXIII* and *United States – Trade Measures affecting Nicaragua* do not support such an interpretation. In the Uruguay case, the Panel considered that measures to protect human health contrary to Article XI but justified under Article XX(b) could nevertheless lead to recourse in a non-violation situation. In the Nicaragua case, the whole procedure was based on the premise that the American embargo was justified under Article XXI. Although it would have been correct to claim that an appeal in a non-violation situation cannot be made if a measure is an exception, in the Nicaragua case, the Panel did not carry out an in-depth examination. This question was, however, never raised by the United States and the Panel therefore carried out an examination and published its findings.

3.530 Canada points out that the EC’s reasoning is not substantiated for three additional reasons. Firstly, contrary to the EC’s claim, inasmuch as Article 26:1(b) of the Understanding does not provide for the granting of compensation rather than withdrawal of a measure, recourse in non-violation affects neither “the adoption” nor “the application” of the contested measure. Secondly, the EC’s attempt to develop a category of “measure(s) of a purely commercial nature” in comparison with a “measure which has health-protection related aspects” is, to say the least, dubious. Not only is it purely artificial, but this distinction has no basis, neither in the texts of the WTO Agreement nor in the case law. Nothing in Article XXIII:1(b) provides for such a distinction. Canada claims that a legitimate expectation does not in any way concern a particular measure or series of measures adopted by a Member, but rather legitimate expectations in relation to the opportunities for competition agreed during multilateral trade negotiations on a given product. Canada notes that the reasoning of the United States cited by the EC suffers from the same confusion. Lastly, the European reasoning is wrong because it does not concur with what was expressed during the preparatory work on the GATT 1947, as it appears in the theory "(...) one of the principal objectives of the Article [XXIII:1(b)] is to prevent circumvention of the provisions of [the Agreement]. Under this Article, if a Member State utilizes the exceptions provided in the Article [XX(b)] as a means of protection, any other Member may take the matter up with the ITO and obtain satisfaction. It is virtually impossible to foresee exceptions that will not lend themselves to abuse if good faith is lacking. The League of Nations adopted an article along the lines of the Article [XXIII:1(b)] precisely because it was unable to find a formula for exceptions that excluded any possibility of abuse". Canada claims that it has provided detailed proof of the elements set out in the case of *Japan – Measures affecting Consumer Photographic Film and Paper* that make it possible to determine that the French measure, even if consistent with the GATT, nonetheless nullifies or impairs benefits accruing to Canada under the WTO Agreements or the attainment of one of the objectives of the Agreements.

3.531 The European Communities claim that the Decree is not incompatible with the provisions of Article XXIII:1(b) of the GATT and its case law. As they have already pointed out, it is not possible to claim “legitimate expectations” with respect to a measure that is taken to protect human health and can therefore be justified, particularly with regard to Article XX(b) of the GATT or Article 2.2 of the TBT Agreement. While a Member may have legitimate expectations in connection with a purely commercial measure, no such expectations can be pleaded when it comes to measures taken to protect human health. This is also the position taken by the United States in its third party intervention. The EC note that Canada referred to two Panel Reports, one concerning Uruguay (1962) and the other Nicaragua (1986), to claim support for its proposition that a non-violation claim can be made in this case. According to the EC, neither of the two Panel Reports referred to by Canada, nor any other panel report, lends support to such claims. The first Panel has no relevance whatsoever to the issues discussed here. The second, unadopted, Panel Report in the Nicaraguan case is also irrelevant, as the Panel in that case specifically stated that it refrained from taking any position on the non-violation claims made by Nicaragua. The EC note that Canada also argued that the background to Article XXIII:1(b) supports the view that non-violation complaints may and should remain possible to deal with situations of bad faith and abuse in the application of the provisions of the General Agreement.
and of Article XX. This is not correct. Canada takes a selective look at some parts of the preparatory documents. As the Appellate Body stated in the United States - Shrimps case, the conditions laid down in the chapeau of Article XX(b) are meant precisely to address situations in which a Member applies in bad faith and in an abusive manner the exceptions laid down in Article XX. In the EC’s view, this means that the potential problem of abuse and bad faith, alluded to by Canada, is adequately covered by the “chapeau” of Article XX and there cannot be two sets of provisions (non-violation and the chapeau of Article XX) which address the same problem twice. The EC therefore propose that this argument of Canada also be rejected.
NOTES

1 Petit Robert 1, Dictionary of the French language.
2 Journal officiel of 26 December 1996. See Annex I to this report.
3 In connection with this aspect, Article 4 refers back to Decree No. 96-98 of 7 February 1996 on the protection of workers against risks linked to inhaling asbestos dust (Journal officiel of 8 February 1996).
4 In connection with this aspect, Article 4 refers back to Decree No. 88-146 of 28 April 1988 on products containing asbestos (Journal officiel of 8 February 1988).
6 European Commission (G. Lohan, DG III), European Justification of Decree No. 96-1133 to the Canadian authorities (15 April 1997) following the French Notification (G/TBT/Notif.97.55).
8 B. Terracini, Review of Technical and Scientific Documents annexed by Canada to its Submission of April 26, 19 May 1999 (document transmitted to the Panel by the EC).
9 See pp.24-230 of the INSERM report.
11 Canada notes that, until the adoption of the Decree, 90 per cent or more of French imports of chrysotile fibres were used to produce chrysotile cement (see Le Déaut, J.-Y. and Revol, H., L’amiante dans l’environnement de l’homme: ses conséquences et son avenir, Office parlementaire d’évaluation des choix scientifiques et technologiques, National Assembly no. 329/Senate no. 41, 1997). According to the INSERM report, in 1984, for example, French production of chrysotile cement was 600,000 tonnes. In 1991, eight French factories produced 540,000 tonnes (see INSERM, Rapport sur les effets sur la santé des principaux types d’exposition à l’amiante, INSERM joint report, Paris, INSERM publications, 1997, p.21).
14 Canadian Ministry of Natural Resources.
15 INSERM report, p.387.
16 Id. Table Importations d’amiante entre 1938 et 1992, p.189.
17 Chrysotile asbestos evaluated by health experts, Press communiqué (51), WHO, 26 July 1996.
18 See paragraph 3.78 below.
19 For a chronology of facts (from June to September 1996), see Info-Science <http://www.infoscience.fr/travaux/amiante/chrono.html> (access date: 4 April 1999). According to Canada, the controversy over asbestos in France results, in large part, from the situation at the University of Jussieu, where a group has led a very active fight against asbestos use for several years. On the topic of the situation in Jussieu, Claude Alègre, who was Minister for Education, Research and Technology in the Jospin Government, said: “The psychosis of those demanding that asbestos be cleared from all buildings is irrational and hazardous. The radical solution that has been chosen is going to shut down, for four years, France’s foremost research centre (the University of Jussieu). That university is being wrecked for a figment of the imagination … I do not know whether asbestos at low doses causes cancer. It is not impossible, but no one has proven it scientifically.” (Le Point, 12 October 1996. See also Le Figaro, 26 December 1996.)
20 L’amiante dans l’environnement: ses conséquences et son avenir, Office parlementaire d’évaluation des choix scientifiques et technologiques, National Assembly no. 329/Senate no. 41, p.57.
22 Pursuant to Article 249 (formerly Article 189) of the Treaty establishing the European Community, a directive “is binding, as to the result to be achieved, upon each Member State to which it is addressed”, while leaving to the national authorities the choice of form and methods.


Article 41 of the Law on the financing of social security; Decree and Orders of 29 March 1990.

Flocking: application of fibres onto any surface to form a coating with a fibrous, velvety or downy appearance.

Lagging: thermal insulation technique used to avoid heat loss from heating installations, pipes and ventilation shafts.

For values measured during the handling of asbestos-cement materials (source EVALUTIL), see Ministry of Labour, Social Dialogue and Participation, Note setting out the main guidelines of the Senior Council for the Prevention of Occupational Hazards, 3 July 1995.

French Senior Council for Public Health, Opinion on premises flocked with asbestos (Division for the assessment of environmental risks to health), session of 15 September 1994.


44 Table of occupational diseases produced by the Caisse d’assurances maladies des travailleurs salariés (CNAMTS).

45 See Section III.B.7 of this report.

46 Institut National de Recherche en Sécurité (National Safety Research Institute).

47 Canada notes that, according to the World Health Organization, “risk of mesothelioma and bronchial cancer attributable to asbestos in the general population is undetectably low; the risk of asbestosis is practically nil”. (World Health Organization, Environmental Health Criteria 53: Asbestos and Other Natural Mineral Fibres, Geneva, p.135).

48 Id.


52 See, in particular, Commins, B.T., The Significance of Asbestos and Other Mineral Fibres in Environmental Ambient Air, Cummins Associates, Berkshire, U.K., 1990, p. 17. See also Commins, B.T., Estimations of Risk from Environmental Asbestos in Non-Occupational Exposure to Mineral Fibres, IARC Scientific Publication no. 90, Lyon, 1989, pp. 476 and 477. Canada notes that the data obtained in the United Kingdom under situations of very heavy vehicular traffic indicate that the use of asbestos in brake linings does not measurably contribute to atmospheric asbestos concentrations in the urban environment. Even at two heavily-used intersections in the London metropolitan area, concentrations varied from 0.0002 to 0.0004 f/ml (Jaffrey, S., Environmental Asbestos Fibre Release from Brake and Clutch Linings in Vehicular Traffic (1990) 34 Ann. Occup. Hyg., Vol.34, p.529.)


55 According to Canada, the studies concluded that there is no health risk linked to the ingestion of asbestos. See, in particular, Commins, B. T., Estimations of Risk from Environmental Asbestos in Non-Occupational Exposure to Mineral Fibres, IARC Scientific Publication no. 90, Lyon, 1989, pp. 476-478.


57 Canada notes that natural concentrations of asbestos fibres are found in spring water, whether or not asbestos-cement pipes are used. The possible contribution to asbestos concentration in water after the use of asbestos-cement pipes was studied in several countries. For example, in the State of Illinois, in the United States, in 15 public water supply networks, some of which were 50 years old, there was no difference in fibre concentrations between samples of drinking water before and after they passed through asbestos-cement pipes. (Hallenbeck, J. et al., Is Chrysotile Asbestos Released from Asbestos-Cement Pipes into Drinking Water? (1978) Amer. Water Works Ass., Vol. 70, no. 2, p.97). See also World Health Organization, Letter to T.A. Jafri, Asbestos in Drinking Water/Amiante dans l’eau de boisson, Helmer, D.R., WHO, Geneva, 5 April 1989.


59 Commins, B.T., The Significance of Asbestos and Other Mineral Fibres in Environmental Ambient Air, Cummins Associates, Berkshire, U.K., 1990, p. 62. According to Commins, “[A]ny risk for future exposures to asbestos are of course likely to be lower still because of improving control measures.” (p. 64.)
Académie nationale de Médecine (Étienne Fournier), Amiante et protection de la population exposée à l’inhalation de fibres d’amiante dans les bâtiments publics et privés, Bulletin de l’Académie nationale de médecine, Vol. 180, no. 4, p. 5.


64 Canada indicates that, over the last 15 years, ambient air concentrations of asbestos in the institutions where chrysotile was being extracted in Quebec never exceeded 0.02 f/ml. In 1986, A. Churg (See Churg, A., Lung Asbestos Content in Long-Term Residents of a Chrysotile Mining Town, (1986) 134 Amer. Rev. Respirat. Diseases 125) concluded that residents of the mining region of Thetford Mines in Quebec showed no sign of disease due to asbestos and that repeated epidemiological studies did not document any abnormal incidence of respiratory diseases in persons that had never worked in this industry. Canada also draws attention to the work of Dr. Camus et al (Camus, M., Siemiatycki, J., Meek, B., Nonoccupational Exposure to Chrysotile Asbestos and the Risk of Lung Cancer, (1998) 338, New England Journal of Medicine 1565) They published a vast study on women in chrysotile mining communities in Quebec, many of whom were exposed to very high levels of fibres between 1920 and 1975. These women were subjected to exposure of 0.0107 f/ml, higher than the current exposure limits in France, and literally thousands of times higher than the levels measured in public buildings. Nonetheless, Canada notes that no excess in lung cancer was detected in this population. According to the study’s authors, this is particularly important in light of the current French situation. In fact, applying the risk model adopted by France for the exposure studied results in a forecast of approximately 100 lung cancer deaths, while in reality there are none. Likewise, use of the French risk model would have resulted in estimates of approximately 250 and, at any rate, no less than 50 deaths from mesothelioma, whereas the preliminary results of the study in question show only 10 cases, some of which may be associated with exposure to amphiboles. Research continues, particularly with an analysis of the work history of each individual in order to determine the exact link, if any, between these cases of mesothelioma and on-the-job exposure, as well as exposure to amphiboles.


66 See paras. 3.120 et seq below.

67 Canada points out that in Germany, for example, a study revealed the low degree of chrysotile fibre emissions in the environment from chrysotile-cement roofing materials, even when these were in a state of advanced corrosion. These chrysotile fibre concentrations, which were measured in urban areas, were well under 0.001 f/ml, i.e. the concentration found acceptable by the German health authorities. (Teichert, U. (1986) 46 Staub Reinhaltung der Luft 432, p. 7 of the English translation.) In Austria, after having compared chrysotile fibre concentrations between areas with and without chrysotile-cement roofing (< 0.0001 f/ml), it was concluded that there was no significant link between the use of chrysotile-cement based materials and the chrysotile fibre concentrations found in the areas concerned. (Felbermeyer, W. S., Ussar, M.B., Environmental Pollution by Atmospheric Effects on Asbestos Cement Sheets, Institute for Environmental Protection and Clean Air, Leoben Austria, 1980). Canada notes that, in Australia, the possible contribution of chrysotile-cement roofing materials on school buildings to environmental asbestos concentrations has been analysed. Most of the concentrations were found to be equal to or less than 0.0002 f/ml. (Safety and Welfare of Western Australia, Asbestos Cement Products, Report of the Working Party, 1990).


74 Id.
79 See also the INSERM report, Chapter 8 (pp.154-190), which analyses in detail how the frequency of mesotheliomas has evolved internationally. INSERM report, pp.172-180.
80 See paragraph 3.222 below.
83 The EC note that Canada seems to believe that this figure is not correct, because it assumes that 5.7 per cent of cases of lung cancer are attributable to asbestos, a figure which comes from a single study carried out in Scotland (and which Canada considers too high, though it does not propose a different value based on the results of scientific studies). This criticism is not justified. The EC point out that, on page 10 of the INSERM report, there is a table summarizing the proportions of lung cancers attributable to asbestos in all the studies which provided an estimate: it shows that in certain studies this fraction is clearly higher than the 5.7 per cent given, and may considerably exceed 10 per cent. In the part of the report which explains the method of calculating the number of cases of lung cancer (page 180), the higher percentage (7 per cent) observed 10 years earlier in Great Britain has not been used since it was believed that this proportion must have declined. According to the EC, it is therefore clear that the INSERM report consistently uses relatively low estimates of the effects of asbestos on cancer mortality in France.
88 INSERM report, pp.259-266.
91 INSERM report, pp.193-202, particularly Table 2, p.196 and figures 1 and 2, pp.198 and 199.
105 See Chapter 9, pp.193-241 of the INSERM report.  
107 See, for example, paragraph 3.124 below.  
108 See, for example, paragraph 3.53 above.  
114 The EC emphasize that the INSERM report, so disparaged by Canada, also stressed the undetectability of the risks associated with very low exposures which are found in the ambient air of towns and buildings (see pp. 145 and 146, and 224-230).  
115 See in particular INSERM report, p. 213.  
116 According to Canada, the pathogenicity of asbestos fibres varies depending on their dimensions. Hazardous fibres are defined as those having a length of more than 5 microns, a diameter of less than 3 microns and a length-diameter ratio of more than 3:1. See in particular IPCS Environmental Health Criteria 203 on Chrysotile, WHO, 1998, p. 14. The longest and finest fibres pose the greatest health risk.  
117 According to Canada, the lack of epidemiological data demonstrating the risks associated with low-dose exposures makes it possible to state that the exposure levels associated with the current uses of asbestos do not pose any detectable health risk. See in particular IPCS Environmental Health Criteria 203 on Chrysotile, WHO, 1998, p. 144.  
118 Id., p. 51; INSERM report, p. 90 et seq. See also para. 3.188 below.  
119 See IPCS Environmental Health Criteria (203) on Chrysotile, WHO, Geneva, 1998, pp. 69 and 81; INSERM report, Table 2, p. 196; EPA, Integrated Risk Information System, Asbestos, Document No. CASRN 1332-21-4 on-line: EPA, <http://www.epa.gov/natispgm3/iris/subst/0371.htm> (access date: 10 June 1999). Canada points out that, in the large majority of the experimental protocols, the comparisons of the effects were still based on gravimetric data; i.e. the effects were produced by an equivalent mineral mass. In fact, retrospective attempts aimed at converting gravimetric doses into doses of numbers of fibres have indicated that, if based on the number of fibres, pathogenicity studies would show that, fibre for fibre, chrysotile fibres are less pathogenic than the other types of asbestos fibres, indeed even less than certain synthetic fibres.  
121 INSERM report, Table 2, p. 196.  
122 See in particular Wagner, J.C. et al, Correlation Between Fibre Content of the Lung and Disease in East London Asbestos Factory Workers (1988) 45 British Journal of Industrial Medicine 305, according to which: “We believe therefore that chrysotile is the least harmful form of asbestos in every respect and that more emphasis should be laid on the different biological effects of amphiboles and serpentine asbestos fibre.”  
116 Mossman, B.T. and Churg, A., *Mechanisms in the Pathogenesis of Asbestosis and Silicosis* (1998) 157 American Journal of Respiratory and Critical Care Medicine 1666, p. 1669. The authors also state: “Both animal and human studies show that continuing exposure to amphiboles results in continuously increasing amphibole fiber levels recoverable from the lung, whereas continuing exposure to chrysotile is associated with a negligible increase in chrysotile fiber burden over time.” (Id., p. 1669).
118 INSERM report, pp. 395-96.
120 Id.
127 Id.
132 See paras. 3.222 and 3.223 below. Canada draws attention to the statement by the WHO in 1998 that the use of this type of data: “contributes less to our understanding of the effects of chrysotile, due to concomitant exposure to amphiboles.” IPCS *Environmental Health Criteria* (203) on Chrysotile, WHO, Geneva, 1998, p. 107.
133 Canada notes that, in the case of bans on asbestos, the regulatory limits apply to asbestos already in place.
136 See paras. 3.76 et seq above.
137 INSERM report, p. 327.
138 See para.3.76 above.
140 See paragraph 3.83 above.
According to Canada, the linear model predicts risks 100,000 times higher than those predicted by a "log-probit" model at doses 100,000 times lower than those for which risks have been observed. See Brown, C.C., Mantel, N., Models for Carcinogenic Risk Assessment, Science 1978; 202:1105.


157 According to the EC, it is estimated that asbestos causes twice as many cancers of the lung as cancers of the pleura (mesotheliomas), see Stayner et al. Exposure to Chrysotile Asbestos and Cancer Risk: a Review of the Amphibole Hypothesis, American Journal of Occupational Health, 1996, 86:179-186.


160 See para. 3.22 above.


164 See paras. 3.78 et seq. above.

165 See para. 3.59 above.

166 Health and Safety Commission, United Kingdom, (1999), Proposals for Amendments to the Asbestos (Prohibitions) Regulations 1992. See also para. 3.59 above.


171 Id.


175 Fibres minérales artificielles et amianté (Résumé). Report of the Groupe scientifique pour la surveillance des atmosphères de travail (G2SAT), INRS – DMT no. 69.


177 Avis concernant l’amiante chrysotile et les produits de substitution envisageables, Scientific Committee on Toxicity, Ecotoxicity and the Environment (CSTEE), 15 September 1998.

The EC point out that, as soon as the INSERM report on man-made mineral fibres came out, France adopted an action plan aimed at protecting workers exposed to man-made mineral fibres.

The EC note that, in 1994, France imported 35,000 tonnes of asbestos and produced 436,000 tonnes of asbestos-cement (Source: Association Française de l’amiante).

That is, without human intervention, explain the EC.


The EC note that an extensive search of the international bibliographical databases failed to reveal any scientific publication by the author of this report concerned with the study of asbestos-related risks.


Report of the Royal Commission on Matters of Health and Safety arising from the Use of Asbestos in Ontario, Table of Contents and Overview, Toronto, Queen’s Printer, 1984.


WHO, Exposure to Asbestos Limits in the Workplace, Report prepared by a WHO committee, Oxford, United Kingdom, 10 and 11 April, 1989, p.12.


Summary of Symposium on Health Aspects of Exposure to Asbestos in Buildings, Energy and Environmental Policy Center, Harvard University, 14-16 December 1988, pp. 26 and 27.

Id., p.27.


Ruling of the 5th Circuit Court of the United States dismissing the EPA regulation, handed down on 18 October 1991.


Decree No.78-394 of 20 March 1978.


Decree No.96-98 of 7 February 1996 concerning the protection of workers against the risks associated with asbestos dust inhalation. Canada notes that this Decree establishes the principles of prevention in the three work situations in which a worker is exposed to asbestos by virtue of his activity: manufacturing activities, for which the prohibition is not an exceptional and temporary measure; asbestos removal or containment operations; con-
tact with asbestos during maintenance work. The report of the Office parlementaire d’évaluation des choix scientifiques et technologiques emphasizes that, according to the dose-response ratios established by INSERM and in view of the current worker exposure standards (i.e., 0.1 f/ml), “it can therefore be asserted that the measures provided for workers […] appear to be quite appropriate and sufficient”. (WTO translation)

205 Toys, materials or preparations for spraying, finished products in powder form meant for retail sale to the public, merchandise for smokers, catalytic screens and insulation materials in heating equipment using liquefied gas, paints, varnishes, road coating products if the fibre content is greater than 2 per cent, mortar, protective finishes, fillers, sealing products, joining pastes, mastics, glues, decorative powders and wall faces, low-density insulation or soundproofing materials, air filters and filters for the transport, distribution and use of natural gas or city gas, backings for plastic-coated wall and floor liners, finished textiles, bituminized felts for roofing, Toasters, heat distribution devices, ironing boards, ironing board covers, iron holders, mobile heaters, insulation panels for the do-it-yourself market, boxed insulation panels for the professional market and materials destined for heat insulation in heating equipment, pipes and cable jackets.

206 Decree No.96-97 of 7 February 1996 concerning the protection of the population against the health risks associated with exposure to asbestos in buildings.


214 See paragraph 3.423 below.


217 See para. 3.122 above.

218 See paras. 3.59 et seq above.


220 Canada points out that, moreover, the manual to which the EC refer deals not only with responsible use but describes controlled use in detail. This manual is intended for producers only and does not mention work methods at construction sites or for building sub-trades.


230 Canadian solid waste regulations.
232 Id.
234 INSERM report, p.179.
235 See para.3.133 above.
238 Weill, H. and Hughes, J., Letter to the Editor: Mesothelioma.
240 European Commission (G. Lohan, DG III), European justification for Decree No.96-1133 to the Canadian authorities (15 April 1997 following the French notification G/TBT/Notif.97.55).
242 Id., p.24.
244 L’évolution des taux d’empoussièrement dans les mines du Québec, in L’amiante chrysotile, un aperçu, Institut de l’amiante, Montreal.
253 Camus, M., L’amiante et les risques pour la santé, April 1999, pp.9 and 10.
255 See para.3.51 above.


266 See Cossette, M., Substitutes for Asbestos, (December 1998) on the technical properties of the main fibres used to replace chrysotile. The paper also deals with substitute products for products based on chrysotile fibre and notes certain undesirable environmental consequences related to the manufacture of the substitute products. Finally, the paper delves into the economic, energy policy and safety considerations related to the use of substitute fibres and products. See also Anderson, A., Fibers in Friction Materials, (December 1998). A. Anderson is the chairman of the sub-committee on health and the environment (Brake Committee) of the Society of Automotive Engineers and was formerly responsible for friction materials at the scientific laboratories of the Ford Motor Company.


268 INSERM report, p.434.


270 Id., p.2.

271 Id.

272 The full INSERM report entitled Effets sur la santé de substitution à l’amiante was published in November 1999, i.e. after the parties had presented their written rebuttals to the Panel. Canada referred to this report at the second substantive meeting with the parties. See paras. 3.325-3.327.

273 Substitutes – Executive Summary, p.33.


276 See Davis, p. 1.

277 Id., p. 6. Canada notes that the report of the Ontario Royal Commission also cautions against the conversion to substitute fibres: “We conclude that the health hazards caused by exposure to asbestos fibres depend very much upon the dimensions of the fibres, and that long, thin, respirable fibres are of primary concern. This leads us to be extremely cautious about concluding from the evidence (on) substitute fibres (that) they will not present health hazards in the future. […] [W]e believe it would be risky to allow the exposure of workers to respirable fibres longer than 5 microns, with small diameters, of any material, if those fibres are likely to be very durable in the lungs” (p.359).

278 Canada notes that the United States Court of Appeal in the matter of the EPA, (see para.3.126 above) rightly emphasized that “Eager to douse the dangers of asbestos, the agency [EPA] inadvertently actually may increase the risk of injury”, Corrosion Proof Fittings v. Environmental Protection Agency, (5th Circuit 1991), p.35.

279 Substitutes – Executive Summary, pp.15-17.

280 Convention concerning Safety in the Use of Asbestos (Convention 162), ILO.


284 Fibres minérales artificielles et amiante (summary); INRS, Documents pour le médecin du travail n°69.


286 Avis concernant l’amiante chrysotile et les produits de substitution envisageables, Scientific Committee on Toxicity, Ecotoxicity and the Environment (CSTEE), 15 September 1998.

287 The EC point out that polyvinyl alcohol fibres (PVA) have been used since 1936; para-aramid fibres have been on the market for about thirty years.


289 Id., p.5.

290 Source: COPACEL and the French customs.


292 Statement for the Health and Safety Executive (HSE) on Carcinogenic Risks of Three Chrysotile Substitutes, Committee on the Carcinogenicity of Chemicals in Food, Consumer Products and the Environment (COC), July 1998. Canada points out that the EC use expressions such as “for a very long time”, “long period” and “use for many years”, but these do not appear in the CSTEE text.

293 Id.


296 Avis concernant l’amiante chrysotile et les produits de substitution envisageables, Scientific Committee on Toxicity, Ecotoxicity and the Environment (CSTEE), 15 September 1998.

297 Id.


300 See Section IV below.


302 Id.

303 Fibres minérales artificielles et amiante (Résumé), Report of the Groupe scientifique pour la surveillance des atmosphères de travail (G2SAT), INRS – DMT No.69.


306 Id.

307 Fibres minérales artificielles et amiante (Résumé), Report of the Groupe scientifique pour la surveillance des atmosphères de travail (G2SAT), INRS – DMT No.69.

308 Id.


315 Canada notes that the Decree “in no way makes it mandatory to use substitute products”, but it does make it unavoidable, as the EC note.


318 The EC note that, in 1984 and 1985, Canada produced 20 million tonnes of cellulose (source: Alternatives to Asbestos, the Pros and Cons, Society of Chemical Industry, United Kingdom, 1989).


320 Avis concernant l’amiante chrysotile et les produits de substitution envisageables, Scientific Committee on Toxicity, Ecotoxicity and the Environment (CSTEE), 15 September 1998.


322 Fibres minérales artificielles et amiante (Résumé), Report of the Groupe scientifique pour la surveillance des atmosphères de travail (G2SAT), INRS – DMT No.69. INSERM, Substitutes – Executive Summary.


324 INSERM report, p. 239.

325 Royal Society of Canada, A Review of the INSERM Report on the Health Effects of Exposure to Asbestos, Ottawa, 1996 [hereinafter “RSC”]. Canada notes that the RSC formed an international committee of independent experts to review the INSERM report. This committee, which was chaired by Dr. F. Kenneth Fare, Ph.D. (Montreal) was composed of the Canadian Michael Brauer, Sc.D. (Harvard, Berkeley), the American Kenny S. Crump, Ph.D. (Montana State), the Briton John M.G. Davis, MA, Sc.D. (Cambridge), FRC. Path., and Enzo Merler, MD (Padua).


328 Ph.D. Camus, M., L’amiante et les risques pour la santé, April 1999.

329 Académie nationale de médecine (Étienne Fournier), Amiante et protection de la population exposée à l’inhalation de fibres d’amiante dans les bâtiments publics et privés, Paris, 1996.

330 INSERM report, p. 428.

331 Comments reported in Le Monde, 18 October 1997, p.13.

332 Canada notes that, two months before the appearance of the INSERM report, the Académie nationale de médecine considered it crucial that scientific reports on the risks associated with asbestos should be based on real data, not hypothetical data. According to the Académie: “A cautious attitude should undoubtedly always be adopted with regard to the purity of the air inhaled, but speaking or writing on asbestos in general without taking into account the doses or quantities actually present in the air inhaled is at the best a sign of ignorance and at worst an attempt at blackmail. We therefore refer to quantities measured.” (WTO translation) (Académie nationale de médecine (Étienne Fournier), Amiante et protection de la population exposée à l’inhalation de fibres d’amiante dans les bâtiments publics et privés, Paris, 1996, p. 10).
INSERM report, p. 232.

Id., p. 235, see also pp. 234, 237, 238.


Id., p. 4 of the English text.

Id., pp.4 and 12 of the English text. Canada indicates that Dr. Gibbs is of the same opinion as the RSC: “In interpreting the risks provided in the report, it is important to recognize that risk estimates given in the report are, in general, not based on actual or estimated exposures but on the current or proposed ‘legal’ limits in France.” (Gibbs, p.2).

INSERM report, p. 232. Canada notes that this level is the maximum legal limit in France for indoor passive exposure. For a critique of this approach, see Camus, M., L’amiante et les risques pour la santé, April 1999, p.8.

INSERM report, pp.233-38.

Id., pp.428 and 429.

InsERM also emphasizes that “this information is practically non-existent in our country and it should now be collected so that we can have an overall estimate of the number of persons concerned by the various situations of exposure to asbestos and the risks they are running because of this”. (WTO translation)

De Vos Irvine, H. et al., Asbestos in Lung Cancer in the West of Scotland (1993) 306 British Medical Journal, 1503. This study is included in Wilkinson, P. et al., Is Lung Cancer Associated with Asbestos Exposure When There are no Small Opacities on the Chest Radiograph? (1995) 345 Lancet 1074. INSERM applied the rate of 5.7 per cent of cancers linked to asbestos in Glasgow to the total 21,617 cancer cases in France in 1996 to arrive at 1,200 deaths by cancer linked to asbestos (INSERM report, p.250). According to the authors of the study, De Vos Irvine, H. et al.: “A comparison of international cancer registry data shows that the west of Scotland experiences one of the highest incidence rates of lung cancer in the world.” (p.503). The RSC makes a point of the lack of scientific rigour on the part of INSERM with regard to this conclusion. It is of the opinion that “INSERM’S estimates of asbestos-related deaths in France in 1996 [were] based on an estimate from the literature specific for Great Britain without critical analysis of the methodology, or establishment that the estimate was applicable to France (RSC, p.3 of the English text; See also Gibbs, p.13).

INSERM report, p.409. In Canada’s view, INSERM’S error can clearly be seen in the following example. According to INSERM’S conclusions, there would be 543.5 cancers in a population of 10,000 people subject to an occupational exposure of 0.1 f/ml for 40 hours a week over 45 years. However, only 21.5 of these cancers would be linked to exposure to asbestos (INSERM report, p.234). Thus, according to INSERM’S data, a little less than 4 per cent of the cancer deaths (21.5/543.5 = 4 per cent) of a population highly exposed to asbestos (0.1 f/ml for 40 hours a week over 45 years) would be linked to asbestos. Why then did INSERM apply the 5.7 per cent from the Glasgow study to the entire French population, which was not subject to exposure of 0.1 f/ml for 40 hours a week over 45 years? Canada notes that, even if one accepts the 4 per cent rate at which INSERM arrived, the number of cancer deaths in 1996 attributable to past exposure to asbestos would be 864 (4 per cent x 21,617), not 1,200.


Id., p. 9.

InsERM report, p.4. Dunnigan, p. 4.

INSERM report, p.412. The Académie nationale de médecine confirms that amphiboles are “currently considered to be the most dangerous” and that chrysotile is “considered as not very dangerous because it is broken down spontaneously in the human organism […] Chrysotile is a form of asbestos that has only caused mesothelioma in cases of massive and
prolonged exposure. This can be explained by its solubility in the organism.” (WTO translation).


352 Gibbs, p. 6. The Royal Society of Canada adds that “the differences between chrysotile and amphiboles may have been underestimated by INSERM, particularly for mesothelioma”. (RSC, p.6 of the English text).

353 Gibbs, p. 8.

354 RSC, p.12 of the English text.

355 Gibbs, p.9. See also Camus, M., L’amiante et les risques pour la santé, April 1999, p.8.


357 RSC, pp.6 and 9 of the English text. According to J. Peto, expressing his opinion of the comparison between present and past exposure rates: “It might therefore be thought that none of the data that have been collected permit any truly scientific prediction of the likely effects of the limits to exposure that have now been set. This may indeed be the case.” (Doll, R. and Peto, J., Asbestos: Effects on Health of Exposure to Asbestos, Her Majesty’s Stationery Office, U.K., 1985, p. 44).

358 INSERM report, p.420. See also RSC, p.9 of the English text.

359 As emphasized by the RSC on p.9 of the English text.

360 According to INSERM: “At present there are no solid direct epidemiological data that enable an assessment to be made of the health effects linked to urban passive environmental or building exposures. […] Even without recalling that such studies would have little chance of directly noting a health effect, if it is low, it has to be recognized that we do not have any reliable epidemiological data.” (WTO translation) (INSERM report, pp.404 and 405).


362 INSERM report, pp.424 and 400.

363 See in particular the French study of the Laboratoire d’hygiène et de contrôle des fibres minérales: Baujon et Authier, Détermination des concentrations de fibres d’amiante dans l’atmosphère lors de la pose sur chantier de plaques ondulées et d’ardoises en amiante-ciment, Laboratoire d’hygiène et de contrôle des fibres minérales, Paris, July 1993. Canada notes that, according to this study, the use of self-boring screws during the process of installing corrugated panels gives rise to an emission of 0.022 f/ml. Moreover, the installation of roof tiles can give rise to emissions of 0.007 f/ml, depending on the method used. These peak exposure levels, which are well below the exposure standards in effect in France, can be further reduced by dampening the materials. Furthermore, workers can be protected from inhaling the fibres simply by wearing a mask.

364 Gibbs, p.10.

365 Académie nationale de Médecine (Étienne Fournier), Amiante et protection de la population exposée à l’inhalation de fibres d’amiante dans les bâtiments publics et privés, Bulletin de l’Académie nationale de médecine, Vol. 180, no. 4, p. 4. Canada notes that Dr. Jacques Dunnigan characterizes the limited usefulness of the INSERM report as follows: “If it was necessary to ban certain uses of chrysotile (for example, friable materials), this is not the case for other uses of asbestos, in particular chrysotile-cement, in which the fibres are bound in such a way that there is practically no risk of emissions at unacceptable levels, by using known and proven control techniques. We believe that monitoring of hygiene and observance of the standard are a better guarantee of protection than blind use of substitute fibres whose long-term effects are not sufficiently known, as is in fact acknowledged by the authors of the INSERM report.” (WTO translation) (Dunnigan, p.7).

366 Id., pp.404 and 405.

367 Id., 226.

368 Id., pp.234, 235, 237 and 238.

369 Id., p.376.
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371 Id., p. 239 and 414.
372 Id., p. 232.
373 Dunnigan, p.7. Canada notes that INSERM acknowledges that “the level of occupational exposure envisaged is 10 to several hundred times lower than that existing in the cohorts on which the risk models were elaborated”. (WTO translation) (INSERM report, p.233).
374 Dunnigan, p.7.
375 Gibbs, p.10.
376 RSC, p.6 of the English text. The RSC also points to the potential for error associated with the integration of different measurement techniques (p.6 of the English text). Regarding the evaluation of measurement techniques, see Camus, M., L’amiante et les risques pour la santé, April 1999, p.4.
377 INSERM report, pp.376 and 428.
378 See paras.3.174 et seq. above.
379 INSERM report entitled Effets sur la santé des fibres de substitution à l’amiante, published in November 1999
381 Document on the link between tobacco and asbestos.
387 Chrysotile Asbestos evaluated by Health Experts, Press Release 51 of 26 July 1996, WHO.
390 See para.3.54 above.
391 The EC note that the average concentration of asbestos fibres in the air inhaled by workers is measured over eight hours.
395 Id.
396 Article 4 of the Decree. This incorporates and refers back to Decree No.96-97 of 7 February 1996 concerning the protection of the population against the health risks associated with exposure to asbestos in buildings.
397 Order of 24 December 1996 on the declaration form for exemptions from the ban on asbestos.
398 Article 4 of the Decree. This incorporates and refers back to Decree No.88-466 of 28 April 1998 concerning products containing asbestos.
399 Notification G/TBT/Notif. 97.55, 21 February 1997.
401 Opening statement by the representative of the European Communities at the consultation held in connection with this case, Geneva, 8 July 1998 (document submitted to the Panel by Canada).
403 Larousse French dictionary.
404 Id.
407 Canada notes that the EC try to show that the Decree does not define the characteristics of a product because Canada made a mistake by citing the definition of the adjective rather than the noun “characteristic”. Having made the necessary correction, this in no way alters the sense of Canada’s argument.
408 Article 4 of the Decree. This incorporates and refers back to Decree No.96-97 of 7 February 1996 concerning the protection of the population against the health risks associated with exposure to asbestos in buildings.
409 Article 4 of the Decree. This incorporates and refers back to Decree No.88-466 of 28 April 1988 concerning products containing asbestos.
410 Notification G/TBT/Notif.97.55, 21 February 1997.
412 Opening statement by the representative of the European Communities at the consultations held in connection with this case, Geneva, 8 July 1998 (document submitted to the Panel by Canada).
413 The EC note that, on its Web site www.wto.org/eol/e/wto03/wto.3_5.htm, the WTO describes a “technical regulation” as follows: “Technical regulations and standards set out specific characteristics of a product – such as its size, shape, design, functions and performance, or the way it is labelled or packaged before it is put on sale. In certain cases, the way a product is produced can affect these characteristics, and it may then prove more appropriate to draft technical regulations and standards in terms of a product’s process and production methods rather than its characteristics per se.”
415 Canada refers in general to the arguments it puts forward below in relation to Article III:4 of the GATT.
416 Tariff Classification 6811. Source: Eurostat, CD-ROM.
418 United States – Section 337 of the Tariff Act of 1930, adopted on 7 November 1989, BISD 365/345, para.5.11
419 See Section III.B.7 above.
420 INSERM report, p.400 and pp. 419 and 420.
421 European Commission (G. Lohan, DG III), European justification of Decree No.96-1133 to the Canadian authorities (15 April 1997) following French Notification G/TBT/Notif.97.55, p.2. Canada notes that the French Académie nationale de médecine also recognizes that “recent important epidemiological studies, going back for 20 years or more, show that the carcinogenic or co-carcinogenic effect of asbestos (lung cancer) is no longer detected among large protected industrial populations working in the conditions prescribed by the regulatory texts. These conditions are laid down in practical leaflets published by the ministries concerned, Labour and Safety, for use by persons in occupations exposed to at least 1f/ml and in recent decrees, in 1996, concern persons working in buildings containing asbestos”. (WTO translation) Académie nationale de médecine (Étienne Fournier), Amiante et protection de la population exposée à l’inhalation de fibres d’amiante dans les bâtiments

European Commission (G. Lohan, DG III), European justification of Decree No.96-1133 to the Canadian authorities (15 April 1997) following French Notification G/TBT/Notif.97.55, p.2.


Canada notes that the French Académie nationale de médecine wrote “A measure viewed so radically by public opinion as the total ban on asbestos does not in any way change the situation in a country. It does not resolve any of the problems raised by this material and may even mean neglecting the essential measures that should be taken immediately”, (WTO translation), Académie nationale de médecine (Étienne Fournier), Amiante et protection de la population exposée à l’inhalation de fibres d’amiante dans les bâtiments publics et privés, Bulletin de l’Académie nationale de médecine, Vol.180, no.4, 1996, p.8. Also see pp.5 and 6 of the same report: “recent important epidemiological studies, going back for 20 years or more, show that the carcinogenic or co-carcinogenic effect of asbestos (lung cancer) is no longer detected among large protected industrial populations working in the conditions prescribed by the regulatory texts. These conditions are laid down in practical leaflets published by the ministries concerned, Labour and Safety, for use by persons in occupations exposed to at least 1f/ml and in recent decrees, in 1996, concern persons working in buildings containing asbestos”. (WTO translation).


See Section IV below.


See the comments of Sir Leon Brittan, para.3.475 below.


United States – Restrictions on Imports of Tuna, report circulated on 3 September 1991, not adopted, BISD 395/155, para.5.27.

The EC note that this principle was underlined by the Appellate Body within the context of the SPS Agreement, (EC Measures Concerning Meat and Meat Products (Hormones), Report of the Appellate Body, WT/DS26/DS48/AB/R, adopted on 13 February 1998, in particular para.109.


See, in particular, Thailand – Cigarettes, adopted on 7 November 1990, BISD 375/200, para.75. The EC also note that this approach is being followed in practice. The document entitled “Streamlined mechanism for reconciling the interests of contracting parties in the event of trade-damaging acts” provides, in particular, that “A measure taken by an importing contracting party should not be any more severe … than necessary to protect the human, animal or plant life or health involved, as provided in Article XX(b)” Document C/M/236, BISD 365/67, point 1.

Gilg Soit Ilg, A., et al., Estimation of the Past and Future Burden of Mortality from Mesothe-

Peto, J. et al., Continuing Increase in Mesothe-
442 The EC indicate that equally relevant, by analogy, is the position of the Appellate Body in Japan – Measures Affecting Agricultural Products, adopted on 19 March 1999 (WT/DS76/AB/R, para.126), according to which the complaining Member must show that the contested measure is more trade-restrictive than necessary to fulfil a legitimate objective taking into account the risk which non-fulfilment would create.
443 See Section V, reply by Dr. Henderson to question 1(b) from the Panel. [Note: in the context of this dispute, the Panel consulted four independent experts in accordance with Article 13 of the Understanding. The written replies by the experts to the questions from the Panel can be found in Section V of this report. Annex VI contains a transcription of the meeting held by the Panel on 17 January 2000 with the experts and the parties].
444 Id.
446 Id., p.144.
447 See Section V, replies by Dr. Musk and Dr. de Klerk to question 3 from the Panel.
448 Id., reply by Dr. Henderson to question 1(e) from the Panel.
449 INRS, Rapport du Groupe scientifique pour la surveillance des atmosphères de travail (G2SAT), 1997, p.47.
451 Id., p.v.
452 Id., p.411.
453 Id., pp.178-181.
454 Id., p.398.
455 See United States Department of Labor (OSHA), Synthetic Mineral Fibers: Hazard Description, on line: OSHA <http://www.osha.gov/oshinfo/priorities/synthetic.html> (access date: 22 June 1999). The OSHA Report states that: “Several epidemiologic studies have demonstrated statistically significant elevations in the risk of lung cancer and other respiratory system cancers among workers employed in fibrous glass and mineral wool manufacturing facilities.”
460 Thailand – Cigarettes, BISD 37S/200, report adopted on 7 September 1990, para.75.
461 See Canada’s arguments in Section III.B.7 of this report.
462 INSERM report, p.428.
463 See Section V, reply by Dr. Henderson to question 1(b) from the Panel.
464 INSERM Report: “the vast majority of these deaths can indisputably be explained by occupational or para-occupational exposure” (WTO translation) (pp.419 and 420).
465 See Section V, reply by Dr. Henderson to question 1(d) from the Panel.
466 See Dr. Henderson’s comments, Section V.C.1(i), citing IPCS Environmental Health Criteria (203) on Chrysotile, WHO, Geneva, 1998. See also the EC’s comments on the replies by the experts, Section V.D.2.
467 See in particular paras.3.155 et seq.
468 European Commission (G. Lohan, DG III), European justification of Decree No.96-1133 to the Canadian authorities (15 April 1997) following French Notification G/TBT/Notif.97.55.
Canada notes that, in the area of public works, chrysotile-cement pipes are used in the construction of piping systems. These pipes are machined into varied lengths that can meet the specifications of buyers and are equipped with couplers that can ensure the linkage between two pipes without the need for cutting or grinding. In the event that the cutting or grinding of chrysotile-cement pipes nonetheless turns out to be inevitable, there are simple techniques, described by standard ISO-7337 that can guarantee the safety of workers on sites. Standard ISO-7337 was designed to be applied at the time of the cutting or grinding of any hard chrysotile-cement product. For example, the cutting of plates or tiles for roof covering, is not a source of emission if the simple techniques of standard ISO-7337 are followed. These techniques consist, in particular, of the use of chains that break the pipes by the effect of pressure, low-speed saws and saws equipped with a vacuum dust extractor, as well as the moistening of the materials prior to any action. The cutting or grinding of a cement pipe, whether or not it contains chrysotile, can also release silica in the air. The International Agency for Research on Cancer (IARC) rates silica among group 1 carcinogens (proven for humans), like asbestos. A worker who cuts any cement pipe therefore has an interest in following the instructions recommended in standard ISO-7337. See also the studies Baujon, Authier & Thomazo, Détermination de concentrations de fibres d’amiante dans l’atmosphère au voisinage de construction en amiante ciment, (October 1993), and Détermination de concentrations de fibres d’amiante dans l’atmosphère lors de la pose sur chantier de plaques ondulées et d’ardoises en amiante-ciment, (July 1993), Laboratoire d’hygiène et de contrôle des fibres minérales, Paris.


Id., Article 12.

Article 10 of Convention 162 reads as follows:
“Where necessary to protect the health of workers and technically practicable, national laws or regulations shall provide for one or more of the following measures:
(a) replacement of asbestos or of certain types of asbestos or products containing asbestos by other materials or products or the use of alternative technology, scientifically evaluated by the competent authority as harmless or less harmful, whenever this is possible;
(b) total or partial prohibition of the use of asbestos or of certain types of asbestos or products containing asbestos in certain work processes.”

Paragraph 12 of Recommendation 172 reads as follows:
“12. (1) The competent authority, wherever necessary for the protection of the workers, should require the replacement of asbestos by substitute materials, wherever
possible. (2) Before being accepted for use in any process, all potential substitute materials should be thoroughly evaluated for their possible harmful effects on health. The health of workers exposed to such materials should be continuously supervised, if judged necessary."

487 European Commission, Directorate General III (Industry), Office A, Industrial Policy to the Canadian authorities as further information for Notification G/TBT/Notif.97.55 made by France to the WTO Committee on Technical Barriers to Trade on 21 February 1997.


489 Id., para.163.


494 See Canada’s arguments in Section III.B, as well as the comments on the replies by the experts, especially those concerning question 3 from the Panel.

495 See Section III.B.6 above.

496 Glass fibre, cellulose fibre, para-aramid fibre, and PVA fibre.

497 Opinion on a study commissioned by Directorate General III (Industry) of the European Commission on Recent Assessments of the Hazards and Risks posed by Asbestos and Substitute Fibres, and Recent Regulation on Fibres World-Wide (Environmental Resources Management, Oxford (opinion expressed on 9 February 1998).


501 Canada’s comments on the replies by the experts to question 6(b) from the Panel.

502 See Annex II to this report, section B.6, para.344.


507 See, for example Canada – Importation, Distribution and Sale of Certain Alcoholic Drinks by Provincial Marketing Agencies, BISD 395/155, paras.5.8-5.16.


510 See, for example, the judgement of the Court of Justice of the European Communities, 11 June 1992, cases C-149/91 and C-150/91, Sanders Adour/Directeur des Services Fiscaux des Pyrénées-Atlantiques, Rep. 1992, p.I-3899, and in particular paras.15 and 19.

511 Korea – Taxes on Alcoholic Beverages, adopted on 17 February 1999, WT/DS57/DS84/R, report of the Panel, in particular point 10.81.

512 In this connection, the EC note that the non-cumulative application of Articles 12 and 95
of the EC Treaty (see above) has been forcefully recalled by the European Court of Justice, which has stated that:

“It is settled law … that the provisions relating to charges having equivalent effect and those relating to discriminatory internal taxation cannot be applied together, so that under the system of the Treaty the same imposition cannot belong to both categories at the same time.”


516 Working Party Report Border Tax Adjustments, BISD 18S/97, para. 18: “... the interpretation of the term should be examined on a case-by-case basis. This would allow a fair assessment in each case of the different elements that constitute a ‘similar’ product. Some criteria were suggested for determining, on a case-by-case basis, whether a product is ‘similar’: the product’s end-uses in a given market, consumers’ tastes and habits, which change from country to country, the product’s properties, nature and quality.” This Report was adopted by the contracting parties in 1970 and has been taken up many times since, notably by the Appellate Body in Japan – Taxes on Alcoholic Beverages, adopted on 1 November 1996, WT/DS8/AB/R; WT/DS10/AB/R; WT/DS11/AB/R, pp.18 et seq.


520 Canada notes that, on the contrary, in the Report Japan – Customs Duties, Taxes and Labelling Practices on Imported Wines and Alcoholic Beverages, it is recalled that products may be deemed like products according to the criterion of end-use only: “Past GATT practice has clearly established that ‘like’ products in terms of Article III:2 are not confined to identical products but cover also other products, for instance if they serve substantially identical end-uses ...” Report adopted on 10 November 1987, BISD 34S/83. See also United States – Taxes on Petroleum and Certain Imported Substances, adopted on 17 June 1987, BISD 34S/136, para.5.1.1.

521 Japan – Customs Duties, Taxes and Labelling Practices on Imported Wines and Alcoholic Beverages, adopted on 10 November 1987, BISD 34S/83, para.5.6: “The Panel agreed in this respect with the arguments submitted to it not only by the European Communities but also by other important producing countries of wines and distilled spirits that gin, vodka, whisky, grape brandy, other fruit brandy; certain ‘classic’ liqueurs, still wine and sparkling wine, respectively, were recognized not only by governments for purposes of tariff and statistical nomenclature, but also by consumers to constitute ‘each in its end-use... a well defined and single product intended for drinking.’” See also Spain – Tariff Treatment of Unroasted Coffee, adopted on 11 June 1981, BISD 285/102, p.112, para.4.7.

522 Canada notes that this interpretation is consistent with the precedent of the GATT 1947, in particular the aforementioned Panel Report on Japan – Customs Duties, Taxes and Labelling Practices on Imported Wines and Alcoholic Beverages. It was clearly established therein that “like” products under Article III:2 are not limited to identical products [...]. (para. 5.5(d)). The Panel judged in this case, that gin, vodka, whisky, brandy, wine and sparkling wine constituted like products for the purposes of Article III.

523 For the purposes of Canada’s argument with regard to Article III:4 of the GATT and Article 2.1 of the TBT Agreement, the term “fibro-cement” means a mix of cement to which PVA, cellulose fibre or fibreglass is added. The term fibro-cement includes “glass cement”. Canada notes that only four products made of chrysotile fibre have no equivalent and are therefore exempt from the ban until 2000 or 2002.

524 See Japan – Customs Duties, Taxes and Labelling Practices on Imported Wines and Alcoholic Beverages, adopted on 10 November 1987, BISD 34S/83, para.5.6: “The Panel agreed in this respect with the arguments submitted to it not only by the European Communities but also by other important producing countries of wines and distilled spirits that gin, vodka, whisky, grape brandy, other fruit brandy, certain ‘classic’ liqueurs, still wine and sparkling wine...”
wine, respectively, were recognized not only by governments for purposes of tariff and statistical nomenclature, but also by consumers to constitute ‘each in its end-use... a well defined and single product intended for drinking.’"


528 Id., p.23.

529 Article 2 of the Decree.

530 World Customs Organization, Category 68.11, which states that: “This heading covers hardened articles consisting essentially of an intimate mixture of fibres (for example, asbestos, cellulose and other vegetable fibres, synthetic polymer, glass or metallic fibres) and cement or other hydraulic binders, the fibres acting as strengthening agents. These articles may also contain asphalt, tar, etc. These products are generally manufactured by pressing together thin layers of a mixture of fibres, cement and water or by moulding (possibly under pressure), by pressing or by extruding. The heading includes all sizes and thicknesses, obtained as described above, and also articles made by cutting these sheets or by pressing, moulding or bending them before they have set, e.g. roofing, facing or partition sheets and tiles; sheets for making furniture; window sills; sign-plates; letters and numbers; barrier bars, corrugated sheets; reservoirs, troughs, basins, sinks; tubing joints; packing washers and joints; panels imitating carving; ridge tiles, gutters, window frames; flower-pots; ventilation or other tubing, cable conduits; chimney cowls, etc. All these articles may be coloured in the mass, varnished, printed, enamelled, decorated, drilled, filed, planed, smoothed, polished or otherwise worked; they may also be reinforced with metal, etc.”


532 Explanatory notes of the World Customs Organization, Harmonized System, note under heading HS 25.24 “Asbestos”.


538 Id.

Source: World Customs Organization.


Canada’s oral submission, 1 June 1999, para.290, See also para. 3.439 above.


Id., para.4.3.

Measures Affecting Alcoholic and Malt Beverages, adopted on 19 June 1992, BISD 395/206, paras.5.73-5.75.


See paras. 3.442-3.449 above.

See para. 3.429 above.

See Section V of this report, the comments by Canada on the replies by the experts to question 6 of the Panel.


Canada notes that the principle of the application of Article XI:1 in such circumstances has been established by several Panels under the GATT 1947, notably in United States – Manufacturing Clause, BISD 315/74, adopted on 15/16 May 1984, para.34; and in Japan – Trade in Semi-Conductors, BISD 355/116, adopted on 4 May 1988, paras.102 et seq.; and under the GATT 1994, in United States – Shrimps, Report of the Panel, WT/DS58/R, adopted on 6 November 1998, paras.7.11-7.17.

See the EC’s arguments in paras.3.395-3.400 and 3.403-3.407.

See paras.3.467 and 3.468 above.

See footnote 565 above.

See para.3.398 above.

Id.


Id., para.4.24.

Id., para.4.26.

See the EC’s arguments in paras.3.395-3.400 and 3.403-3.407 above.


656
583 Id.
584 See also Canada – Administration of the Foreign Investment Review Act, adopted on 7 February 1984, BISD 365/140, para.5.20 and United States – Section 337 of the Tariff Act of 1930, adopted on 7 November 1989, BISD 365/345, para.5.27.
590 Thailand – Cigarettes, adopted on 7 November 1990, BISD 375/200, para.73.
591 United States – Restrictions on Imports of Tuna, circulated on 3 September 1991, not adopted, BISD 395/155, para.5.27.
592 The EC note that the Panel report on United States – Restrictions on Imports of Tuna refers to the Panel report on Thailand – Cigarettes, adopted on 7 November 1990, BISD 375/200, paras.73 and 74.
594 Thailand – Cigarettes, adopted on 7 November 1990, BISD 375/200, para.73.
595 INSERM report, p.401.
596 Id., p.388.
598 Canada notes that the Decree is not aimed at the use of amphiboles and the manufacture of friable asbestos materials as these uses had already been banned in France:
599 European Commission (G. Lohan, DG III), European Justification of Decree No.96-1133 to the Canadian authorities (15 April 1997) following the French notification G/TBT/Notif.97.55, p.2.
600 See INSERM report, p.182.
602 Id.
603 See Académie nationale de médecine (Etienne Fournier), Amiant et protection de la population exposée à l’inhalation de fibres d’amianté dans les bâtiments publics et privés, Bulletin de l’Académie nationale de médecine, vol.180, no.4, April 1996.
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Uruguay Round of Multilateral Trade Negotiations, Legal Instruments restating the results of the Uruguay Round of Multilateral Trade Negotiations effected in Marrakesh on 15 April 1994, volume 19, List LXX – European Communities. After the 1996 binding, the numbers of the tariff items in this case, reproduced on the list CXL – European Communities are: 2524.00.30 (initial negotiating right), 2524.00.80 (initial negotiating right), 6811.10.00, 6811.20.00, 6812.10.00, 6812.20.00, 6812.30.00, 6812.40.00, 6812.50.00, 6812.60.00, 6812.70.00, 6812.90.10, 6812.90.90, 6813.10.10, 6813.10.90, 6813.90.10, 6813.90.90.


Note DPPR/SDPD/BGTD/LT/LT No. 97-320 of 12 March 1997, Section III.

Canada notes that Article 26.1 of the Understanding provides that: “Where the provisions of Article XXIII:1(b) of the GATT 1994 are applicable to a covered agreement, a panel… may only make rulings and recommendations where a party to the dispute considers that any benefit accruing to it directly or indirectly under the relevant covered agreement is being nullified or impaired or the attainment of any objective of that Agreement is being impeded as a result of the application by a Member of any measure, whether or not it conflicts with the provisions of that Agreement.” The provisions of Article XXIII:1(b) apply to WTO Agreements (because the General Agreement is an integral part of the WTO Agreements under Article II.2 of the WTO Agreement) and to the General Agreement.


Id., para.10.38.
The EC note that Article XX states that, “Subject to the requirement that such measures are not applied in a manner which would constitute a means of arbitrary or unjustifiable discrimination between countries where the same conditions prevail, or a disguised restriction on international trade, nothing in this Agreement shall be construed to prevent the adoption or enforcement by any contracting party of measures... (b) necessary to protect human, animal or plant life or health.”


Id., para.10.82.


Id., para.10.82.


Id., para.10.82.


Id., para.10.82.


Id., para.10.82.


IV. ARGUMENTS PRESENTED BY THIRD PARTIES

A. Brazil

1. Introduction

4.1 Brazil explains that the proceeding challenges the WTO-consistency of France’s 1 January 1997 Decree, which bans the manufacture, processing, sale and possession for sale, importation, exportation, domestic marketing, offer and transfer of all varieties of asbestos fibres and products containing them (the Decree or the ban).1 The ban has four narrow exceptions that apply where no substitutes exist for chrysotile products. The substitute products that do exist generally are more expensive than chrysotile products. Thus, the ban clearly operates to create a commercial advantage for substitute products. According to Brazil, a ban is the most trade restrictive of measures. Therefore, the justification for any ban must be subject to the strictest scrutiny, especially as applied to a developing country such as Brazil. The ban has ended Brazilian exports of uncontaminated chrysotile to France. In 1994 and 1995, France imported from Brazil 1,100 and 1,500 metric tonnes of uncontaminated chrysotile, respectively. Since the ban took effect in 1997, France has not imported any chrysotile from Brazil.

4.2 According to Brazil, the importance of this proceeding extends far beyond the French ban – the proceeding is a test case. Will other WTO Members be allowed to ban products of developing countries that can be safely used with appropriate, tested precautions simply to appease the public? Modern economies use hundreds of products that present health risks if they are misused, but that present no risks if they are used properly. Uncontaminated chrysotile is one of them; if properly used, uncontaminated chrysotile presents no health risk. Similar products include organic fibres, man-made fibres, benzene, mercury, ammonia, nearly all forms of pesticide, etc. Societies regulate these products to ensure they are used safely so as to protect the health of workers handling them directly and of the general population which is exposed to them indirectly. The same treatment is appropriate for uncontaminated chrysotile. Uncontaminated chrysotile—the only asbestos fibre Brazil mines and exports – is the safest by far of all asbestos fibres. In particular, it is much safer than amphibole, the asbestos responsible for current health problems from past exposure. All of the asbestos that Brazil mines, produces and exports is uncontaminated chrysotile. For this reason, Brazil’s chrysotile products are among the safest in the world. The medical explanation for these facts is set forth in detail in a recent bio-persistence study by Dr. David S. Bernstein, an expert in fibre toxicology (indeed, the EC often seeks his expertise on this topic).2

4.3 Brazil asserts that the primary issue in this proceeding is not – as the EC would suggest - whether asbestos can be hazardous to human health. It can. Years of misuse and unsafe utilization of the most hazardous form of asbestos – amphibole - have caused significant damage to health. All countries, including Brazil, regret the harm to human health caused by decades of exposure earlier this century to amphibole produced and used worldwide. Brazil understands well the basis of the public outcry, experienced in many countries (including Brazil), that led the French Government to commission the INSERM Report3 (a study focusing on the health effects of earlier, unsafe uses of amphibole asbestos) and then to ban asbestos. France imposed the ban only one day after INSERM released its Report. The Report was commissioned and released to provide a scientific “cover” for a political decision that had already been taken. However, as a review of the INSERM Report demonstrates, the causes of asbestos-related health problems in France are past uses, especially in the spraying of brittle amphibole on to fireproof buildings and, until quite recently, warships (flocking). Given the long latency period between exposure to amphibole and the onset of any related diseases, workers who were victims of heavy exposure with virtually no protection 30 years ago are experiencing serious health problems today. The INSERM Report is based on analyses of these workers’ health. The INSERM Report does not focus on data from studies of modern uses of chrysotile. Moreover, in the Report, INSERM concedes that it was unable to produce “scientifically certain” conclusions, but could present only an “aid to understanding” based on “plausible, though uncertain, estimates.”4 Quite simply, the INSERM Report is an inadequate basis for the ban.
4.4 Brazil argues that it has a deep appreciation of the desire - indeed, the need - for the French Government to address public concern and protect public health. Brazil also understands the frustration of being unable to remedy or even mitigate the health consequences of past exposure from unsafe use of amphibole, and the frustration of being unable to take measures to remedy or decrease exposure from flocked amphibole asbestos that is already in French buildings (because disturbing flocking increases exposure). However, when France approved the WTO Agreement, it agreed not to restrict trade merely to appease domestic sentiment, no matter how strong. Brazil cannot accept France’s adoption of a politically motivated measure that will neither (i) make those already sick from asbestos exposure healthy; nor (ii) reduce risk to the healthy beyond existing levels of protection guaranteed by modern, controlled uses of chrysotile. As the European Commission recently stated:

[V]arious national organisations, including the Health and Safety Executive in the United Kingdom, have made very disturbing projections about the numbers of deaths which are likely to be attributable to asbestos over the next few decades. However, it is important to note that these figures relate to past exposures to mixed asbestos types, including the fibres which have already been banned. It would be wrong to use these statistics alone to justify a ban on the marketing and use of chrysotile because such a ban would not lead to a lower risk of exposure for workers to asbestos which is already in place, nor would it reduce the number of deaths which are occurring today as a result of past exposure to asbestos.5

4.5 Modern uses of asbestos are or should be limited to chrysotile, which most parties, including INSERM, agree is safer than other forms of asbestos. Moreover, modern uses are or should be confined to products in which the fibres are bonded in a finished product and, thus, cannot escape, e.g., asbestos-cement products.6 For these and other reasons, modern uses are quite safe; they involve exceedingly low levels of exposure (that often do not exceed even the “natural” levels in ambient air). Chrysotile is used in a very wide variety of products. It is used as a flame retardant, to strengthen friction materials (e.g., truck brakes) and to create cement pipes for carrying water that are far less subject to corrosion, cracking and breaking than traditional cement pipes. In most applications, chrysotile is used because it increases public safety; thus, using other, less-efficient products in its place often decreases public safety. Chrysotile is used primarily in truck brake pads, drum brakes and brake blocks to control heat build-up, thus maximizing friction and stopping power. It is the preferred product for this application. As one of the authors of an American Society of Mechanical Engineers (ASME) study commissioned by the EPA concluded:

(a) The “replacement/substitution of asbestos-based with non-asbestos brake linings will produce grave risks”; and

(b) “the expected increase of skid-related highway accidents and resultant traffic deaths would certainly be expected to overshadow any potential health-related benefits of fiber substitution.”7

4.6 Brazil pleads that chrysotile’s numerous public safety benefits - the many contributions it makes to societies around the world - not be ignored in this proceeding, as they were when France passed its ban. In Brazil’s view, the primary question in this proceeding is quite narrow - is a complete ban necessary to protect public health or can public health be ensured by regulating modern, controlled uses of chrysotile and chrysotile products? The answer arrived at by those countries in the Americas that have examined the issue closely, ranging south from Canada, to the United States, to Brazil, is that public health can be ensured by regulating modern controlled uses. France may, of course, take measures that are designed to, and actually do, protect its citizens. However, the ban does not meet even this very generous characterization of the general rule set forth in the WTO Agreement on Technical Barriers to Trade (the TBT Agreement). France must not be allowed to
impose a ban on imports and safe, modern uses of chrysotile as a response to public pressure. That the ban does not apply to man-made fibres produced in France, which the available scientific data show present greater risks when their use is not controlled and which have not been proven safer, confirms that the basis for the ban may be political and economic, but is not scientific or medical.

4.7 Brazil argues that in many respects, the French reaction is identical to that of the United States Environmental Protection Agency (the EPA) promulgated in 1989, when it banned asbestos under pressure from panicked U.S. public opinion. The EPA was unable to justify its ban scientifically to the United States Court of Appeals for the Fifth Circuit. After lengthy legal proceedings, the Fifth Circuit ordered the EPA to reverse its decision and to acknowledge publicly that modern products containing chrysotile enclosed in a matrix of cement or resin do not pose any detectable risk to public health.\(^8\) (Today, although amphiboles are prohibited in the United States, a number of products containing non-brittle chrysotile are permitted, including the products manufactured by Brazil and previously manufactured in France from Brazilian chrysotile.) Unfortunately, France has adopted a measure that unnecessarily and to no good effect impedes international trade.

4.8 Brazil makes the following claims regarding the ban: (1) the ban is inconsistent with Article 2.2 of the TBT Agreement because it creates unnecessary obstacles to trade and is more trade restrictive than necessary; (2) the ban is inconsistent with Article XI of the General Agreement on Tariffs and Trade 1994 (GATT 1994) because it is a quantitative restriction that is not excused by the exceptions in Article XI:2 or Article XX; (3) the ban is inconsistent with Article 2.8 of the TBT Agreement because it applies to asbestos but not to man-made fibres or other substitute products and thus operates as a technical regulation setting forth an unnecessary design or descriptive characteristic; (4) the ban is inconsistent with Article 2.4 of the TBT Agreement because international standards for producing and using chrysotile and chrysotile products exist and France should have used them; (5) the ban is inconsistent with Article III:4 of the GATT 1994 and Article 2.1 of the TBT Agreement (national treatment) because it does not apply to domestic man-made fibres and other substitute products, which are like products to chrysotile; and (6) the ban is inconsistent with Article I:1 of the GATT 1994 and Article 2.1 of the TBT Agreement (MFN) because, insofar as it bans imports of chrysotile and chrysotile products, but not imported like product substitutes, it improperly discriminates among imports.

2. **Factual Aspects**

4.9 Brazil concurs with practically all aspects of Canada’s presentation, agreeing (i) that the French ban was passed in response to public outcry in France over the deaths associated with the intensive exposure to amphibole that had taken place early on in the century; (ii) with the circumstances and risks of exposure presented by Canada. In particular, it agrees with the statement that exposure, even in asbestos product plants, has decreased significantly and that, apart from existing flocked amphibole, current exposure is limited, or could be limited, entirely to chrysotile: in contrast, past exposure and current exposure from past uses (e.g. flocking) included exposure to amphibole; (iii) that current levels of exposure to modern uses of chrysotile are not significant and are not associated with substantial health risks; (iv) with the fact that current controlled-use policies and standards which are accepted internationally are sufficient to ensure the health of chrysotile workers and others exposed to chrysotile and to guarantee their safety; and, (v) with Canada’s argument that the INSERM Report has many defects and that it was not the reason for France’s ban on modern, controlled uses of chrysotile and chrysotile products.

4.10 Brazil considers that a “battle of experts”, with one side presenting experts who support banning chrysotile and the other presenting experts who oppose banning chrysotile would be, in this case, both uninformative and unnecessary because the INSERM Report and the Synthesis, as a matter of law, not fact, cannot support the ban.\(^9\) This Report and the Synthesis have several defects that render them utterly incapable of supporting the ban.\(^10\) INSERM has not conducted original research, but merely based itself on existing studies and, furthermore, it has not examined all existing studies as it has deliberately
excluded those that have established a distinction between chrysotile and amphiboles. More specifically, the shortcomings of the INSERM Report include the following. First, the Report completely fails to examine the modern uses of chrysotile and chrysotile products and, thus, ignores the current state of the industry. Instead, it focuses on the health effects of exposure to amphiboles that took place in previous decades. INSERM concedes that it does not have “direct, certain scientific” data on the health risks associated with current levels of exposure to the modern uses of any form of asbestos, much less chrysotile. In short, INSERM does not examine current uses and exposure levels and does not distinguish among the different levels of risk associated with the different types of asbestos fibres (chrysotile, the only type produced and exported by Brazil, as well as used in it, is accepted as being the safest of asbestos fibres, even by INSERM itself).13

4.11 Second, the INSERM Report fails to examine the efficiency of the ways in which worker exposure has been reduced through the use of air filters in mines and plants and employing masks, laundry services, etc. Third, it does not even compare the risks of the past to the risks associated with man-made fibres and substitute products (such as ductile iron or polyvinyl chloride (PVC) pipes). By the time INSERM began to examine substitutes, the ban had already been in effect for 1.5 years and, in any case, INSERM issued only a synthesis and not a complete report on these substitutes. INSERM concedes in its Report that it lacked the data required to recommend the banning of chrysotile and only to allow its substitutes. INSERM emphasizes that because it is the structure (size and shape) of fibres that makes them toxic when inhaled, any substitute fibre must be viewed as dangerous to human health. Finally, INSERM concedes that, although the health data it applied to chrysotile are from past, massive and prolonged exposure to amphibole, the data being collected for substitutes is based on much lower levels of exposure, replicating modern conditions. Most telling is that INSERM states that toxicity levels for “asbestos” as a whole (and not merely for chrysotile) would yield similar results to those obtained for substitutes if similar testing conditions had been used.

4.12 Brazil further argues that INSERM uses a linear risk model to assume illogically and without any evidence that a threshold does not exist for safe exposure. France and INSERM are forced to commit this methodological error (the assumption) due to the fact that they had data from past prolonged exposure to amphiboles but not to current, much lower exposure to chrysotile. To justify the ban on the modern uses of chrysotile, France/INSERM had to assume that significant risk is present at all levels of exposure, even at those that are insignificant, out of political self-interest. INSERM adopted the linear risk model despite the fact that studies cited by the European Communities (hereinafter “EC”) themselves indicate that “bricoleurs” are not at risk. The study conducted by Iwatsubo et al. indicates that low, sporadic, intermittent and cumulative exposure of up to 0.5 fibres/ml-years does not present increased risk of mesothelioma. In commenting upon the results of an earlier study, the authors note that “no significant risk was observed for those whose exposure was intermittent”.

4.13 Brazil argues that a close examination of the INSERM Report reveals that: (i) prolonged exposure to amphiboles (its past uses) is associated with severe health problems (a proposition with which everyone agrees); (ii) substitute fibres have similar structures and, thus, when subject to scientific scrutiny, are expected to have similar health effects at similar levels of exposure; (iii) insufficient data exists on the health effects of current levels of exposure to chrysotile and substitute fibres, but the available data suggests that their health effects would be the same; and (iv) the Report does not purport to be as conclusive as France would have all believe; rather, to overcome (iii) above, INSERM extrapolated from the data used in (i), which as it itself concedes “does not produce scientifically certain knowledge, but only an aid to understanding the implications for risk management.” Brazil contends that the ban has been based on the irrelevant data described above. France employs the linear risk model as a tool to make data on past uses relevant to the imposition of the ban. However, INSERM researchers themselves recognize the limitations of this model and clearly state that it cannot produce “scientifically certain knowledge,” but can only serve as an “aid to understanding,” based on “plausible, though uncertain, estimates.” These “conclusions” do not support significant trade restrictions, much less the ban. Rather, they are merely a call for further research.
4.14 Brazil argues that recent research focusing on uncontaminated chrysotile demonstrates why it presents no health risks whatsoever. According to Dr. David Bernstein’s medical explanation, the serpentine (braided) structure of chrysotile leads it to unravel in the lungs (whereas the tubular structure of amphibole and substitute fibres does not allow them to unravel and is unchanging); and once unravelled, the smaller and thinner particles are more easily and rapidly enveloped by macrophages and/or expelled from the lungs. Dr. Bernstein’s research demonstrates that uncontaminated Brazilian chrysotile of less than 20 microns (the length that has been associated with pathogenicity for all fibres) is very quickly cleared. The clearance half-time is 1.3 days (and is 2.4 days for fibres of a length of 5-20 microns). He concludes that, once in the lung, chrysotile fibres defibrillate (or unravel), breaking down into shorter fibres. According to Dr. Bernstein, this result “is in stark contrast to amphibole asbestos where a portion of the fibres longer than 20 [microns] remains indefinitely or with synthetic mineral fibres where even very soluble fibres are removed by dissolution in the lung with half-times greater than this.” He concludes that uncontaminated chrysotile’s lack of bio-persistence suggests that it has “little if any toxicological effect.” However, it is of course a fact that if used improperly, uncontaminated chrysotile could be dangerous, but that would be the case for virtually all products in existence and not just chrysotile.

4.15 Brazil indicates that it mines, produces and exports only uncontaminated chrysotile and chrysotile products, and subjects mining, production and use to strict regulations. In 1990, it signed the ILO Convention and Recommendation Concerning Safety in the Use of Asbestos (Convention 162 and Recommendation 172). To ensure safety in the mining, manufacture and use of chrysotile and chrysotile products and to meet its ILO obligations, Brazil passed a primary law and decree on asbestos. In addition, the production and use of chrysotile and chrysotile products is governed by “national tripartite (government-industry-workers) agreements”. These set exposure limits, and processes of production and safety procedures to be used to guarantee worker safety. Finally, the Brazilian Asbestos Association (ABRA), a watchdog organization comprised of asbestos producers and sellers, further regulates the safety of, and trade in, chrysotile and chrysotile products.

4.16 Brazil explains that the ILO Convention and Recommendation are international standards that establish safety procedures for the handling of chrysotile and chrysotile products. They follow the ILO Code of Practice on Safety in the Use of Asbestos. The goal of the Code is to prevent the risks of exposure to asbestos and its harmful effects and to provide practical control procedures for its use. Convention 162 and Recommendation 172 recommend the controlled and safe use of asbestos. Their wording clearly indicates that the replacement of asbestos fibres should only take place when it is established that this is necessary to protect worker health and when replacement is technically feasible. The replacement of chrysotile asbestos fibres contained in modern materials or products (i.e. where it is sealed in a matrix and cannot be released into the environment) is not necessary since these products do not pose any detectable health risks. International standards, such as Convention 162 and Recommendation 172, recommend the regulation of asbestos on the basis of the type of asbestos fibre employed, the products in which certain fibres are included, and their planned use. Thus, Convention 162 and Recommendation 172 stipulate the prohibition of crocidolite and materials containing friable asbestos for flocking, but permit many uses of chrysotile, including those central to this dispute (asbestos-cement and friction products). They allow countries to prohibit other specific uses if national authorities deem this necessary for worker protection, but only on condition that substitute products be subjected to a thorough scientific examination of their health effects.

4.17 In 1995, Brazil passed Law No. 9055 to discipline the extraction, industrialization, use, commercialization and transportation of asbestos and of asbestos-containing products, as well as of natural and synthetic fibres of any source used for the same purpose. The Law (i) bans the processing and use of all types of asbestos, except chrysotile and chrysotile-containing products; (ii) bans the crushing and spraying (flocking) of all types of asbestos, including chrysotile, and of all substitute fibres; (iii) provides the framework...
for the tripartite agreements in that it sets deadlines for the government’s confiscation of the operating licences of companies that do not execute the tripartite agreements, establishes medical inspection requirements for workers, and sets exposure limits for those who work with chrysotile and substitute fibres subject to annual reduction. (In compliance with Article 2.4 of the TBT Agreement, the exposure limits are determined based, in part, on the recommendations of “international entities which are scientifically accredited”); (iv) prohibits miners and wholesalers from supplying chrysotile or substitute fibres to companies that do not comply with all provisions of the Law; (v) applies special restrictions to the use of chrysotile and substitutes in products currently considered to be the riskiest, such as textiles; (vi) calls for research into the health effects of chrysotile and substitute fibres and provides financing for the effort; and (vii) provides for prompt Department of Justice action against infractions.

4.18 Brazilian Decree No. 2350 implements the Law, and (i) requires that prior to marketing, all products containing chrysotile of imported or national origin bear a “seal of compliance to the Brazilian System of Certification”, and provides for the development of the certification system; (ii) requires research into and confirmation of the health effects of chrysotile and its substitutes; (iii) establishes additional requirements for the tripartite agreements which apply to all mines and companies producing chrysotile and chrysotile products; (iv) establishes requirements for monitoring and controlling the use of chrysotile and its substitutes, and ensures that a record is kept of the exposure measurements made by companies while guaranteeing access to them; and (v) establishes a permanent National Commission on Amianthus (NCA) to ensure the safety of workers involved in the chrysotile or substitute fibre industry. The Decree also establishes certain bodies, such as the NCA, composed of government and industry officials as well as workers, to ensure worker safety.

4.19 The tripartite agreements (otherwise known as The National Agreements for the Furtherance of the Safe Use of Asbestos) are required by both the Law and Decree. They are executed by the Federal Government of Brazil, the industries involved (e.g. the mining or asbestos-cement industry) and the workers in the industry (through their unions). They establish mandatory medical procedures inspection and safety measures, as well as exposure limits. They also give workers certain rights, both individual and collective, within their industries. Their objective is to continuously work towards improved worker safety and to decrease exposure limits as well as actual exposure. First, tripartite agreements set the maximum permissible exposure limits to 0.30 f/cm³, with 50 per cent of all measurements being below 0.10 f/cm³ (and without there being any constant exposure above 0.3 f/cm³, even when the workers exposed have special breathing equipment). Second, they require the use of specific “collective protection” procedures to protect workers. The procedures are to include the installation of air filter and exhaust systems, the use of wet processes in the handling of chrysotile (which reduces dust release and, thus, exposure), the sealing of workspaces and processes to limit exposure, the demarcation of areas of exposure for warning, the prohibition of dry sandpapering processes, the implementation of a daily programme for the washing, wetting or vacuum cleaning of production sites, and provisions for a change of work clothes (which may not be taken off site), of laundry services and showers for employees. Third, the agreements require employers to provide workers with individual protection equipment that complies with relevant standards. Fourth, they also require them to conduct regular and detailed environmental evaluations of working conditions as well as to medically inspect their employees. All results are to be filed with the Control Commission on the Safe Use of Asbestos and with the Brazilian Asbestos Association, known as ABRA. The Control Commission is comprised of plant workers elected by their peers. Fifth, they require the provision of worker education programmes to communicate the health risks of exposure to chrysotile, the measures which can be taken to reduce exposure, and the “multiplier effect” which tobacco smoking has on exposure. Sixth, they make ABRA responsible for providing the companies with technical assistance regarding controls and preventive measures.

4.20 Founded in 1984, ABRA is an industry watchdog group composed of companies from Brazil’s asbestos industry. Its main goal is to oversee industrial activity in order to
ensure that ABRA members comply with the Law, the Decree and the tripartite agreements, as well as to educate workers, wholesalers and end-users of chrysotile asbestos and asbestos products on safe use. To accomplish this goal, ABRA has an extensive, independent, monitoring programme. Biannually, it conducts spot measurements at the facilities of its members. It maintains an ISO 9000-certified laboratory and sends control samples once a year to independent laboratories in Edinburgh (AFRICA) and Paris (LHCF) to ensure the accuracy of its measurements. If the company that has been tested fails to meet the applicable exposure limits, ABRA sends it a letter and informs its suppliers. It then provides the company with a maximum number of days in which to comply, and instructs its suppliers to withhold chrysotile and/or chrysotile products from the company until it is able to notify its compliance. The agreement restates the requirements of both the Law and Decree, and develops certain safety procedures. In exchange for compliance (and dues), ABRA serves as a low-cost repository for state-of-the-art safe-use technologies, covering areas such as plant, air filter and process design. It attempts to encourage as well as facilitate safe use, with its overarching objective being to regulate the industry in such a manner as to render additional government regulation unnecessary. The regulatory regime (consisting of the Law, the tripartite agreements and ABRA itself), aligns the self-interests of the industry with those of its workers. The industry and the workers individually, as well as through Safety Commissions and Unions, cooperate to reduce health risks. The result of this cooperation has been the creation of an extremely safe workplace with very low exposure levels. In general, this system encourages individual plants to exceed applicable requirements in order to guarantee worker and user safety. At the Capivari Asbestos Cement Plant, which is the largest chrysotile-cement plant in South America, the on-site doctor did not report a single case of asbestos-related disease among the employees whose contact with asbestos had been limited to the plant.

4.21 With respect to regulation of asbestos in the United States, Brazil asserts that, in response to public outcry based on sensationalist media reports on the dangers of asbestos, the Environmental Protection Agency (EPA) banned asbestos in 1989. It prohibited “at staged intervals, the future manufacture, importation, processing, and distribution in commerce of asbestos in almost all products […]”. In reaction, a United States company that manufactured asbestos pipes, Corrosion Proof Fittings, filed suit against EPA arguing that the ban was not based on scientific and medical information. In a 1991 decision, the United States Court of Appeals for the Fifth Circuit called for the lifting of the ban and ordered EPA to issue new rules grounded in science. The Fifth Circuit concluded that EPA had presented “insufficient evidence to justify its asbestos ban.” Specifically, it found that the EPA had failed to (i) consider all of the necessary and relevant evidence, and (ii) “give adequate weight to statutory language requiring it to promulgate the least burdensome, reasonable regulation” that would protect human health. Similarly, France has failed to (i) examine existing evidence on the modern, controlled uses of chrysotile, (ii) assess the danger associated with substitute products, and (iii) impose a regulation that is not more restrictive than necessary. In 1993, EPA lifted the ban and issued new provisions regulating the production and use of asbestos and asbestos products. Based on a thorough scientific and medical review, EPA then authorized more asbestos products (18) than it banned (6). None of the uses that are banned are at issue in this proceeding. Of the authorized uses, two are central to Brazil’s exports to France and had previously been allowed (they include chrysotile-cement products and chrysotile friction materials). Under existing regulations, the United States produced 6,890 metric tonnes of chrysotile and imported 20,900 metric tonnes in 1997. In the same year, it consumed nearly 21,000 metric tonnes of chrysotile, exported unmanufactured fibre for a total value of US$5,690,000 and manufactured products for a total of US$197,000,000. Public health has not suffered in the United States and public outcry did not resume. United States regulations ban the dangerous uses of asbestos, and regulate those that are safe.

3. Legal Aspects

4.22 Brazil argued that, as it turns to Brazil’s legal arguments, the Panel should recall Brazil’s complex system of regulation that ensures public safety. The Panel should serve the same role regarding France’s political decision which the Fifth Circuit served regard-
ing EPA’s political decision - that of a neutral arbitrator. Brazil understands that the Corrosion Proof decision does not in the least bind the Panel - the procedures, legal standards and status of the parties are quite distinct. However, the court there faced similar circumstances and issues, and in the face of contrary public sentiment, issued a very focused, well-reasoned opinion, which is precisely what Brazil seeks here.

(a) The Agreement on Technical Barriers to Trade

(i) Article 12 of the TBT Agreement

4.23 Brazil argues that a prohibition of the trade and use of a product, such as France’s ban, is the most restrictive of all possible trade measures and must be closely scrutinized by the Panel. It requests the Panel to devote particular attention to the ban on imports from Brazil, which is a developing country (and from Zimbabwe, a least-developed country). In general, WTO agreements provide for the special and differential treatment of developing and least-developed country exports. In the context of the TBT Agreement, special provisions are set forth in Article 12, which obliges Members that are developing technical regulations and standards to consider the special needs of developing and least-developed countries and to provide them with differential treatment. Article 12.2 obliges France to “take into account the special development, financial and trade needs” of developing countries and of least-developed countries, when developing its technical regulations. France did not meet this obligation. Rather, it adopted an outright ban that advantages French producers of substitute fibres and products to the detriment of Brazil’s chrysotile and chrysotile product producers (and to Zimbabwe’s detriment as well). Moreover, the ban has not contributed to improving public health in France.

4.24 France violated Article 12.3 which covers the “preparation and application” of technical regulations and standards. Article 12.3 requires France to ensure that its technical regulations “do not create unnecessary obstacles to exports” from developing countries such as Brazil (and from least-developed countries such as Zimbabwe). However, France’s ban applies to Brazilian (and Zimbabwean) exports and creates, to say the least, an “obstacle” to their trade. The obstacle is “unnecessary” because it does not contribute to the supposed objective of increasing safety. The only trade permissible under the ban is that of chrysotile and chrysotile product substitutes. The risks associated with substitute fibres are unknown, but they are suspect. Meanwhile the risks associated with the modern, controlled uses of chrysotile, are zero.

(ii) Article 2.2 of the TBT Agreement

4.25 Brazil argues that the ban is inconsistent with Article 2.2 of the TBT Agreement because it is more trade restrictive than necessary to fulfil a legitimate objective. Once it is established that the Decree is a “technical regulation”, the EC must demonstrate (and have the burden of proving) that four different conditions have been met if they are to argue that the ban is in fact consistent with Article 2.2.41 To defend the ban, the EC must demonstrate to the satisfaction of the Panel that (i) the objective of the ban is “legitimate”, (ii) that it “fulfils” this legitimate objective, (iii) that it is not “more trade restrictive than necessary” to fulfil the legitimate objective, and (iv) that France evaluated the health effects (i.e. “the risks non-fulfilment would create”) on the basis of “available scientific and technical information”. According to Brazil, the ban meets only the first of these four conditions.

4.26 Brazil argues that the Decree is a “technical regulation” within the meaning of the TBT Agreement. The ban sets out certain (i) product characteristics, (ii) process and production methods, (iii) administrative provisions, as well as (iv) packaging, marking and labelling requirements with which compliance is mandatory. Article 1 of the Decree prohibits the production, importation, exportation, manufacture, transformation, sale and offer for sale of all types of asbestos fibres and asbestos-containing products (except those temporarily excepted from the ban by virtue of Article 2.1). Thus, the ban is explicitly directed at product characteristics (asbestos and asbestos-containing products) and at process and production methods (all forms of production, manufacture and transformation of
asbestos and asbestos-containing products). Both the prohibition imposed by Article 1 and the procedures for implementing and reviewing the entitlement to the exceptions set out in Articles 2.11 and 3 of the Decree are “applicable administrative provisions” relating to product characteristics and process and production methods. Article 4 of the Decree prescribes certain marking and labelling requirements for those few asbestos-containing products excepted under Article 2. Compliance with the ban is mandatory and violations are penalized under Article 5. Brazil argues that both France and the EC have conceded that the Decree is a technical regulation. In WTO document G/TBT/Notif.97.55, dated 21 February 1997, the French Government notified the ban to the TBT Committee as a technical regulation. Paragraph 3 of the Notification indicates that the ban was being notified under Articles 2.9.2 and 2.10.1 of the TBT Agreement, both of which establish notification obligations for technical regulations. The European Commission has also recognized that the ban is a technical regulation both in a 15 April 1997 document justifying the French ban and during the 8 July 1998 consultations on this dispute. Therefore, both France and the EC concede that the ban falls within the scope of paragraph 1 of Annex 1 of the TBT Agreement and is a technical regulation.

4.27 Brazil does not contest that the objective of protecting the health of French workers and consumers is a “legitimate objective” within the meaning of Article 2.2 of the TBT Agreement. However, it argues that the ban imposed by the Decree creates an unnecessary obstacle to trade. It does not in reality fulfil its stated objective, and is more trade-restrictive than necessary to protect the health of French workers and consumers. In using the word “fulfil” (as in the requirement that “technical regulations shall not be more trade-restrictive than necessary to fulfil a legitimate objective”), the text of Article 2.2 requires the existence of a rational link between the regulation and its stated objective. However, this rational link is absent as the ban does nothing to accomplish its objective. It does not make those who are now sick healthy and removing it would not make any of those now healthy, sick. The lack of a rational link between the ban and its purported objective is demonstrated by the following: (i) that asbestos-related health risks are due to old and already prohibited uses of asbestos; (ii) that there are no detectable health risks associated with modern uses of chrysotile; and (iii) that health risks associated with substitute fibres remain unknown and are suspect.

4.28 Brazil asserts that the health risks addressed in the INSERM Report are based on past exposure to high levels of asbestos fibres (largely amphiboles) and to exposure to old uses of asbestos, such as flocking. In prohibiting future importation and sale of chrysotile and modern chrysotile-containing products, the ban does nothing to address the effects (today) of exposure between 1940 and the early 1960s to extremely high levels of asbestos, mainly amphibole fibres. It does not cure workers who now suffer because of long-term exposure in the past to amphibole, the use of which was banned in France in 1994, or to unregulated concentrations of fibres that are “50,000 times” higher than the modern-day internationally recognized controlled-use level of 1 f/ml. Likewise, prohibiting the future importation and sale of chrysotile and modern chrysotile-containing products does nothing to address the effects of exposure to (or the disturbance of) friable asbestos, mainly amphibole, in French buildings prior to the 1978 French ban on flocking. This was recognized by European Commissioner, Mr. Bangemann, who, in response to a question posed by the European Parliament, responded that “[I]t is important to mention that a new ban would not lead to a lower risk of exposure to existing asbestos for workers, nor would it reduce the number of deaths from past exposure to asbestos.”

4.29 Brazil maintains that there are no detectable health risks associated with modern uses of chrysotile. There is no rational link between the ban and its purported objective because modern uses of uncontaminated chrysotile are safe. Prior to the ban, more than 90 per cent of the chrysotile imported into France was used in the manufacture of chrysotile-cement products. Currently, the chrysotile is bound to the cement and encapsulated in it, without there being any loose or friable fibres. Furthermore, most chrysotile-cement products are produced in such a way that sawing or drilling are unnecessary, and in the few instances when either or both are required, widely recognized and well-established procedures have been developed for these tasks which prevent fibre release. Similarly, in all
other modern uses of chrysotile, the fibres are sealed, bonded or encapsulated in the product. In no instance are loose, friable fibres allowed to be. Brazil contends that France does not have credible evidence to suggest that (i) sealed, bonded or encapsulated chrysotile poses a health risk, (ii) concentrations of chrysotile fibres at or below the internationally-recognized controlled use level of 1 f/ml present a health risk, and (iii) controls do not eradicate all risk throughout a product’s life-cycle (from mining to manufacture, distribution, sale and use, and eventual disposal). On the other hand, much science-based research concludes that the level of chrysotile encountered in the workplace today, or in buildings, presents no detectable health risk. After an exhaustive study of the existing scientific literature, the Health Effects Institute concluded in 1991 that the health hazards created by asbestos at the levels commonly encountered today are “unlikely to be large enough to be actually observed and measured.” This conclusion (which was reached by an independent United States health watchdog), confirmed the 1984 findings of the Ontario Royal Commission. Similarly, in the case brought by Corrosion Proof Fittings against the EPA, the Fifth Circuit made the following comment on the risk of asbestos products relative to toothpicks:

“As the petitioners point out, the EPA regularly rejects, as unjustified, regulations that would save more lives at less cost. For example, over the next 13 years, we can expect more than a dozen deaths from ingested toothpicks - a death toll more than twice what the EPA predicts will flow from the quarter-billion-dollar bans of asbestos pipe, shingles, and roof coatings.”

Brazil concludes that, because there are no detectable risks attributable to modern uses of chrysotile, there is no rational link between the French ban and its purported objective.

4.30 Brazil asserts that the French ban induces consumers to use chrysotile substitutes, whose health risks are unknown, in place of chrysotile, whose risks are known. In his 1998 paper on the biological effects of substitute fibres, Dr. J. M. G. Davis concluded that “replacement [of chrysotile by substitute fibres] is premature in the present state of our knowledge .... The need for full toxicology testing of new fibre products is recommended before these products are marketed.” This conclusion was shared by the European Communities Directorate General for Consumer Protection Policies, which stated that “there is no significant epidemiology base to judge the human health risks [of substitute fibres] ... hence the conclusion that [the uses of] specific substitute materials pose a substantially lower risk to human health, particularly public health, than the current use of chrysotile, is not well founded ... “. The INSERM Report itself acknowledges that the risks associated with substitute fibres are unknown. INSERM “urgently” cautions against their use until further scientific tests are conducted. It states that “[T]he absence of epidemiological data concerning the long-term safety of these substitute products should not obscure the results of experimental systems indicating the possibility that pathological modifications could result. It is urgently important that suitable research into this area be conducted prior to the widespread use of substitute fibres.” Despite this urgent warning from its own experts, the French Government banned chrysotile and not its substitutes the day after it received the INSERM Report. Thus, the French Government knowingly shifted consumption from chrysotile used in modern ways, and for which there is no detectable health risk, to substitute fibres for which “experimental systems indicate the possibility that pathological modifications could result”. Brazil concludes, therefore, that the ban does not “fulfil” a legitimate objective as required by Article 2.2 of the TBT Agreement. The rational link between the ban and its stated health objective does not exist because, as demonstrated above, (i) asbestos-related health risks are due to old, already prohibited, uses of asbestos, and not to the modern uses of chrysotile, (ii) no detectable health risks are associated with the modern uses of chrysotile, and (iii) substitute fibres, whose health risks are unknown, will replace chrysotile.

4.31 Brazil further argues that even if there were a rational link between the ban and the purported objective, the French ban would nonetheless be inconsistent with Article 2.2 of the TBT Agreement because it is “more trade-restrictive than necessary to fulfil a legiti-
mate objective, taking account of the risks non-fulfilment would create. A ban is the most trade-restrictive measure possible. It could be justified only if France were able to prove that there was no reasonably available, less trade-restrictive, alternative. France cannot do so. Controlled use policies demonstrably fulfil the objective of protecting the health and safety of French workers and consumers. In assessing whether the ban is more trade-restrictive than necessary, within the meaning of Article 2.2, the Panel should examine both the risks of non-fulfilment and whether a less trade-restrictive measure is available to fulfil the objective.

4.32 Brazil argues that available scientific and technical information does not support the imposition of the ban. Article 2.2 provides that, in assessing the risk that a technical regulation is meant to address, Panels should consider, inter alia, relevant scientific and technical information, related processing technology and intended end uses of products. The risk to be avoided in the dispute at hand is the risk of illness resulting from exposure to (a) modern uses of chrysotile and chrysotile-containing products and (b) the disturbance of previously installed friable asbestos (largely amphibole) in buildings. Illnesses associated with old, previously banned, uses of asbestos are not relevant to this analysis. Moreover, they cannot be addressed through the present ban on trade, domestic sale and use. The INSERM Report, which provides the supposed scientific justification for the ban, does not assess the health effects of current levels of exposure to modern uses of chrysotile. To determine the health risk associated with exposure to low levels of bonded, sealed and encapsulated chrysotile in chrysotile-cement and other modern applications, it applies the same risk of exposure associated in previous decades with higher levels of exposure to friable asbestos (largely amphibole). There is no scientific logic for such an extrapolation. The INSERM Report itself concedes that its conclusions are not “scientifically certain” but are merely “plausible, though uncertain, estimates”. Several other scientific reports concur that there is no detectable health risk from modern uses of chrysotile.

4.33 Brazil explains that all modern-day uses of chrysotile involve bonding, sealing or encapsulation. Such uses, or modern products, do not contain loose, friable chrysotile fibres – which were the cause of past asbestos-related illnesses. The risk associated with modern use is undetectable. Most modern products are manufactured to specifications well known in the building and public works trades, so that sawing or drilling operations are seldom necessary. When sawing or drilling are necessary, there are well-established procedures to ensure that workers are not exposed to fibre release. Thus, Brazil argues that neither available scientific information, intended end-uses nor processing technology, necessitate a ban on chrysotile. Brazil argues that while the term “necessary” has not yet been interpreted in the context of Article 2.2 of the TBT Agreement, the interpretation provided by the Panel in the case of Section 337 of the Tariff Act of 1930 is instructive:

”[I]t was clear to the Panel that a contracting party cannot justify a measure inconsistent with another GATT provision as ‘necessary’ in terms of Article XX(d) if an alternative measure which it could reasonably be expected to employ and which is not inconsistent with other GATT provisions is available to it. By the same token, in cases where a measure consistent with other GATT provisions is not reasonably available, a contracting party is bound to use, among the measures reasonably available to it, that which entails the least degree of inconsistency with other GATT provisions.”

4.34 Brazil argues that the focus is on the range of measures “reasonably available” to France. Just as under Article XX(d), under Article 2.2 of the TBT Agreement, the French ban cannot be justified as “necessary” since a less trade-restrictive measure that fulfils the legitimate objective is available. There are numerous examples of controlled use that are both readily available and effective in addressing the health risks associated with the modern uses of chrysotile. First, the ILO Convention and Recommendation Concerning Safety in the Use of Asbestos (Convention 162 and Recommendation 172) establish procedures to ensure safety in the handling of chrysotile and chrysotile products. Second, the ILO’s 1990 Code of Practice on Safety in the Use of Asbestos details appropriate controlled use proce-
dures to ensure worker safety with respect to all chrysotile-containing products currently in use. Third, Brazil, the United States and Canada have demonstrated that controlled-use policy is effective in eliminating the health risks attributable to the modern uses of chrysotile. Controlled use policy is less restrictive than a ban. Trade and sales are permitted as long as appropriate safety measures are employed in the manufacture, installation and use of chrysotile-containing products. While complying with safety regulations could be expensive for firms, the decision of whether or not to use chrysotile or substitutes under the safety regulations should be determined by the marketplace and not by government. Given the availability of controlled use policy and its effectiveness in addressing the legitimate public health objective which France wishes to achieve, the ban is inconsistent with Article 2.2 in that it is more trade-restrictive than necessary to fulfill its objective.

(iii) Article 2.4 of the TBT Agreement

4.35 Brazil contends that the French ban is inconsistent with Article 2.4 of the TBT Agreement because it ignores appropriate and effective international standards. Article 2.4 obliges France to base its technical regulations on existing international standards, or on any “parts of them”, that would be effective and appropriate in any given circumstance. France violated this obligation when it banned chrysotile and chrysotile products, ignoring existing international standards that would have been both appropriate and effective. To establish that France has not violated Article 2.4, the EC must show that: (i) there are no international standards that apply to asbestos; (ii) if international standards exist, that the ban is consistent with them; or (iii) if international standards exist and the ban is inconsistent with them, that the international standards would not have been an effective or appropriate means of accomplishing France’s stated objective. The EC cannot make such arguments.

4.36 Brazil argues that a number of international standards apply to chrysotile and chrysotile products, including ILO Convention 162 and Recommendation 172, on the types of asbestos that can be used (only chrysotile) and how, and the International Organization for Standardization’s (ISO) 7337 standard, entitled Chrysotile Cement Products Guidelines for On-Site Work, regarding the proper installation and use of chrysotile-cement products. The fact that the ISO 7337 standard is an applicable international standard is beyond doubt. Annexes 1 and 3 to the TBT Agreement expressly recognize the authority and status of ISO as an international standard-setting body, and the ISO 7337 standard directly governs the primary chrysotile product group. Each of these documents state that chrysotile products may be manufactured and used, but only under controlled conditions and in modern applications. Each of the standards sets out specific controls to guarantee the safety of workers and end-users. They have been incorporated into Brazilian legislation as well as that of many other countries, including the United States and Canada. The ban is inconsistent with these international standards because it bans all imports, manufacture, use, etc., of chrysotile and chrysotile products, whereas these permit their use in modern applications. They only subject them to safety controls. Current international standards provide an effective and appropriate means of fulfilling France’s stated objective and the EC cannot argue otherwise. ILO and ISO standards are “appropriate” for France’s stated objective since they were specifically drafted to protect the health of industrial workers, the general public and others who may come into contact with asbestos. ILO and ISO standards would also be “effective” in achieving France’s stated objective since they have successfully protected human health in economies as diverse as those of Brazil, the United States and Canada. The EC would be hard pressed to provide evidence of a deterioration in the health of citizens from Brazil, the United States or Canada due to adherence to ILO or ISO standards.

4.37 Brazil states that a closer examination of the term “ineffective and inappropriate means” is justified. The text of Article 2.4 clarifies that this exception is to be quite narrowly construed and applied. Were it not to be so, Article 2.4 would be rendered useless. Members would all too easily claim that the applicable international standard was “inappropriate”. Second, the Article provides examples of the situations in which exceptions to the use of international standards are allowed. These include when an international stand-
ard would be ineffective or inappropriate due to fundamental climatic or geographical factors or fundamental technological problems. Thus, a Member may ignore an international standard only if the standard will not achieve the results it seeks because of its unique conditions in terms of climate, geography, or its economy (i.e. level of technological development). No such conditions exist in France. The EC would be unable to present any evidence to suggest that different conditions apply to France so as to make the standards followed by Brazil, the United States and Canada inappropriate or ineffective for it. France ignored ILO and ISO standards because it wished to ban chrysotile to appease public opinion and advantage domestically-produced and substitute products.

(iv) Article 2.8 of the TBT Agreement

4.38 Brazil argues that the French ban is inconsistent with Article 2.8 of the TBT Agreement because it establishes design requirements for products. France’s ban is inconsistent with this obligation because, by prohibiting chrysotile and its use in any product, the ban sets out an impermissible “design or descriptive characteristic”. To establish that France has not violated Article 2.8, the EC must demonstrate that (i) the ban is a performance requirement; or, in this case, (ii) that adopting a performance requirement would not have been “appropriate”. The Communities can demonstrate neither of these points. The ban sets out an impermissible “design or descriptive characteristic” because it regulates on the basis of the content and description of a product. France has banned chrysotile and products containing it but has not banned competing fibres and products that contain them. Therefore, the ban advantages French-produced substitute fibres, products which are “like”57 chrysotile, and chrysotile containing products. The ban does not contain regulations based on the performance of a product. Rather, it states that certain products may be imported and sold only if they do not contain a certain input, namely chrysotile. Article 2.8 obliges France to adopt a performance requirement “whenever possible”. In the case of chrysotile, France could have adopted any of a number of performance requirements that would have enabled it to achieve its stated objective.

4.39 According to Brazil, France could have adopted, for example, detailed regulations regarding the importation, production, modern use and disposal of chrysotile and substitute fibres and their products (as France had previously done and as do Brazil, the United States, Canada and many other countries). Alternatively, France could have established a single, never-to-be-exceeded exposure level to apply to the manufacture, use and disposal of chrysotile and substitute fibres, and to their products. France could, and should, have adopted a performance requirement for chrysotile and the products which contain it. Instead, if adopted a design or descriptive requirement and violated Article 2.8 of the TBT Agreement. Were any other findings to be reached by the Panel, it would allow Member countries to take the much easier route of banning, rather than regulating, products which they claim create health risk. The TBT Agreement is based on the assumption that certain products present risks and that those risks are to be managed through standards. To allow a Member to ban, instead of to regulate, products due to perceived risks would render the Agreement meaningless.

(b) The General Agreement on Tariffs and Trade

(i) Article XI of the GATT

4.40 Brazil submits that the ban is also inconsistent with GATT Article XI because it is a quantitative restriction that is not permitted by the WTO. The ban includes (i) a prohibition of the sale in France of chrysotile and chrysotile products, which is a violation of GATT Article III:4, and (ii) a prohibition of the importation of chrysotile and chrysotile products. In fact, paragraphs I and II of Article 1 of the ban prohibit “the import […] of all kinds of asbestos fibres […] whether or not these substances are incorporated into materials, products or devices”. The latter aspect violates Article XI. In reference to paragraph 1 of Article XI, Brazil argues that the ban on importation is not a “duty, tax or other charge,” but is a “prohibition or restriction” that France has instituted and maintained on the importation of chrysotile from Brazil. Indeed, the ban is the most restrictive of all quantita-
tive restrictions in that it sets a quota at the level of zero imports. Therefore, the portion of the Decree that bans imports is inconsistent with Article XI:1. Brazil further argues that none of the three exceptions contained in paragraph 2 of Article XI apply to the ban. Simply put, the ban is an outright prohibition of all imports, supposedly imposed to protect public health. It is not a “standard or regulation for the classification, grading or marketing” of chrysotile or chrysotile products. Moreover, whether chrysotile is a “commodity” within the meaning of this exception is questionable. All three exceptions only relate to agricultural products.

(ii) Article III of the GATT and Article 2.1 of the TBT Agreement

4.41 Brazil argues that the ban is inconsistent with France’s national treatment obligations under GATT Article III:4 and Article 2.1 of the TBT Agreement. The national treatment obligations of Article III:4 and TBT Article 2.1 are violated when a law, regulation or requirement (or a technical regulation) that affects the internal sale, offering for sale, purchase, transportation, distribution or use of any imported product, accords less favourable treatment to the imported product than that accorded to “like” domestic products. Each of these criteria are satisfied with respect to the ban. The ban is indisputably a law and its three implementing “Arrêts” are regulations. For the purpose of Article 2.1 of the TBT Agreement, the ban is a technical regulation. Article 1 of the Decree bans, among other things, the manufacture, processing, sale, offer for sale, distribution and use of all varieties of asbestos fibres and all asbestos-containing products (except for the few temporary exceptions permitted by its Article 2). Thus, it indisputably meets the second criterion for the application of GATT Article III:4 and TBT Article 2.1. It provides less-favourable treatment to chrysotile and chrysotile-containing products (which, prior to the ban, were imported from Brazil), than that which it does to French products used as asbestos substitutes (and which are not banned).

4.42 Finally, the ban itself recognizes that the so-called “substitute fibres”, and products incorporating them, are “like” chrysotile and chrysotile-containing products. The few exemptions permitted by Article 2 of the ban apply when no substitute fibre is equivalent in terms of its end-use to chrysotile. In other words, whenever a French substitute fibre can replace chrysotile, chrysotile is banned. There can be no more convincing proof that chrysotile and substitute fibres are “like” products. Even if the ban did not by its own terms prove the “likeness” of French substitute fibres to imported chrysotile, analysis of the precedents set under GATT demonstrate that chrysotile and substitute fibres are indeed alike, as are chrysotile and substitute fibre-containing products. Using the criteria identified by the Appellate Body in the case on Taxes on Alcoholic Beverages for establishing “likeness”, it is self-evident that the end uses of chrysotile and substitute fibres are the same. The fibres are used solely because they emulate the desired characteristics of chrysotile in particular products. With regard to “consumers’ tastes and habits”, chrysotile and substitute fibres are not consumer goods. They are used solely as inputs in certain products (primarily in various cement products today). Industrial consumers purchase substitute fibres rather than chrysotile based on considerations of cost and availability. They can do so because substitute fibres are intended to emulate chrysotile’s characteristics.

4.43 Brazil asserts that the same reasoning applies to the assessment of the products’ properties, nature and quality. Substitute fibres are “like” chrysotile precisely because they emulate chrysotile’s characteristics. An additional criterion to determine likeness was added after Border Tax decision – tariff classification. As previously noted, almost all chrysotile is used as an input into various cement products. Chrysotile and other fibre-containing cement products are classified under the same Harmonized Tariff System heading (which is number 68.11). In all instances, the six and eight digit classification of chrysotile and other fibre-containing cement products are the same. Therefore, France’s conduct violates GATT Article III:4 and the TBT Agreement’s Article 2.1, and is inconsistent with France’s national treatment obligation.
(iii) Article I of the GATT and Article 2.1 of the TBT Agreement

4.44 The most-favoured-nation obligations of Articles I:165 and 2.1 are violated “with respect to all matters referred to in paragraphs 2 and 4 of Article III” (or, for purposes of the TBT Agreement, Article 2.1), whenever any “advantage, favour, privilege or immunity” is granted to a product from one country and is not “accorded immediately and unconditionally” to a “like product” from other WTO Members. This happens to be the case with the French ban. As previously demonstrated, Brazil argues that the ban violates GATT Article III:4 and, for the purposes of Article 2.1 of the TBT Agreement, is a technical regulation. The fact that substitute fibres may be imported into France while chrysotile imports are banned constitutes an “advantage, favour, privilege or immunity.” This advantage is accorded to imported substitute fibres but is denied to imported chrysotile, which is banned.

(iv) Article XX of the GATT

4.45 Brazil argues that the General Exceptions of Article XX do not excuse the Decree. To obtain an exception under Article XX, the EC must establish that (i) the ban does not “constitute a means of arbitrary or unjustifiable discrimination between countries where the same conditions prevail”, (ii) that it is not a “disguised restriction on international trade”, and (iii) that it is “necessary to protect human life or health.” The EC cannot argue that the ban meets these conditions. As demonstrated above, the ban discriminates between like products, without advancing its stated objective. Therefore, it is a “means of arbitrary or unjustifiable discrimination”. Also, it disadvantages imports of chrysotile, but not imports of man-made fibres. Countries like Brazil (and Canada) produce both chrysotile and substitute fibres. Therefore, the criterion of “discrimination between countries where the same conditions prevail” is obviously satisfied. Similarly, as previously demonstrated, the ban is a “disguised restriction on international trade”. Although it masquerades as a measure designed to protect public health, it is an outright product ban that is designed to quell public outrage and advantage domestic and European manufacturers of substitute fibres and products. Moreover, it cannot be argued that the ban is “necessary” to protect human life or health. For these reasons, the EC should not be granted recourse to Article XX.

B. United States

1. Introduction

4.46 The United States’ submission first discusses the facts concerning the health risks of exposure to chrysotile asbestos, and reduction of these risks through regulation. In this connection, the United States supplies information correcting certain errors and mischaracterizations in Canada’s description of the United States regulation of asbestos and the former United States ban and phaseout on asbestos-containing products. United States regulations are not at issue in this proceeding. The United States’ submission nevertheless seeks to set the record straight because of Canada’s assertions concerning United States policy. Following a factual discussion, it addresses the legal provisions that the Panel has been asked to interpret.

4.47 In the view of the United States, chrysotile asbestos is a toxic material that presents a serious risk to human health. Chrysotile asbestos is no less toxic than other forms of asbestos. A regulatory approach that treats all forms of asbestos on a par with each other is scientifically justified. France, like all other Members of the WTO, has the right to set its own desired level of protection against risks arising from exposure to asbestos, and its regulation on asbestos appears neither discriminatory nor unnecessarily trade restrictive in ensuring that level of protection. The United States currently relies on specified work practices and other controls (including a limited ban) to reduce the risk to human health from asbestos exposure. However, the United States does not consider its approach to be the only appropriate one for regulation of asbestos. Specification of work practices and other controls does not avoid all the risks associated with a hazardous material such as chrysotile asbestos. First, “controlled use” does not eliminate all the risks associated with asbestos. Although it is generally true that asbestos contained in a cement matrix does not
present substantial risks while that product is intact, the same is not true during the production, installation, maintenance, removal, or disposal of that product. Second, in many cases a matrix containing asbestos does not remain intact during its useful life. Moreover, while the bulk of Canada’s submission focuses on cement-matrix applications, it also acknowledges that chrysotile asbestos is currently used in brake linings and spun fibres for the production of insulating tissues or cords. Significant health risks attend the manufacture and repair of such substances. Finally, even the best work practice is effective only to the extent that it is followed; accidents, use of improper techniques, and intentional non-compliance are virtually inevitable in the use of these products. For these reasons, France’s ban on the manufacture, processing, distribution in commerce, export, import and sale of asbestos and its products appears to be a WTO-consistent response to the risks posed by the use of asbestos.

4.48 As for the legal issues: In the view of the United States, Canada has not met its burden of proof with respect to any violation by the French Decree of provisions of the GATT or the Agreement on Technical Barriers to Trade (“TBT Agreement”). In particular, Canada has not shown that imported asbestos and asbestos-containing products are “like products” with respect to substitute fibres and products containing them which are of French origin. As a finding that these products are not “like products” eliminates any violation of Article III:4, and Article XI:1 is simply irrelevant to the analysis of this measure, there does not appear to be any violation of the GATT 1994. With respect to the TBT Agreement, the United States disagrees with the EC position that the TBT Agreement is inapplicable to the French Decree. The Panel should reject the EC’s position and find that the Decree is a “technical regulation” within the meaning of Annex 1 of the Agreement; any other result will open up a loophole which could entirely nullify the TBT Agreement. Nevertheless, in the United States view Canada has not proven any violation of Articles 2.2, 2.4, 2.8 or 2.1 of the Agreement. Finally, Canada has not met the particularly high burden of proof for cases of non-violation nullification and impairment.

2. Factual Aspects

4.49 The United States argues that asbestos - whether chrysotile or in other forms - is a toxic substance. In United States lexicography, it is a “Class A carcinogen”, meaning a substance whose carcinogenic properties have been proven conclusively. The IPCS has reached the same conclusion: “[E]xposure to chrysotile asbestos poses increased risks for asbestosis, lung cancer and mesothelioma in a dose-dependent manner.” The IPCS Report also concludes: “[C]ommercial grades of chrysotile have been associated with an increased risk of pneumonia, lung cancer and mesothelioma in numerous epidemiological studies of exposed workers.” In regulating asbestos, the United States treats chrysotile asbestos the same as any other recognized form of the substance. The findings presented by Stayner et al. support the decision not to distinguish between chrysotile and other forms of asbestos. This study concluded that it is prudent to treat chrysotile with virtually the same level of concern as the amphibole forms of asbestos, based on the evidence of a significant lung cancer risk, the fact that workers are generally exposed to a mixture of fibres, and the lack of conclusive evidence for the “amphibole hypothesis.” More recent confirmation of the hazardous nature of chrysotile was provided by Landrigan, concluding on the basis of an epidemiological study undertaken in Quebec that “chrysotile asbestos is still indisputably a human carcinogen.” Concerning exposure to asbestos, IARC stated in 1976 that “at present it is not possible to assess whether there is a level of exposure in humans [to asbestos] below which an increased risk of cancer would not occur.” The IPCS reaffirmed this conclusion specifically with regard to chrysotile asbestos in 1998, stating: “[N]o threshold has been identified for carcinogenic risks” with regard to chrysotile asbestos. That means that it cannot be assumed that any exposure, no matter how small, to asbestos is safe. Canada questions France’s scientific approach, attacking the use of a “linear risk model”. The United States takes issue with Canada’s criticism of the INSERM Report’s use of a linear dose-response model to estimate cancer risk. The use of such a model is entirely appropriate when it comes to estimating the risk of cancer from exposure to asbestos.
4.50 The United States notes that it is not in a position to draw definitive conclusions concerning the regulatory process in France and the actual factual basis for the French Decree. However, generally speaking, regulatory decision-making relating to carcinogens involves two components: risk assessment and risk management. Risk assessment defines the adverse health consequences of exposure to toxic agents. Risk management combines the risk assessment with the directives of regulatory legislation, together with socio-economic, technical, political, and other considerations, to reach a decision as to whether or how much to control future exposure to the suspected toxic agents. Risk assessments are carried out independently from considerations of the consequences of regulatory action. A risk assessment involves, among other things, quantitative and/or qualitative estimation of risks associated with low levels of exposure to carcinogens. While it is always preferable to rely on human data, epidemiological studies are often either not available or sufficiently definitive, particularly regarding the specific exposure levels involved, and thus often cannot be relied upon as the sole basis for a risk assessment. In addition, because testing of thousands of animals would be necessary in order to have the sensitivity to detect any but large effects, it generally is not practical to measure risks at low exposure levels directly in animal experiments. Accordingly, a number of mathematical models have been developed to extrapolate from high dose animal studies to low human doses.

4.51 In the United States, models or procedures that incorporate low-dose linearity have been adopted when data and information are limited and when there is uncertainty regarding the mechanism of carcinogenic action. While low-dose linearity may not be appropriate for all carcinogenic risk assessment, it is commonly used in the United States as a default methodology. This methodology is supported by scientific studies and is a reasonably protective approach in the face of uncertainty. The use of a linear model is appropriate for a quantitative estimation of the risks associated with low levels of exposure to asbestos because of the observed linearity of the response in occupational studies. The United States has adopted this approach, in addition, because of the incomplete understanding of how asbestos causes diseases in humans. In assessing the risk from asbestos, EPA notes that "direct evidence for linearity of response with asbestos exposure is available from seven studies (two of the same plant) that compared lung cancer mortality to the cumulative total dust exposure in asbestos workplaces". Similarly, the limited data that exist for mesothelioma also indicate a linear relationship. The IPCS states: "there was a clear dose-response relationship, with crude rates of mesotheliomas (cases/1000 person-years) ranging from 0.15 for those with cumulative exposures less than 3530 million particles per cubic meter years … to 0.97 for those with exposures of more than 10,590 mpcm-years […]." After identifying and defining the adverse effects of asbestos through the risk assessment process, the next step is to make risk management decisions. A risk management decision, while taking into account the scientific findings of the risk assessment process, also includes a country’s choice on whether and how much to regulate a toxic agent. It is at this stage that a country selects measures and regulations that will achieve its chosen level of protection relating to the health of its people.

4.52 In its arguments, Canada has referred to United States regulations concerning asbestos. Because its description of the U.S. regulatory approach is substantially inaccurate, the United States proceeds to set the record straight. The U.S. regulatory approach at present includes a mix of control measures, including bans and required work practices. This approach involves a number of complex statutes, some of which require the consideration of cost, feasibility and other factors besides human health. Almost all control measures are designed to protect workers and building occupants from exposures resulting from contact with asbestos in installed products. Although France’s approach to the same problem is different, this different approach is also reasonable under the circumstances.

4.53 Canada references the 1989 rule promulgated by the EPA prohibiting the future manufacture, importation, processing and distribution in commerce of asbestos in almost all products ("the Asbestos Ban and Phase-Out Rule"). Several of Canada’s statements on this point are factually inaccurate. According to the United States, the Asbestos Ban and Phase-Out Rule was in large part vacated and remanded to EPA by the United States Court of Appeals for the Fifth Circuit in a case entitled Corrosion Proof Fittings vs. Environmental
Protection Agency, based on the court’s view that EPA had not appropriately addressed cost-benefit issues. Contrary to the claims made by Canada that EPA was incapable of scientifically justifying its ban, and that the risks posed by asbestos were not supported by scientific facts, the court specifically agreed with EPA’s scientific judgment by recognizing that “[a]sbestos is a toxic material, and occupational exposure to asbestos dust can result in mesothelioma, asbestosis, and lung cancer.” Indeed, in the record of rulemaking, EPA provided, among other things, a number of scientific studies and reports on the health risks of asbestos, including: Airborne Asbestos Health Assessment Update, Report to the United States Consumer Product Safety Commission by the Chronic Hazard Advisory Panel on Asbestos, Asbestiform Fibres: Non-occupational Health Risks and “Short-term asbestos work exposure and long-term observation.” These studies and reports were discussed in the preamble to the Rule. The court based its decision on procedural flaws in the EPA rulemaking process and on the court’s own interpretation of the applicable United States statutory risk/benefit balancing standard for promulgating such rules, not on any disagreement with EPA’s findings concerning the health hazards posed by asbestos. After the judicial remand, EPA imposed a more limited ban on asbestos-containing products, including a ban on any new uses of asbestos. That ban remains in place.

4.54 The United States notes that Canada makes much of the argument that asbestos entrained in a cement matrix does not present “detectable risk.” However, Canada’s focus ignores the risks presented by asbestos products throughout their life-cycle. The most significant sources of exposure to asbestos, and therefore risk from asbestos-containing products derive from their manufacture, installation, repair, removal, and disposal, including disposal of products containing asbestos in a cement or resin matrix. Moreover, while Canada appears to acknowledge France’s concern about protecting health in general terms, it appears to disagree about how protective France should be. By arguing that certain small risks are equivalent to zero risks, the Canadian submission implicitly questions the sovereign authority of a WTO Member to determine the appropriate level of protection for its citizens. What Canada or the United States might consider to be adequate protection in a particular context is not necessarily what other countries must choose. To put it another way, Canada concedes that a ban is acceptable for “certain uses where exposure cannot be controlled to an acceptable degree”. The United States agrees, but submits that it is up to each Member to determine what that “acceptable degree” is. Canada indicates that among the most important commercial applications of asbestos are as part of brake linings or clutches and in the form of spun fibres for the production of insulating tissues or cords. In connection with its analysis of “friction products”, which include brake linings and clutches, the United States court in the Corrosion Proof Fittings case recognized that “[w]orkers are exposed to asbestos during the manufacture, use, repair and disposal of these products” and that, in the asbestos ban and phase-out rule, “EPA demonstrates that the population exposure to asbestos in this area is great.” The court agreed with EPA’s determination that friction products containing asbestos pose a risk to human health.

4.55 With respect to Canada’s argument that a major commercial application for asbestos is as a reinforcement material for cement, plastic, or rubber, the United States asserts that, in the Asbestos Ban and Phase-Out Rule, EPA made certain determinations respecting worker exposure to these products that were not questioned by the Corrosion Proof Fittings court. EPA determined that the manufacture, installation, repair, and disposal of flat and corrugated asbestos-cement sheet expose workers to asbestos. Similarly, EPA determined that the manufacture and installation of asbestos-cement pipe provide “primary routes of exposure” of workers to asbestos from these products, and workers may also be exposed during the removal of asbestos-cement pipe. The United States generally agrees with Canada’s assertion that as long as asbestos is held within a cement or resin matrix in an undisturbed state, there is minimal exposure to the fibres - but only while the matrix retains its integrity. Much asbestos has been installed in United States buildings. Because of the high health risk from disturbed building materials and the reduced risk from intact asbestos-containing materials, EPA has issued guidance recommending management of asbestos-containing materials in place. Unfortunately, cement and resin matrices do not remain undisturbed. Putting aside significant releases that occur during the manufacturing process, releases of asbestos can occur when, for example, asbestos-
cement pipes are installed (which requires cutting the material), and when asbestos-containing material (such as cement) deteriorates, as through peeling, cracking, or crumbling. Fibre release could also result when the material is dry, has the capacity to be crumbled, pulverized or reduced to powder by hand pressure, or is subject to sanding, grinding, cutting or abrading.103

4.56 The United States asserts that a cement matrix in which asbestos is bound can undergo a natural process of erosion or degradation resulting in asbestos fibre release: “[T]he release of fibres from external asbestos-cement products [such as siding] due to weathering can be an important external source of asbestos contamination that can be carried into or can infiltrate into the building environment”. This has been acknowledged by the 1991 Health Effects Institute-Asbestos Research (HEI-AR) report, *Asbestos in Public and Commercial Buildings: A Literature Review and Synthesis of Current Knowledge*.104 As reported in the HEI-AR document, researchers found that weathered asbestos-cement sheet products washed from gutters and onto walkways were an important source of chrysotile carried by foot or wind into a classroom.105 The report also cited a research finding of increased ambient air concentrations in the vicinity of buildings with asbestos-cement products on their exterior.106 Mere maintenance of some asbestos bound in a matrix may disturb the matrix and thereby create additional exposures to the asbestos fibres. For example, in an EPA study107 conducted at 17 schools in New Jersey involving spray buffing of resilient floor tile containing asbestos, airborne asbestos concentrations were approximately five times higher during than before spray-buffing with high-speed machines, whereas spray-buffing with low speed machines showed a two-fold increase. For school maintenance workers, the maximum estimated eight-hour time-weighted average exposure concentration was 0.093 f/cc. Similarly, routine spray-buffing and wet-stripping as well as ultra-high speed burnishing and wet-stripping of asbestos-containing resilient floor tiles can result in elevated levels of airborne asbestos.108 Part 2 of the Kominsky study demonstrates that the ultra high-speed burnishing and wet-stripping procedures were associated with a maximum estimated eight-hour time-weighted average exposure concentration of 0.275 f/cc to operations and maintenance staff.109 Likewise, the HEI-AR Report states that: “buffing, wax stripping, and other abrasive treatments may cause the release of particulate material from the surface of the floor tile”.110

4.57 The United States’ regulatory programme on asbestos is largely aimed at controlling exposure to asbestos when it is no longer bound in a matrix. United States regulations address renovation and demolition of buildings111 and identification and management of asbestos-containing material in schools.112 Regulations of the Occupational Safety and Health Administration (OSHA) of the Department of Labor address worker exposures to asbestos including manufacture, installation, renovation, removal and custodial work where workers come into contact with asbestos-containing material. OSHA has established a permissible exposure limit and mandated extensive work practice controls, enclosures, hazard communication, training, medical and other industrial hygiene practices to protect both workers contacting asbestos and other workers who are nearby. Enforcing and complying with these regulations entail a significant commitment of resources by both the public and private sectors. Even within a cement or resin matrix, the risk from asbestos is not negligible. According to an analysis conducted in 1991 by the Health Effects Institute-Asbestos Research113, janitors, custodians and maintenance workers exposed to ambient asbestos fibre levels of 0.1 f/ml (the permissible exposure limit currently allowed by OSHA regulations) were subject to an estimated increased risk of death from cancer of 2 in 1,000. The same analysis estimated that building occupants (school children and office workers) exposed to airborne asbestos fibres from asbestos-containing materials (presumably many of which, as building materials, would have been encased in cement or resin) have a lifetime cancer risk from such asbestos exposure of 4 to 60 per 1,000,000. Contrary to Canada’s suggestion, such risks are not equivalent to zero. Each country must determine for itself what level of protection from risks of exposure to asbestos it wishes to achieve, i.e. what risks to its population it is willing to accept. It is not for another country to tell France that certain risks to its population are not significant. By way of illustration, the United States regulates risks on the order of 1 in 100,000 (1 x 10^-5) or 1 in a million (1 x 10^-6) in a number of instances. Canada concedes that “[t]he principle of controlled use also
means that certain uses for which exposure cannot be controlled to an acceptable degree would be banned”. The discussion above demonstrates that exposure to asbestos even in a cement or resin matrix cannot be controlled sufficiently to eliminate all risk.

4.58 The United States argues that both of the parties have mischaracterized the substitutes for asbestos and asbestos-containing products. Some products that currently contain asbestos may be manufactured simply by removing the asbestos, thus eliminating any substitution of risk. Alternatively, a wide variety of fibrous substances are being used commercially as replacement materials for asbestos-containing products. These include the man-made mineral fibres (consisting of glass fibres, rock wool, slag wool, refractory ceramic fibres), selected organic fibres (e.g., aramid, carbon/graphite, polyolefin), and several naturally occurring mineral fibres other than asbestos (e.g., wollastonite, sepiolite, palygorskite). The potential health effects for these non-asbestos fibres have been evaluated by EPA,\textsuperscript{114} the IARC\textsuperscript{115} and the IPCS.\textsuperscript{116} Although limited health effects information exists for many of these fibres, available data do not indicate that these fibres are as toxic as chrysotile asbestos. For example, none of these fibres have been found to cause either malignant or non-malignant respiratory diseases similar to those associated with asbestos exposure in humans. Unlike asbestos fibres, these substitute fibres have not been classified as carcinogenic to humans or known human carcinogens. The only fibre that has been shown to be more hazardous than asbestos fibres is erionite. Erionite, however, is not known to be available in commerce at this time.\textsuperscript{117}

4.59 The United States notes that Canada continually urges that “controlled use” will bring the risk associated with chrysotile asbestos to “undetectable” levels. It also compares the risk from asbestos to that of other products and activities, concluding that many are more risky than asbestos. Canada overstates the efficacy of “controlled use”. For example, Canada states that where it is necessary to cut chrysotile-cement materials on site, “the use of tools that almost entirely eliminate emissions (low-speed saws, with water injection or equipped with suction units), and the wearing of a mask by the operator guarantee their safety”. United States regulations recognize, however, that masks may not be sufficient in some situations and require the use of a supplied-air respirator which is obviously more cumbersome and costly.\textsuperscript{118} When a government makes its choice whether to ban a product or to opt for controlled use, it must necessarily take into account the anticipated effect of the regulations on its population. Beyond the obvious point that “almost” eliminating emissions of asbestos fibres is not the same as eliminating them, it must be acknowledged that 100 per cent compliance with “controlled use” of asbestos is not a realistic expectation due to the burdensome nature of certain work practices concerned. Even the best work practice is effective only to the extent that it is followed; accidents, use of improper techniques, and international non-compliance are virtually inevitable in the use of these products.\textsuperscript{119}

4.60 The United States argues that Brazil has made a number of unjustified and inaccurate assertions concerning the U.S. Environmental Protection Agency’s former ban on all forms of asbestos and asbestos-containing products in the United States, and a domestic court decision concerning the EPA ban. Of course, this U.S. domestic court did not rule on the consistency of the EPA ban with the TBT Agreement or the GATT. It only dealt with whether EPA had complied with the risk/benefit balancing requirements of the U.S. Toxic Substances Control Act (TSCA). The risk/benefit standard in this U.S. statute is irrelevant to the Panel’s present task of determining whether France’s ban on asbestos is consistent with the WTO Agreement. France has not adopted such a risk/benefit balancing standard as the basis for a ban. First, Brazil misrepresents the domestic court decision and the situation following, repeating errors made by Canada. The court in 1991 upheld EPA’s determination that “[a]sbestos is a toxic material, and occupational exposure to asbestos dust can result in mesothelioma, asbestosis, and lung cancer”. The court based its decision not on any disagreement with EPA’s findings concerning the health hazards posed by asbestos, but instead on procedural flaws in the EPA rulemaking process and on the Court’s own interpretation of the statutory risk/benefit balancing required by TSCA. Because the court agreed with EPA on the health effects of asbestos, EPA did not, after the court decision, need to carry out a “thorough review of scientific and medical data” as a basis to
authorize or ban the products Brazil has listed. All EPA did, pursuant to the court's instructions, was to determine which product categories were no longer manufactured, imported, or processed when the rule was issued. For all such products the ban was maintained. EPA also banned new uses of asbestos.120

Second, contrary to what is alleged by Brazil, the United States has not determined that controlled-use policy effectively eliminates the health risk attributable to modern-day uses of chrysotile. Third, Brazil has downplayed the health risk from asbestos by lifting a quotation out of context from the Health Effects Institute report, in a manner that does not accurately reflect the discussion of mathematical models found in this section of the HEI report.121 The United States addressed these issues as described above at paragraph 4.40. Finally, concerning Brazil's discussion of the French Decree, the United States notes that Brazil concedes that the objective of the Decree - protection of the health of workers and the public - is a legitimate objective within the meaning of Article 2.2 of the TBT Agreement. The United States agrees with Brazil on this point but would also note that France has a right to set its "legitimate objective" to establish the protection of the health of French workers and consumers at the level it deems appropriate. However, when alleging that there is no "rational link" between the French Decree and health protection, stating that the Decree will not make those now sick healthy, and removing it would not make any of those now healthy sick, Brazil conveniently omits the function of the Decree in preventing future exposure and future disease that would result from that exposure.

3. Legal Aspects

For the reasons below, the United States suggests that the Panel should find that Canada has failed to meet its burden of proof that the French Decree violates any provision of the WTO Agreement. The Panel should also find that Canada has failed to make a showing that the French Decree gives rise to non-violation nullification or impairment of benefits accruing to Canada.

(a) The General Agreement on Tariffs and Trade

(i) Article XI of the GATT

With respect to Canada's argument that the Decree violates Art. XI:1 because it imposes an absolute prohibition or restriction on imports, the United States agrees with the EC that Article XI is simply not relevant to these proceedings, and the Decree should be analyzed under Article III instead. The Decree regulates characteristics of asbestos and asbestos-containing products. It applies to all asbestos, and is applied to imported products "at the time or point of importation," in the words of the Note Ad Article III.

(ii) Article III of the GATT

With respect to Canada's allegation that the Decree violates the national treatment obligations embodied in GATT Article III:4, the United States argues that, to show a violation of Article III, there must be discrimination - that is, unlike treatment of like products. Yet the relevant domestic and imported products here are not "like products" for the purposes of Article III:4. As the EC has noted, the classic statement of the factors relevant in determining what constitutes a "like product" is found in the Working Party report on Border Tax Adjustments of 1968, which defined like products in terms of "the product's end-uses in a given market; consumers' tastes and habits, which change from country to country; the product's properties, nature and quality",122. The United States generally agrees with the EC's analysis that the properties, nature and quality of asbestos and asbestos-containing products on the one hand, and substitutes on the other, are not "like". The substitutes are by definition substitutable for asbestos and asbestos-containing products for certain uses, but that does not mean that they are "like products".

According to the United States, Canada has not made the correct product comparison for the purpose of determining whether the relevant products are "like products"
under Article III:4. In considering a regulation that bans asbestos and requires the use of substitutes, the relevant products to compare are the following: (i) asbestos must be compared to substitute fibres; and (ii) products containing asbestos must be compared to products that do not contain asbestos but which perform the same function. Where the asbestos elements of a product were inessential, the substitute product may consist of the same product minus the asbestos element (e.g., a kitchen hot pad with thick cotton padding but no asbestos); or the same product redesigned to eliminate the need for asbestos; or a similar product which uses different fibres (e.g., a hot pad made of glass fibre); or a similar product made of what the EC describes as “classic materials” (e.g., a trivet made of cast iron, ceramic or plastic). The physical correspondence between the two classes of products is therefore considerably weaker than Canada has assumed. Canada has failed to show that asbestos and asbestos-containing products and the substitutes have the same “properties, nature and quality”. The known severe adverse human health effects of asbestos are another reason why asbestos-containing products are not “like” the substitutes for which adverse health effects have not been demonstrated. The substitute fibres differ considerably in physical structure and properties from chrysotile asbestos and thus cannot be considered “like products.” For example, while chrysotile is a naturally occurring mineral which is crystalline in nature, the man-made mineral fibres (MMMF) are amorphous (non-crystalline) silicates which are produced from a liquid melt of different starting materials (e.g., slag, natural rock, glass, clays). Furthermore, unlike chrysotile asbestos, MMMF do not split longitudinally into smaller fibrils of smaller diameter, but may break transversely into shorter segments.123

(iii) Article XXIII:1(b) of the GATT

4.66 The United States argues that Canada has failed to meet the special burden imposed by Article 26:1 of the Dispute Settlement Understanding on parties making claims of non-violation nullification or impairment. The United States has been one of the strongest proponents of the non-violation remedy, as an essential safeguard for bargained-for market access rights against frustration by government actions. But the requirements of the non-violation remedy are not satisfied here. As the EC have noted, the text of Article XXIII:1(b) establishes three elements that a complaining party must demonstrate in order to make a cognisable claim under Article XXIII:1(b): (i) application of a measure by a WTO Member; (ii) a benefit accruing under the relevant agreement; and (iii) nullification or impairment of the benefit as a result of the application of the measure. Canada, as the complaining party, has the burden of presenting detailed evidence in support of all three elements. In the present case, there is no dispute that the French Decree is a measure of a Member. The question is simply whether Canada has a legitimate expectation of benefits accruing to it. The precedents are clear that for expectations to be legitimate, they must take into account all measures of the party making the concession that could reasonably have been anticipated at the time of the concession.

4.67 The United States considers that, as a matter of principle, the Panel should reject the possibility of a finding of non-violation nullification or impairment with respect to health and safety regulations that respond to the development of scientific knowledge concerning health risks. Members do not have a legitimate expectation that regulatory measures will stay static in the face of expanding scientific knowledge concerning health risks, and changing societal decisions concerning the level of acceptable risk. Canada is also in a poor position to argue that an asbestos ban was unforeseeable at the time it negotiated the tariff concessions on asbestos. The dangers posed to human health by asbestos are notorious and have been so for many years. Pliny, the ancient Roman author, described the “diseases of slaves” as including exposure to the textile processes of preparing and weaving asbestos, and even referred to the use of a transparent bladder skin as a respirator to avoid inhalation of dusts by slaves.124 As of the time of the first GATT negotiating round in 1947, asbestosis had already (in the 1920s) been identified as a distinct condition caused by asbestos.125 By 1935 asbestosis was widely recognized as a mortal threat affecting a large fraction of those who regularly worked with the material.126 In addition, by the mid-1940s there were indications that exposure to asbestos in animals and humans was associated with lung tumours.127 Thus Canada should have reasonably expected subsequent regula-
tory action (such as a ban) by a GATT contracting party as a result. As of the Dillon Round of 1960-61, Canada had even more reason to foresee the possibility of restrictive regulations of asbestos. An international symposium of experts on the pathogenesis of lung cancer in 1953 published its conclusions and recommendations in a journal which editorialised: ‘‘[I]t seems to be beyond discussion that cancer of the lung is sometimes caused by occupational exposure to asbestos.’’ In addition, two major studies had been published in 1955 on cancer in the textile industry demonstrating the relationship between asbestosis and lung cancer. Since the early 1960s, the hazards posed by asbestos - and particularly chrysotile asbestos - have become even more widely known and documented.

(b) The Agreement on Technical Barriers to Trade

4.68 With respect to Canada’s arguments that the French ban on asbestos is inconsistent with a number of provisions in the Agreement on Technical Barriers to Trade (TBT), the United States argues that Canada has misread the relevant TBT articles. The interpretation of the TBT Agreement on which Canada’s arguments are based attempts to read into the Agreement obligations that do not exist. The United States urge the Panel to reject that interpretation. The EC, on the other hand, have argued that the French Decree is not a “technical regulation” because it is a categorical ban on asbestos and products containing asbestos. They have argued that general bans, and specifically this product ban, are not “technical regulations” because they allegedly do not “lay down product characteristics or their related processes and production methods” within the meaning of paragraph 1 of Annex 1 of the TBT Agreement. The United States disagree with the EC’s view on this point. In this instance, the Decree lays down “product characteristics […] with which compliance is mandatory”. The characteristics in question are that the product may not contain any asbestos if it is to be marketed, offered for sale, imported, exported, etc. in France. Compliance with the exclusion of asbestos is mandatory except if the French Government has accorded a derogation, in which case adherence to the terms of the derogation is mandatory. In any event, the French Decree is a technical regulation within the meaning of the TBT Agreement and is subject to the substantive rules of the TBT Agreement. The EC’s interpretation of Annex 1 would open up a loophole of potentially huge dimensions in the TBT Agreement. Measures having a very significant impact on trade - for instance, regulations limiting the characteristics of spreadable butter or wool - could simply be redefined as product bans. Similarly, the EC argument would mean that a regulation on the safety of infant toys that excluded any parts below a certain size (to prevent choking) would not be a “technical regulation” nor would regulations excluding water from being added to ham. The provisions of the TBT Agreement would then be rendered a nullity. Such a reading of the TBT Agreement is impermissible as a matter of treaty interpretation, and undesirable as a matter of trade policy. This does not mean that the French Decree does not satisfy the requirements of the TBT Agreement, however. As discussed below, although the TBT Agreement applies to the French Decree, Canada has failed to make a case that the French Decree violates any of the provisions it has cited.

(i) Article 2.1 of the TBT Agreement

4.69 The United States argues that, for the reasons discussed in relation to Article III of the GATT, these products are not “like products.” Furthermore, since the ban is applied without discrimination as to the source of the product, discrimination between foreign sources is not an issue.

(ii) Article 2.2 of the TBT Agreement

4.70 The United States notes that Article 2.2 provides a key element of the disciplines in the TBT Agreement. From the standpoint of the United States, certain aspects of Article 2.2 are particularly important with respect to health and safety regulation. The first sentence of Article 2.2 is important because it recognizes that in certain circumstances, technical regulations may create necessary obstacles to trade, and the creation of such necessary obstacles is consistent with the TBT Agreement. We note that the “legitimate objectives” enumerated (non-exhaustively) in Article 2.2 specifically include protection of human health
or safety. Article 2.2 also recognizes that in assessing the risks that may arise from non-
fulfilment of a legitimate objective, a government may consider a number of elements,
including available scientific and technical information, related processing technology or
the intended end-uses of a product.

4.71 The obligation in Article 2.2 that technical regulations are not to be more trade
restrictive than necessary to fulfill a legitimate objective should be interpreted in a manner
similar to Article 5.6 of the Agreement on Sanitary and Phytosanitary Measures (SPS). Such
a reading is supported by the sixth clause of the Preamble to the TBT Agreement,
which provides that: “[R]ecognizing that no country should be prevented from taking
measures necessary ... for the protection of human, animal or plant life or health, [or] of
the environment [...] at the levels it considers appropriate[...].” The United States argues
that the preamble is part of the context of Article 2.2 in the sense of Article 31 of the Vienna
Convention on the Law of Treaties, and provides an authoritative indication of the
TBT Agreement’s object and purpose for the purposes of treaty interpretation. Thus, in
order for a Member to show that a government’s technical regulation is more trade-restric-
tive than required, it would need to show that there is another measure that is reasonably
available, fulfils the regulating Member’s legitimate objectives, and is significantly less
restrictive to trade. Accordingly, the complaining party should be required to identify a
specific alternative measure that is reasonably available - as a Member is not required to do
what is unreasonable. Furthermore, the alternative measure must make a significant dif-
ference from a trade perspective. There should be no need to adopt an alternative measure
if it makes only an insignificant difference in terms of trade. Most importantly, the com-
plaining party must demonstrate that the alternative measure fulfils the government’s ob-
jectives. Canada has failed to demonstrate that its preferred alternative to the French ban
– i.e. “controlled use” of asbestos and asbestos products – fulfils the French Government’s
stated “legitimate objective” of protection of human health.

4.72 Canada alleged that the Decree does not address the “true problem” of asbestos
in France which Canada identifies as the flocking of asbestos. Yet it is not for Canada to
determine what France’s “true problem” is. It is up to France to determine what level of
protection to afford its citizens. Second, Canada alleges that the French Decree violates
Article 2.2 because it fails to acknowledge the “scientific reality” that chrysotile encapsu-
lated in a matrix is harmless. Yet as discussed above and as demonstrated by the EC,
encapsulated asbestos is not harmless at all, as the encapsulation can easily be breached,
and is likely to be breached during the product’s life cycle, resulting in release of fibres and
elevated risk to human health. Canada alleges that the French Decree violates Article 2.2
because it replaces use of chrysotile – an allegedly harmless product – with substitutes
whose health risks are unknown. The United States fundamentally objects to this reading
of the TBT Agreement. Canada is implicitly arguing that any regulatory action negatively
affecting trade in a product has to be tested against the hypothetical risks engendered by
use of likely alternative products. This test has no basis whatsoever in the TBT Agreement.

(iii) Article 2.4 of the TBT Agreement

4.73 Canada has asserted that Article 2.4 requires a panel to determine: (i) whether a
technical regulation on chrysotile is required; (ii) whether there are international stand-
ards concerning chrysotile; (iii) whether the international standards are effective and ap-
propriate to achieve the objective; and (iv) whether the Decree is based on international
standards. Under this analysis, Canada concludes that France adopted the most restrictive
measure possible despite the fact that the international community has developed stand-
ards representing a less restrictive approach (i.e., controlled use). This analysis misreads
Article 2.4. First and fundamentally, Article 2.4 does not contemplate that a panel deter-
mine whether a technical regulation is or is not required. The burden of proof is on Canada
to demonstrate that international standards exist and are relevant. In regard to the ILO
standard, both the ILO Convention 162 and Recommendation 172 allow participating coun-
tries to choose the approach they find to be appropriate to protect workers from asbestos
hazards. Indeed, the Provisional Record to the 72nd Session of the International Labour
Conference, which adopted Convention 162, states concerning Article 10 of Convention
162 (which Canada now claims to condition a ban on a finding concerning the risk posed by product substitutes): “The Government member of Canada saw nothing in Article 10 that would prevent any country from doing whatever it wanted with respect to asbestos.”

4.74 The United States argues that Brazil’s interpretation of Article 2.4 of the TBT Agreement ignores the fact that climate, geography and fundamental technological problems are listed as examples, not an exhaustive list, of reasons that an international standard may be an “ineffective and inappropriate means” of achieving a Member’s legitimate objective.

C. ZIMBABWE

1. Introduction

4.75 As an important producer and exporter of chrysotile (white) asbestos fibre and products containing chrysotile asbestos and also as a developing country in need of foreign exchange, Zimbabwe argues that it has a substantial interest in the outcome of this proceeding. In fact, the present dispute is of such importance to Zimbabwe’s asbestos industry and indeed its whole economy that the Government of Zimbabwe has decided, for the first time ever, to have recourse to the dispute settlement mechanism of the WTO. Zimbabwe is of the view that the ban of chrysotile asbestos and products containing chrysotile asbestos by France is unjustified and contrary to relevant rules of the World Trade Organization (WTO). The ban should therefore be lifted without delay. Zimbabwe believes that it is not incumbent upon it as a third party to this dispute to set out in full the case against the responding party, i.e. the EC. Accordingly, Zimbabwe will limit itself in this submission to the Panel to addressing a number of factual and legal aspects of this dispute that it feels are of particular importance to the outcome of this proceeding. Zimbabwe argues that the complaining party in this case, i.e. Canada, has made a compelling case with respect to both the factual and legal issues in dispute as to why the ban on chrysotile asbestos and products containing chrysotile asbestos is inconsistent with relevant WTO rules and must be withdrawn immediately.

2. Factual Aspects

4.76 Zimbabwe asserts that the chrysotile asbestos industry is of great economic importance to its economy. Zimbabwe ranks among the world’s largest producers of chrysotile asbestos. In Africa, Zimbabwe is the number one producer of chrysotile asbestos. It produces a high-quality chrysotile asbestos fibre and has sufficient underground reserves for at least another 25 years and infrastructure to continue operations for many more years to come. Chrysotile asbestos currently accounts for about 18 per cent of Zimbabwe’s mineral production index of volume and value. Crocidolite (blue) and amosite (brown) asbestos are not mined in Zimbabwe. As a developing African country, Zimbabwe relies primarily on natural resource products and other primary products for much of its export revenue. In terms of the revenue it generates, chrysotile asbestos is second only to gold as far as the mining sector is concerned. As much as 95 per cent of the country’s total asbestos fibre production is exported. In 1998, for example, 150,000 tonnes of chrysotile asbestos were exported out of a total production of some 175,000 tonnes, generating foreign exchange in excess of ZWS1.5 billion. In addition to the export of chrysotile asbestos fibres, more than 7,500 tonnes of asbestos-cement products, valued at over ZWS30 million, were exported. Zimbabwe’s sole producer of chrysotile asbestos fibre is African Associated Mines. The European Union in general, and Spain and France in particular, have traditionally been important export markets for African Associated Mines.

4.77 Zimbabwe argues that African Associated Mines suffered a dramatic (more than 50 per cent) drop in its sales to France in 1996. The setback suffered by African Associated Mines in the French market is directly attributable to the French Government’s actions. It should be pointed out in this connection that in mid-1996 the French Government announced its intention to ban asbestos. Before that, i.e. towards the end of 1995, the French Government had already announced a programme to reduce the risks associated with exposure to
asbestos. There is therefore clear evidence that the French ban on asbestos and products containing asbestos has had a direct and damaging impact on Zimbabwe’s asbestos industry. The significance of the asbestos industry to Zimbabwe cannot be overstated. The country has immensely benefited from its existence. African Associated Mines directly employs about 6,000 people in Zimbabwe, which amounts to about 20 per cent of total employment in the mining industry. The industry indirectly sustains more than 70,000 people in and around the mining towns of Zvishavane and Mashava. There are no other industries in these towns, meaning that a decline of the asbestos industry would cause dislocation with all its attendant social consequences. It should be borne in mind in this context that the Zimbabwean economy has faced considerable difficulties in the past decade and has not been able to create a sufficient number of new jobs. Out of a labour force of 5 million people, only 1.4 million people are gainfully employed. Apart from generating revenue for the Government of Zimbabwe, the asbestos industry has injected dynamism into the country’s economy. In addition to the salaries and wages paid by the companies engaged in the mining and marketing of asbestos and asbestos products, the asbestos industry’s more than 300 suppliers of goods and services receive payments of around ZW$600 million each year, including over ZW$150 million for the state-owned Zimbabwe Electricity Supply Authority (ZESA) and the National Railways of Zimbabwe.

4.78 It is apparent from the foregoing that a ban on asbestos would have severe repercussions for the Zimbabwean economy. In fact, as has been demonstrated, the ban on asbestos by France has already impacted negatively on the Zimbabwean economy. It must be mentioned in this connection that Zimbabwe views with great concern the potentially wider implications of the French ban on the use of chrysotile asbestos. While it is true that most countries, including the United States, still do not generally prohibit the use of chrysotile asbestos or products containing chrysotile asbestos, there is the probability that other governments may be tempted to follow the French example if the French measure were upheld by the WTO. Indeed, the European Union has just announced - without awaiting the outcome of a WTO ruling - that it will move to ban the use of chrysotile asbestos in all its member States. Zimbabwe wished the WTO to be aware of the wider implications of the decision it would render in this dispute.

4.79 Zimbabwe argues that the risks involved in the use of chrysotile asbestos can be adequately controlled. It appears that the concerns that governments have with respect to the use of chrysotile asbestos relate to airborne asbestos dust or respirable asbestos fibres, as they may have an effect on human health. For this reason, the United Nations Environment Programme (UNEP), the International Labour Organization (ILO), and the World Health Organization (WHO), within the Framework of an Inter-Organization Programme for the Sound Management of Chemicals, commissioned a Task Group of international experts to make an evaluation of the risks for human health from exposure to chrysotile asbestos and to make recommendations for health protection and further research. The report of the Task Group was published in 1998. One of the Task Group’s main conclusions was that “[e]xposure to chrysotile asbestos poses increased risks for asbestosis, lung cancer and mesothelioma in a dose-dependent manner”. The Group acknowledged, however, that it was not possible to provide quantitative estimates of risks to humans given the dearth of information and data. Furthermore, the Group cautioned that there was a need for further epidemiological studies of populations exposed to pure chrysotile so as to clearly and reliably be able to distinguish between chrysotile and amphibole exposure. In other words, there is a possibility that the available data may actually overestimate the risks to humans from exposure to chrysotile asbestos. What is quite clear from the Task Group’s conclusion - and this is crucial - is that the risks to humans are conditional on exposure as well as on doses or concentrations. The key objective for any responsible government must therefore be to reduce exposure. This said, it should be borne in mind that chrysotile asbestos is a natural product. It is present in the air we breathe and in the water we drink. Exposure is therefore inevitable, and no ban can change that. With these facts at hand, the question arises as to whether the French ban of asbestos is justifiable given the information within the public domain. Zimbabwe believes that what is at the heart of this dispute is the risk of occupational exposure to cement containing chrysotile
4.80 It is the submission of Zimbabwe that transportation and storage of imported chrysotile asbestos fibre do not entail a risk of exposure provided that there is proper packaging. Another possible activity involving a risk of exposure is the production of chrysotile asbestos-cement itself. In Zimbabwe, this risk has been contained, as demonstrated by the monitoring done by a group of independent experts for Turnall Fibre Cement Company Limited, which is a Zimbabwean company engaged in the manufacture of chrysotile asbestos-cement. The focus of the research has been on the health hazards related to asbestos during the process of manufacture. This research has been going on for more than ten years and so far there have been no reported cases of risks to human life. It should be mentioned here that the EC has adduced as relevant evidence a very recent study by the U.K. Health and Safety Commission which is alleged to demonstrate that notwithstanding the application of control measures, “primary users” of chrysotile asbestos fibres, i.e. workers in asbestos-cement factories, showed a higher mortality rate in relation to asbestos-related lung cancer and mesothelioma. Zimbabwe views this study with considerable scepticism in view of the fact that there are long latency periods involved in the above-named diseases and that the current “cases” go a long way back to a time when the control measures implemented were far less sophisticated than they are now.

4.81 Zimbabwe asserts that a risk of exposure may also be incurred by workers or any person, for that matter, during installation, maintenance and repair of asbestos-containing products. The risks involved in the use of asbestos-containing products can be adequately controlled, even taking into account France’s high level of protection against health risks, thus making a ban unnecessary. In fact, the 1998 Task Group report supports this conclusion when stating that “[n]on-friable products and appropriate technological controls greatly reduce fibre release.” It can thus be said that the risk of occupational exposure (i) is a function of the nature of the product and (ii) the risk inherent in that product can in any event be further reduced through appropriate control measures. Regarding the products at issue, i.e. products made from asbestos-cement, the first thing that should be noted is that asbestos-cement does not contain friable asbestos. Moreover, and equally importantly, products made from asbestos-cement are products of high density and thus chrysotile asbestos fibres are firmly blended into the final product. This reduces to a minimum the likelihood of fibres being released into the air and thereby posing a health hazard to human beings. The ILO came to the same conclusion in a report released in 1985: “[l]a manipulation de produits contenant de l’amiante dans lesquels les fibres d’amiante sont solidement fixées dans un liant de telle sorte qu’il ne puisse pas se former de poussières ne présente pas de danger pour la santé.”

4.82 It emerges therefore that when products made from asbestos-cement are used and handled properly, the risks associated with their use are minimal. The recommendation of the 1998 Task Force was to the same effect. It recommended that appropriate control measures be implemented wherever occupational exposure might occur. Among the control measures which might be used to minimize exposure to chrysotile asbestos are engineering controls, special work practices (including workplace hygiene), and protective equipment, such as technical appliances which eliminate or minimize the formation of asbestos dust, as well as protective respiratory equipment or special protective clothing. That risk control is in fact an effective means of dealing with chrysotile asbestos-related health concerns is borne out by the following passage taken from the report of the 1998 Task Group: “[d]ata from industries where control technologies have been applied have demonstrated the feasibility of controlling exposure to levels generally below 0.5 fibres/ml. Personal protective equipment can further reduce individual exposure where engineering controls and work practices prove insufficient.” In light of the foregoing, it is the contention of Zimbabwe that the combined use of high-density products made from asbestos-cement, which inherently are low-risk products, coupled with adequate risk control measures minimize the risk of exposure to asbestos dust. Whatever residual risk may remain does not, in Zimbabwe’s view, justify an outright ban on chrysotile asbestos.
3. Legal Aspects

4.83 It is the submission of Zimbabwe that the French ban of chrysotile asbestos is contrary to WTO rules and should be lifted without any delay. It is the view of Zimbabwe that the French Decree constitutes a technical regulation within the meaning of the Agreement on Technical Barriers to Trade. As such, it must be in accordance with Article 2.2 of the TBT Agreement and hence must not be “more trade-restrictive than necessary to fulfil a legitimate objective”. By totally banning the import of chrysotile asbestos, the French legislation contravenes the express language of this Article. Furthermore, in the event of the French Decree being found to fall outside the ambit of the TBT Agreement, the Decree contravenes the provisions of GATT Article III:4, as it discriminates against imported asbestos in favour of other like products which are used in France for the same purpose. In the same vein, the French Decree cannot be justified under the terms of GATT Article XX(b), as claimed by the EC.

(a) The Agreement on Technical Barriers to Trade

4.84 Zimbabwe disagrees with the view of the EC that the Decree does not fall within the scope of the TBT Agreement. For a mandatory measure to come within the scope of the TBT Agreement, it must be a “technical regulation”. The Decree clearly is a mandatory measure. Notwithstanding the EC’s claim to the contrary, it is the submission of Zimbabwe that the Decree, to the extent that it applies to products containing chrysotile asbestos, qualifies as a technical regulation within the meaning of Annex 1 of the TBT Agreement. The argument of the EC that for the TBT Agreement to be applicable, the Decree should have specified which particular products were covered by the ban is without any merit. It is the view of Zimbabwe that such an interpretation is overly restrictive. Annex 1 of the TBT Agreement talks about “product characteristics” in general. Nowhere does it state that the national legislator should adopt only product-specific regulations. Even ignoring this point, Zimbabwe fails to understand why a Member should be precluded from laying down horizontal rules applicable to a group or groups of products which call for the same regulatory approach. In fact, it appears that there would be little merit in forcing Members to specifically enumerate all products covered by a particular regulation when it is in the nature of things that new products would regularly have to be added to the list due to, for example, technological developments. From a public policy perspective, this would seem to be a rather inefficient and costly approach to adopt.

4.85 Zimbabwe argues that the second reason advanced by the EC in support of its argument that the TBT Agreement is not applicable in this case is also without any merit. According to the EC, the ordinary meaning of the noun “characteristic” supports the view that, for the TBT Agreement to be applicable, product characteristics must be positively defined. Applying this reading of the TBT Agreement to the present case, the EC argues that “not containing chrysotile asbestos” should not be seen as the equivalent of a product characteristic. Zimbabwe finds this reasoning of the EC very tenuous. According to the Shorter Oxford English Dictionary, the noun “characteristic” designates a “distinguishing quality or peculiarity”. Zimbabwe believes that, without doing injustice to these terms, a product’s “distinguishing quality or peculiarity” can lie in the fact that it does not contain asbestos. The absence of any trace of asbestos clearly sets apart a product in terms of its qualities from another product which contains asbestos. In any event, Annex 1 does not actually require positive product characteristics. Zimbabwe submits that its interpretation of Annex 1 is also in conformity with the relevant context of Annex 1 of the TBT Agreement. All the Agreements annexed to the WTO Agreement are part of the relevant context. Thus, Article 2(f) of the Agreement on Rules of Origin obliges Members to ensure that “their rules of origin are based on a positive standard”. From this it follows that where Members wanted to give a special meaning to a term - in this case, to the term “standard” - they used appropriate language to reflect their intention. Members did not adopt that approach as far as Annex 1 of the TBT Agreement is concerned.

4.86 Given the object and purpose of Annex 1 of the TBT Agreement, Zimbabwe wonders what would be the rationale of a rule which compels Members to define product
characteristics positively when all they care about is a negative characteristic. Why, for example, should France have to positively define the characteristics of a host of products when its only regulatory concern is with the asbestos contained in those products? It is the submission of Zimbabwe that its interpretation is also in conformity with the jurisprudence of the WTO Appellate Body. Thus, according to the Appellate Body, the term “measure” as it appears in various WTO agreements is to be understood to include a government’s failure to act. In other words, a “negative” measure, i.e. a failure to act, counts as a measure no less than a “positive” measure. By token of the same reasoning, the term “characteristics” should encompass negative characteristics. In view of the above reasons, Zimbabwe joins Canada in believing that the general term “product characteristics” lends itself to an interpretation which includes negative characteristics.

Having demonstrated that the Decree qualifies as a technical regulation under the TBT Agreement to the extent that it bans products containing chrysotile asbestos, Zimbabwe now turns to show that the same is true also with respect to the Decree’s ban on the use of chrysotile asbestos fibres as such. The EC has expressed the view that the French ban on the production and importation of chrysotile asbestos fibre is not a technical regulation within the meaning of Annex 1 of the TBT Agreement because, just like the ban on asbestos-containing products, the ban on asbestos fibres is general (rather than specific) and lays down negative characteristics (rather than positive ones) or, for that matter, does not lay down any characteristics. As the issues of specificity and of “positive vs. negative standards” have already been discussed, the following submissions will focus on whether or not the French Decree lays down product characteristics with regard to the ban on asbestos fibres. Zimbabwe contends that the matter is more complex than the EC makes it out to be. To be sure, an independent and isolated ban on sales, say, of all cigarettes would not normally be considered a technical regulation. Yet the situation as it presents itself in this dispute is quite unlike that. As Canada rightly pointed out, unlike cigarettes, asbestos fibres per se, i.e. as products in their own right, serve no useful purpose. It is the products containing asbestos fibres which have commercial use and value. By necessary implication, when it comes to dealing with the health hazards of asbestos, the concern of policymakers and the law should be with products containing asbestos fibres, not asbestos fibres, per se. If the products containing asbestos disappear, so will asbestos fibres.

Zimbabwe considers that the Decree is fully consistent with this straightforward principle. The EC does not contest this. On the contrary, the EC sets out the objective of the Decree as follows: “[l]’interdiction de l’amiante, en France et dans d’autres pays, n’a pas pour objectif de supprimer les quelques 0,0002 fibres/ml qui existent ‘naturellement’ dans l’air. L’interdiction vise simplement à protéger l’ensemble des travailleurs et des utilisateurs de l’amiante qui sont souvent exposés à des valeurs très supérieures [...] pour des opérations courantes d’intervention sur des matériaux contenant de l’amiante-ciment.” The EC explains the rationale of its asbestos-control policy in the following terms: “[l]a politique adoptée en France en 1996 vise en tout premier lieu au remplacement des matériaux contenant de l’amiante par d’autres matériaux sans danger [...]”. It clearly emerges from these two quotes that the Decree aims at asbestos-containing products, not at asbestos fibres per se. The inference that can be drawn from this is that the import ban - just like the corresponding ban on domestic production - does not perform an independent function, but a subsidiary one. Indeed, the EC expressly states that nothing would change if the import ban - and, by implication, the ban on domestic production - were lifted. Imported and domestically produced asbestos fibres could still not be sold on French territory - because no products containing them could be sold. The following sentence pinpoints this underlying logic of the French ban: “[l]e but est donc bien d’arrêter la diffusion d’amiante le plus en amont possible”. The ban on asbestos fibres is thus based on considerations of administrative efficiency, which is arguably only a secondary objective pursued by France. Again, this is confirmed by the EC: “[l]’interdiction d’importation a simplement pour but de rendre plus efficace, en termes de contrôle, l’interdiction d’utilisation [which is France’s primary goal]”.

Zimbabwe asserts that, for the foregoing reasons, it should be readily apparent that the ban on chrysotile asbestos fibre is very closely related to the ban on asbestos-
containing products. Assuming the ban on chrysotile asbestos fibres could be viewed in isolation, it could possibly be argued that it does not, *stricto sensu*, lay down product characteristics. As Zimbabwe has demonstrated, however, such a line of reasoning is unwarranted and misses the point. The ban on asbestos fibres is an integral part of the Decree. In fact, it is part and parcel of the same Article of the same Decree. Zimbabwe therefore submits that for purposes of this proceeding there is one single, indivisible regulatory package - the Decree - whose consistency the Panel needs to examine with the TBT Agreement. Zimbabwe is of the view that the Decree falls within the ambit of the TBT Agreement. This view is buttressed by the reasoning of another Panel which faced a comparable situation.

In the *Kodak/Fuji* film case, the Panel had to decide whether a measure that had not been directly brought up under Article 4 of the DSU could nevertheless be within the Panel’s terms of reference. The Panel found that such a measure was not within the Panel’s terms of reference, unless it was “subsidiary” or “closely related” to the measure that was properly before the Panel.155 By way of analogous reasoning, Zimbabwe argues that the French ban on asbestos fibres is “subsidiary” and “so closely related” to the ban on asbestos-containing products - which, as shown, qualifies as a technical regulation within the meaning of the TBT Agreement - that it can reasonably be found to form an integral part of the latter, and thus constitute a technical regulation in, and of, itself.156

4.90 Zimbabwe further submits that treating the ban on asbestos fibres and the ban on asbestos-containing products as separate and “unrelated” could give rise to unreasonable results. Such a situation could in fact arise in the present case. It could be envisaged, for instance, that the ban on asbestos fibres might be found to be consistent with the provisions of the GATT, while the ban on asbestos-containing products might be found to violate the provisions of the TBT Agreement because - to use but one example - it is more trade-restrictive than necessary to fulfil a legitimate governmental objective. Zimbabwe submits that such an outcome would be unreasonable and could undermine the practical effectiveness of the TBT Agreement. Taken to its logical conclusion, such a situation would imply, on the one hand, that France could not produce asbestos-containing products domestically as a result of the ban on imported or domestically-produced asbestos fibres. On the other hand, France would be required to lift its ban on imports of asbestos-containing products and adopt instead a less trade-restrictive measure which, in practice, would mean that a certain quantity of asbestos-containing products would cross the border into French territory. France would thus have no choice but to idly sit and watch as other countries take advantage of the business opportunities offered by the French domestic market. Zimbabwe is of the view that the drafters of the TBT Agreement did not and could not have intended such a result.

4.91 It is therefore the submission of Zimbabwe that the TBT Agreement applies to the French Decree in its entirety, i.e. with regard to the ban on asbestos-containing products as well as the ban on asbestos fibres. The French legislation does not meet the requirements of Article 2.2 of the TBT Agreement, as amply demonstrated by Canada. Zimbabwe adopts the arguments presented by Canada in this connection and would like to support the views expressed therein by also relying on the arguments presented below on whether or not the French measure is necessary within the meaning of GATT Article XX(b).

(b) The General Agreement on Tariffs and Trade

(i) *Article III of the GATT*

4.92 Zimbabwe argues that, in the alternative, and in addition to the claimed violations of the TBT Agreement, the Decree violates GATT Article III:4. Zimbabwe submits that chrysotile asbestos fibres and, at a minimum, cellulose fibres, aramid fibres and glass fibres are “like products” within the meaning of Article III:4. The EC confirms that cellulose and aramid fibres count among those fibres which are most frequently used to substitute asbestos fibres in the manufacture of cement.157 Cellulose, aramid and glass fibres are all produced in France.158 Whereas they may lawfully be sold in that country, the importation and sale of asbestos fibres is prohibited. There is thus no doubt that asbestos fibres are accorded “less favourable treatment” than cellulose, aramid and glass fibres, despite the fact that they are “like products”159.
Zimbabwe notes that the EC contests that asbestos fibres, cellulose, aramid and glass fibres are “like products” within the meaning of Article III:4. It is well established in WTO jurisprudence that the determination of whether or not products are “like products” must be made in accordance with such criteria as the products’ physical characteristics and the products’ end-use.\(^{160}\) It is equally clear from WTO jurisprudence that any such determination can only be made on a case-by-case basis, i.e. taking into account the specific and unique circumstances of each case.\(^{161}\) Regarding the first criterion, i.e. physical characteristics and properties, the EC claims that cellulose, aramid and glass fibres are not sufficiently similar to asbestos fibres in that their chemical composition is different. Zimbabwe wishes to recall that the EC has acknowledged that the chemical composition of all varieties of asbestos fibres is different as well. This did not preclude the EC, however, from concluding that chrysotile asbestos fibres and amphibole asbestos fibres were “like products”. Zimbabwe submits that the same logic applies and extends to cellulose, aramid and glass fibres.

Even ignoring the inconsistency in the reasoning of the EC, Zimbabwe does not believe that the differences pointed out by the EC are significant enough to make the relevant products “unlike” within the meaning of Article III:4. Zimbabwe wishes to recall, first of all, that “likeness” does not require that products be “identical in all respects”.\(^{162}\) The second thing that should be noted is that the significance that is attached to differences in physical characteristics depends on the particular circumstances of each case. In this case, as has previously been stated, the starting-point of any analysis must be the fact that chrysotile asbestos fibres, as products in their own right, serve no useful purpose.\(^{163}\) Chrysotile asbestos fibres are predominantly used as “inputs” in the manufacture of fibre-cement products. It follows that substitute fibres like cellulose, aramid or glass fibres, on the one hand, and asbestos fibres, on the other hand, should not be compared to each other as products in their own right. Instead, asbestos fibres and the relevant substitute fibres should be compared to each other as products incorporated into cement. It is obvious that if this approach is adopted, as it should be, the differences identified by the EC become minor ones and irrelevant. The EC essentially makes the point that cellulose and aramid fibres are, on average, less fibrillose and larger in diameter than asbestos fibres and that only asbestos fibres are internationally recognized as “category I” products, i.e. as products that have been shown to cause cancer. With regard to these alleged varying degrees of health risk associated with the fibres at issue, it should be noted that whatever differences exist between the relevant products become far less relevant when the fibres are blended with other materials to produce cement and other related products.\(^{164}\) As explained by Zimbabwe, any remaining risks arise from improper handling and manipulation of cement-products and not from the cement-products themselves. Beyond that, Zimbabwe is not convinced that much significance should be attached to the fact that only asbestos fibres are listed by the WHO as a “category I” product. In fact, even the EC concedes that there is a lingering uncertainty about the risks involved in the use of alternative fibres. Zimbabwe submits that the fact that there are to date no known negative effects on human health from the use of alternative fibres does not necessarily mean that they are risk-free.\(^{165}\) Zimbabwe notes that the EC shares that view, for it expressly acknowledges that “… un risque indétectable n’est pas égal à une absence de risque”.\(^{166}\)

With regard to the second criterion, i.e. commonality of end-uses, Zimbabwe submits that asbestos, cellulose, aramid and glass fibres serve “substantially identical end-uses”.\(^{167}\) Their chemical resistance and reinforcing capabilities make them almost perfect substitutes for asbestos fibres. It is therefore not the case that chrysotile asbestos fibres are unique products, as the EC would have the Panel believe. As previously noted, the EC, in fact, acknowledges that cellulose and aramid fibres are commonly used substitutes for asbestos fibres.\(^{168}\) Moreover, like Canada, Zimbabwe believes that the structure of the Decree is at least suggestive of the substitutability of asbestos fibres with other fibres. This becomes clear if the French Decree is seen in terms of the functioning of the political process. If very close substitutes had not been available to the principal users of asbestos fibres at the time the Decree was signed into law, it is reasonable to assume that they would have lobbied the French Government and in all likelihood would have secured a broader exception (allowing the continued use of asbestos fibres) than the one that is now in the De-
In light of the foregoing considerations Zimbabwe believes that asbestos fibres and cellulose, aramid and glass fibres should be regarded as “like products” within the meaning of GATT Article III:4.

(ii) Article XX of the GATT

Zimbabwe argues that the Decree is not justified under paragraph (b) of Article XX because it is not “necessary to protect human […] health”. More particularly, the Decree does not satisfy the necessity requirement. GATT 1947 case law has established that a measure qualifies as “necessary” within the meaning of Article XX if there is “no alternative measure consistent with the General Agreement, or less inconsistent with it, which [a Member] could reasonably be expected to employ to achieve its […] policy objectives”. Zimbabwe believes that it is sufficient for it to establish that - even assuming that asbestos fibres posed more of a health risk to humans - there are less trade-restrictive measures available to France to achieve its health objective. The EC claims that in order for France to achieve its health policy objective there was no measure reasonably available to it other than an outright ban on chrysotile asbestos fibres. In particular, the EC submits that control measures used to minimize exposure to chrysotile asbestos fibres are not sufficient to ensure that France reaches its high level of protection. It also argues that control measures are impracticable in the case of the large group of “secondary users” of asbestos fibres, i.e. those workers and do-it-yourself people who, in the absence of control measures, may be exposed to chrysotile asbestos dust during installation, maintenance and repair of products containing chrysotile asbestos. The problem is compounded, according to the EC, by the fact that in many instances “secondary users” do not have any information as to whether they are dealing with products that contain asbestos. The EC submits that even if they were given that information, control measures are costly and turn what would otherwise be a simple operation into a costly, complicated and awkward one. Furthermore, the EC believes that “une fois mis sur le marché, il n’existe plus aucun moyen raisonnable de contrôler l’usage de l’amiante et, en particulier, de contrôler des opérations banales (découpage, sciage …) que de nombreuses personnes peuvent être amenées à réaliser”.

Zimbabwe is not convinced by the arguments of the EC. First of all, regarding the effectiveness of control measures, Zimbabwe believes that the observance of certain work practices and the use of technical appliances in accordance with the ISO standard 7337, for example, would be sufficient to meet the maximum exposure level acceptable to France. The EC argues that, even where special technical equipment is used when high-risk activities are undertaken, peak exposure levels to asbestos would still exceed the French maximum level. What the EC fails to mention, however, is that, as argued by Canada, the wearing of protective respiratory equipment and humidification of the materials during those activities could significantly reduce the exposure - so much so, in fact, that the respect of the French maximum level of exposure would be ensured. Regarding the argument of the EC that mandatory control measures are impracticable because they are too costly, Zimbabwe contests the relevance of such considerations. After all, whether or not these costs are too high, is a matter to be left to the dictates of the market. If the producers of asbestos-cement face insufficient demand for their products because of expensive control measures imposed on their customers, they will go out of business or diversify into the production of cement using alternative fibres. Likewise, Zimbabwe does not see any merit in the argument that control measures make certain work procedures complicated and awkward. Where certain practices are imposed by law, the question of whether they are appreciated by those who must follow them becomes meaningless. It certainly does not in itself provide a rationale for trade-restrictive measures.

While Zimbabwe recognizes that it may not be readily apparent to an inexperienced person whether or not he/she is handling a product containing asbestos fibres, it is by no means justification for instituting a far-reaching ban on products which might contain asbestos fibres. It is the contention of Zimbabwe that it would be possible under the WTO legal framework for Members to impose a disclosure requirement, which would enable purchasers to make informed decisions as to whether or not they purchase prod-
ucts containing asbestos fibres. Where the materials have already been installed or incorporated, say, in a building, Zimbabwe does not see why there could not be, for instance, an asbestos warning message next to the evacuation instructions on a notice board of that building. Moreover and specifically with respect to the work of plumbers, electricians and the like, Zimbabwe does not see why the owner of an installation or building could not be required to make available some sort of map which would document in which parts of the installation asbestos is present. With reference to the concern of the EC that the use of asbestos-containing products cannot sufficiently be controlled, especially when it comes to "secondary users" of such products, Zimbabwe again does not think that banning all imports of such products would solve the problem. In fact, it would raise more problems than it would solve. To begin with, if indeed the French Government is so concerned about do-it-yourself users of asbestos-containing products, it could have easily banned the sale of such products in all do-it-yourself outlets. Furthermore, as a supporting measure, it could have also restricted the handling of asbestos-made products to certified experts, thus eliminating contact with asbestos by inexperienced people. The protection of workers, such as electricians and plumbers, could also have been ensured relatively easily. The French Government could have, for example, required certification, which would only be bestowed upon an individual once he/she had successfully followed information and training courses on the use and handling of asbestos-containing products. The French Government could also have laid out the precise work practices and technical appliances that must be used in all contacts with asbestos-containing products. To ensure compliance, the regulations could authorize the imposition of heavy fines or a custodial sentence in the event of a wilful disregard of the government's regulations. Needless to say, it is also open to a Member to run information campaigns, so as to raise awareness among workers of the risks of asbestos fibres and the procedures to be observed in all contacts with such fibres. It is clear from the foregoing that the French Government had a number of alternative measures at its disposal which would have interfered less with trade and at the same time would have assisted in realizing its overriding objective of protecting the health and safety of its citizens.

V. PANEL’S CONSULTATION WITH SCIENTIFIC EXPERTS

A. DETERMINATION OF THE PROCEDURE

5.1 The Panel noted that the dispute before it raised scientific and technical issues. At the first substantive meeting, the Panel informed the parties of its intention to seek the opinion of individual scientific experts except where, in the light of the parties’ written rebuttals, it concluded that such a procedure was not necessary. The areas in which the Panel wished to obtain information included the circumstances of exposure to chrysotile asbestos and the associated risks, as well as the effectiveness of the controlled use of chrysotile. The Panel invited the parties to submit their comments to it in writing, particularly regarding the areas on which the experts were to be consulted, the possible approaches to such a procedure and the international or other bodies that could usefully be consulted in order to identify suitable experts.

5.2 In a letter to the Panel dated 14 June 1999, Canada proposed, in regard to the possible approaches to a procedure for consultation with individual experts, that five requirements should be met, each one intended to ensure observance of the right of the parties to be heard at all stages of the procedure: (i) the Panel should consult the parties on the choice of scientific experts; (ii) the Panel should seek the opinion of the parties concerning the formulation of the questions to be put to the experts; (iii) the Panel should provide the parties with an opportunity to make written comments on a draft report by each of the experts; (iv) the parties should be able to question each of the experts on the content of his final report at a meeting with the Panel; (v) the parties should be given the possibility to make written comments on the conclusions set out in the final report of each expert and their legal implications. Like the Panel, Canada also believes that the areas on which the scientific experts should be consulted ought to include the circumstances of exposure to chrysotile asbestos and the risks associated with present applications as well as risk management by the controlled use of chrysotile asbestos. The experts should also be consulted.
in two other areas, namely the comparative toxicity of the different types of asbestos fibres and substitute fibres, and risk assessment methods, including the question of whether there are exposure thresholds below which the risk is undetectable in practice. In Canada’s opinion, there are four specializations that in one way or another cover the above-mentioned areas and from which the experts should be drawn. These are toxicology, epidemiology, risk analysis and occupational health. Given the scientific characteristics of the dispute, Canada would wish that each question be submitted to more than one expert, and that each expert submit an individual report. As regards the international institutions that could usefully be approached in order to identify suitable experts, Canada believes that they should be consulted in order to come up with a sampling of experts in the above-mentioned domains. The main selection criteria and hence the best guarantee of impartiality should be that experts must have conducted recognized and independent research into chrysotile asbestos. The international organizations that could be approached included the World Health Organization, the International Labour Office and the International Organization for Standardization. Once a list of prospective candidates has been drawn up with the help of the international organizations, the parties should then be able to submit their own list of names of specialists who could act as scientific experts in the areas mentioned above.

5.3 In a letter dated 14 June 1999, the European Communities were of the opinion that the scientific issues raised in this dispute were simple and clear. The DSU rules on the burden of proof also provided the Panel with sufficient guidance in dealing with the factual and scientific issues raised by the parties to the dispute. With respect to the general selection procedures and criteria, the European Communities believed that the Panel’s use of experts for obtaining scientific and technical advice should respect the general principles of law. In particular, it should be transparent, avoid conflicts of interest, reinforce the integrity of the dispute settlement mechanism and foster public confidence in the outcome of the dispute. In the view of the European Communities, the Panel can in this case establish only an expert review group under the terms of Appendix 4 to the Dispute Settlement Understanding. Indeed, the measure at issue in the present dispute is one that must be examined strictly in terms of the GATT 1994, to the exclusion of the Agreement on Technical Barriers to Trade. Article 13:2 of the DSU provides as follows: ‘… With respect to a factual issue concerning a scientific or other technical matter raised by a party to a dispute, a Panel may request an advisory report in writing from an expert review group. Rules for the establishment of such a group and its procedures are set forth in Appendix 4’. The establishment of an expert review group is the only option provided under the DSU for panels wishing to obtain information on scientific matters. The first sentence of Article 13:2 applies only to situations in which a Panel wishes to obtain factual or technical but not scientific information. In their context, the ordinary meaning of the terms, as well as the object and purpose of Article 13:2, first and second sentences, clearly lead to the conclusion that panels are not authorized to deviate from the procedure laid down by Appendix 4 to the Dispute Settlement Understanding. Whether the request comes from a party or arises at the initiative of the Panel itself makes no difference. Strictly scientific matters cannot be resolved by means and/or procedures other than those envisaged in Appendix 4 to the Dispute Settlement Understanding. The chapeau to Appendix 4 to the Dispute Settlement Understanding also confirms this interpretation by providing that the rules and procedures set forth in the Appendix ‘... shall apply to expert review groups established in accordance with the provisions of paragraph 2 of Article 13’, that is, regardless of whether the Panel bases itself on the first or second sentence of that Article. This interpretation is supported by the fact that, if the Agreement on Technical Barriers to Trade (TBT) should be applicable (which is not the case), Article 14:2 of that Agreement explicitly prescribes that panels establish only a technical expert group (which is equivalent to an expert review group). In such a case, the procedural rules set forth in Annex 2 to the TBT Agreement must apply. Annex 2 to this latter agreement and Appendix 4 to the Dispute Settlement Understanding are almost identical. Moreover, by virtue of Article 1:2 and Appendix 2 to the DSU, only Article 14:2 of the TBT Agreement is applicable.

5.4 The European Communities also point out that the previous cases in which panels requested the opinion of scientific experts all came under the Agreement on the appli-
cation of Sanitary and Phytosanitary Measures, which is not applicable in this case. Those previous cases are therefore irrelevant to the present dispute. The dispute concerning Shrimp is the only other case for which the opinion of scientific experts was requested under the GATT 1994. But this example per se is not enough to set a valid precedent applicable to all cases, especially because the parties to the Shrimp dispute apparently did not request the exclusive application of Appendix 4 to the Dispute Settlement Understanding. The result is that, in the present case, should the Panel decide to seek the scientific opinion of external experts, it can do so only under Article 13.2, second sentence of the Dispute Settlement Understanding or under Article 14.2 of the TBT Agreement.

5.5 According to the European Communities, Appendix 4 to the DSU and/or Annex 2 to the TBT Agreement lay down almost identical rules on the establishment of an expert review group. These rules must all be observed in this dispute. Moreover, to ensure that the aforementioned principles are respected, the European Communities believe that the Panel should observe the following specific criteria when choosing scientific experts: (i) the experts should not be citizens of the parties to the Dispute; (ii) the Panel should select scientific experts in different areas of specialization in order to ensure coverage of all the areas identified by it. These areas are: the human health hazards posed by asbestos, especially chrysotile asbestos; the inapplicability of a threshold; the circumstances of exposure and the question as to whether what is known as “controlled use” can eliminate the potential hazards to human health; (iii) the European Communities believe that if the Panel decides to request information, it should consult at least five experts so that more than one expert will have the requisite expertise and provide answers to the questions in the various areas identified by the Panel. In the light of the number of experts that the Panel should consult, only scientists with proven expertise in the realm of asbestos should be selected; (iv) the experts should be drawn mainly if not exclusively from the International Agency for Research on Cancer (IARC), a specialized agency of the WHO. The IARC has studied asbestos from all possible angles and should therefore be well placed to propose experts covering all the areas in which questions could be posed. The Panel should also explore the possibility of consulting the International Labour Office (ILO) in the event that the IARC is unable to cover all the areas in question; (v) the experts chosen must have no link whatsoever, present or past, with the industry producing asbestos or substitute products. They must furthermore clearly demonstrate the lack of any conflict of interest. The parties should receive at the outset the Curricula Vitae of all the candidates proposed and should have at least ten working days in which to verify the skills, expertise and possible conflicts of interest of the candidates; (vi) the Panel should also request the opinion of the parties as to the aim of the consultation with experts, the type and nature of the questions to be put to them; (vii) the aim of the consultation should be to further the knowledge of the scientific considerations germane to this dispute. Therefore, and in accordance with the provisions of the Dispute Settlement Understanding, the questions to be put by the Panel must have a direct and strict bearing only on the scientific aspects of the case. The questions may not relate to legal problems nor to any problem of interpretation of any WTO Agreement under examination.

5.6 Having taken cognizance of the comments from the parties, the Panel decided to consult the experts on an individual basis, pursuant to paragraphs 1 and 2, first sentence, of Article 13 of the Understanding on Rules and Procedures Governing the Settlement of Disputes. The Panel convened the parties to a meeting on 10 July 1999 to acquaint them with the procedure it intended to follow and to give them the opportunity to state their opinions on the matter. The Panel recalled Article 13 of the Dispute Settlement Understanding which, among other things, provides that:

“Each panel shall have the right to seek information and technical advice from any individual or technical body which it deems appropriate.” […]

“Panels may seek information from any relevant source and may consult experts to obtain their opinion on certain aspects of the matter.”
5.7 At that meeting, the Panel told the parties that, in its opinion, Article 13 of the Dispute Settlement Understanding empowered it to seek such information and technical advice as it deemed fit in a given matter; in particular, a panel was free to determine whether it was necessary or appropriate to establish an expert review group. In the case at hand, the consultation of experts acting in their own right seemed to it to be the most appropriate form of consultation. The Panel intended to seek information concerning the circumstances of chrysotile exposure and the attendant hazards. In the circumstances, the Panel indicated that it would structure its questions around the following main topics: the pathogenicity of chrysotile, the relative pathogenicity of amphiboles, chrysotile and substitute products; the assessment and management of risks associated with the use of chrysotile; the effectiveness of controlled use of chrysotile.

5.8 The Panel then presented to the parties the procedure that it intended to follow, which is the same used by previous panels that had consulted experts selected on an individual basis:

- The experts will be placed under the authority of the Panel. They will be consulted on a personal basis and not as representatives of a government or organization. Their opinion will be strictly in the nature of advice; it will not be binding on the Panel;
- the number of experts to be chosen by the Panel will be decided depending on the number of matters on which an opinion will be sought, as well as the number of matters on which each expert can give an opinion;
- the Panel intends to request names from the World Health Organization (WHO), the International Labour Organization (ILO), the International Programme on Chemical Safety (IPCS), the International Agency for Research on Cancer (IARC), the International Organization for Standardization (ISO), and from the parties;
- the Panel does not intend to appoint experts who are citizens of one or other of the parties to the dispute, unless the parties consent to their appointment or the Panel believes that it would otherwise be impossible for it to secure the specialized scientific advice needed;
- the Secretariat will request the persons suggested to submit a curriculum vitae. The curricula vitae will be transmitted to the parties. The parties may not establish contact with the experts suggested;
- the parties will have an opportunity to make comments and to state any major objections they may have to any expert under consideration. The Panel will inform the parties of the experts it chooses;
- the experts will receive all the relevant elements of the communications on a confidential basis;
- the Panel will prepare draft questions for the experts. They will be communicated to the parties. The parties will have the opportunity to comment on the questions proposed or to suggest additional questions before they are sent to the experts. The Panel will then draw up a definitive list of questions which will be sent to the experts and simultaneously to the parties;
- each expert will receive all the questions. He will be requested to reply to the questions falling within his sphere of competence and, if necessary, to indicate the areas on which he does not feel competent to
reply. The experts will be invited to provide written answers; copies of those answers will be transmitted to the parties. The parties will have an opportunity to make written comments on the replies from the experts and the replies will be included in the Panel’s final report;

- should the Panel deem it fitting, either on its own initiative or at the request of a party, a meeting may be held with the experts immediately before the second substantive meeting. Before the meeting, the Panel will ensure that: (i) experts are made privy to the parties’ comments on their replies; (ii) the experts each receive the replies of the other experts to the Panel’s questions;

- the minutes of the meeting with the experts will be submitted to the parties and to the experts so that they may make corrections. The corrected version will be attached to the Panel’s final report.

5.9 The Panel gave the parties the opportunity to transmit their written comments to it.

5.10 In a letter dated 19 July 1999, Canada recalled all the points that it had notified to the Panel in its letter of 14 June 1999. Canada agrees with the Panel as to the nature of the information and advice that it intends to seek from the scientific experts. It nevertheless believes that the experts best qualified to reply to the Panel’s questions concerning the circumstances of exposure to chrysotile and the associated hazards are to be found in the areas of toxicology, epidemiology, risk assessment and occupational safety. In addition to the opportunity given to the parties to make written comments on the experts’ replies, the Panel should also provide for the possibility of a final written submission by the parties following the second substantive meeting. As regards the stipulation that the scientific experts may not be citizens of any of the parties to the dispute, Canada believes that this procedural rule, established in Appendix 4 to the Dispute Settlement Understanding, normally applies only to the establishment of an expert review group. In the Hormones case, the Appellate Body stated in that connection: “… once the Panel has decided to request the opinion of individual scientific experts, there [was] no legal obstacle to the Panel drawing up, in consultation with the parties to the dispute, ad hoc rules for those particular proceedings”177. As the agreement of the two parties to the dispute is required if the selection of citizens of one of the parties is to be allowed, Canada is surprised at the refusal of the European Communities to allow the selection of their citizens. Canada is prepared to consider the selection of experts who are citizens of the European Communities despite the refusal of the European Communities to consider experts from Canada. In this dispute, if the citizens of the parties are automatically excluded, the Panel risks facing a situation in which it will be unable to select the experts with the best scientific knowledge considering the nature of the advice being sought. Canada therefore requests the European Communities and the Panel to reconsider their decision with regard to the non-participation of citizens of the parties.

5.11 Moreover, Canada cannot accept that, as demanded by the European Communities, the experts must clearly demonstrate the absence of any conflict of interest. It is not incumbent upon a prospective expert to prove his impartiality, instead he is merely required to fill out a disclosure form concerning his interests, relationships and any matters that may affect his independence. This form is provided for in the document entitled Rules of Conduct for the Understanding on Rules and Procedures Governing the Settlement of Disputes.178 Once the persons approached as potential experts have filled out their disclosure forms, the parties to the dispute may oppose any candidate who has disclosed an interest, relationship or matter that may place him in a situation of conflict of interest. The Panel is empowered to decide whether the information disclosed in the form really places a candidate expert in a situation of conflict of interest and to uphold a party’s objection to an expert’s candidature. The approach taken by the Panel in the Shrimp case should be followed in this instance. Having noted that in their disclosure forms three of the experts approached had disclosed what might be considered as potential conflicts of interest, the
Panel nevertheless decided to confirm their appointments “being of the view that the disclosed information was not of such a nature as to prevent the individuals concerned from being impartial in providing the scientific information expected of them. The Panel has also taken into account the disclosed information when evaluating the answers provided. The Panel underlined that, in making its choice, it had been guided primarily by the need to gather expertise of the best quality and covering as wide a field as possible. In [the circumstances specific to this case], it was difficult – if not impossible – to reconcile this need with an agreement by all the parties to the dispute on each and every individual concerned”.179 Canada is surprised at the European Communities’ insistence on the absence of any link between the experts and producers of chrysotile asbestos but not between the experts and anti-asbestos pressure groups. No one opposes the principles of independence and impartiality of experts or the observance of the rules on conflicts of interest. The single pertinent consideration remains the way in which these principles should be applied in this particular instance.

5.12 In a letter dated 19 July 1999, the European Communities took note of the Panel’s decision to consult individual scientific experts pursuant to Article 13:1 of the Dispute Settlement Understanding. The European Communities contest the legal basis of the Panel’s decision. Under the international customary principles of treaty interpretation, a systematic interpretation of Articles 13:1 and 13:2 of the Dispute Settlement Understanding suggests that as far as scientific matters are concerned, the preferred option in the Dispute Settlement Understanding is the establishment of an expert review group. The term “scientific matter” appears only in the second sentence of Article 13:2 of the Dispute Settlement Understanding, which envisages only the constitution of an expert review group. The drafting history of the WTO Agreements also confirms this interpretation.180 The three previous cases in which panels sought the opinion of scientists in their own right all had to do with matters arising under the SPS Agreement, Article 11:2 of which expressly mentions “scientific” matters and envisages the possibility of consulting experts individually.181 Canada furthermore requests that the TBT Agreement be applied to the measure at issue here. It is worth noting that Article 14:2 of the TBT Agreement provides only for the possibility of consulting a technical expert group. This Agreement contains no provision equivalent to Article 13:1 of the Dispute Settlement Understanding or to Article 11:2 (first sentence) of the Dispute Settlement Understanding and Article 11:2 of the SPS Agreement. This difference is not accidental.182 It denotes the clear intention of the WTO Members to settle scientific or technical matters in the framework of the TBT Agreement only by establishing an expert review group. The decision of the Panel to consult experts on a personal basis is also contrary to Article 1:2 of the Dispute Settlement Understanding, which provides as follows:

“To the extent that there is a difference between the rules and procedures of this Understanding and the special or additional rules and procedures set forth in Appendix 2, the special and additional rules and procedures in Appendix 2 shall prevail.”

5.13 As explained above, there is a clear difference between Article 13:1 and 13:2 (first sentence) of the DSU, invoked in this case by the Panel, and Article 14:2 of the TBT Agreement. The special rules and procedures mentioned in Appendix 2 to the DSU, namely Article 14:2 of the TBT Agreement, which provides for the establishment of a technical expert group, should thus be applied in the present case, should the Panel judge the TBT Agreement to be applicable.183 Therefore, the European Communities consider the Panel’s decision contrary to the letter, object and purpose of Article 14:2 of the TBT Agreement (if the latter is applicable), in conjunction with Article 1:2 of the DSU, and to Article 13:2 (second sentence) of the DSU. Besides, from a systematic point of view, the Panel’s decision renders useless and obsolete the provisions of the Dispute Settlement Understanding and of the TBT Agreement regarding expert review groups, which are clearly the option preferred by WTO Members and the only one for which rules of procedure have been drawn up in the WTO for the settlement of “scientific” questions.184 At this stage, the
European Communities are therefore obliged to reserve all their rights on this issue. They would also request the Panel, in keeping with current WTO practice and for the sake of transparency and due process, to state in writing the criteria and the reasons for its decision to call on individual scientific experts and the reasons for which it has not entertained the arguments put forward by the European Communities, and to communicate this information to the parties to the dispute.

5.14 As regards the type of scientific background and specializations, the European Communities take the view that the experts should be cancer specialists, in particular in lung cancer and mesothelioma. They should also be epidemiologists experienced in the area of asbestos and cancer. The European Communities are not clear as to what type of scientific discipline would encompass those persons who would be required to provide advice regarding “risk evaluation and management in the use of chrysotile” and “the effectiveness of the controlled use of chrysotile”, nor what type of technical expertise they should have. If such experts exist, they should be able to provide information about all the categories of persons who could come into contact with asbestos and asbestos-containing products, such as those working in maintenance, repair and construction (for example, carpenters, plumbers, heat repairers, workers in insulating materials, do-it-yourself enthusiasts, etc.). The European Communities believe that the scientists chosen should also have expertise in the inspection of houses, buildings and factories for the presence and possible removal of asbestos. Obviously, such experts cannot be allowed to have any link, whether direct or indirect, with the industries producing asbestos or those producing the equipment for reducing the risk of asbestos fibre inhalation. Such a link would seem particularly possible if the experts were to be designated by the ISO. The European Communities consider that at least two experts should be designated for each scientific domain and each area of questions. That is a minimum prerequisite for a balanced view and for not being entirely dependent on the views of just one person. At all events, the overall number of experts should not be less than six.

5.15 The European Communities have expressed their wish to receive copies of the letters to be sent by the Panel to the aforementioned institutions under this point and of their replies. The experts appointed should not be nationals or residents of the parties to the dispute. The European Communities consider that all the candidates must submit a detailed curriculum vitae in time so as to enable the parties to verify their scientific credentials, experience and independence. The candidates must therefore clearly indicate in their curriculum vitae whether in the course of their professional life they have worked for or provided advice, in whatever form, to the industries producing asbestos, asbestos-containing products and substitute products or to the industry producing “controlled use” equipment. In addition, the selected experts must complete a disclosure form concerning potential conflict of interest, pursuant to the Rules of Conduct for the Understanding on Rules and Procedures Governing the Settlement of Disputes adopted (WT/AB/WP/3, Annex II, page 16, 28 February 1997). The disclosure form must contain all the information indicated in the illustrative list appearing in Annex II to the Rules of Conduct mentioned above. It should also explicitly contain information as to whether the expert has done any type of paid or unpaid work (scientific research, consulting, expert advice, participation in the board of directors or board of management, etc.) for the enterprises engaged in the extraction, production, processing of or trade in asbestos, asbestos-containing products or substitute products, or for enterprises producing the equipment intended for “controlled use”.

5.16 It is the opinion of the European Communities, that the Panel should, for example, request that the disclosure form further indicate: (i) the expert’s professional situation (job in an enterprise or institute connected to the asbestos, substitute products or “controlled use” equipment industries); (ii) whether the expert is a member of the board of directors, board of management or any other supervisory body within an enterprise, association, institution or interest group linked with the industries producing asbestos, substitute products or equipment for “controlled use”; (iii) whether he has conducted scientific research or provided expert advice at the request of or under contract to an enterprise, association, institution or interest group connected with the industries producing asbestos,
substitute products or a “controlled use” equipment. If the aforementioned clarifications and information are not given in the curriculum vitae and in the disclosure form, the parties will not be in a position to exercise their rights and make the type of comments being requested of them by the Panel. Therefore, the European Communities consider that the issue of the scientific credentials, experience and, in particular, that of the independence and impartiality of the experts, are of paramount importance and that they need to be reflected in the Panel’s decision on the selection and consultation of scientific experts. They therefore wish to reserve their rights until completion of the selection procedures. The parties should be allowed sufficient time to enable them to make effectively known to the Panel their views on the above issues. Specifically, they should be given sufficient time to make known their views on the list of potential experts to be chosen by the Panel and to submit their comments on the written replies from the experts to the questions put to them by the Panel.

5.17 In a letter to the parties dated 2 August 1999, the Panel confirmed its intention to consult experts individually, in application of Article 13 of the DSU. The Panel carefully examined the arguments advanced by the parties concerning the expert consultation procedures, in particular, the European Community argument that Article 13.2 of the DSU Agreement requires the constitution of a technical expert group as envisaged in Appendix 4 to the DSU for the purposes of consultation with experts on scientific matters. Article 13 of the DSU provides, among other things, that “each Panel shall have the right to seek information and take advice from any individual or body which it deems appropriate” and that “Panels may seek information from any relevant source and may consult experts to obtain their opinion on certain aspects of the matter”. In addition, Article 13.2 prescribes that panels “may” request an advisory report in writing from an expert review group specifically though not exclusively to examine a factual issue concerning a scientific matter. The Panel deems this text to allow for the establishment of such an expert group, while not ruling out consultation of experts on an individual basis, both with regard to a scientific matter “or other technical matter”. This interpretation of Article 13.2 of the DSU seems to the Panel to be perfectly in line with the text of this provision, interpreted in accordance with Article 31 of the Vienna Convention on the Law of Treaties, and with the interpretation given by the Appellate Body that Article 13 of the DSU does not prevent panels from consulting with individual experts and leaves to the sound discretion of a panel the determination of whether the establishment of an expert review group is necessary or appropriate.

5.18 The Panel also considered the European Community’s argument that, if the measure at issue should be deemed to fall under the TBT Agreement, which the Communities contest, Article 14.2 of that Agreement would require the establishment of an expert review group for any scientific or technical matter, and the EC position that pursuant to Article 1:2 of the DSU, that provision would prevail over those of Article 13 to the DSU. Article 14.2 of the TBT Agreement is among the provisions mentioned in Appendix 2 to the DSU and which, under Article 1:2 of that Understanding, will prevail over the provisions of the Understanding to the extent that there is a difference between the two. The Panel notes, however, that it is only “to the extent that there is a difference” between the rules and procedures of the Understanding and a special or additional rule or procedure in Appendix 2 to the DSU that the latter will prevail. Yet, as stated by the Appellate Body, it is only where the provisions of the DSU and the special or additional rules of Appendix 2 cannot be read as complementing each other that the special or additional provisions will prevail over those of the DSU, that is, in a situation where the two provisions would be mutually incompatible. In the present case, Article 14.2 of the TBT Agreement provides that a panel “may” establish a technical expert group. Like Article 13.2 of the DSU, this text envisages the possibility of establishing a technical expert group and lays down the procedures that would be applicable in the event. Nevertheless, it does not exclusively prescribe the establishment of a technical expert group, and this possibility, in our opinion, is not incompatible with the general authorization given under Article 13 of the DSU to consult with individual experts. The two provisions can be read as complementing each other.
The Panel believes that in this case the consultation of experts on an individual basis is the more appropriate form of consultation, inasmuch as it is the one that will better enable the panel usefully to gather opinions and information on the scientific or technical issues raised by this dispute. Considering in particular the range of areas of competence that might be required, it is appropriate in this case to gather information and different individual opinions rather than asking for a collective report on the various scientific or technical matters in question. In the light of the foregoing, the Panel wishes to underline that its decision to consult experts on an individual basis is without prejudice to the applicability of the TBT Agreement to the measure in question, on which the parties disagree.

**B. SELECTION OF EXPERTS**

The Panel has requested the assistance of five institutions in identifying experts. The institutions concerned are the World Health Organization (WHO), the International Labour Organization (ILO), the International Programme on Chemical Safety (IPCS), the International Agency for Research on Cancer (IARC) and the International Organization for Standardization (ISO). The parties have also submitted names to the Panel. The Secretariat then requested those of the proposed experts who were prepared to participate to submit to it a detailed *curriculum vitae*. Those *curricula vitae* were forwarded to the parties, who were able to convey to the panel their comments concerning the potential experts and to indicate, where appropriate, whether they had any major objections to any of them. Upon careful examination of the *curricula vitae* and the comments of the parties, the Panel accepted the following four experts, whose nominations were not opposed by the parties:

- Dr. Nicholas H. de Klerk, Senior Research Fellow, Department of Public Health, University of Western Australia, Australia;
- Dr. Douglas W. Henderson, Professor of Pathology, Head of the Department of Anatomical Pathology, Flinders Medical Center and The Flinders University of South Australia, Australia;
- Dr. Peter F. Infante, Director, Office of Standards Review, Health Standards Programme, Occupational Safety and Health Administration, Washington D.C., United States;
- Dr. Arthur W. Musk, Clinical Professor of Medicine and Public Health, University of Western Australia, and Physician, Department of Respiratory Medicine, Sir Charles Gairdner Hospital, Nedlands, Australia.

The experts were asked to acquaint themselves with the *Rules of Conduct for the Understanding on Rules and Procedures Governing the Settlement of Disputes*[^188], paying special attention to Annex 2 (Illustrative list of information to be disclosed). No expert has disclosed any circumstance that could be considered as the potential source of a conflict of interest.

In consultation with the parties, the Panel prepared precise questions which it submitted to each expert individually. The experts were requested to answer only those questions that they considered to be within their domain(s) of competence. Written communications by the parties, transcriptions of their oral statements, as well as the references they submitted to the Panel were transmitted to the experts for their information. The written answers from the experts have been forwarded to the parties, who have had a chance to comment on them. The questions posed by the Panel and the answers given by the experts are contained in section V.C. The observations of the parties are reproduced in section V.D.

On 17 January 2000, the experts were invited to discuss with the Panel and the parties their written answers to the questions and to provide additional information. Annex VI to this report contains the minutes of the meeting.
C. Questions by the Panel and Comments by the Scientific Experts

5.24 The Panel requested the experts to comment on the areas of difference between the parties highlighted in the first paragraph of each question, as well as to address the specific points listed. The Panel encouraged the experts to indicate, to the extent possible, key points on which they considered that (i) there is scientific proof, (ii) there is broad agreement among experts, (iii) there is uncertainty and/or a range of divergent opinions among experts.

1. Introductory Comments by Dr. Henderson

(a) Introduction

5.25 This introduction sets out a general summary of prevailing knowledge and uncertainties on asbestos-related disorders, with emphasis on mesothelioma and lung cancer, together with discussion of both the amphiboles and commercial chrysotile, patterns of exposure, and some brief details of \textit{in vivo} and \textit{in vitro} experimental studies.

5.26 This introduction has two purposes: (i) to provide a general background and broad perspective to the questions and answers that follow; and (ii) to correct some inaccuracies and errors in the documentation supplied already to the WTO. In so doing, I have tried to broaden the perspective beyond the classical Canadian studies on the Quebec chrysotile miners and millers, and beyond the INSERM Report. A number of the general discussions in this introduction have been truncated after the issue has been put into context, and some of these discussions are then continued and amplified in my specific responses to the questions. This has produced some iteration of some points, but I believe that the advantages - avoidance of the potential for distortion created by answers without adequate background information - outweigh any disadvantages. The division of my report into these sections also provides an opportunity to indicate the relative importance of epidemiological studies versus \textit{in vivo} or \textit{in vitro} experimental models in the formulation of my opinions and answers.

5.27 At the outset, I emphasize that Australia (including Western Australia) is no longer an asbestos producer. Production of crocidolite at the Wittenoom blue asbestos industry stopped in 1966. None has been produced or exported since. Crocidolite was used in asbestos-cement products in Australia until 1966 when its use was discontinued, but imported amosite was used in these products until 1984 [NICNAS 99][189]. The use of chrysotile in fibro-cement products was discontinued in 1987.

5.28 As stated repeatedly in the documentation provided to the WTO, asbestos has the capacity to induce at least five benign pleuropulmonary disorders, and two cancers: parietal pleural fibrous plaques; benign asbestos pleuritis with effusion; diffuse pleural fibrosis; rounded atelectasis; asbestosis; primary lung cancer; malignant mesothelioma. The essential characteristics of these disorders are discussed in the documentation submitted to the WTO and lie beyond the scope of this report; if further details are required, standard texts should be consulted [26-30]. There is no persuasive or compelling evidence that asbestos of any type causes cancers other than lung cancer and mesothelioma, with the arguable exception of cancer of the larynx. At this stage, it is sufficient to point out that:

\textit{... there is an exposure-response relationship for all chrysotile-related diseases. Reduction of exposure through introduction of control measures should significantly reduce risks. Construction and demolition operations may present special control problems"}. [EHC 203, p 141].
Malignant mesothelioma is a cancer of the mesothelial cells that line the serosal membranes of the major body cavities, namely the pleura, the peritoneum, the pericardium and the tunica vaginalis testis; the constituent neoplastic cells characteristically express the phenotype of a recognized pattern of differentiation of the mesothelium, whether epithelioid, sarcomatoid or both (biphasic), as revealed by conventional light microscopy, mucin immunohistochemistry, immunohistochemistry or electron microscopy, or a combination of these techniques [31-33]. Like other forms of cancer, mesothelioma has the capacity for local invasion of tissues such as the chest wall or lung, with confluent serosal spread in most cases but not all, and in some instances distant metastasis [31], with an almost invariably fatal outcome. Mesothelioma is resistant to conventional cancer therapies (e.g. radiotherapy or chemotherapy), but some long-term survivals have been recorded following radical surgery (pleuropneumonectomy) in patients in good physical condition and with early-stage disease [34-43]; radical surgery of this type is not a treatment option for the majority of mesothelioma patients.

Most mesotheliomas encountered in the 1990s are a consequence of prior occupational exposure to asbestos [24], including bystander exposure. The relationship between asbestos - especially one or more of the amphibole varieties - and mesothelioma is accepted by virtually all authorities as causal. In this respect, asbestos fulfils all The Bradford Hill Criteria for the establishment of causality (e.g. please see Stolley and Lasky [44]). The following points about asbestos and mesothelioma are worth emphasis:

(i) Inhalation of asbestos fibres represents the overwhelming cause of mesothelioma in industrialized societies, so much so that the incidence of mesothelioma is usually considered to be an index of those societies’ past usage of asbestos.

According to Peto et al. [24]:

“The great majority of mesotheliomas are caused by asbestos, and the much higher incidence in men indicates that most are due to occupational rather than environmental exposure. The incidence continues to rise approximately as the third power of time since first exposure to asbestos for many decades after exposure has ceased (Peto et al., 1982), and most patients are men first exposed 30 or more years ago. A country’s mesothelioma rate is therefore a quantitative indicator of its population’s past exposure — mainly occupational — to asbestos.” [p 666].

Boffetta [15] claims that:

“Asbestos is the only established risk factor of mesothelioma. Because of the rarity of the disease and the specificity of the causal association, all cases occurring among asbestos-exposed workers are attributed to this exposure.” [p 476; please see following discussion].

The asbestos exposure may take the following forms: (i) direct or indirect occupational exposure (including bystander exposure); (ii) domestic exposure: e.g. household contacts of asbestos workers, such as wives who washed the asbestos-contaminated work clothes of their husbands [45-47]; (iii) environmental exposure: this category includes those who lived downwind of asbestos industries or in townships contaminated by asbestos [45-47]. For example, ≈ 27 mesotheliomas have been recorded among those who lived at Wittenoom as children (the roads, airstrip and school yards were surfaced with crocidolite tailings from the mine, and children often played in the mine tailings).
The tumor [mesothelioma] is more often seen in workers who have only moderate or small amount of asbestos in their lungs, and who show little, if any, clinical or radiologic evidence of pulmonary fibrosis. This amount of asbestos may be inhaled not only by professional asbestos workers, but also by those who handle products containing only a small proportion of asbestos, those who do not handle asbestos at all but merely work alongside asbestos workers such as craftsmen employed in the building industry — carpenters, electricians, etc. — those who have relatives who carry asbestos home in their workclothes and those who live close to asbestos plants.” [47] [p 295].

5.34 No history of asbestos exposure is obtainable in about 15-25 per cent of mesothelioma cases [31, 48]. Nonetheless, absence of a history of exposure does not equate to absence of exposure, and evidence indicates that many of these mesotheliomas are in reality attributable to asbestos inhalation — e.g. remote, brief or forgotten exposure, or alternatively, the individual may be unaware that he (the male:female ratio is about 8:1) was in fact exposed to asbestos: (i) from my own series of mesotheliomas, 79 per cent of the request forms that accompanied the biopsies on which the diagnosis was made gave a positive history of past asbestos exposure; clinical review of the remaining 21 per cent yielded a history of asbestos exposure in a substantial proportion, including some for whom the original history stated that there was no exposure, so that my estimate of the proportion for whom a positive history of exposure was eventually obtained is = 85-90 per cent. This estimate is in reasonable agreement with figures in the 1999 Report for the Australian Mesothelioma Register [AMR 99], where 85 per cent of mesotheliomas had a history of asbestos exposure; (ii) Leigh et al. [49] found measurable asbestos fibre levels (> 200,000 fibres per gram dry lung tissue) in 81 per cent of the 28 per cent of Australian mesothelioma cases that had no history of occupational or environmental exposure to asbestos.

5.35 In the reply to Question 3 from the European Communities, Canada makes the following statements:

"... Canada wishes to inform the European Communities of the considerable body of evidence contradicting their statement that asbestos in all forms (amphiboles and chrysotile) is the only known factor that can cause mesothelioma or pleural cancer. ... A number of studies suggest other potential risk factors that may have been under-estimated in epidemiological studies in industrialized countries. ... A number of artificial fibres cause mesothelioma when they are injected into the pleura and peritoneum of laboratory animals. Note also that the International Agency for Research on Cancer (IARC) has classified refractory ceramic fibres as probable carcinogens, partly because of instances of mesothelioma induced by inhalation and injection in animal experiments. The SV40 virus readily induces mesothelioma when injected into animals; studies suggest that the virus contaminated anti-polio (poliomyelitis) vaccines from 1955 to about 1963 and may induce mesothelioma with or without the help of asbestos fibres. Some studies of humans report the presence of the simian SV40 virus in the biological tissue of mesothelioma victims. Ionizing radiation used in cancer therapy and perhaps occupational exposure to radiation have induced mesothelioma. ... [E]rionite has been shown to be even more toxic than crocidolite in causing mesothelioma; it has killed large numbers of villagers in Turkey. Erionite is a mineral fibre but does not belong to the asbestos family.”

5.36 Possible factors other than asbestos implicated as contributory or causative for rare mesotheliomas are tabulated below:
TABLE 1: PUTATIVE OR POSSIBLE RISK FACTORS AND MEDIATORS OF RISK OF MESOTHELIOMA OTHER THAN ASBESTOS

<table>
<thead>
<tr>
<th>Factor</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Erionite</td>
<td>Very high incidence of mesothelioma due to environmental exposure in Turkey (restricted geographic localization only).</td>
</tr>
<tr>
<td>Chronic Inflammation</td>
<td>Pleural scars (tuberculosis, pleurisy, therapeutic pneumothorax, familial Mediterranean fever); see following discussion.</td>
</tr>
<tr>
<td>Radiation</td>
<td>Single cases after Thorotrast injection or radiotherapy, causality unproven. One case in atomic bomb survivor.</td>
</tr>
<tr>
<td>Beryllium</td>
<td>Two doubtful cases described.</td>
</tr>
<tr>
<td>Vegetable fibres</td>
<td>No proof in humans.</td>
</tr>
<tr>
<td>Hereditary factors</td>
<td>Familial cases (explicable by common asbestos exposure ± unidentified genetic susceptibility factors, including association with other cancers in first-degree relatives).</td>
</tr>
<tr>
<td>Immunological factors</td>
<td>Rapidly progressive cases in patients with HIV infection; very rare — single case(s) only.</td>
</tr>
<tr>
<td>Dietary factors</td>
<td>Provitamin A, ß-carotene may decrease the risk (unproven).</td>
</tr>
<tr>
<td>Viruses</td>
<td>Mesotheliomas in animals. Simian virus 40 (SV40) DNA sequences reported in mesotheliomas; see following discussion</td>
</tr>
</tbody>
</table>

Modified from Hillerdal [20].

5.37 There are anecdotal reports of mesothelioma following radiation, including radiotherapy for childhood cancer such as Wilms’ tumour [50-56]. In addition, excess rates of mesothelioma have been reported among both Danish and German patients exposed to radioactive thorium dioxide (Thorotrast) for radiological procedures [57, 58], although a similar but smaller Japanese study found no such excess [59]. Neugut et al. [60] investigated women with breast cancer and patients with Hodgkin’s disease, many of whom had been treated by radiotherapy (RT):

“The authors performed a retrospective cohort study utilizing 251,750 women registered with breast carcinoma in the Surveillance, Epidemiology, and End Results Program of the U.S. National Cancer Institute from 1973-1993, 24.8% of whom received RT as part of their initial management, and 13,743 people with Hodgkin’s disease, 50.6% of whom received RT as part of their initial management. RESULTS: Six cases of malignant pleural mesothelioma were found: two in breast carcinoma patients treated with RT and four found in women not treated with RT. No cases occurred in the patients with Hodgkin’s disease. The overall estimated relative risk for malignant pleural mesothelioma after RT was 1.56 (95% confidence interval, 0.18-5.63). CONCLUSIONS: To the authors’ knowledge, this is the first controlled study to investigate thoracic radiation exposure and malignant pleural mesothelioma, and no association was found.” [abstract].

5.38 I am also aware of at least one mesothelioma in a patient with HIV infection (AIDS) [61]. Other mesotheliomas have occurred many years after chronic inflammatory lesions of the pleura — e.g. chronic empyema or packing of the pleural cavity with leukite spheres as treatment for tuberculosis [62, 63], and there are a few reports of an association between familial Mediterranean fever (FMF) and mesothelioma (about eight cases only; possibly related to recurrent FMF serositis [64-67]). However, cases of this type are exceptional and most cases of “post-inflammatory” mesothelioma with a short interval between inflammation and tumour (e.g. = 2-3 years by analogy with the criteria for the diagnosis of benign asbestos pleuritis [33, 68, 69]), are probably mesotheliomas that presented with a burst of inflammatory activity, followed by a period of quiescence [70].

5.39 In addition, background asbestos exposure represents a confounding factor for some cases associated with radiation and immunodeficiency: (i) in one report on mortality among 260 plutonium workers, all six mesotheliomas occurred in individuals who had also sustained asbestos exposure [71]; “… no apparently elevated causes of death except for six cases of mesothelioma and six cases of astrocytoma glioblastoma multiforme. The
mesothelioma cases had a documented occupational exposure to asbestos...” [extract from abstract]; (ii) in one of my own cases, the patient had been treated for Hodgkin’s disease by mantle radiotherapy 10 years before the diagnosis of his primary pericardial mesothelioma, but he also had a background of occupational exposure to asbestos; (iii) in another case — a pleural mesothelioma in a renal transplant recipient — the patient had also sustained earlier occupational exposure to asbestos.

(iii)  Erionite and mesothelioma in Turkey

5.40  Erionite (a fibrous zeolite) represents a naturally occurring fibrous mineral implicated in the induction of mesothelioma in certain villages (notably Karain and Tuskoy) in the Cappadocian region of Turkey [72, 73], and in Turkish emigrants [74]. So far as I am aware this represents a restricted geographic pocket of mesothelioma cases induced by erionite used as stucco or whitewash in buildings, so that the inhabitants were exposed to high concentrations of erionite fibres from birth. Erionite has no relevance to the broader mesothelioma problem in Western Europe, North America, and Australia. Nonetheless, in its physical properties erionite has similarities to the amphibole varieties of asbestos and it has been suggested that its greater mesotheliomagenicity is related to a greater surface area (200 m² per gram) than crocidolite (8-10 m² per gram), due to the presence of pores in the crystal lattice (see Roggli and Brody [75]); such differences in surface topography might correlate with differences in free radical generation at the surface of fibres.

(iv)  Simian virus 40 (SV40) and mesothelioma

5.41  Recently, a voluminous literature has grown rapidly on the detection of SV40 DNA in up to 60 per cent of human mesotheliomas [76-87] and some other tumours, such as papillary carcinoma of the thyroid [88], osteosarcomas and brain tumours [83, 89-91]. These observations followed an initial finding that SV40 could induce mesothelioma in hamsters when injected into the pleural cavity [92], and the later demonstration that SV40 could inactivate the tumour suppressor genes p53 and the retinoblastoma gene (Rb) via the large T antigen (TAG) [80, 82, 93, 94]. For humans, early poliomyelitis vaccines contaminated with SV40 were a potential source for the SV40 DNA [82-84]. The following points on this interesting association are also worth emphasis:

- It has been suggested that the presence of SV40 might explain: (i) why mesothelioma only develops in a relatively small proportion of asbestos-exposed individuals (usually < 10 per cent); and (ii) why no history of asbestos exposure is obtainable on a sizeable minority of mesotheliomas [95]. However, almost all the mesotheliomas in which SV40 DNA has been found were asbestos-associated; to the best of my knowledge, there is no reported case-control analysis of SV40-associated mesotheliomas where asbestos fibre counts were not elevated above reference values, with the exception of a recent study by Mayall et al. [96] (please see following discussion). Therefore, the existing data do not adequately address either (i) or (ii): there are many other possible explanations for these observations.

- In other studies, SV40 or TAG could not be detected within mesotheliomatous tissue [97-99]. Galateau-Sallé et al. [100] found that SV40 was present not only in mesotheliomas but also in benign inflammatory disorders of the pleura and non-neoplastic lung tissue. In an as yet unpublished investigation carried out in collaboration with Prof. Alec Morley in the Department of Haematology-Oncology at the Flinders University, we have also identified SV40 in mesotheliomas, and in non-neoplastic pleural lesions, normal tissues and colon cancers, casting doubt on the specificity of the association.

- Two epidemiological studies have shown no increase in the incidence of bone or brain tumours — or mesotheliomas — 30 years after the
The evidence so far only points to SV40 as a possible co-factor for asbestos in the genesis of mesothelioma [96]. For example, Mayall et al. [96] detected SV40 sequences in five of seven asbestos-associated mesotheliomas, but none of four mesotheliomas that were not asbestos-related (investigated by fibre burden analysis of lung tissue, using electron microscopy). However, the evidence in favour of SV40 as a co-factor for mesothelioma induction is still inconclusive and non-persuasive, and in humans the SV40 may represent an innocent bystander or passenger: the criteria for causality [44] have not been fulfilled.

“It remains to be shown whether the presence of SV40 contributes significantly to malignant transformation or whether certain human neoplasms provide a microenvironment that favors viral replication in humans with latent SV40 infection.” [91] [last sentence of abstract].

5.42 The point of these comments is that the evidence for a role of SV40 in the development of mesothelioma is inconclusive, and most of SV40-associated cases still represent asbestos-associated mesotheliomas. Although the literature contains anecdotal reports of mesothelioma following radiation, some of these cases (e.g. among plutonium workers) are complicated by coexistent asbestos exposure and it is worth emphasizing that these cases are rare: together they add up to only a small fraction of 1 per cent of the total burden of mesotheliomas in industrialized societies, for which asbestos remains the overwhelming cause. As emphasized already, there is general agreement that the incidence of mesothelioma in various nations is a reflection of the past usage of asbestos by those societies.

5.43 Hillerdal [20] comments along similar lines:

“... SV40 might be a cofactor to asbestos in some patients with mesothelioma, but the [findings] have not been confirmed and are still disputed. ... In summary, then, as far as is known today, factors other than mineral fibres can only explain a very small proportion of mesotheliomas, and can for practical purposes be disregarded [i.e. when approaching the causation of mesothelioma among large cohorts or populations]. Thus, a malignant mesothelioma can be regarded either as caused by asbestos or belonging to a normal background level — that is a spontaneously occurring tumour.” [p 506].

(v) Male: female ratio for mesothelioma

5.44 Asbestos-induced mesothelioma affects males more often than females in a ratio of about 8:1, as a reflection of occupational exposure.

(ii) Anatomical distribution of mesothelioma

5.45 With the exception of one series in which 44 per cent of mesotheliomas were peritoneal [104], there is general agreement that primary asbestos-induced mesothelioma affects the pleura more often than the peritoneum, in a ratio of at least 3:1 or even up to = 11:1 [31, 33] (see also AMR 99). In Australia, = 91 per cent of mesotheliomas arise in the pleural cavities, whereas about 7 per cent represent primary peritoneal mesotheliomas and = 1 per cent affect the pericardium or tunica vaginalis testis [33]. This predominance of pleural mesotheliomas in comparison to the peritoneum appears to correlate with gender differences in the frequency of occupational exposure to asbestos (the same high ratio of pleural to peritoneal tumours is also encountered in the United States). In females, a smaller proportion of mesotheliomas arises in the pleura, and in one study of Swedish
insulation workers, all seven mesotheliomas arose in the peritoneum [105] (please see following discussion).

5.46 One report [106] that included cases notified to the Australian Mesothelioma Register from 1986 through 1988 gave figures for the anatomical sites affected in men and women: 676 of 723 men had a pleural mesothelioma (93 per cent), whereas 38 were peritoneal tumours (5 per cent) and nine occurred in other sites (1 per cent). In contrast, 84 mesotheliomas in 101 women were pleural in location, whereas 17 per cent had a peritoneal mesothelioma.

5.47 Presumably, this difference in anatomical distribution between sexes is a reflection of different rates of occupational exposure to asbestos. On theoretical grounds, one would expect mesotheliomas entirely unrelated to asbestos to occur with about equal frequency in the pleura and peritoneum, or more often in the peritoneum because of the greater surface area of the peritoneal cavity.

5.48 A partial list of the factors that might explain the higher proportion of peritoneal mesotheliomas in some series and in women includes the following [33]:

- The high proportion of pleural mesotheliomas in men is presumably a reflection of asbestos exposure, with deposition of asbestos fibres in lung tissue, followed by translocation of the fibres to the pleura; on this basis, asbestos inhalation appears to skew the proportional distribution of mesothelioma towards the pleura in comparison to other sites. In contrast, fibres presumably follow a more circuitous route from the lung to the pleura, across the diaphragm and into the peritoneal cavity, to induce peritoneal mesothelioma; higher inhaled doses of asbestos might be necessary for the requisite number of fibres (whatever that is) to reach the peritoneum via the pleura, in order to induce peritoneal mesothelioma.

- The high proportion of peritoneal tumours in some series may be a consequence of patterns of referral for cases that constitute problems in diagnosis, because the diagnosis of peritoneal mesothelioma is, in general, more difficult than for pleural mesotheliomas. This may explain the higher proportion of peritoneal mesotheliomas among cases referred to the US-Canadian Mesothelioma Panel [107], because many of these represented problems in diagnosis, whereas the Australian Mesothelioma Surveillance Program (AMSP) captured all mesotheliomas throughout Australia [48].

- Genuine biological differences in the inhaled dose, deposition or transport of different asbestos fibre types in some groups of workers, notably insulation workers [108] and former Wittenoom workers [109] — as a consequence of heavy occupational exposure — and in women [106, 110].

5.49 In answer to questions posed by the European Communities (Question 3, see Annex II), the following comment is made:

"... malignant diffuse mesothelioma is a cancer of the mesothelial cells of the pleura, the pericardium and the peritoneum. Furthermore, peritoneal mesothelioma is an even more typical result of exposure to amphiboles than pleural mesothelioma."

5.50 From the preceding discussion on the proportions of mesotheliomas arising in the pleural cavities versus the peritoneum, it is evident that this proposition is not correct: use of the term typical in this context is inappropriate. In reality, pleural mesothelioma is a more typical or usual outcome of asbestos exposure, whereas asbestos-induced peritoneal...
mesotheliomas are usually associated with more prolonged and heavier exposures than pleural mesotheliomas, so that the proportion of patients with asbestosis is higher than for pleural mesothelioma [111]. It has also been claimed that peritoneal mesotheliomas are almost always a consequence of amphibole exposure (as opposed to chrysotile only) [112]. Nonetheless, although some of the peritoneal mesotheliomas in my own series of cases followed high-dose exposures to asbestos that included one or more of the amphiboles, a few followed lower cumulative exposures, and Neumann et al. [111] have reported peritoneal mesotheliomas as a consequence of exposure in the building trades and metal industries, in addition to asbestos industries; Rogers et al. [3] recorded peritoneal mesotheliomas in whom only chrysotile fibres were detected on lung fibre analysis (see Table 9, paragraph 5.137).

(vii) Latency intervals (lag-times)

5.51 In all reported studies, mesothelioma is a disease of long latency between exposure to asbestos and the subsequent diagnosis of the mesothelioma. In the AMSP [48], the mean latency interval (lag-time) was 37 years, with a reported range of 4-75 years; the lag-time was reported to be < 10 years in only four of 499 asbestos-associated mesotheliomas (0.8 per cent). Many authorities set a minimum lag-time of 10 years (e.g. The Helsinki Criteria [113]), and for most patients the lag-time is in the range of 20-40 years. When the lag-time is < 10 years, it is likely that the proximate exposure was coincidental, and that there were one or more earlier exposures.

(c) Spontaneous or background mesothelioma: does it exist?

5.52 The rare occurrence of mesothelioma in childhood and even as a congenital malignancy supports the existence of a background of spontaneous mesotheliomas unrelated to asbestos (in addition, mesothelioma has been reported in fish (trout) [114], where inhalation of airborne asbestos fibres cannot be invoked). However, in epidemiological studies on adult populations, it is virtually impossible to separate spontaneous mesotheliomas from those that are arguably attributable to environmental exposure to asbestos [70, 115]. The incidence of mesothelioma in women is sometimes used as an index of the background or spontaneous rate: the crude incidence rate for women in Western Australia is about 2.6 per million person-years at age = 15 years [115]. The incidence rate in other populations is listed in Table 2 (following page).

5.53 In the answer to the WTO Panel’s questions to Canada (Question 9, see Annex II), the following statement is made:

“Recent analyses of Canadian data on mesothelioma in Canada, British Columbia and Quebec all agree that the incidence rate of mesothelioma has been stable among women of all age groups since 1984. The rates are 70% higher in Quebec than in the rest of Canada, presumably as a result of more frequent and more intense exposure in the workplace.”

5.54 The statistics for Australia differ on this point (Table 2): mathematical modelling of Western Australian data suggests that the incidence rate in women has risen about two-fold from the 1970s until the 1980s, which might be explicable by increased general environmental exposure to asbestos, plus some occupational exposures among women [70, 115] (please see also AMR 99 — i.e. the graph for the Age Specific-Incidence Rates of Malignant Mesothelioma in Australia Women, 1986-1995, especially for ages 50-64 and 65-79). This increased incidence among women presumably reflects direct or indirect occupational exposure, domestic exposure or environmental exposure [115]. In this respect, it is worth emphasizing that domestic (household contact) exposure to asbestos — e.g. among wives laundering the dust-laden workclothes of an asbestos-exposed husband — is not necessarily low-level exposure, and analysis of the asbestos fibre content of the lungs in a small number of such patients indicates that this type of exposure can approach occupational levels [116].
5.55 The often-cited background or spontaneous rate of mesothelioma of 1-2 per million person-years [10, 117], has in part also been derived from backward extrapolation of the incidence rates in men, to the point where the estimated incidence rates for men and women diverged from each other (i.e. linear extrapolation to the point where the sex ratio = 1:1) [117]. Hillerdal [20] suggests that this incidence probably represents a high estimate and comments in the following terms:

“... there seems to be a small spontaneous basal or background incidence of the tumour [mesothelioma] ... However, it is of course possible that some of these background cases might in fact be due to occupational, domestic, or even environmental exposure, unknown to (or forgotten by) the patients themselves. ... There are authors who claim that the presumed background levels must be very low, and retrospective searches for the tumour in the medical literature reveal no convincing cases of mesothelioma before 1946, although such negative evidence is of questionable value.191 McDonald and McDonald, in a recent review, estimated the background level to be 1-2/million per year; they came to this figure by extrapolating backwards from epidemiological studies from various countries. ... It is nevertheless possible that there is a background level of mesothelioma, — that is, that the tumour can occur even in the complete absence of asbestos (or erionite) fibres. However, the data reviewed here indicate that if so, this background level must be very low — probably much < 1 case/million people/year. This figure comes from studies of industrialized countries, where background exposure to asbestos is unavoidable. What the true figure is can only be guessed ...”. [p 507].

5.56 De Klerk [115] and Comin et al. [70] have commented that in the absence of specific exposure to asbestos, the final estimated rate for both men and women in Australia is 2.6 per million person-years — higher than the equivalent figure of 1.6 for Los Angeles [115]. This difference may lend some support to the proposition that general environmental exposure to asbestos may have produced an increase in the mesothelioma rate in Western Australia [115]. However, it is difficult or impossible in general to draw firm conclusions from differences between different studies, because of variation in the accuracy of diagnosis and differences in the ways that data are collected.

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**TABLE 2: INCIDENCE OF MORTALITY OF MESOTHELIOMA IN VARIOUS COUNTRIES AND AREAS OVER TIME, 1960s TO 1994 (PER MILLION INHABITANTS PER YEAR)**

<table>
<thead>
<tr>
<th>Country or area</th>
<th>Year</th>
<th>Males</th>
<th>Females</th>
</tr>
</thead>
<tbody>
<tr>
<td>United States</td>
<td>1968-81</td>
<td>2.1</td>
<td>0.8</td>
</tr>
<tr>
<td>North America</td>
<td>1972</td>
<td>2.8</td>
<td>0.7</td>
</tr>
<tr>
<td>Texas</td>
<td>1976-80</td>
<td>5.6</td>
<td>2.1</td>
</tr>
<tr>
<td>Selected cities, United States</td>
<td>1970s</td>
<td>4.4-11.1</td>
<td>1.2-3.8</td>
</tr>
<tr>
<td>United States</td>
<td>1986</td>
<td>7.13</td>
<td>1.2</td>
</tr>
<tr>
<td>Nantes-Saint-Nazaire, France</td>
<td>1956-74</td>
<td>5.2</td>
<td>0.2</td>
</tr>
<tr>
<td>Nantes-Saint-Nazaire, France</td>
<td>1979-83</td>
<td>1.2</td>
<td>0.3</td>
</tr>
<tr>
<td>Nantes-Saint-Nazaire, France</td>
<td>1985-92</td>
<td>19.4</td>
<td>4.0</td>
</tr>
<tr>
<td>Great Britain</td>
<td>1968-71</td>
<td>8.4</td>
<td>2.3</td>
</tr>
<tr>
<td>Great Britain</td>
<td>1972-76</td>
<td>12.6</td>
<td>2.8</td>
</tr>
<tr>
<td>United Kingdom</td>
<td>1983</td>
<td>17.5</td>
<td>3.2</td>
</tr>
<tr>
<td>Great Britain</td>
<td>1986-87</td>
<td>20.7</td>
<td>4.3</td>
</tr>
<tr>
<td>Great Britain</td>
<td>1982-86</td>
<td>30.8</td>
<td>4.9</td>
</tr>
<tr>
<td>Great Britain</td>
<td>1987-91</td>
<td>44.0</td>
<td>6.4</td>
</tr>
<tr>
<td>Australia</td>
<td>1982-86</td>
<td>28.3</td>
<td>3.3</td>
</tr>
<tr>
<td>Australia</td>
<td>1994</td>
<td>49.9</td>
<td>4.8</td>
</tr>
<tr>
<td>Denmark</td>
<td>1978-80</td>
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<td>7.0</td>
</tr>
<tr>
<td>Barcelona, Spain</td>
<td>1983-90</td>
<td>8.3</td>
<td>4.7</td>
</tr>
<tr>
<td>Finland</td>
<td>1990-94</td>
<td>10</td>
<td>2.9</td>
</tr>
</tbody>
</table>

Modified from Hillerdal [20].
In response to questions from the WTO Panel (Question 9, see Annex II) the Canadian document also observes that:

“The incidence of [mesothelioma] among men levelled off after 1984 in British Columbia ... and seems to have levelled off in Quebec after 1990 ... Finally, analysis of Canadian rates between 1973 and 1992 ... estimates that the risk is four times greater for men born before 1940 than for men born between 1951 and 1955. Those analyses therefore suggest that the incidence of mesothelioma has levelled off in Canada, is declining in British Colombia, and has levelled off in Quebec ...”

5.58 In response to these observations, I emphasize the following: (i) the incidence of mesothelioma among Australian males shows little evidence of levelling off, and has continued to rise until 1994-1995 and thereafter (please see Table 2 and the 1998 and 1999 Reports for the Australian Mesothelioma Register); (ii) from the recent report by Peto et al. [24], it is also evident that the incidence of mesothelioma in Western Europe continues to rise, with particular emphasis on males born between 1945 and 1950 who used asbestos-containing products in the 1960s and the 1970s (and the early 1980s).

(d) Magnitude of the mesothelioma problem

5.59 Malignant mesothelioma continues to represent a major health problem within industrialized societies, and together with lung cancer it represents the most important occupational cancer among so-called blue-collar workers [121-124].

5.60 It has been estimated that across Western Europe, North America and Australia (combined population ~ 800,000,000), around 10,000 mesotheliomas and 20,000 asbestos-induced lung cancers occur annually, related mainly to occupational exposure (about one mesothelioma for every 200 tons of asbestos produced, taking into account the prolonged lag-times) [125]. Steenland et al. [126] estimate that approximately 9000-10,000 men and 900-1900 women develop lung cancer each year in the United States because of past exposure to occupational carcinogens, and more than half of these lung cancers are related to asbestos (this overall estimate is considered probably to be conservative). Predictions of asbestos-related diseases in Australia (population ~ 18,000,000) indicate that about 13,000 cases of mesothelioma (range 8000-20,000), about 40,000 cases of lung cancer (range 30,000-76,000) and 1000 cases of asbestosis are likely to occur between the years 1987 and 2020 [70, 127].

5.61 More recently, Peto et al. [24] have predicted that about 190,000 mesothelioma deaths are likely to occur throughout Western Europe (Britain, France, Germany, Italy, the Netherlands and Switzerland) over the next 35 years. If one adds in lung cancer at a ratio of one lung cancer for every mesothelioma death, this figure would rise to 380,000 deaths, and if the ratio of lung cancers to mesotheliomas is 2:1 the figure rises to 570,000 deaths.

5.62 Overall, asbestos may have caused approximately 5,000,000 deaths across industrialized societies so far. When future deaths in so-called developing nations are added, the final toll is almost certain to be substantially higher, especially because occupational exposures in those countries are likely to be heavier (e.g. China). Estimates of this magnitude are likely to engender alarm among those who set social policy. Even so, it is important that this problem, like others (e.g. atomic energy), is approached with common sense, rationality and prudence, taking into account population-based risk estimates: it would be irrational to swap one risk for another higher risk if the two risks were equally serious.

(e) Some general observations on approaches to risk assessment on society and on epidemiological studies of asbestos-related cancers

5.63 The documentation supplied to the WTO includes estimates of risks from low-level exposure to chrysotile in proportion to various other risk factors in society: in fact, the relative risk of mesothelioma from low-level exposure to asbestos in place is the focus of considerable controversy. Clearly, detailed analysis of this issue is beyond the scope of
this report, but the excess risk of mesothelioma from very low-level exposure to asbestos — e.g. simple occupancy of public buildings or schoolrooms where average asbestos fibre concentrations are about < 0.001-0.02 fibres per litre — appears to be very slight: about = 5.5 mesotheliomas per million lifetimes of 80 years, or < 1 case per 10,000,000 per year.

5.64 The estimated mesothelioma risk for a 10-year exposure to low levels of airborne asbestos in schools (age start 7-8 years; fibre concentration 0.00065-0.001 fibre per ml) is in the range 6.6-20 per million lifetimes (0.0825-0.25 per million person-years). Estimates of this type are predicated on linear dose-response models with no threshold, and these have been the subject of argument and criticism. The occupational groups from which they were derived were exposed to mixtures of asbestos types, but one might expect the risk to be even less or “undetectably low” in nations where only chrysotile was used. At this point, it is sufficient to emphasize that - even if one accepts for a moment the non-threshold linear dose-response relationship - calculations indicate that a single asbestos fibre (the so-called “one fibre” hypothesis) would only have a 50-50 chance of producing a single excess mesothelioma among all the humans who have ever inhabited Planet Earth.

5.65 These observations on mesothelioma risk estimates for very low-level exposure to asbestos in buildings do not contradict the earlier suggestion in this report that Western Australian and Australian increases in mesothelioma incidence among females were a possible consequence of general environmental exposure to asbestos: the two-fold rise in incidence in Western Australia could be due to environmental fibre levels higher than those recorded in public buildings elsewhere.

5.66 If it exists, the risk of mesothelioma from very low-level environmental exposure to asbestos needs to be considered in proportion to other risks of death in society. The lowest death-risk at any age occurs in girls aged 4-14 years and is ~ 100 per million per year, but the risk in late teenage increases by 300-400 per million per year — attributable largely to increased travel by motor vehicle. A 40-year-old man at risk of mesothelioma from low-level exposure to asbestos during childhood (excess mesothelioma risk < 1 per million person-years) has an annual risk of death from all causes of about 2000 per million. In 1990, de Klerk [128] had this to say on the subject:

“In the US the acceptable (or possible litigation-proof) lifetime risk seems to be one per million. The FDA have this as a set policy; the EPA approximates to it, and other agencies seem to employ similar figures. In the UK the Royal Society has set a higher range with an annual risk of one per million considered negligible, with any form of control unjustified; one per 100,000 considered low (“very few would consider action necessary” — e.g. 16,000 km air or rail travel); one per 10,000 moderate (few would commit their own resources to reduce risk” — e.g. 16,000 km car driving, working as a coalminer); one per 1,000 high (e.g. age 30-39, 16,000 km motor cycling); and one per hundred unacceptable (e.g. age 55-59, smoking 20 cigarettes per day, heavy exposure to crocidolite).”

5.67 Within this context, one might ask what constitutes a negligible as opposed to an acceptable (or unacceptable) risk? In the context of the foregoing discussion, one might argue that a negligible risk is a statistical and scientific concept: a risk so slight that it does not require preventive or remedial steps in comparison to other risks in society (although some might argue about the dividing line between negligibility and unacceptability in this context). With acceptability or unacceptability other factors come into play: they include, for example, social, political and industrial considerations — and the likelihood of litigation over any situation wherein the theoretical or estimated risk is elevated to a background level, no matter how slightly. Accordingly, a risk, though slight or even negligible, might still be considered unacceptable in legal or sociopolitical terms.

5.68 Others may dissent from the acceptability of the Royal Society approach to low, moderate and high risks discussed above. Conclusions about the acceptability or unacceptability of risk will also vary according to the seriousness of the risk (e.g. the approach to a lethal risk such as mesothelioma would be quite different from a factor that caused a large proportion of the population to sneeze once or twice): these assessments
will also vary according to the avoidability of the risk, the individuals making the assessment, and the question of informed consent by those at risk.

5.69 Furthermore, society abounds with inconsistencies and contradictions over the relativity of various risks. For example, some societies that regulate or propose a ban on chrysotile asbestos make extensive use of radioactive materials — e.g. in nuclear power stations and the production of radioactive isotopes for medical purposes. Even so, the use of fissionable materials for these purposes may be justified and justifiable within those societies, because: (i) the risk of morbidity or death from well-publicized mishaps at nuclear reactors is still substantially less than the risk of death from alternative energy sources (e.g. higher mortality rates among coalminers); and (ii) because the materials in question can be regulated and controlled so that they are accessible to only a small fraction of society (i.e. workers who can be trained in the controlled use of radioactive substances); and (iii) nuclear power does not contribute significantly to air pollution or greenhouse gas emissions in comparison to the burning of fossil fuels.

5.70 In addition, Nicholson [129] places the problem into the perspective of voluntary versus involuntary risks:

“Rather than compare asbestos risks with voluntary risks (smoking, school football) or risks that remain high despite expenditures of substantial public and private money (aircraft and highway accidents), it is worthwhile to compare them with other involuntary environmental risks that are controlled by regulatory agencies (pesticide exposures, drinking water contamination). In a review of regulatory actions taken by the FDA ... and the EPA it was found that for estimated population risks exceeding one death/year, the individual lifetime risks were usually regulated if they exceed 1/1,000,000 for a lifetime exposure. Only eight of 31 carcinogenic exposure circumstances that exceeded this level were not regulated. They involved saccharin, aflatoxin, formaldehyde and polycyclic organic matter ... ” [p 81].

5.71 In fact, it is my view that over-reaction to the low risks produced by asbestos in place may lead to a greater risk — i.e. the carcinogenic risks imposed by asbestos removal programmes. Two mesotheliomas encountered in my own practice during 1999 occurred not in asbestos removal workers, but in others who sustained bystander exposure as a result of this activity: (i) pleural mesothelioma in a lecturer who had walked to and from her classroom at an Australian university each day over a period of weeks, through a building where an asbestos insulation removal programme was being carried out; (ii) pleural mesothelioma in a fireman who attended fires in buildings that contained asbestos-cement products and who participated in clean-up operations thereafter; about once a month for some years, he also attended and examined buildings where fire alarms had been activated by high atmospheric concentrations of asbestos fibres produced by removal programmes. (In addition, a recent survey in Finland found “occasional high fibre concentrations even inside personal protectors during asbestos removal work” [130]).

5.72 Two other important points are worth emphasis. First, because they usually focus on specific cohorts or groups of workers, epidemiological studies may fail to identify a small but real risk, because of low statistical power. In this respect, the documents submitted to the WTO state that it may be impossible to prove a negative (absence of risk), but one can also state that absence of proof does not constitute proof of absence. For example, a number of investigations have failed to identify a statistically significant increase in the relative risk (RR) of cancer among individuals with parietal pleural fibrous plaques. In an extensive review of asbestos and lung cancer, Henderson et al. [131] commented along the following lines in relation to pleural plaques and lung cancer:

“Nurminen and Tossavainen [132] also emphasized the issue of statistical power; they calculated the RR for plaque-associated lung cancer in the general population to be as low as 1.1, given the prevalence of 4.6% among unlikely exposed and 15.0% among probably exposed men with an estimated twofold risk of lung
cancer. Detection of this RR at a level of statistical significance would require a population sample of about 300 000.” [p 102].

5.73 In a discussion of the Hughes-Weill study [133] on radiological asbestosis and lung cancer in New Orleans asbestos-cement factory workers — one of the three key investigations that proposed an obligate intermediary step of pulmonary fibrosis for the induction of lung cancer by asbestos — Henderson et al. [131] also commented in the same review:

“... the number of lung cancer cases [in the Hughes-Weill investigation] was small. What number of workers would be required in such a study to detect an increase in risk of, say 1.4, 1.56 or 2.0, as opposed to the risk in workers with chest x-ray opacities? ... person-years of follow-up equivalent to 20-50 expected cases would be required to have any reasonable chance of detecting RRs of 1.4 to 1.6 at a level of significance of 0.05. ... The power level for the actual sample of 420 ... to detect a risk of 1.5 would be about 40%. That is, a true effect would be falsely declared ‘non-significant’ 60% of the time. ... The low power of the Hughes-Weill study is exemplified by the fact that ... lung cancer risk was not significantly associated with duration of employment or cumulative exposure (there was a fairly restricted range of employment periods) and even the association of lung cancer with fibrosis was only marginally significant.” [pp 93-94].

5.74 The point is that a low, non-significant or undetectable risk in a small cohort may nonetheless translate into a substantial body of disease when spread over a large population: e.g. an RR of 1.1 representing an increase in risk of 10 per cent may require a population size of 300,000 to be detectable at a level of statistical significance of 0.05, whereas this 10 per cent increase in a common disease such as lung cancer may amount to a substantial burden of disease when spread across a population of, say, 1,000,000, 10,000,000, or 100,000,000. (Please see also later discussion on mesothelioma among brake mechanics: answer to Question 2.)

5.75 Another point is that a high frequency of a cancer such as mesothelioma in a small population may be overshadowed in absolute numbers by a lower occurrence rate for the same disorder spread over a large population. For example, among non-smoking former Wittenoom workers, mesothelioma is now the most common cause of death [70] (in most cohorts exposed to amphibole asbestos, < 10 per cent will develop mesothelioma). Nonetheless, mesotheliomas among the Wittenoom cohort constitute only 5-6 per cent of the total burden of mesothelioma across the Australian population [AMR 99]. For example, the 1999 Report for the Register records 189 mesotheliomas among the former Wittenoom population with only a single exposure to asbestos, in comparison to 187 mesotheliomas among carpenters/joiners with only a single exposure to asbestos; the point is that the lower risk of mesothelioma from asbestos exposure among carpenters has produced almost the same number of cases, because carpenters among the Australian workforce constitute a much larger occupational group than the entire Wittenoom cohort of about 6000.

5.76 This observation also applies to the numbers of mesotheliomas among chrysotile miners and millers in Quebec, in proportion to other cases among the general population of Quebec. Bégin et al. [134] divided Quebec mesotheliomas into three groups, as shown in the following Table:

<table>
<thead>
<tr>
<th>Group</th>
<th>Type of asbestos exposure</th>
<th>Number of cases</th>
<th>Average age</th>
<th>Average duration of exposure</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Chrysotile miners and millers, Thetford and Asbestos, Quebec</td>
<td>49</td>
<td>62 ± 8.1 yrs</td>
<td>30.5 ± 13.7 yrs</td>
</tr>
<tr>
<td>2</td>
<td>Manufacturing, industrial insulation, shipbuilding yards of Quebec</td>
<td>50</td>
<td>56.7 ± 8.6 yrs</td>
<td>21.4 ± 14.5 yrs</td>
</tr>
<tr>
<td>3</td>
<td>General construction / building maintenance industries of Quebec</td>
<td>21</td>
<td>57.2 ± 7.2 yrs</td>
<td>27.2 ± 7.2 yrs</td>
</tr>
</tbody>
</table>

From Bégin et al. [134].
In this study, Bégin et al. [134] also commented that “the incidence of pleural mesothelioma in chrysotile miners and millers, although not as high as in the crocidolite workers, is well above the North American male rate”. They also observed that “asbestos exposures in Group 3, although difficult to quantify on the basis of the record, appear to be often very low intensity”. Bégin et al. also commented in the following terms:

“The present study documents an increasing incidence of malignant mesothelioma in chrysotile miners and millers of the eastern townships of Quebec, with 49 cases in the last 23 years and a rate of 2.5 cases per year in the last 10 years in the primary industry, as compared with a rate of 0.3 per year in the years prior to 1969 ... To put these rates into perspective, a comparison of the incidence for the combined population of the Asbestos and Thetford townships of Quebec of some 40,000 adult males or the maximal estimated workforce of 10,00-15,000 men [sic; surely this is a typographic error in the original, and it should be 10,000-15,000], 20 years ago and currently at risk, reveals that the incidence of mesothelioma in the chrysotile mining townships of Quebec would give an annual incidence rate of 62.5 cases per million per year for the 1980-1990 period, or in chrysotile miners and millers of Quebec, would give an incidence rate of 150-250 cases per million per year for the 1980-1990 period. These values are well below the annual incidence rate of the crocidolite mining townships of South Africa, estimated at 542 cases per million per year, and well above the rate for the North American population, estimated at between 2.5 to 13 cases per year per million adult males for the 1970-1980 period, and 14.1 cases per year per million adult males in 1984 and 15 cases per million for 1980 and projected to increase for the 1990s. ...

Thus, our observations add information of interest to the on-going debate regarding the relative carcinogenicity of different types of asbestos fibers. Our data suggest that some of the cases of malignant mesothelioma in Quebec chrysotile miners and millers may not be necessarily attributable to amphibole and could be chrysotile-induced. Lung tissue burden analyses, a better indication of exposure than tumour tissue burden, will be done on these cases to further investigate this point. ....

Finally, our data strengthen the view that a substantial number of malignant mesothelioma cases have a relatively short asbestos exposure, particularly seen in Group 3. In our study, 25% of all cases are in such a category” [pp 539-541].

In one of the documents submitted to the WTO, it is argued that evaluation of, and actions on, risks should be based on probability rather than mere possibility. This proposition is open to dispute. For example, action is often taken to avoid the possibility of harm — by regulation or prohibition — even though the likelihood of injury is remote, because of the seriousness of the potential outcome. In medical ethics, this is the principle of first, do no harm (primum non nocere). Two examples follow: (i) the antibiotic chloramphenicol was known to be highly effective in the treatment of various infections, including typhoid fever, but on rare occasions it induced bone marrow aplasia; despite the low likelihood of this side-effect — about 1 in 250,000 — the use of chloramphenicol was restricted to only a few life-threatening infections (e.g. typhoid fever), and it is now almost never used because safer effective alternatives are available; (ii) over recent years, there has been a flood of publicity over global warming and greenhouse gas emissions. A causal or direct relationship between greenhouse gases (such as CO2 and methane) and climate change is open to argument, and Earth undergoes repeated cycles of natural cooling and warming; in this respect, there is also evidence that melting of the Antarctic ice cap has been going on for some thousands of years, and global warming for over 100 years. Nonetheless, the consequences of inaction over greenhouse gas emissions are potentially so serious that strategies to reduce the release of these gases into the atmosphere are entirely appropriate, despite uncertainty over the link between them and global warming.
(f) General observations on induction of mesothelioma by asbestos, especially the amphibole varieties of asbestos such as crocidolite and amosite.

(i) The linkage between the amphibole varieties of asbestos, and commercial chrysotile, and the subsequent development of malignant mesothelioma is well established and is not in dispute.

5.79 This link is generally accepted as causal; in this respect, asbestos fulfils all The Bradford Hill Criteria for the establishment of causality [44].#

(ii) There is a dose-response relationship between cumulative exposure to asbestos and the subsequent incidence of mesothelioma in asbestos-exposed cohorts or populations; the incidence is also related to time since exposure, so that early exposures are more significant for mesothelioma induction than later exposures, other factors being equal.

5.80 This relationship is expressed by the Peto model and its various modifications:

$$ I = K^*F^*(T^p - [T - D]^p) $$

where $I$ = incidence; $K$ depends on fibre type, mix, size and other site-specific variables; $F$ = intensity of exposure in f/ml; and $D$ = years of exposure. For the purposes of modelling, $T$ can be replaced by $(T - 10)$ to build in a minimum 10-year lag-time, and the cubic power of time $(T^3)$ is often used, so that:

$$ I = K^*F^*([T - 10]^3 - [T - 10 - D]^3) $$

An important aspect of this model is that early exposures are more significant for mesothelioma induction than later equivalent doses.

5.81 Of the variables $D$, $F$ and $K$, it is $D$ that is the most accurately measurable, whereas the values for $K$ and $F$ are often unknown, though some estimates of $F$ can be made from the type of work activity. When there are multiple periods of employment for which the type of work is similar for each, one can assume that the value for each of $F_1$, $F_2$, $F_3$ ... $F_n$ remains constant, which also applies to $K_1$, $K_2$, $K_3$ ... $K_n$, so that:

$$ I \mu ([T - 10]^3 - [T - 10 - D]^3) $$

In practice, a simpler equation can be used: $I = ct^k$

where the constant $c$ is dependent on exposure, usually taken as proportional to the intensity of exposure multiplied by its duration (i.e. cumulative exposure), with weightings for different fibre types; the power $k$ remains about 3.5, or 3 for short periods of exposure. As de Klerk and Armstrong [135] state:

“The model predicts that risk is increased after each increment of exposure by an amount proportional to the level of exposure and the cube of time after that. In terms of the multistage model of cancer, it implies that asbestos acts at the first stage of a 4-stage process. ... The model predicts that incidence is much more dependent on early or low levels of exposure and increases less rapidly as exposure continues to increase, depending mainly on time since first exposed.” [p 232].

5.82 When one is faced with multiple exposures to asbestos, the following points emerge, specifically for mesothelioma induction, provided that the characteristics and time for each exposure are appropriate for a biological effect: (i) it is not valid to point to one exposure among the others and incriminate it as the sole cause of a mesothelioma, with exoneration of the other exposures; (ii) it is not valid to point to one exposure among the others and exonerate it from a causative role in the development of a mesothelioma, and to incriminate all the others; (iii) when there are multiple episodes of exposure as a background to a mesothelioma, it is often the case that each exposure in isolation would be
sufficient for attribution of the mesothelioma to asbestos, with the provisos mentioned above (characteristics and times of exposures). When each exposure among others is appropriate for mesothelioma induction if the particular exposure occurred alone, it is not logical to state that this exposure — which could have a biological effect in isolation — has no effect when in combination. In such circumstances, it is not the presence or absence of an effect that is in question, but the magnitude of each effect in proportion to the others.

5.83 A dose-response relationship has been observed with both estimates of airborne exposure to asbestos [136], and quantitative and qualitative fibre burden analysis of the asbestos content in human lung tissue of mesothelioma patients [3, 25, 137, 138]: e.g. see Rogers et al. [3], and, more recently, Williams et al. [138], who noted in 1997 that:

“It was shown there that while the relative risk of all three diseases [i.e. asbestosis, mesothelioma and lung cancer] increased with increasing exposure, the relative risk of malignant mesothelioma is greater at low levels of exposure when compared with the risk of asbestosis but is lower at very high levels of exposure.” [p. 39].

5.84 In their study on the relationship between lung asbestos fibre type and the lung tissue concentration of asbestos versus the relative risk of mesothelioma, Rogers et al. [3] made the following comment:

“Fiber content in the lung depends on both the amount of fiber deposited and the amount cleared. The amount deposited depends on duration and intensity of exposure in the occupational or general environment. Clearance rate is thought to be dependent on the amount deposited at any point in time, i.e., clearance is exponential. Thus, the same fiber content in the lung at death or time of resection may be achieved from a high initial deposition, followed by absence of deposition and absence of clearance over a long period of time, or by a continuous deposition at a lower level, with or without clearance. Since detailed mechanisms of mesothelioma initiation and progression are not known, ‘dose’ as estimated by final lung fiber content may not relate to the ‘dose’ required to produce mesothelioma. It is thus possible that a high lung fiber content in a mesothelioma case may represent continuing accumulation of fibers after a lower level of fibers had produced malignant change. It is more likely, however, that the malignant change did not occur until the fiber content reached a sufficiently high level.” [p 1913].

(iii) The dose-response relationship between the amphiboles and mixtures of asbestos types is linear at high exposures [15]

5.85 For example, please see EHC 203 and Table 4.

(iv) This dose-response relationship between asbestos exposure and the risk of mesothelioma has also been detected at low levels of exposure, which overlap with environmental exposures.

5.86 A recent case-control study [136] from France on the dose-response relationship between low levels of asbestos exposure and the odds ratio (OR) for mesothelioma showed a clear dose-response relationship between estimated cumulative asbestos exposure and the OR for pleural mesothelioma. In the final paragraph of the article, the authors stated:

“We found a clear dose-response relation between cumulative exposure to asbestos and pleural mesothelioma in a population-based control study, with retrospective assessment of exposure. A significant excess of mesothelioma was observed for levels of cumulative exposure that were probably far below the limits adopted in many industrial countries during the 1980s.” [last sentence of abstract].
Although some concerns have been expressed about this type of investigation [139], it is my opinion that these points were addressed in the original paper [136], and they are common and intrinsic to epidemiological studies of this type — e.g. see Camus et al. [140, 141]. This study [136] found an OR for mesothelioma of 4.2 [95 per cent CI 2.0-8.8] at estimated cumulative exposures of 0.5-0.99 fibre-year, with elevation of the OR at about 0.5 fibre-year. 

In a fibre burden study on mesothelioma patients, Rödelsperger [137] observed that:

“...A significantly increased OR [for mesothelioma] is obtained even within the very low concentration range of 0.1-0.2 F/µg [i.e. concentrations in the range of 100,000-200,000 fibres per gram dry lung tissue], which may be expected for about 5% of the population.” [p III] (which also corresponds to an estimated cumulative exposure in the range of about 1-2 fibre-years).

In a more recent study on mesothelioma cases (N = 66) and controls (N = 66), Rödelsperger et al. [25] found an OR for mesothelioma of 4.5 at fibre concentrations of 100,000 to < 200,000 per gram dry lung tissue (for fibres > 5 µm in length; 95 per cent CI 1.1-17.9). These authors also recorded an OR = 2.4 at concentrations of 50,000 to < 100,000 fibres per gram dry lung tissue (95 per cent CI 0.8-7.6). The controls for this study — surgical lung resections mainly for lung cancer — would be expected to bias the OR towards 1.0 (i.e. underestimate the effect) [25], and hence the OR of 2.4 probably represents a genuine doubling of risk or more at these low fibre concentrations.

From de Klerk and Armstrong [135].

### TABLE 4: INCIDENCE OF MESOTHELIOMA IN OCCUPATIONALLY EXPOSED GROUPS BY FIBRE TYPE AND TIME SINCE FIRST EMPLOYED

<table>
<thead>
<tr>
<th>Fibre type</th>
<th>Industry</th>
<th>Years since first employed</th>
<th>Rate per million person–years</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mixed: crocidolite, amosite and chrysotile</td>
<td>Manufacture textiles and insulation</td>
<td>20-24</td>
<td>1520</td>
</tr>
<tr>
<td></td>
<td></td>
<td>25-30</td>
<td>1710</td>
</tr>
<tr>
<td></td>
<td></td>
<td>30+</td>
<td>3180</td>
</tr>
<tr>
<td>Mixed, mainly amosite</td>
<td>Insulation workers</td>
<td>30-24</td>
<td>290</td>
</tr>
<tr>
<td></td>
<td></td>
<td>25-29</td>
<td>1550</td>
</tr>
<tr>
<td></td>
<td></td>
<td>30-34</td>
<td>2760</td>
</tr>
<tr>
<td></td>
<td></td>
<td>35-39</td>
<td>6300</td>
</tr>
<tr>
<td></td>
<td></td>
<td>40-44</td>
<td>6330</td>
</tr>
<tr>
<td></td>
<td></td>
<td>45+</td>
<td>8110</td>
</tr>
<tr>
<td>Mixed: crocidolite and chrysotile</td>
<td>Fibrous cement manufacture</td>
<td>20-24</td>
<td>2200</td>
</tr>
<tr>
<td></td>
<td></td>
<td>25-29</td>
<td>6300</td>
</tr>
<tr>
<td></td>
<td></td>
<td>30-34</td>
<td>9600</td>
</tr>
<tr>
<td>Chrysotile, some crocidolite</td>
<td>Textile manufacture</td>
<td>20-24</td>
<td>108</td>
</tr>
<tr>
<td></td>
<td></td>
<td>25-29</td>
<td>143</td>
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<tr>
<td></td>
<td></td>
<td>30-34</td>
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<td></td>
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<td>35-39</td>
<td>493</td>
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<td></td>
<td></td>
<td>40+</td>
<td>1774</td>
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<td>Insulation manufacture</td>
<td>20-24</td>
<td>734</td>
</tr>
<tr>
<td></td>
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<td>25-29</td>
<td>2623</td>
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<tr>
<td></td>
<td></td>
<td>30-34</td>
<td>5078</td>
</tr>
<tr>
<td></td>
<td></td>
<td>35+</td>
<td>1842</td>
</tr>
<tr>
<td>Mixed</td>
<td>Dockyards</td>
<td>20-24</td>
<td>120</td>
</tr>
<tr>
<td></td>
<td></td>
<td>25-29</td>
<td>410</td>
</tr>
<tr>
<td></td>
<td></td>
<td>30-34</td>
<td>220</td>
</tr>
<tr>
<td></td>
<td></td>
<td>35-40</td>
<td>370</td>
</tr>
<tr>
<td></td>
<td></td>
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</tr>
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<td>45-49</td>
<td>1510</td>
</tr>
<tr>
<td>Crocidolite</td>
<td>Mining and milling</td>
<td>20-24</td>
<td>900</td>
</tr>
<tr>
<td></td>
<td></td>
<td>25-29</td>
<td>2200</td>
</tr>
<tr>
<td></td>
<td></td>
<td>30-34</td>
<td>3000</td>
</tr>
<tr>
<td></td>
<td></td>
<td>35-39</td>
<td>7000</td>
</tr>
</tbody>
</table>
“Even within the concentration interval of 0.1-0.2 f/µg dry weight [i.e. 100,000 to 200,000 fibres per gram dry weight], a significantly increased odds ratio of 4.5 was obtained. Previously, the same method of tissue analysis was used to estimate a 95 percentile of the amphibole fibre concentration of 0.11 f/µg dry weight for persons without detectable exposure to asbestos at the workplace. Therefore, within the range of the normal background level [up to 300,000 fibres per gram dry lung in Germany], a positive dose response is observed.” [p 191].

This study did not detect an increase in the OR for chrysotile or for other mineral fibres.

5.89 The risk detected by Rödelsperger et al. [25] appears to correlate reasonably well with the French case-control study reported by Iwatsubo et al. [136], which found an OR for mesothelioma of 4.2 at estimated cumulative exposures of 0.5-0.99 fibre-year, with elevation of the OR at about 0.5 fibre-year.

(v) No lower threshold (minimum) level of asbestos exposure has been delineated, below which there is demonstrably no increase in the risk of mesothelioma.

5.90 This observation is expressed by Hillerdal [20] in the following terms: “There is no proof of a threshold value — that is, a minimal lower limit below which asbestos fibres cannot cause the tumour [i.e. mesothelioma] — and thus it is plausible that even such low exposure can cause mesothelioma (even if the risk is extremely low). Patients with mesothelioma whose lungs show fibre concentrations within the normal range cannot be dismissed as background cases, — that is, not due to asbestos. ... The only way to prove such a hypothesis would be to compare the incidence of mesothelioma in a group with such background exposure with the incidence in a truly non-exposed group. This is not possible, as no such group can be found.” [i.e. the lung tissue of virtually all mammals contains some asbestos fibres derived from natural, environmental or occupational sources] [p 510].

5.91 Points worth emphasis in Hillerdal’s review [20] include reports of mesothelioma among school teachers (9/487 patients with mesothelioma in one reference), in jewellers, and in individuals exposed to asbestos insulation at home (6/262 patients with mesothelioma according to one reference). Hillerdal [20] also makes the point that low-level asbestos exposure “more often than not contains peak concentrations which can be very high for short periods” (e.g. airborne asbestos fibre concentrations of up to 78 f/ml from sweeping asbestos from the floor [New Caledonia]).

(vi) The dose-response relationship for commercial Canadian chrysotile and mesothelioma incidence is also linear at high levels of exposure.

5.92 For example, among the Quebec chrysotile miners and millers it has been noted that:

“All the observed 38 cases were pleural with the exception of one of low diagnostic probability, which was pleuro-peritoneal. None occurred in workers exposed for less than two years. There was a clear dose-response relationship, with crude rates of mesotheliomas (cases/thousand person-years) ranging from 0.15 with cumulative exposure < 3530 million particles per m³ (mpcm)-years (< 100 million particles per cubic foot (mpcf)-years) to 0.97 for those with exposures of more than 10 590 mpcm-years (> 300 mpcf-years).” [EHC 203, p 8].

(vii) So far as I am aware, there are no observational data on dose-response relationships between chrysotile only at low exposure levels and mesothelioma incidence; in this respect, estimates are based on extrapolation of a linear dose-response line from high exposures down to low exposures.
“Overall, the available toxicological data provide clear evidence that chrysotile fibres can cause fibrogenic and carcinogenic hazard to humans. The data, however, are not adequate for providing quantitative estimates of the risk to humans. This is because there are inadequate exposure-response data from inhalation studies, and there are uncertainties concerning the sensitivities of the animal studies for predicting human risk.” [EHC 203, p 7].

5.93 Because of the lack of such data, no definite threshold for chrysotile in relation to mesothelioma and lung cancer has been delineated: According to EHC 203 (p 144):

“(a) Exposure to chrysotile asbestos poses increased risks for asbestosis, lung cancer and mesothelioma in a dose-dependent manner. No threshold has been identified for carcinogenic risks.”

5.94 In summary:

TABLE 5: ASBESTOS-RELATED DOSE-RESPONSE RELATIONSHIPS FOR MESOTHELIOMA

<table>
<thead>
<tr>
<th>Exposure Level</th>
<th>Amphiboles</th>
<th>Chrysotile</th>
</tr>
</thead>
<tbody>
<tr>
<td>Heavy exposure</td>
<td>Dose-response effect; linear</td>
<td>Dose-response effect; linear</td>
</tr>
<tr>
<td>Low-level exposure</td>
<td>Dose-response effect</td>
<td>No data</td>
</tr>
<tr>
<td>Threshold</td>
<td>No threshold delineated</td>
<td>No threshold delineated</td>
</tr>
</tbody>
</table>

(viii) To the best of my knowledge, there are no observational data on the interactive effect of low (or for that matter, high) concentrations of inhaled chrysotile fibres only, when these are superimposed later and separately upon a pre-existing amphibole ± chrysotile burden within lung tissue (?superimpositional additive or multiplicative effect).

5.95 For example, it has been estimated that up to 15-20 per cent of men in industrialized societies may have sustained occupational exposure to asbestos (chrysotile/amphiboles). Rödelsperger et al. [137] indicate that fibre concentrations of 100,000-200,000 amphibole fibres per gram dry weight lung tissue may be expected for about 5 per cent of the population in Germany. We do not know what the effect of subsequent chrysotile fibre inhalation on top of this type of amphibole burden might be.

“Data were analysed on a case-referent basis, to relate relative risks of mesothelioma to dose of fibre, as measured both by lung content and estimated airborne exposure. Multivariate analysis of cases found a dose response relationship for lung fibre content of crocidolite, amosite and chrysotile and the development of mesothelioma. Either a multiplicative or additive model could be used to fit the relative risk/dose coefficients for the various asbestos types. A progressive increase in relative risk with increasing fibre content was reported for all fibres ... Tests for trend were highly significant in all cases.” [NICNAS 99, p 61].

(ix) There is a long lag-time between asbestos exposure and the subsequent diagnosis of mesothelioma (10 years as a minimum; usually in the range of 20-40 years). It follows that the mesotheliomas encountered in the 1990s and the incidence of mesothelioma in various nations are a consequence of exposures, especially occupational exposures, sustained from the 1940s through to the 1970s and even beyond.

5.96 Exposures from the 1940s through to the 1980s usually involved one or more of the amphibole varieties of asbestos. For example, asbestos-cement building products used in Australia usually contained one or more of the amphibole varieties of asbestos, namely crocidolite or amosite, or both, at different times; in this respect, the use of crocidolite in the products was discontinued in 1966, and amosite in 1984.

5.97 Peto et al. [24] make this point in the following terms:
“The extraordinarily high mesothelioma incidence throughout Western Europe in men born around 1945-50 reflects the extent of asbestos use in the 1960s and 1970s at the beginning of their working lives. Annual raw asbestos imports to European Union countries peaked in the early to mid 1970s and remained above 800,000 tonnes per year until 1980, falling to about 100,000 tonnes by 1993 (European Commission, 1996). Increasingly stringent exposure limits were enforced in the manufacture of asbestos-containing products over this period, but exposure to users of such materials, particularly in the building industry, remained virtually uncontrolled in many countries. Chrysotile asbestos products are still widely used in several European countries, and maintenance or demolition work on older buildings may result in substantial exposure to amphiboles as well as to chrysotile. We have not included men born after 1955 in our projections, but the effects of asbestos exposure during the 1980s and 1990s, although not yet apparent, may prove considerable.” [p 670].

(x) Properties of asbestos fibres that determine carcinogenicity

5.98 As indicated in the documents submitted to the WTO Panel, the properties of asbestos fibres implicated for mesothelioma induction (and, possibly, lung cancer and other disorders), can be summarized as the three Ds:

5.99 Dose: this issue is covered in the preceding sections [(f)(ii)to (vi)].

5.100 Dimensions: according to The Stanton Hypothesis, the carcinogenicity of asbestos fibres appears to reside primarily in long thin fibres (length > 5 µm and especially > 8 µm, and in the range of 10-20 µm, and diameters < 0.25 µm) — e.g. see Pott [142]. On the other hand, shorter fibres appear to be less carcinogenic, although data indicate that tremolite fibres > 4 µm in length and < 1.5 µm in diameter produce malignant mesenchymal tumours when implanted into the pleural cavities of rats [2]. On the other hand, very short-length fibres appear to have little carcinogenic activity, although Churg [143] comments on fibre dimensions in the following terms:

“The has been extensive investigation of the relation of mesothelioma induction and fiber size in experimental models. Using intrapleural inoculation of different types of fibres with different size distributions, Stanton et al. concluded that long, thin (i.e., high aspect ratio) fibres were much more powerful mesothelial carcinogens than were short, thick fibres and that fiber type was less important. The exact size of fiber that qualifies as long and thin is unclear: fibers ... longer than 8 µm and widths narrower than 1.5 µm are usually cited from Stanton’s work, but the same experiments show that fibers with lengths greater 4 µm and widths less than 0.25 µm were also effective carcinogens. The Stanton hypothesis has been supported by animal inhalation experiments using size-separated fibers: few mesotheliomas were found with either amosite or chrysotile prepared to contain few fibers longer than 5 µm.

Human data on the question of fiber length and mesothelioma are equivocal. The tremolite found as a natural constituent of chrysotile ore is a relatively short, thick fiber compared with commercial amosite or crocidolite, and if one attributes ‘chrysotile-induced’ mesotheliomas in man to the tremolite component, the differences in mesothelioma do correlate with fiber size. However, attempts to prove this proposition directly have produced equivocal results ... McDonald et al. concluded that the number of fibers longer than 8 µm explained most mesotheliomas and that chrysotile played no role. However, Rogers et al. found that fibers both longer and shorter than 10 µm, including chrysotile fibers, played a role, although long fibers were generally more important. The problem with both of these studies is that most patients with mesothelioma have had occupational asbestos exposure, and fibers in lungs from those with occupational exposure are always longer than fibers in the general population; thus the same result would have been obtained if the test group were exposed but had no disease or
some disease other than mesothelioma. My colleagues and I have attempted to circumvent this problem by comparing fiber sizes in a chrysotile mining and milling cohort and a cohort with heavy amosite exposure, using exposed workers with no disease as the control group. In neither cohort could we show that fibers in mesothelioma cases were significantly longer and thinner than those in the other disease categories or even in the disease-free workers.” [p 353].

5.101 In other words, it is possibly the bio-persistence of amphibole fibres that is important for mesothelioma induction, rather than precise fibre dimensions.

5.102 Durability (bio-persistence): the greater mesotheliomagenic (mesothelioma-producing) potency of the amphiboles in comparison to chrysotile is widely ascribed to greater persistence of the amphiboles in tissues, with significantly longer half-lives than chrysotile (please see later discussion, Section (g)(v)). On the other hand, it is conceivable that the same effect might be achieved by sustained inhalation of chrysotile over a prolonged time interval or, possibly, shorter, but more intense exposures so that the chrysotile fibres persist despite shorter half-lives than the amphiboles.

(xi) There is general though not universal agreement of a differential potency between the amphiboles versus chrysotile for mesothelioma induction

5.103 In this respect, the amphiboles are substantially more potent, with estimates ranging from 2-4X, to 10X, to 12X on a fibre-for-fibre basis, to 30X, to a 30-60X greater potency, or more (e.g. please see EHC 203). A minority view that the amphiboles in chrysotile have roughly equal mesotheliomagenicity is not supported by the prevailing evidence for humans. Although acknowledging the greater potency of the amphiboles for mesothelioma induction, some argue that chrysotile is of equal or greater importance overall, because chrysotile accounts for > 95 per cent of world asbestos production. According to this perspective, commercial chrysotile is a weaker carcinogen on a fibre-for-fibre basis, but this lesser potency is multiplied across a much greater tonnage, leading to an overall equivalent or greater effect [144].

(xii) Tobacco smoke plays no role in the development of mesothelioma at any anatomical site — unlike the synergy between asbestos and tobacco smoke for the causation of asbestos-related lung cancer (see section (i)(i) below).

(g) Commercial Chrysotile and Mesothelioma Induction

(i) There is general agreement that commercial chrysotile has the capacity to induce mesothelioma in experimental animals and humans

5.104 There is dispute, however, over which fibres in commercial chrysotile are implicated (i.e. the predominant chrysotile or the trace quantities of fibrous tremolite).

(ii) Canadian chrysotile contains trace amounts of tremolite, including fibrous tremolite, as a contaminant [2, 10, 13, 14, 145-148]

5.105 The amount of tremolite appears to vary from one sample to another, but is generally < 1 per cent (please see EHC 203).

(iii) It has been argued that the occurrence of mesotheliomas among the Quebec chrysotile miners and millers is a consequence — not of the chrysotile per se — but of the coexistent trace quantities of tremolite (a non-commercial amphibole).

5.106 Analysis of the asbestos fibre content of lung tissue from this cohort demonstrates disproportionately high concentrations of tremolite in comparison to chrysotile; this appears to represent a bio-accumulation phenomenon whereby chrysotile is cleared from lung tissue more rapidly than the tremolite, so that the tremolite not only persists but increases in proportional concentration. In this respect, the tremolite content of the lung tissue can be used as an index on the past chrysotile exposure and some claim that the incidence of mesotheliom-
(iv) It is known that fibrous tremolite has the capacity for mesothelioma induction

5.107 Mesotheliomas related to the use of tremolite in whitewash or stucco have been reported in Turkey, Greece, Cyprus and Corsica [149-152] (for additional references, see Hillerdal [20]).

“Tremolite asbestos, a minor component mineral of commercial chrysotile, has also been shown to be carcinogenic and fibrogenic in a single inhalation experiment and an intraperitoneal injection study in rats. Exposure/dose-response data are not available to allow direct comparison of the cancer potency of tremolite and chrysotile.” [EHC 203, p 6].

5.108 Tremolite has also been implicated in lung cancer and mesothelioma induction in a group of vermiculite miners in Montana [2, 16, 153, 154]. It appears that these miners were exposed only to tremolite-actinolite fibres. The group was shown to have a:

“... very high lung cancer incidence (standard mortality ratio [SMR] 285 ...), as well as four cases of mesothelioma and eight of pneumoconiosis. Examination of sputum samples from all but three (170/173) current workers demonstrated asbestos bodies (AB) in 75%, the numbers showing a close parallel with cumulative exposures in fibre-years.” [2] [p 493].

5.109 Case [2] has extensively reviewed the biohazards of tremolite, including epidemiological investigations in humans and experimental data on animal models. In his review, he emphasized the pathogenicity of the tremolite found in Quebec chrysotile samples, especially at Asbestos and in the Thetford mine:

“Tremolite was not identified in Montreal air, was just detectable (0.2 fibres/l) in Asbestos, and was one order of magnitude higher in Thetford mines (still only 1.5 fibres/l or 0.0015 fibres/cc ...).” [pp 496-497].

5.110 He also favoured the expression “chrysotile/tremolite” for Quebec chrysotile:

“As to the separate issue of ‘chrysotile vs. tremolite’, few would dispute the abilities of both to produce lung cancer and asbestosis, again in sufficient exposure dose. The weight of epidemiological, animal, and, especially, lung internal-dose biomarker studies leads to the inevitable conclusion that it is the tremolite ‘component’ of Quebec chrysotile which causes mesothelioma [but please see later discussion in this report]. It is unfortunate that adequate terminology for tremolite-contaminated chrysotile has not been introduced: I for one would favour the simple compound phrase ‘chrysotile/tremolite’. ” [p 500].

5.111 Case [2] also states:

“... it becomes important to know to what degree ‘chrysotile-in-place’ is really ‘chrysotile/tremolite-in-place’. No easy answer can be expected: both bulk analyses and air sampling, even with analytical electron microscopy, can miss very low levels of tremolite. Studies in the Quebec mining district indicate that, at the very least, such low levels (roughly 0.0015 fibres/cc) can induce biological effects (i.e., pleural plaques). Unfortunately, only expensive in vivo animal bioaccumulation assay systems can truly answer the question: the alternative is to wait 40 to 50 years for the next wave of asbestos disease — which is likely to occur mainly among present-day asbestos abatement workers and to some degree in custodial personnel and other tradesmen” .... [p. 500].

(v) Clearance of chrysotile from lung tissue

5.112 It is well known that chrysotile fibres are cleared more rapidly than amphiboles,
especially in long-term studies [145]. Clearance of amphibole fibres does occur and the clearance mechanisms appear to be more effective for short fibres (for both chrysotile and the amphiboles) so that the mean length of retained fibres increases over time. Chung and Vedal [155] calculated a half-life in lung tissue of about 20 years for amosite. Estimates of the tissue half-life for crocidolite fibres have been somewhat shorter (in the order of 5-10 years) [156-158], and de Klerk et al. [158] could find no difference between the clearance rates for long and short fibres. Oberdörster [159] estimates human clearance half-times to be about 90-110 days for chrysotile and 200-1500 days for crocidolite fibres > 16 µm in length, based on extrapolated rat and primate inhalation data.

5.113 It has been claimed that chrysotile is cleared from lung tissue within 28-48 hours of inhalation. This claim seems extraordinary and begs the question: why, if chrysotile is cleared from lung tissue so rapidly, is it still demonstrable in human lung tissue many years or decades after cessation of inhalation of commercial chrysotile (or mixtures of asbestos types)? For example, in one of my recent referral cases — an elderly man with lung cancer who sustained exposure from mixing loose asbestos and sweeping up dried insulation materials — an asbestos fibre analysis carried out on lung tissue resected 16 years after his exposure stopped showed a total asbestos fibre count of 8,440,000 fibres/g of dry lung (> 1 µm in length; aspect ratio = 3:1), made up by 6,250,000 chrysotile fibres + 940,000 tremolite fibres + 940,000 anthophyllite fibres + 310,000 crocidolite fibres (the 24 year lag-time is enough for a carcinogenic effect).

(vi) The Quebec chrysotile cohort

4.114 In an analysis of mesotheliomas among the Quebec chrysotile miners and millers, up to 1997, McDonald et al. [13, 14] reported 38 mesotheliomas, and most of these occurred after prolonged and heavy exposure, especially at the mine where the greatest concentrations of trace tremolite occurred (Thetford). For example, these authors [13] recorded the breakdown of the mesotheliomas shown in Table 6 (below).

5.115 McDonald et al. [13] identify two main reasons for the low mesothelioma rate from the five smallest mines (1 case only among 6010 person-years, equivalent to 166 cases per million person-years): firstly, workers within this sub-group were younger than the remainder of the cohort; secondly, these mines had been opened recently so that “there were inadequate periods of latency”. A single additional mesothelioma shortly after completion of the study would erase the difference in incidence rates between the five smallest mines and the main complex. McDonald et al. [13] go on to indicate that the other rates are “reasonably comparable”. In comparison to the Thetford main complex, there were relatively few mesotheliomas among workers at the asbestos mine and mill (23 versus 8), despite nearly equivalent person-years of observation; in addition, asbestos fibre analysis on lung tissue demonstrated crocidolite and amosite in five out of the eight cases from the mine and mill at Asbestos and in two out of the five mesotheliomas from the Asbestos factory (Table 7, below). In focussing on the Thetford mines group, it was noted that most of the mesotheliomas came from the five central mines (Area A; Group C) as opposed to the 10 peripheral mines (Area B; Group P), so that the odds ratio for mesothelioma for Group C plus employees who had jobs in both Area A and Area B (Group M) was 2.50 (based on net service; 20 adjusted years), in comparison to an odds ratio of 0.80 for Group P.

<table>
<thead>
<tr>
<th>Number of mesothelioma deaths</th>
<th>Subject-years (000s)</th>
<th>Rate (per 100,000 subject-years)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Thetford Mines: Main complex and the oldest of the smaller mines</td>
<td>23</td>
<td>65.14</td>
</tr>
<tr>
<td>The five smallest mines</td>
<td>1</td>
<td>6.01</td>
</tr>
<tr>
<td>Asbestos: Mine and mill</td>
<td>8</td>
<td>60.64</td>
</tr>
<tr>
<td>Factory</td>
<td>5</td>
<td>10.84</td>
</tr>
</tbody>
</table>

From McDonald et al. [13].
The clear implication of this complex and sophisticated study is that the risk of mesothelioma was related strongly to years of service in the central area at Thetford where geological factors “in Area A would probably result in tremolite, some in fibrous form, being mined with the ore”. In addition, the mesothelioma rate for miners and millers was > 2.5 times higher at Thetford mines (excluding the smallest mines) than at Asbestos, and this difference was also attributed to differences in the amount of fibrous tremolite in the ores. Despite these differences within the cohort for the distribution of mesothelioma related to chrysotile and tremolite (and also to crocidolite and amosite at the Asbestos factory and the Asbestos mine and mill), the results indicate that Quebec chrysotile — on average contaminated by fibrous tremolite in small amounts — is capable of mesothelioma induction: the Abstract describes 25 mesotheliomas from the Thetford mines, representing a mesothelioma rate of 337 per million person-years, which is substantially (almost 20X) higher than the mesothelioma incidence rate of about 17 per million per person-years for men in British Columbia and the USA in 1982 and 1973-1984 respectively, and well above the background rate for spontaneous mesotheliomas of 1-2 per million person-years.

**Table 7: Asbestos fibre concentrations in lungs at autopsy from 21 mesothelioma cases among Quebec chrysotile miners and millers**

<table>
<thead>
<tr>
<th>Place of employment</th>
<th>No. of cases</th>
<th>Chrysotile</th>
<th>Tremolite</th>
<th>Crocidolite</th>
<th>Amosite</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mines and mills</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Thetford Mines</td>
<td>14</td>
<td>12.8</td>
<td>104.1</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Asbestos</td>
<td>5</td>
<td>4.3</td>
<td>7.5</td>
<td>1.7</td>
<td>0.3</td>
</tr>
<tr>
<td>Factory</td>
<td>2</td>
<td>2.1</td>
<td>0.5</td>
<td>6.4</td>
<td>0.3</td>
</tr>
</tbody>
</table>

Table from McDonald et al. (1997): Table 2 in the original reference. See also Table 1 in the original.

In calculating geometric means, a zero count has been replaced by half the detectable limit.

For crocidolite and amosite, all counts were zero: i.e. below the detectable limit.

For fibre counts/g lung tissue, multiply the figures by 106.

In the final two paragraphs of the paper, McDonald et al. [13] comment as follows:

“The tremolite hypothesis, if correct, has several important implications. First, it supports the widely but not universally held view that most, if not all, asbestos-related mesotheliomas are caused by amphibole fibres. This in turn points to fibre durability and biopersistence as critical factors in aetiology ... a point of even greater relevance in assessing the safety of man-made mineral fibres. Second, it implies that uncontaminated chrysotile carries very little risk of mesothelioma. In Asbestos, exposures were not to uncontaminated chrysotile, but also to some tremolite and crocidolite, yet among the miners and millers only five deaths from a total of over 3,300 can be confidently attributed to their work. At present-day levels of dust control the mesothelioma risk must be vanishingly small. Even so, it remains desirable to minimise, perhaps by screening, the contamination of commercial chrysotile by amphibole fibres, however difficult this may be.” [p 718].

Despite the importance of this study by McDonald et al. [13], the following comments can also be made:

- The number of mesotheliomas in all groups except for the Thetford main complex was small (1, 8 and 5 mesotheliomas respectively; please
see Table 6 above). In this respect, misdiagnosis or misclassification of the mesotheliomas according to the places worked could significantly affect the results, although there is no evidence that this happened; however, the probability for the diagnosis of mesothelioma also varied, with a high probability in 19 cases, moderate probability in 14, and a low probability (though considered more likely than not) in five; of these 38 cases, only 18 had been coded on the death certificate to ICD 163, and the rest to a variety of other diagnostic codes. Furthermore, in analysing the mesotheliomas according to Area A versus Area B at the Thetford mines (groups C, M and P), the numbers were 104 for group C, 69 for group P and 35 for group M; McDonald et al. noted that the odds ratio for group P was unstable as shown by the “very wide confidence intervals, and as the point estimate is well below unity it is quite unrealistic.”

• The low incidence of mesotheliomas in the Quebec chrysotile cohort appears to parallel similar low incidence rates for asbestosis and lung cancer for the same cohort

[160, 161]: the incidence rates for lung cancer and mesothelioma appear to be different in other chrysotile-exposed cohorts.

5.119 For these reasons and because of the different rates of various asbestos diseases (asbestosis, lung cancer and mesothelioma) between the Quebec cohort and other groups of workers, I would be reluctant to recommend national policies from the findings in this cohort in isolation, and I would look for coherence of the evidence across different cohorts and studies.

5.120 In relation to the Quebec cohort, there is an important error in the Canadian reply to Question 4 (see Annex II) from the European Communities, where the following statement is made:

“Regarding asbestos-related mesothelioma, a number of studies have demonstrated cogently that this type of cancer is almost exclusively linked to exposure to amphiboles. Cases of mesothelioma in chrysotile asbestos miners in Quebec are quite rare — in a cohort of 11,000 workers who were very carefully tracked (in the McDonald study), there were no more than 50 or so cases over several decades. Exhaustive research on their employment history revealed that most of the cases were related to short-term exposure to commercial amphiboles. For example, during World War II, some of the miners with mesothelioma had worked in plants manufacturing products for the Allied Forces and amphiboles imported into Canada had been used to make a variety of products, including gas masks, to assist in the War effort.”

5.121 The statement that “most of the cases were related to short-term exposure to commercial amphiboles” is incorrect and misleading. As demonstrated in the study by McDonald et al. [13], most of the mesotheliomas occurred among chrysotile miners who worked at the Thetford main complex, without exposure to commercial amphiboles such as crocidolite or amosite. This is clearly shown in Table 7 (above), slightly modified from the paper by McDonald et al. [13] where fibre burden analysis on lung tissue from 14 mesothelioma cases from the Thetford mines showed both chrysotile and a high concentration of tremolite, with a zero count for the commercial amphiboles crocidolite and amosite. The point to be emphasized is that the mesotheliomas from the Thetford mines were not related to commercial amphiboles such as crocidolite or amosite, but to chrysotile with its content of fibrous tremolite.

(vii) As discussed earlier, a dose-response relationship between the incidence of mesothelioma and cumulative asbestos exposure has been demonstrated for commercial chrysotile.

5.122 Mesotheliomas have also been produced in experimental animals by implants-
tion and inhalation of chrysotile (presumably also containing trace amounts of tremolite). Mesotheliomas can also be induced in rats by intraperitoneal injection of chrysotile, with evidence of a dose-response effect [1] (see also bibliography for EHC 203).

“In non-inhalation experiments (intraperitoneal and intraperitoneal injection studies), dose-response relationships for mesothelioma have been demonstrated for chrysotile fibres.” [EHC 203, p 5].

(viii) Chrysotile is also known to be toxic to a variety of cell lines in vitro, with induction of a variety of chromosomal alterations (e.g. please see EHC 203, pp 69-102).

(h) Other Chrysotile-Exposed Cohorts and Studies

5.123 In addition to the Quebec chrysotile miners and millers, mesotheliomas have also been reported among other workforces apparently exposed only to chrysotile, with no significant tremolite.

(i) Russia

5.124 Chrysotile from the Urals region (Uralasbest) in Russia [162, 163] is said to represent pure chrysotile. Although precise figures for the mesothelioma incidence in this area are difficult to procure, Kogan [164] makes the following comment in a recently-published textbook on occupational lung diseases:

“In the Middle Ural mountains, the main asbestos mining region in Russia, only chrysotile asbestos is produced. In the 50 districts of this region, the mortality from mesothelioma over a 10-year period was six-fold higher than the average rate in the Sverdlovsk region, an area of negligible asbestos mining. Most with mesothelioma had worked at the asbestos mining and milling plants, or had lived in an adjacent town near old and very ‘dusty’ mills.” ... [p 251].

5.125 Because it is difficult to equate exposure levels in the Russian chrysotile industry with other industries (e.g. the airborne fibre concentrations at Uralasbest are usually expressed as gravimetric measurements), and I have been unable to ascertain the numbers of cases relative to exposure levels, I consider this evidence to be weak in comparison to other studies.

5.126 One might expect data on mesothelioma incidence in Central and Eastern European nations to be of interest, from an assumption that some of these countries would have imported mainly chrysotile from Russia until the breakup of the Soviet Union. Unfortunately, it is difficult to evaluate national mesothelioma statistics, because a number of these nations also imported amphibole asbestos. For example, in Slovenia, the total consumption of asbestos (1947-1995) was 580,000 tonnes, of which crocidolite accounted for 37,133 tons, until its use was stopped in 1992 [165]. Similarly, the annual usage of asbestos in Bulgaria during the 1970s and 1980s reached approximately 32,000 tons of chrysotile (mainly from Russia and Canada), together with about 1000 tons of crocidolite from Africa and 6000-7000 tons of Bulgarian amphibole material (anthophyllite and tremolite) [166]. In Poland, total consumption of asbestos for the manufacture of asbestos-cement products between the end of the Second World War until 1993 was about 1.4 million metric tons which included about 8500 metric tons of amosite and approximately 86,000 metric tons of crocidolite [167].

(ii) Germany

5.127 The former German Democratic Republic (GDR): Sturm et al. [5, 7] have published data on asbestos-related diseases and asbestos types in the German State of Saxony-Anhalt. These authors pointed out that:

“All asbestos-based products were made from raw asbestos which was primarily imported from the former Soviet Union, particularly from the Kiemba min-

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ing area in the Ural mountains (said to represent pure chrysotile). Small quantities of long-fibred grades came from Canada (2,990 tonnes in 1989) and were mainly used for the manufacture of asbestos-cement pressure pipes free of amphibole asbestos. This was a share of approximately 7% in total imports. We never obtained any information about the Canadian mines from which the asbestos processed in the former GDR originated. ... However, several analyses carried out by the GDR Central Institute for Industrial Medicine confirmed that both the Canadian and the Russian asbestos were pure chrysotile. In addition to these imports of chrysotile asbestos, smaller quantities of amphibole asbestos were imported. For example, in the period from 1980 to 1985, some 90 tonnes of anthophyllite were imported annually from Mozambique. This anthophyllite was used exclusively by a Berlin manufacturer and was of acid-proof products, similar to the way crocidolite had been used in previous years to produce filters, seals and acid and lye-proof plastic materials. In Saxony-Anhalt, our region of work, these amphibole imports did not have any significance from the point of view of industrial medicine” ... [p 318/173].

5.128 Between 1960 and 1990, a total of 1082 mesotheliomas was recorded in Saxony-Anhalt, and these included 843 “proven asbestos-accepted mesotheliomas”; Table 8 from Sturm et al. [5, 7] gives a breakdown of 812 cases for which adequate data were available: 67 were said to follow exposure to chrysotile only, and 331 were associated with “chrysotile; possible amphiboles”.

(iii) Italy

5.129 Two mesotheliomas have now been recorded among more than 900 workers employed at the Balangero mine and mill in Italy [168, 169]. EHC 203 gives the following summary:

“The cohort of chrysotile production workers employed at the Balangero mine and mill ... was almost exactly one tenth the size of the Quebec cohort. At the end of 1987, when 427 (45%) of the cohort had died, there were two deaths from pleural mesothelioma, both in men employed for more than 20 years with cumulative exposure estimated respectively at 100-400 and > 400 f/ml years. One diagnosis was confirmed histopathologically, and one was based on radiological findings and examination of pleural fluid. Fibrous tremolite was not detected in samples of chrysotile from this mine, but another fibrous silicate (balangeroite), the biological effects of which are not known, was identified in low proportions by mass (0.2-0.5%). At a comparable stage in the evolution of the Quebec cohort, mesothelioma accounted for 10 of 4547 deaths, a lower but not dissimilar proportion.” [p 112].

<table>
<thead>
<tr>
<th>TABLE 8: MESOTHELIOMAS ACCORDING TO TYPES OF EXPOSURES</th>
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</thead>
<tbody>
<tr>
<td>Amphiboles</td>
</tr>
<tr>
<td>Age at beginning of exposure</td>
</tr>
<tr>
<td>Duration of exposure</td>
</tr>
<tr>
<td>Lethal period (years)</td>
</tr>
<tr>
<td>Age of person dying of mesothelioma</td>
</tr>
<tr>
<td>Number of mesotheliomas</td>
</tr>
</tbody>
</table>

Note: All types of application of asbestos with common addition of chrysotile fall under the heading “Chrysotile. Amphiboles possible” when previous admixture of amphiboles cannot be definitely excluded. From Sturm et al. [5, 7].

(iv) China

5.130 At the XV International Scientific Meeting of the International Epidemiological
Association (Florence, September 1999), Yano et al. [170] presented a paper on lung cancer incidence in a cohort of 515 male asbestos workers heavily exposed to chrysotile containing < 0.001 per cent tremolite, in Chongqin; two mesotheliomas over 11,850 person-years of observation occurred in this cohort (discourse to the paper; assuming this rate to be representative, it would amount to 170 mesotheliomas per million person-years).

5.131 In a retrospective cohort mortality study of 1227 men employed at a chrysotile mine in Hebei Province of China before 1972, Zou et al. found three deaths from mesothelioma (please see EHC 203, p 120).

(iii) United States

5.132 Two mesotheliomas have also been observed among the cohort of South Carolina chrysotile textile workers — who used Canadian chrysotile — studied by Dement et al. [171, 172] (please see EHC 203, p 115).

(vi) Australia

5.133 There is also some indication of an increased frequency of mesothelioma among Australian brake mechanics who were potentially exposed only to chrysotile from grinding of brake blocks that contained Canadian chrysotile (please see later discussion on friction products, and NICNAS 99 and AMR 99).

(vii) Zimbabwe

5.134 One pathologically confirmed case of mesothelioma has been recorded in association with occupational exposures to asbestos in the Zimbabwe mines and/or mills, with one other case said to resemble mesothelioma radiologically (EHC 203, p 121).

(viii) Fibre burden studies on human lung tissue from mesothelioma patients

5.135 Fibre burden analyses also support the notion that some mesotheliomas occur in association with, or as a consequence of, inhalation of pure chrysotile.

5.136 Morinaga et al. [173] detected asbestos fibres in 19 of 23 mesothelioma studied; amphibole fibres were found in 13 cases, but six were found to have only chrysotile fibres (five pleural mesotheliomas and one peritoneal mesothelioma). Nonetheless, the methodology for this study seems unimpressive, with relatively small numbers of fibres analysed.

5.137 The 1991 paper by Rogers et al. [3] recorded a substantial number of mesothelioma patients in whom the only detectable type of asbestos was chrysotile (Table 9), with evidence of a dose-response effect as reflected in a trend to an increasing odds ratio (OR) at a relatively low fibre concentration of = 10^6 fibres per gram dry lung tissue (log_{10} = 5.5–6; OR = 8.67).

| Fibre burden studies on human lung tissue from mesothelioma patients |

<table>
<thead>
<tr>
<th>TABLE 9: DISTRIBUTION OF FIBRE CONCENTRATION: TRANSMISSION ELECTRON MICROSCOPIC ANALYSIS, CHRYSOTILE ONLY (ALL LENGTHS)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mesothelioma cases</td>
</tr>
<tr>
<td>No.</td>
</tr>
<tr>
<td>---</td>
</tr>
<tr>
<td>f/g</td>
</tr>
<tr>
<td>log_{10} (f/g)</td>
</tr>
<tr>
<td>5.5-6</td>
</tr>
<tr>
<td>6-6.5</td>
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<tr>
<td>6.5-7</td>
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<tr>
<td>7-8</td>
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<tr>
<td>&amp;</td>
</tr>
<tr>
<td>From Rogers et al. [3]. CI: confidence interval; f/g: fibres per gram of dried lung tissue.</td>
</tr>
</tbody>
</table>

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Finally, fibre burden studies have demonstrated that both chrysotile fibres and amphibole fibres can translocate from lung parenchyma to reach the pleura; EHC 203 summarizes these findings in the following way:

"In a study of asbestos fibres in the lung parenchyma and the parietal pleura of 29 asbestos workers, Sebastien et al. (1980) found that chrysotile fibres predominated in the pleura and that amphibole fibres could not be detected. A similar result was reported by Dodson et al. (1990). Kohyama & Suzuki (1991) found short chrysotile fibres in pleural plaques and in mesothelial tumours. In contrast, Boutin et al. (1993) found 0.21 x 10^6 fibres per g of parietal pleura and 1.96 x 10^6 in samples of lung parenchyma. Fibre concentrations were higher in subjects with a history of asbestos exposure and most of the fibres were amphiboles. Chung (1994) reported detection of chrysotile fibres in the subpleural parenchyma in chrysotile miners and millers." [pp 64-65].

(ix) Other Observations

Nicholson and Raffn [8] analysed mesothelioma risk over 40 studies for which little or no exposure information was available, using the excess numbers of lung cancers as a measure of exposure and comparing the ratios of mesotheliomas to excess lung cancers across these studies. They suggested that:

"... the ratio of mesothelioma to excess lung cancer is the same for exposures to 100% chrysotile (presumably Canadian chrysotile), 97%+ chrysotile, 100% amosite, and mixtures of chrysotile, amosite and crocidolite, within statistical uncertainty. Only 100% crocidolite exposures appear to have a greater ratio, about two to four times that of predominantly chrysotile. This relatively small difference in the potential for crocidolite to produce mesotheliomas compared with other fibre exposure cannot explain the high risk seen in chrysotile exposures accompanied by a very small crocidolite exposure. The data speak strongly that much of the mesothelioma risk in predominantly chrysotile exposures is from the chrysotile." [p 402].

In other words, these authors appear to argue, like Smith and Wright [144], Stayner et al. [11], and Landrigan et al. [21] that although chrysotile may be cleared more rapidly from lung tissue than tremolite — and that tremolite can be used as an indicator of past chrysotile — it may not be valid to ascribe all the mesothelioma risk to the tremolite and to ignore the far more numerous chrysotile fibres. Nonetheless, I do not find Nicholson and Raffn’s argument to be persuasive, taking into account the K values for different industries.

Therefore, it is my perception that epidemiological and experimental evidence clearly demonstrates that Canadian chrysotile with its trace amounts of fibrous tremolite has the capacity for mesothelioma induction. Although the tremolite may have a disproportionately large effect, it is my perception that the evidence does not allow one to conclude that the chrysotile has no effect on mesothelioma induction: there is evidence from other cohorts and studies that chrysotile per se can also induce mesothelioma, even when tremolite is undetectable, and in experimental models in animals, chrysotile is as carcinogenic as, and more toxic than, the amphiboles. However, there is also general agreement that in humans, chrysotile is substantially less carcinogenic for the mesothelium than the amphiboles, and my estimate is that it has a potency 1/10^6 – 1/30^6 the carcinogenicity of crocidolite, with amosite being less mesotheliomagenic than crocidolite but more carcinogenic than chrysotile on a fibre-for-fibre basis. Amosite is an important factor in the incidence of mesothelioma in the United States, because of its widespread use in insulation materials from the 1960s [155, 174-176].

Some salient features of asbestos-associated lung cancer include the following:

(i) Synergy between asbestos and tobacco smoke

Historically, most asbestos workers have also been cigarette smokers, and the lung cancer rate in virtually all cohorts is an outcome of the combined and synergistic effects of tobacco smoke and asbestos. Vainio and Boffetta [179] emphasize that asbestos and tobacco smoke are complex carcinogens that can affect multiple steps in the multistage process of cancer evolution, and that the combined effects will depend on the relative magnitude of each carcinogen at each stage; the interactive effect ranges from less than additive to supramultiplicative, but the model for insulation workers approximates a multiplicative effect (reviewed in Henderson et al. [131]). If the multistage model of carcinogenesis holds, and asbestos and smoking act at different stages, then a multiplicative relationship follows [180]. Leigh et al. [178] have reviewed various models for the apportionment of fractional contributions from cigarette smoke and asbestos towards the development of lung cancer.

(ii) Lung cancer incidence rates for asbestos-associated lung cancer vary greatly from one cohort to another

Please see following discussion.

(iii) Asbestos fibre type and lung cancer risk

The greater carcinogenicity of the amphiboles for the mesothelium in comparison to chrysotile appears not to extend to the induction of lung cancer [11]. In this respect, chrysotile is implicated in one of the lowest rates of asbestos-associated lung cancer (in Quebec chrysotile miners and millers), but also the highest rate (in South Carolina asbestos textile workers who used Canadian chrysotile) [171]. The reasons underlying this 30-fold difference in lung cancer risk remain unknown (reviewed recently by McDonald [161]; please see also EHC 203). The risk of lung cancer in other asbestos-exposed cohorts is intermediate between these two extremes [15].

(iv) Dose-response relationship

In most studies, there is a direct and linear relationship between the relative risk of lung cancer and cumulative exposure to asbestos, including chrysotile and the amphiboles.

Accordingly, EHC 203 gives the following account:

“The slopes of the relationship between cumulative exposure to chrysotile and the relative risk of lung cancer are summarized in Table 23 for those studies that reported this information. These studies all expressed this relationship using the following linear relative risk (RR) model:
$RR = 1 + B \times E$

where $B$ is the slope and $E$ is the cumulative exposure to chrysotile asbestos expressed in f/ml-years.

The slopes from the studies of the mining and milling industries (0.0006 to 0.0017), the latter having been estimated on a subset of the cohort on which the former was based, and the friction production industries (0.0005 to 0.0006) are reasonably similar. Hughes et al. (1987) in a study of cement workers (section 7.1.2.1b) reported a similar slope (0.0003) in one plant (plant 1) that only used chrysotile, and a nearly 20-fold higher slope (0.007) among workers only exposed to chrysotile in another plant (plant 2).

The slopes of 0.01 and 0.03 reported for the two studies of the chrysotile-exposed textile workers conducted on overlapping populations, as well as the slope of 0.007 from one of the two plants (plant 2) of cement workers in the study of Hughes et al. (1987), were an order of magnitude greater than those reported for the other cohorts. It should be noted that the two textile cohorts were identified from the same textile facility, but were based on different cohort definitions. Hence, it is not surprising that the slopes from these two studies were similar. The slopes in the studies of chrysotile-exposed textile workers are also remarkably similar to those reported in other studies of textile workers with mixed fibre exposures (Peto, 1980; McDonald et al., 1983b; Peto et al., 1985). This similarity in findings provides some support for the validity of the slopes reported in the chrysotile-exposed textile cohorts.

The reason for the much higher slopes observed in studies of textile workers is unknown, although several possible explanations have been suggested. The first is that these differences might be attributed to errors in the classification of exposures in these studies. Particular concern has been raised about errors in the exposure assessment related to conversions from mpcm (mpcf) to fibres/ml that were performed, particularly in the mining and milling studies (Peto, 1989). Sebastien et al. (1989) conducted a lung burden study specifically designed to examine whether the differences in lung cancer slopes observed in the Charleston chrysotile textile cohort and the Quebec mining industries could be explained by differences in errors in exposure estimates. Lung fibre concentrations were measured in: (a) 32 paired subjects that were matched on duration of exposure and time since last exposure; and (b) 136 subjects stratified on the same time variables. Both analyses indicated that the Quebec/Charleston ratios of chrysotile fibres in the lungs were even higher than the corresponding ratios of estimated exposures. This finding was interpreted by the author as being clearly inconsistent with the hypothesis that exposure misclassification could explain the large discrepancy in the lung exposure-response relationships observed in the two cohorts. “[pp 118-119].”

Boffetta [15] expresses the relationship in the following terms:

“A large number of studies have been conducted on lung cancer risk following asbestos exposure. The interpretation of their results is complicated by several factors: (i) dose, geological type of fibres and industry are all important determinants of risk and are strictly correlated; (ii) the biologically relevant exposures occur 20 or more years before appearance of the disease, and their quantitative assessment is imprecise; and (iii) the role of potential confounders, in particular, tobacco smoking, can hardly be evaluated. In general, the risk of lung cancer is smaller in studies of miners and friction product manufacturers, is intermediate in studies of asbestos-cement and asbestos product manufacturers, and is highest in studies of asbestos textile workers. This likely reflects a stronger carcinogenic effect of individual, long and thin fibres, like those occurring in the
textile industry, as compared to grouped, short and coarse fibres, like those occurring in mining.

Several cohort studies provide sufficient details to allow a quantitative evaluation of the risk of lung cancer from cumulative asbestos exposure. In all cohorts, the empirical relationship fits well a linear correlation with no threshold, which can be expressed as:

$$RR_1 = 1 + K_1 \times CE,$$

where $RR_1$ is the relative risk of lung, $CE$ represents cumulative asbestos exposure, expressed as $fb/ml$-yrs, and $K_1$ is the industry-specific slope of the relationship (RR for the increase in 1 $fb/ml$-year of exposure) for lung cancer and varies across cohorts. Similarly, the risk difference ($RD_1$) can be expressed as

$$RD_1 = K_1 \times CE \times Exp,$$

where $Exp$ is the number of expected cases of lung cancer. In other words, the number of cases of (or deaths from) lung cancer attributable to asbestos exposure depends on the number of expected cases (deaths), the cumulative exposure, and the intrinsic carcinogenic potential of the exposure circumstance. The value $K_1$ varies from 0.05-0.01 in cohorts of insulation and asbestos textile workers to 0.001-0.0005 in friction manufacturers and miners, while cohorts with mixed exposure have, in most cases, intermediate values. ... While all estimated values of $K_1$ are positive, the type of asbestos does not seem to be correlated to lung cancer risk.

In the interpretation of these results, however, one should consider several limitations. Most studies are based on a small number of cases or deaths: for example, the risk estimate of 100 $fb/ml$-yrs for the cohort of asbestos textile workers presented by McDonald and colleagues (RR 2.4) has a 95% confidence interval from 1.7 to 3.8. Another source of uncertainty, and possibly bias, relates to the estimate of cumulative exposure: in the same cohort of asbestos textile workers, the range of RRs based on the extremes of the distribution of possible exposure values is 1.3-6.7. For these reasons, several governmental and scientific committees have suggested to adopt an ‘average’ value of $K_1$, independent from fibre type and circumstance of exposure ...: the most widely accepted value is 0.01 which corresponds to an increase of 1% of the risk of lung cancer for each $fb/ml$-yr of exposure. ...

Tobacco smoking is the main cause of lung cancer, and this applies also to the cohorts of asbestos-exposed workers. Despite the limitations of the available studies, which limit the precision of the estimates of the combined effect of the two carcinogens, the risk from tobacco smoking seems to act synergistically with that of asbestos exposure, according to a multiplicative model. ... The available data are consistent with the most widely accepted model of quantitative dose-response between cumulative exposure to asbestos and lung cancer risk, which assumes a linear relationship with no threshold. Alternative models, however, would also be consistent with the data. In particular, as no precise data are available for cumulative exposures below 1 $fb/ml$, a model with a threshold at low exposure cannot be rejected.” [pp 473-475].

(v) Histological types of lung cancer

5.150 Although some studies have shown a relative excess of adenocarcinomas in proportion to other histological types of lung cancer, all of the major histological types occur among asbestos workers in proportions equivalent to, or only slightly different from, those
in the general population [112]. Therefore, the histological type of a lung cancer has no value in ascertaining whether or not asbestos has contributed significantly to the genesis of the cancer (reviewed by Henderson et al. [131]).

(vi) Lobar distribution and the central versus peripheral distribution of asbestos-related lung cancer

5.151 Some studies have reported a reversal of the upper lobe:lower lobe ratio for lung cancers in asbestos workers, in comparison to a reference non-exposed population. Recently, Lee et al. [181] addressed the lobar distribution of lung cancer in asbestos-exposed individuals and found that the tumours were predominantly located in the upper lobe (i.e. they did not find a reversal of the upper lobe to lower lobe ratio). The lobe of origin for a cancer has no value in ascertaining whether the cancer is likely to be asbestos-related. The distribution of lung cancer between the central versus the peripheral airways does not differ significantly in asbestos workers from a control non-exposed population (please see Henderson et al. [131]).

(vii) Asbestos and lung cancer risk

5.152 Cumulative exposure versus fibrosis (asbestosis): as discussed already, most epidemiological studies dealing with lung cancer risk in asbestos workers have reported a direct correlation between the relative risk of lung cancer and cumulative asbestos exposure, although the slope of the dose-response line varies from one cohort to another. Most of the documents submitted to the WTO appear to agree on this relationship, the main area of uncertainty or dispute being the question of whether a threshold exists or not.

5.153 However, Canada’s answers to questions from the Panel and the European Communities appear to resurrect the fibrosis-cancer hypothesis, which postulates that asbestos does not induce lung cancer per se, but only through an obligate intermediary step of pulmonary fibrosis (asbestosis), so that fibrosis becomes the determinator of lung cancer risk, not cumulative exposure:

"1. Canada does not disagree that chrysotile causes lung cancer. However, the way in which exposure to chrysotile asbestos may increase the risk of lung cancer has not yet been fully explained; it could be just an indirect cause. ...

2. The risk may become detectable in cases of long-term exposure to high levels, but it is by no means certain that chrysotile acts as a direct carcinogen or that it acts in the form of pulmonary fibrosis, which would be a precursor to neoplasia. In other words, exposure must be intense and long enough to induce pulmonary fibrosis, which predisposes the pulmonary parenchyma to a high risk of cancer.”

5.154 It is my perception that the fibrosis?cancer hypothesis represents a minority opinion: with some prominent exceptions, most authorities in this area reject the fibrosis?cancer theory and focus instead on the asbestos fibre burden in lung tissue as the main determinator for lung cancer risk, as discussed earlier in this report.

5.155 The fibrosis?cancer hypothesis is predicated upon three key but flawed studies:

- In the investigation reported by Kipen et al. [182], there was major problem with case selection (only 138 cases out 450 — 31 per cent — had a tissue specimen with sufficient non-malignant tissue for assessment of fibrosis); in addition, the histological criteria used for the diagnosis of asbestosis are unacceptable to most pathologists — i.e. no asbestos bodies in some cases; fibrosis restricted to the subpleural zone considered to be asbestosis — so that this study seems to have suffered from an over-diagnosis of asbestosis [183, 184].
As discussed in Section (e) above, the Hughes-Weill study [133] on chest X-ray opacities related to lung cancer mortality in New Orleans asbestos-cement workers had low statistical power, so that it had only a 40 per cent chance of detecting a significant lung cancer standardized mortality ratio (SMR) of 1.5. Other studies based on X-rays have shown an increase in risk or mortality for lung cancer in the absence of radiological asbestosis (e.g. Wilkinson et al. [185], Finkelstein [186] and de Klerk et al. [187]).

The autopsy study on South African crocidolite miners reported by Sluis-Cremer and Bezuidenhout [188] was also bedevilled by problems of selection (black people excluded; autopsies on 36.7 per cent of deaths only; autopsies on cases for which compensation was sought). Analysis of the findings indicates that the effect of duration of exposure (the most accurately measurable of the exposure variables) was still significant even after adjustment for the grade of asbestosis and other variables. This indicates that exposure to asbestos still had an independent effect on lung cancer mortality even after adjustment for the grade of asbestosis, as in the study reported by Wilkinson et al. [185]. In subsequent correspondence, Sluis-Cremer and Bezuidenhout [189] conceded that when they carried out a logistic regression analysis, allowing for the grade of asbestosis, years of exposure accounted for most of the variation, but the degree of asbestosis still emerged as a highly significant risk factor.

Recently, Case and Dufresne [190] have commented as follows:

"... Hughes and Weill go much further in stating that asbestosis is a prerequisite for lung cancer attribution in those with asbestos exposure. This statement goes beyond the known facts and relies on mechanistic speculation. The authors believe that asbestosis is produced by a mechanism or mechanisms that will also lead to lung cancer. Their hypothesis requires that the mechanism(s) always be intermediate in that lung cancer always follows asbestosis. Finally, the speculation requires that lung cancer occurring without asbestosis can never be caused by asbestos exposure alone (or in synergy with cigarette smoking) regardless of the level of that exposure, and that no mechanism can occur that does not involve intermediate fibrosis. The biological fallacy of this argument has been well documented ... one must remember that lung cancer originates in the large airways, while asbestosis is a disease of the lung parenchyma at and beyond the respiratory bronchioles. ... To ignore our knowledge of indices of exposure other than the simple presence or absence of asbestosis is simplistic and biologically naive." [p 1118].

The case-control studies carried out on South Carolina asbestos textile workers by Dement et al. [171] clearly undermine the fibrosis-cancer hypothesis and, in this respect, they constitute Pepper’s Black Swan factor:194 Dement and his colleagues clearly identified a lung cancer SMR > 2.5 at 2.7-6.8 fibre-years of exposure (well below the exposure level necessary for histological asbestosis in the same cohort [191]).

The first written submission from Canada also refers to the study reported by Camus et al. [140] on non-occupational exposure to chrysotile asbestos in Quebec and the risk of lung cancer:

"It is also interesting to note the work of Dr Camus et al. (see Camus, M., Siemiatycki, J. Meek, B., Nonoccupational Exposure to Chrysotile Asbestos and the Risk of Lung Cancer, (1998) 338, New England Journal of Medicine 1565). They
published a vast study on women in chrysotile mining communities in Quebec, many of whom were exposed to very high levels of fibres between 1920 and 1975. These women were subjected to exposure of 0.0107 f/ml, higher than the current exposure limits in France, and literally thousands of times higher than the levels measured in public buildings. Nonetheless, no excess in lung cancer was detected in this population. According to the study’s authors, this is particularly important in the light of the current French situation. In fact, applying the risk model adopted by France for the exposure studied, results in a forecast of approximately 100 lung cancer deaths, while in reality there are none. Likewise, use of the French risk model would have resulted in estimates of approximately 250 and, at any rate, no less than 50 deaths from mesothelioma, while the preliminary results of the study in question show only 10 cases, some of which may be associated with exposure to amphiboles. Research continues, particularly with an analysis of the work history of each individual in order to determine the exact link, if any, between these cases of mesothelioma and on-the-job exposure, as well as exposure to amphiboles.”

In fact, Camus et al. [140] investigated the relative risk of death from lung cancer among 2242 deaths between 1970 and 1989 among women = 30 years of age who lived in two chrysotile asbestos-mining areas that comprised eight towns of which three (Thetford mines, Black Lake and Asbestos) contained nearly all the asbestos mines and mills. Eighty percent of the women lived within 4 km of a mine or mill, and all lived within 10 km.

The estimated average cumulative level of exposure was 25 fibre-years (range 5-125 fibre-years) made up by neighbourhood exposure (16.0 fibre-years), household exposure of 7.8 fibre-years and occupational exposure of 1.2 fibre-years, making a total of 25.0. The authors of this study pointed out that:

“... The lower limit of 5 fibre-years per ml corresponds, for example, to 50 years of exposure to asbestos at a level of 0.1 fibre per ml (the actual mean ambient airborne asbestos level in the area in 1974); the upper limit of 125 corresponds for example, to 50 years of exposure to 2.5 fibres per ml — a relatively low exposure level in local asbestos-mining and asbestos-milling industries before 1960.”

This investigation found a standardized mortality ratio of 1.0 in comparison to the reference population (i.e. no observed excess of lung cancer mortality). However, seven deaths from “pleural cancer” were observed (RR = 7.64; p < 0.05).

A few points about this study are worth emphasis:

- The Quebec chrysotile miners and millers have a low risk of lung cancer in comparison to other cohorts, such as the South Carolina chrysotile textile workers, for whom the frequency of lung cancer is at least 30 times higher. Therefore, it is not surprising that the low risk of lung cancer in the chrysotile miners and millers of Quebec extends across residents exposed environmentally to the same ore. In other words, the absence of a detectable increase in lung cancer mortality in female residents of this region of Quebec may not apply to other groups exposed environmentally to asbestos from other asbestos industries.

- The study reported by Camus et al. [140] stimulated considerable correspondence in the columns of the same journal (NEJM), and at least two of the correspondents (Churg [193] and Case [192]) emphasized that the seven-fold increase in mesothelioma mortality (seven cases) among the women was probably explicable by occupational exposure to amphiboles from manufacture of gas masks, repair of bags that contained imported asbestos, and, possibly in one case, domestic exposure to “tremolite brought home on miners’ clothes”.
In his letter to the editor, Case [192] also pointed out that "[T]hese women were exposed to levels of chrysotile as high as 1 fibre per ml of air as recently as one month in 1984."

I have some misgivings over the exposure estimates for this female population, and the figure of 25 fibre-years from environmental exposure in the general neighbourhood or vicinity of the Quebec chrysotile industry seems high in comparison to neighbourhood or environmental exposures from other industries. For example, ECH 203 (p 35) reproduces a Table of asbestos fibre concentrations in Quebec chrysotile mining towns, where the fibre concentration in 1984 is in the vicinity of 0.005 fibre/ml and the concentrations in 1973 and 1974 are given as 0.08 fibre/ml. In other words, Case’s figure of 1 fibre/ml for one month in 1984 [192] may be doubtful, unless there were some catastrophic event in the industry, with a burst of asbestos into the general environment. Unless earlier environmental airborne fibre concentrations were substantially above the 1973/1974 concentrations, it is difficult to see how a cumulative exposure of 25 fibre-years would come about; e.g. Camus et al. [140] state that residence in the area for 50 years at a mean fibre concentration of 0.1 fibre/ml would lead to the lower estimate of 5 fibre-years.

In addition, the estimate of 25 fibre-years seems high in comparison to data on environmental airborne fibre levels related to the Zimbabwean and Russian chrysotile industries. For example, EHC (p 47) states:

“There are some data concerning fibre levels in the air close to chrysotile mines. Baloyi (1989) found fibre levels around the Shabani mine (Zimbabwe) to range from below the limit of detection of the method” (less than 0.01 f/ml) to 0.02 f/ml of air, assayed by PCOM. [PCOM = phase contrast optical microscopy].

Scherbakov et al. [163] also give a comparable environmental airborne fibre concentration in Asbest City of 0.1 mg/m³ (comparative data for the same industry [194] suggest that the gravimetric measurement of mg/m³ is very roughly equivalent to the same number of fibres/ml).

The point is that if the estimate of cumulative asbestos exposure in the Quebec female population is high, this would lead to underestimation of lung cancer risk or mortality. For example, no detectable increase in lung cancer mortality among the 2242 deaths would be expected at the low cumulative estimate of = 5 fibre-years.

In addition, in their reply to the Letters to the Editor, Camus and Siemiatycki [141] state that “[W]e agree ... that the study had low statistical power to detect small risks; this was conveyed by the wide confidence intervals for our risk estimates ...”, although they go on to indicate that the Quebec study should have detected a risk of the magnitude predicted by the Environmental Protection Agency [EPA].

(ix) The Helsinki Criteria

5.163 This set of criteria deals with attribution of lung cancer to asbestos for the individual patient [113]:

736
"Because of the high incidence of lung cancer in the general population, it is not possible to prove in precise deterministic terms that asbestos is the causative factor for an individual patient, even when asbestosis is present. However, attribution of causation requires reasonable medical certainty on a probability basis that the agent (asbestos) has caused or contributed materially to the disease. The likelihood that asbestos exposure has made a substantial contribution increases when the exposure increases. Cumulative exposure, on a probability basis, should thus be considered the main criterion for the attribution of a substantial contribution by asbestos to lung cancer risk." [p 314; emphasis in original].

5.164 The Helsinki Criteria set an exposure level of = 25 fibre-years of exposure; however, it should be emphasized that this level of cumulative exposure is required for the individual patient as an index for an asbestos-attributable relative risk of lung cancer of = 2.0 (which, in the individual patient, equates to a probability of causation or material contribution of = 50 per cent — the civil standard of proof). Intended as a criterion for individual compensation, this exercise is clearly different from population-based relative risks relevant to the dispute before the WTO.

5.165 In summary:

<table>
<thead>
<tr>
<th>TABLE 10: ASBESTOS-RELATED DOSE-RESPONSE RELATIONSHIPS FOR LUNG CANCER</th>
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<tbody>
<tr>
<td>Chrysotile or Amphiboles</td>
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<tr>
<td>Heavy exposure</td>
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<tr>
<td>Dose-response effect; linear</td>
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<td>Low-level exposure</td>
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<tr>
<td>Dose-response effect for South Carolina textile workers (chrysotile)</td>
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<td>Threshold</td>
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<td>No threshold delineated</td>
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(i) Some General Observations on Experimental Models of Asbestos Carcinogenesis, including in vivo and in vitro Systems

(ii) In vivo experimental models

5.166 Although animal models of asbestos carcinogenesis - especially induction of mesotheliomas in animals such as rats - are of value to demonstrate the capacity of different fibres to induce tumours and to elucidate the mechanisms underlying carcinogenesis, they are not strictly comparable to carcinogenesis in humans, for a number of reasons:

- The airborne fibre concentrations to which experimental animals are exposed for inhalation experiments are substantially higher than in workplace or environmental situations for humans.

- The routes of administration of asbestos or other fibres - e.g. injection or direct implantation into the pleura or peritoneum - are not comparable to the human situation, with the exception of inhalation experiments.

- High concentrations of asbestos or other fibres are necessary to reduce latency intervals so that a reasonable yield of mesotheliomas or other cancers is obtainable within the life span of the animal used. In other words, the latency intervals are not comparable to the human model.

- There are known to be marked differences in the susceptibility of different species to asbestos carcinogenesis.

5.167 For example, in a review of asbestos and lung cancer, Henderson et al. [131] state the following:
“The dose of asbestos delivered by inhalation or installation over a short time interval in experimental animals, the lag-times, and the histological spectrum of the tumors also make it difficult or impossible to extrapolate the findings from such models to humans. The exposure to asbestos in positive inhalation experiments seems to have been so high that fibrosis was an unavoidable association with an increased cancer risk (exposure to at least 100 f/ml, > 1,000 f/ml for some groups, 5 x 7 hours per week, up to 12 months or more). Wagner et al. remarked on a number of ‘surprising’ results in their study (e.g. no differences in carcinogenicity or fibrogenicity between chrysotile and the amphiboles). ...”

The sensitivity of humans to the carcinogenic effects of asbestos is about 100-fold greater than that of rats. ...

... Experimental studies of this type address asbestos inhalation in isolation, instead of asbestos combined with tobacco smoke [for the study of lung cancer]. Hence, they are of questionable relevance to most lung cancers in asbestos workers, for which tobacco smoke is an important co-factor.

For the reasons stated above, we consider that the existing literature on tumorigenesis by inhalation of asbestos in laboratory animals allows no conclusions on the asbestos-asbestosis-lung cancer controversy in humans.” [p 96].

5.168 Davis [195] comments in the following terms:

“In experimental inhalation and injection studies, however, chrysotile has repeatedly produced as many mesotheliomas as other asbestos types. This finding probably indicates that the carcinogenic potential of chrysotile to cells is as high as the other asbestos types, and it is just sufficiently durable to exert its maximum effect in rats, although it is unable to survive long enough to do so in humans.” [p 201; but see discussion in this report on chrysotile clearance from lung tissue, Section A.(g)(v)].

(ii) In vitro systems

5.169 It is obvious that the effects of asbestos and other fibre types on isolated cell lines used for in vitro studies are not comparable to the induction of mesothelioma or lung cancer in humans. In vitro studies of this type are of most value in showing that asbestos and other fibres can induce chromosomal injury, oncogene expression or mutations similar to those induced by other known carcinogens.

5.170 Detailed discussion of the voluminous literature on this topic lies beyond the scope of this report. Henderson et al. [131] give some details of the effects of asbestos on cell lines in vitro; more extensive reviews are given in EHC 203 (pp 69-102), Both et al. [196], and Mossman et al. [197-202], and Bielefeldt-Ohlmann [203]. Only a few recent studies on chrysotile follow:

- “In the study by Haugen et al. [204], chrysotile was about 10 times more cytotoxic than amosite or crocidolite (as assayed by inhibition of clonal growth rate) and > 100-fold more toxic than glass fibres; epithelial cells were 10-15 times more sensitive to the cytotoxic effects of asbestos fibres than bronchial fibroblasts from the same human. We can find no comparison with mesothelial cells in this paper [204], despite at least one claim to this effect [197] ...” [p 97].

- “Harrison et al. demonstrated synergy between the lung carcinogen N-nitrosodimethylamine (NMDA) and chrysotile in the production of hyperplastic epithelial lesions in the lungs of rats, with a dose-response relationship for NMDA, augmented by chrysotile. Neoplastic
lesions (adenoma and adenocarcinoma) were found only in animals treated with both NHMI and asbestos, but the number of such tumours was small (N = 6 among 115 rats studied)” [p. 118; see Henderson et al. [131] for references].

- “Hei and Piao reported on malignant transformation of a human papillomavirus-immortalized human bronchial epithelial cell line (BE2PD) by a single 7-day treatment with chrysotile: the cells so treated evolved through a series of sequential steps to become tumorigenic, with the formation of progressively growing tumours in nude mice” [p. 118; see Henderson et al. [131] for references].

- In an investigation of the capacity of different asbestos fibre types to induce loss of heterozygosity [LOH] mutations in lymphocytes and diploid mesothelioma cells that were heterozygous for the HLA A2/A3 histocompatibility complex studied (in collaboration with Dr David Turner at the Department of Haematology-Oncology at the Flinders University), it was found that chrysotile was more toxic to the cell lines used so that few viable cells remained, making it difficult to evaluate LOH mutations, in contrast to UICC South African crocidolite.

- More recently, Dr. Turner and I have investigated the effects of UICC South African crocidolite injected into the peritoneal cavity of mice, for the investigation of somatic intrachromosomal recombinational events in mice that are transgenic for the gene that encodes the enzyme β-galactosidase; using PCR [the polymerase chain reaction], we detected a 5-fold reduction in SIR within only a few days of administration of the crocidolite. This finding parallels the results obtained with other carcinogens (e.g. cytotoxic drugs used for cancer chemotherapy) and may be explicable by a reduction of SIR because the asbestos produces an increase in other classes of mutation (e.g. point mutations or deletions), or because of impairment of DNA repair mechanisms.

5.171 The picture now emerging on asbestos carcinogenesis is a prolonged multistage parametric process [205], in which asbestos fibres may participate in both the initiation and promotion phases [196]. Some classes of mutation potentially inducible by asbestos - e.g. loss of heterozygosity mutations - are implicated in the initiation or progression phases of cancer development in humans, thought to be related to loss of tumour suppressor genes (e.g. retinoblastoma, astrocytoma, and colonic, gastric, prostatic and breast cancer) [206-211].

5.172 Free radicals — generated either from the surface of the fibres themselves [205, 212-215] or via macrophages [213, 216-218] — have been shown to have genotoxic or clastogenic properties [205, 212-214, 217, 219, 220], and are also implicated in asbestos carcinogenesis.

2. Questions by the Panel and Comments by the Scientific Experts

Question 1:

High-density chrysotile products (i.e. products where chrysotile fibres are bound in a matrix, such as chrysotile-cement, as opposed to “friable” products, such as flocking and heat insulation) represent the main use of chrysotile asbestos. The parties to this dispute disagree as to the circumstances of exposure to chrysotile and the risks to human health associated with such products. In this context, various questions arise with respect to the risks to human health associated with the use of high-density chrysotile products, in particular chrysotile-cement (of particular concern are installation, modification, repair, maintenance, demolition and disposal).
Canada argues that workers who are at greatest risk of exposure to chrysotile asbestos are, in descending order: (i) chrysotile miners and workers employed in the processing (milling) industry; (ii) workers in the chrysotile textile industry; (iii) workers involved in the production of friction materials (such as brakes, clutches); (iv) workers involved in the manufacturing of chrysotile-cement products; (v) workers involved in the removal of asbestos from buildings; and (vi) workers involved in construction, renovation, maintenance and the heat insulation of buildings. Furthermore, according to Canada, the last two categories are likely to be exposed to amphiboles. On the other hand, the European Communities argues that, in France, the secondary users, which includes installation, maintenance, repair, insulation, waste management and “handyman” type persons, etc. are at the greatest risk of exposure and that they are mainly exposed to chrysotile asbestos, since, for some fifty years, chrysotile has represented about 97 per cent of asbestos consumption in that country. Could you comment on these contrasting views, with a special focus on current uses and products?

Dr. de Klerk:

This question is rather curious and is either irrelevant or the wrong words have been used: trying to elucidate an ordering of working groups according to their “risk of exposure to chrysotile asbestos”. The risk of an event is the probability that it will occur. The event in question here is that a worker will come into contact with chrysotile asbestos. It is certain that the workers in groups (i) to (v) are exposed to chrysotile so their risks are all the same and equal to 1.0. Workers in group (vi) may not come into contact with chrysotile so their risk of exposure is less. The more relevant question here is: who is likely to receive the most exposure and therefore have the greatest risk of disease? In general, workers in well-regulated industries, where government inspection is mandatory, where there is a long history of efficient industrial hygiene practices, will have less risk of disease than those in the smaller less well-regulated industries. A good example can be found with silicosis: the majority of cases now occurring in both the USA and Australia arising from small unregulated industries with no awareness of risks or hygiene practices. Similar examples from own experience are: witnessing (in 1992) use of Russian asbestos in an asbestos-cement factory in Czechoslovakia (as it was then) where all the warnings on the bags of asbestos were in English; passing by demolition in progress of an old asbestos cement factory building in Sydney last month where no observable precautions of any kind were being taken. (Note added later: It has struck me that the misunderstanding with this question could be due to the relative imprecision of the French language, where “de” means both “of” and “from”, words with quite different meanings, especially in this context!)

Dr. Henderson:

In past historical terms, the Canadian proposition about the classes of workers at risk of exposure to chrysotile is correct — provided that this risk is expressed in terms of a numerical value for the risk per person-years of observation (e.g. per 100,000 or 1 million person-years). However, this situation has changed over recent years, as airborne fibre concentrations have been reduced in the mining and milling industries and during the production of friction products. As one example, NICNAS 99 points out that manufacture of friction products (brake linings and gaskets) in Australia is a completely closed operation, with low airborne fibre concentrations.

EHC 203 refers to this reduction in airborne fibre concentrations:

"Based on data mainly from North America, Europe and Japan, in most production sectors workplace exposures in the early 1930s were very high. Levels dropped considerably to the late 1970s and have declined substantially to present day values. In the mining and milling industry in Quebec, the average fibre concentration in air often exceeded 20 fibres/ml (f/ml) in the 1970s, while they are now generally well below 1 f/ml. In the production of asbestos-cement in Japan, typical mean concentrations were 2.5-9.5 f/ml in 1970s, while mean con-
In contrast, the risk per million person-years of observation may be less in building construction, renovation and maintenance workers, but this smaller risk is spread across a substantially larger workforce (i.e. there are many more carpenters/joiners, builder’s labourers, electricians, plumbers and other tradespeople in Western societies than the numbers of workers engaged in the mining, milling or production of high-density asbestos-containing materials such as asbestos-cement sheets and pipes or brake blocks).

5.177 According to EHC 203:

“It should be recognized that although the epidemiological studies of chrysotile-exposed workers have been primarily limited to the mining and milling, and manufacturing sector, there is evidence, based on the historic pattern of disease associated with exposure to mixed fibre types in western countries, that risks are likely to be greater among workers in construction and possibly other user industries.” [EHC 203, p 9].

“Past uncontrolled mixed exposure to chrysotile and amphiboles has caused considerable disease and mortality in Europe and North America. Moreover, historical experience to mixed fibre types in European countries has clearly indicated that a larger proportion of mesotheliomas occurs in the construction trades than in production. Far larger quantities of chrysotile than of other types of asbestos were used in most construction applications. Epidemiological studies that contribute to our understanding of the health effects of chrysotile conducted to date and reviewed in this monograph have been on populations mainly in the mining or manufacturing sectors and not in construction or other user industries. This should be borne in mind when considering potential risks associated with exposure to chrysotile.” [EHC 203, p 137].

“Few data on concentrations of fibres associated with the installation and use of chrysotile-containing products were available to the Task Group, although this is easily the most likely place for workers to be exposed.” [EHC 203, p 138].

“There is potential for widespread exposure of maintenance personnel to mixed asbestos fibre types due to the large quantities of friable asbestos materials still in place. In buildings where there are control plans, personal exposure of building maintenance personnel in the USA, expressed as 8-h time-weighted averages, was between 0.002 and 0.02 f/ml. These values are the same order of magnitude as exposures reported during telecommunication switch work (0.009 f/ml) and above-ceiling work (0.037 f/ml), although higher concentrations have been reported in utility space work (0.5 f/ml). Concentrations may be considerably higher where control plans have not been introduced. For example, in one case, short-term episodic concentrations ranged from 1.6 f/ml during sweeping to 15.5 f/ml during cleaning (dusting off) of library books in a building with a very friable chrysotile-containing surface formulation. Most other values, presented as 8-h timed-weighted averages, are about two order of magnitude less.” [EHC 203, p 139].
5.178  These points are also borne out by the 1999 Report for the Australian Mesothelioma Register [AMR 99], where the broad spread of prior occupations among mesothelioma victims is plain. For example, the number of mesotheliomas from the former Wittenoom blue asbestos industry (189 mesotheliomas related to a single exposure only; 25 additional mesotheliomas as a consequence of multiple exposures; total = 214) is less than the numbers of mesotheliomas as a consequence of asbestos exposure in different occupations (e.g. carpenters/joiners: 187 mesotheliomas from a single exposure; 33 additional mesotheliomas due to multiple exposures; total = 220; for builders/builders’ labourers the corresponding numbers are 150 + 27 = 177). In other words, mesotheliomas among the former Wittenoom cohort constitute a relatively small number (214) in comparison to the aggregate numbers of mesotheliomas from asbestos exposures in other occupations (2371 other asbestos-associated mesotheliomas; no exposure data for 717 cases, and no apparent exposure for 443; aggregate total = 3745).

5.179  NICNAS 99 makes the same point (p 59):

“Occupation/industry classification of the mesothelioma cases on the register are based on the Australian Bureau of Statistics ‘Industry and Occupation Codes’. The percentage of overall cases of mesothelioma (January 1986 to March 1995) according to exposure category are: repair and maintenance of asbestos material (13%), shipbuilding (3%), asbestos cement production (4%), railways (3%), power stations (3%), boilermaking (3%), mining (Wittenoom) (5%), wharf labour (2%), para-occupational, hobby, environmental (4%) carpentry (4%), building (6%) navy (3%), plumbing (2%) brake linings (manufacture/repair) (2%) and combinations of the above (multiple) (12%) (Leigh et al., 1997). Leigh (1994) reported that the pattern of exposure is shifting away from the older traditional industries towards product, domestic and environmental exposure. An analysis of 16 years data in 1996 by Yeung et al. (1997) showed more cases (on a number of cases basis) in more recent years in the asbestos user industries and from occupations such as plumbers, carpenters, machinists and car mechanics.”

5.180  Similar patterns of exposure — and resultant diseases (lung cancer; mesothelioma) — have been recorded in the United Kingdom (EHC 203, pp 123-124):

“Based on analyses of mortality of workers with mixed exposures to chrysotile and amphiboles in the United Kingdom, by far the greatest proportion of mesotheliomas occurs in users of asbestos-containing products, rather than those involved in their production. ...

1. Asbestos exposure caused approximately equal numbers of excess deaths from lung cancer (749 observed, 549 expected) and mesothelioma (183 deaths) within the occupations covered by the 1969 and 1984 Regulations ...

2. Only a few (5%) of British mesothelioma deaths were among workers in regulated occupations (Peto et al., 1995). The majority of deaths occurred in unregulated occupations in which asbestos-containing products are used, particularly in the construction industry. The risk was particularly high among electricians plumbers and carpenters as well as among building workers.”

5.181  As shown by the literature cited in this discussion, it is my perception that there is broad agreement among experts on these patterns of exposure.

Dr. Infante:

5.182  The relative exposure categorization of the six job situations mentioned in the question depends on the nature of controls being used in each situation. In general, exposures are more easily controlled in manufacturing and more difficult to control in construction, maintenance, repair, demolition and disposal activities. Today, exposures would be
more easily controlled in mining and milling because of awareness of the hazard and the clear identification of the operations as sources of asbestos exposure. Quite often workers involved in maintenance, repair and handyman type activities do not know whether asbestos is present or not. In the absence of such knowledge, workers usually do little, or nothing to protect themselves from exposures to asbestos in these situations. As a result, workers involved in these activities are most likely to be the most heavily exposed in the occupational setting today. These types of activities often result in asbestos being carried home on the workers’ clothing. A typical scenario that comes to mind is a situation whereby a worker is in a crawl space and encounters asbestos insulation. There is no active supervision in this situation and the asbestos most likely is not labelled. Thus, the worker cuts through the insulation to get to the area needing to be repaired without knowledge of the hazard and without having the appropriate personal protective equipment. In the latter scenario, even when workers do wear respirators, they are often dust masks, which do not provide a proper face seal and the filter medium is not adequate i.e., HEPA filters are not part of the filtration material on these masks. In the repair trades particularly, it is common practice for workers to use dust masks that do not provide HEPA filtration. As a result, the respirator is inadequate for filtering out the fibres of dimensions that are thought to lead to cancer and other asbestos related diseases. Furthermore, even in situations where the appropriate respirators may be worn, comprehensive respiratory fit-testing programmes may not be included as part of the industrial hygiene programme and as a result, the respirators leak because of the inability to achieve a proper face seal. In situations where workers may be drilling, sawing, crushing, or sanding asbestos cement products the only appropriate respirator may be a supplied air respirator, but it may not be used because it is too cumbersome for the job situation. In my opinion, scenarios (v) and (vi) are usually the most dangerous in current times because the workers are not aware of the presence of asbestos and they are more likely not to have received training and education about the hazards of asbestos exposure.

5.183 The risk of exposure should be considered not only by level of exposure, but also by the extent of the populations exposed to chrysotile asbestos. The large number of mesotheliomas associated with secondary and tertiary users of chrysotile asbestos (maintenance workers, electricians, bystanders, etc.) is a reflection of the large number of individuals in the population exposed in these situations. Thus, in terms of the risk of disease from chrysotile exposure, one must consider not only the intensity of exposure in the various work situations, but also the extent of the population exposed. One study (Begin et al., 1992) reports that 33 per cent of mesothelioma cases identified among maintenance workers, electricians, bystanders, etc. were the result of exposure for less than five years, and that the incidence of these occasionally exposed cases was increasing more rapidly than in the primary industries (mines and mills), or in the secondary industries (manufacturing, daily handling of asbestos).

Dr. Musk:

5.184 The term “risk of exposure” is taken to mean who is most likely to receive the most exposure and therefore be at the greatest risk of developing asbestos-related disease. This would depend on the nature of the industry in the locality and the type of asbestos being produced or used or otherwise encountered. Those workers likely to receive the most exposure would be those in industries where regulations are most permissive or compliance with them is poorest from absence of supervision or means of personal protection. It would also depend on the conditions of work such as indoor versus outdoor etc. Canada’s “argument” could be settled by monitoring of exposure! The “arguments” do not seem to be incompitable.

1(b) Should we consider that the risk to human health associated with the various uses of chrysotile throughout its life-cycle is a workplace issue or does this risk affect a larger part of the population?
Dr. de Klerk:

5.185 The risk of disease from chrysotile affects everyone. The risk of disease depends on intensity of exposure, the duration of exposure and the time since exposure. The population who do not work with asbestos will still come into contact with it, albeit at a much lower intensity, however this population is much larger and hence the burden of disease may be greater. There are numerous examples of asbestos-related disease arising in people living in the vicinity of asbestos works or living with asbestos workers.

Dr. Henderson:

5.186 From my perspective, this is overwhelmingly a workplace issue (e.g. construction workers). The risk of cancer for the larger general population from exposure to asbestos in place has been discussed in an earlier part of this report (see above section C.1.e)). Please see also my answer to the preceding question.

5.187 For example, asbestos-cement roofs are common in Germany where corrosion by acid rain represents a potential problem. Measurements carried out by Spurný et al. [221-224] on airborne asbestos fibre concentrations in the vicinity of such buildings consistently reveal levels in the order of 0.0002-0.0012 f/ml, in comparison to fibre concentration in other urban environments that range up to 0.1 f/ml (but generally = 0.001 f/ml).

5.188 Measurements have also been made on airborne fibre levels related to asbestos-cement roofing in schools in Western Australia [128], with only one asbestos fibre detected in each of two schools (air monitoring at 9 sites over 720 hours). Based on the findings, it was estimated that airborne fibre concentrations would be unlikely to exceed 0.002 f/ml and were likely to be < 0.0002 f/ml. These levels were considered to represent a negligible risk to health; the Western Australia Advisory Committee on Hazardous Substances that carried out this investigation considered that a greater risk to health would arise from: (i) unskilled attempts to clean up the asbestos-cement roofs before application of protective coating; and (ii) trauma to the workers — e.g. falling from or through the roofs.

5.189 It is my perception that there is little or no dispute among experts on this issue.

Dr. Infante:

5.190 In general, workers are at relatively greater risk of exposure to chrysotile and disease, particularly those involved in maintenance, modification, demolition, repair and disposal activities as compared to those exposed in non-occupational situations. A large number of people from the general population, however, also will be exposed to chrysotile and elevated risk of disease when they engage in home repairs that involve manipulating or disturbing asbestos-containing products. The latter individuals usually have little or no education about the hazards of asbestos, nor of the most appropriate means to handle it with the least amount of exposure.) These types of operations will also create some standby exposures (Ascoli et al. 1996). If appropriate controls are not used when handling asbestos insulation in buildings, the building can become contaminated and the occupants will become exposed. Therefore, the major problem with asbestos exposure is related to occupational situations though a much larger population is exposed beyond the occupational setting to relatively lower levels. Reports of cases of mesothelioma among non-occupationally exposed individuals document non-occupational exposures to asbestos causing disease. Family members of workers involved in the asbestos cement industry (Magnani et al. 1993) as well as children of miners and millers (McDonald and McDonald 1980) have been diagnosed with mesothelioma.

Dr. Musk:

5.191 It is my opinion that the risks resulting from exposure affects all exposed people and depends on the cumulative level of exposure. It is also my opinion that there is not an exposure threshold below which there is no risk. The risks to people not occupationally exposed to asbestos are likely to be much less than the risks to those with occupational exposure because the degree of exposure is likely to be less (though not necessarily always
so). However, while the individual risks may be much less the total burden of disease in the community may not be because it is likely that there are many more people experiencing these risks (albeit lower). For example the burden of disease in the residents of the town of Wittenoom, Western Australia has been significant albeit less than that of the workers. The WA Mesothelioma Registry contains subjects whose only exposure was from neighbourhood industries. Similar cases have been documented in the Quebec areas.

1.(c) Can chrysotile-cement products (for instance in buildings) release fibres, through weathering, corrosion or general degradation, thus presenting a possible risk to human health? Can you quantify this risk?

Dr. de Klerk:

5.192 There is good evidence that both wind and rain cause the release of fibres even from new asbestos cement sheeting. Other possibilities are fires and unwanted demolition. It is hard to quantify the risk which again depends on intensity and duration, but measurements have been made in the vicinity of such buildings which are detectable but low.

Dr. Henderson:

5.193 Please see the preceding answer, and section C.1.1(e). Quantitation of the risk is based on backward extrapolation according to the linear no-threshold model because there are no observational data on the dose-response effects from low-level exposure to chrysotile, and the estimates are, therefore, open to question and dispute, but the risks to health from very low-level environmental exposure appear to be minuscule or negligible.

Dr. Infante:

5.194 Yes, weathered and corroded asbestos cement products are capable of releasing chrysotile fibres into the environment, and most of the fibre is transported by rainwater though some will be released into the ambient air in low concentrations. One study indicates that chrysotile exposure in such circumstances will generally be less than 1,000 fibres longer than 5 microns per cubic meter of air. The fibres released were shown to have the same carcinogenic potency as “standard” chrysotile fibres (Spurny, 1989). Asbestos fibres also will be released into water from cement water pipes. I have not seen any estimates of risk from this type of asbestos exposure. Although the relative risk of disease is considerably less than that from occupational exposures, the population at risk is considerably larger.

Dr. Musk:

5.195 I understand that asbestos-cement products do release fibres as they weather. Release of fibres occurs from both old and new products. Asbestos fibres can also be released when asbestos-cement products are involved in fires. Quantitative estimates of the risks are theoretically possible as airborne concentrations can be measured and dose-response relationships are known.

1.(d) Can interventions on chrysotile-cement and other high-density chrysotile products release fibres, thus presenting a possible risk to the health of the individual making such interventions or to the public in general? Can you quantify this risk?

Dr. de Klerk:

5.196 It is during interventions such as drilling, sawing, sanding, moving in stacks, loading onto transport etc, that concentrations of fibres are greatest, both for the operators and bystanders. The concentrations associated with such operations have been extensively tabulated in the literature. Exposure response relationships can be used to estimate the risk for any combinations of intensity, duration and time after exposure, as shown in the Table below.
5.197 Lifetime risks (to age 85 years) of mesothelioma after exposure to chrysotile, assuming 0.1 f/ml for 10 years from age 20 with competing causes of death at 1992 Western Australian death rates.

<table>
<thead>
<tr>
<th>Assumptions</th>
<th>Expected cases per million lifetimes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Health Effects Institute equation</td>
<td>724</td>
</tr>
<tr>
<td>Wittenoom crocidolite equation 1/12th potency</td>
<td>210</td>
</tr>
<tr>
<td>Wittenoom crocidolite equation, 1/80th potency</td>
<td>32</td>
</tr>
<tr>
<td>Background risk (Peto study of “unexposed” Los Angeles population)</td>
<td>112</td>
</tr>
</tbody>
</table>

Dr. Henderson:

5.198 My answer to the first question is YES. Operations such as drilling or sawing asbestos-cement products release fibres and produce elevated airborne fibre concentrations. (i) asbestos-cement sheets can release respirable fibres in the absence of manipulation, even when new (up to 0.001 f/ml; for references, see de Klerk and Armstrong [135]); (ii) a 1938 report in New South Wales indicated that cutting asbestos-cement products with a power saw could generate 4-5 million particles/cubic foot (roughly equivalent to 12-15 f/ml); cutting with hand saws produced lower concentrations; (iii) as shown in the following Table 11, Sturm et al. [5, 7] reported measurable fibre concentrations from various operations on asbestos-containing materials, including asbestos-cement in the former East Germany, as measured by occupational inspectors.

**TABLE 11: ASBESTOS FIBRE CONCENTRATIONS AT WORKPLACES, WITHOUT SUCTION DEVICES, DETERMINED BY KONIMETRY (FROM UNPUBLISHED REPORTS PREPARED BY OCCUPATIONAL INSPECTORATES)**

<table>
<thead>
<tr>
<th>Type of Work</th>
<th>Fibre Concentration (f/ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Scratching and crushing of asbestos-cement</td>
<td>0.03 to 0.3</td>
</tr>
<tr>
<td>Abrasive cutting of asbestos-cement without dust removal by suction</td>
<td>0.3 to 10.0 approx.</td>
</tr>
<tr>
<td>Drilling asbestos-cement without dust removal by suction</td>
<td>0.5 to 3.4</td>
</tr>
<tr>
<td>Machining of brake linings</td>
<td>0.1 to 13.0</td>
</tr>
<tr>
<td>Replacement of gaskets</td>
<td>0.02 to 0.5</td>
</tr>
<tr>
<td>Punching of gaskets (rubber asbestos)</td>
<td>0.02 to 1.9</td>
</tr>
<tr>
<td>Use of asbestos gloves</td>
<td>0.02 to 0.6</td>
</tr>
<tr>
<td>Replacement of clearing layers</td>
<td>0.06 to 0.5</td>
</tr>
<tr>
<td>Use of talcum for powdering gloves</td>
<td>0.6 to 20.0</td>
</tr>
<tr>
<td>Level limit value (over a whole working day)</td>
<td>1.0</td>
</tr>
</tbody>
</table>

5.199 In 1993, Kumagai et al. [4] in Japan reported on dust levels generated by repair work on asbestos-cement pipes, including use of a high-speed disc cutter both inside holes dug in the ground to gain access to the pipes and outside the holes. The concentration of asbestos fibres > 5 µm in length ranged from 48-170 f/ml inside the hole (average = 92 f/ml) and ranged from 1.7-15 f/ml outside the hole. The Abstract for this paper follows:

“Asbestos cement pipes (ACPs) containing 15 to 20% chrysotile or crocidolite have been used for underground conduits. Even today 16.2% of all conduits in Japan are ACPs, though the production of ACPs was suspended in 1985. When such a conduit is accidentally damaged the workers belonging to the Waterworks Bureau of a local government cut off the damaged conduit using a high-speed disk cutter and replace it with a new conduit. This operation develops a cloud of dust and the workers involved run the risk of asbestos exposure. It was the
aim of the present study to estimate asbestos exposure levels among these workers. First, in the experiment, we established the typical working conditions and requested an experienced worker to cut an ACP using a high-speed disc cutter in a hole dug in the ground as he routinely does. The experiment was repeated three times. During a bout of each experiment, dust was sampled at several points both inside and outside the hole. Second, a self-administered questionnaire survey was conducted to obtain information from the workers regarding their working conditions in cutting ACPs. The subjects of the survey were 1,048 men belonging to conduit repair sections of the Waterworks Bureau of 119 local governments. The results obtained can be summarized as follows. (1) Each bout of cutting ACPs required about five minutes. The concentration of asbestos fibers longer than 5 microns with 3:1 aspect ratio ranged from 48 to 170 fibers/ml (92 fibers/ml on average) inside and 1.7 to 15 fibers/ml outside the hole. The concentration inside the hole exceeded the ceiling limit (10 fibers/ml recommended for asbestos by the Japanese Association of Industrial Health. A concentration of 92 fibers/ml is equivalent to 0.96 fibers/ml as 8-h time-weighted average. (2) The number of subjects with experience of cutting ACPs was 849 (81.0%). The average length of service in conduit repair section was 14.2 yr. Based on the information obtained from each subject regarding the average working days per yr for each decade from 1946, the cumulative days to date expended in cutting ACPs was estimated to average 235 d, that is, 17 d per yr. Only 18.1% of the subjects used a protective respiratory device."

5.200 EHC 203 also gives the following data (p 40):

"Weiner et al. (1994) reported concentrations in a South African workshop in which chrysotile asbestos-cement sheets were cut into components for insulation. The sheets were cut manually, sanded and subsequently assembled. Initial sampling showed personal sample mean concentration of 1.9 f/ml for assembling, 5.7 f/ml for sweeping, 8.6 f/ml for drilling and 27.5 f/ml for sanding. After improvements and clean-up of the work environment, the concentrations were 0.5-1.7 f/ml.

Nicholson (1978) reported concentrations of 0.33-1.47 f/ml in a room during and after sawing and hammering of an asbestos-cement panel."

5.201 It is my perception that there is no dispute among experts on this issue.

5.202 In relation to the second part of this question, apart from stating that there is a risk because of the generation of airborne asbestos fibres from interventions on asbestos-cement and other high-density asbestos products, it is not possible to quantify the risk in a way that would meet with universal agreement or a broad consensus, because few data are available for the risks for this type of operation on chrysotile-cement products: the risk would be related to cumulative exposure, which would vary according to the types of operation carried out, and their frequency. In addition, risk estimates would be dependent on extrapolation from the linear dose-response model that has been called into question by Canada. Therefore, I would expect disagreement among authorities on the magnitude of the risk.

5.203 Table 12 derived from NICNAS 99 gives risk estimates for lung cancer at airborne chrysotile concentrations of 0.1-1.0 f/ml, according to the National Occupational Health and Safety Commission in Australia (NOHSC) and two US occupational health and safety bodies (OSHA and NIOSH).

<table>
<thead>
<tr>
<th>Exposure (yearly average fibre/ml)</th>
<th>Excess risk (per 100,000 persons exposed)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>NOHSC</td>
</tr>
<tr>
<td>1</td>
<td>173</td>
</tr>
<tr>
<td>0.5</td>
<td>86</td>
</tr>
<tr>
<td>0.1</td>
<td>17</td>
</tr>
</tbody>
</table>

Excess risk = Risk coefficient x lifetime exposure (yrs) x average exposure level (f/ml) background risk.*

*A cumulative background risk for lung cancer in the male population was used in these calculations (i.e. 7200/100,000 assuming mixed smoking habits).
However, NICNAS 99 goes on to discuss uncertainties concerning these risk estimates:

“There are several other reasons why there is considerable uncertainty regarding these risk estimates, which include:

1. Past occupational exposures have generally involved exposure to a mixture of asbestos fibres. As it appears likely that different types of asbestos have different degrees of hazard, it is difficult to determine the risk attributable to chrysotile per se. In addition, commercial chrysotile often has low levels of tremolite contamination.

2. Fibre size, such as difference in fibre size between different chrysotile industries, probably influences the degree of hazard and/or potency.

3. There is a long latency between exposure to asbestos and development of lung cancer. Hence, it is not possible to state definitively what fibre type and level of exposure caused the disease. Consequently, risk estimates are related more to duration of employment rather than intensity of exposure.

4. A linear, non-threshold model may not be an appropriate model as there is some evidence suggesting that lung cancer due to chrysotile exposure may have a threshold for effect.

5. Past exposure estimates (both quantitative and qualitative) are subject to considerable error. For example, conversion of historical results in mpcf units to fibres/mL has inherent uncertainties.

6. There is a high background level of lung cancer in the general population due to smoking. Cases of lung cancer attributable to asbestos cannot be distinguished from those due to smoking. Attribution can only be assessed in terms of excess of lung cancers above a control population, hence the choice of control population is critical.

7. The identification of the disease is dependent on medical diagnosis, however autopsies are not always conducted.

The impact of some of these uncertainties can be accounted for to some extent. For example, it is considered that (1) and (2) are largely accounted for by basing risk estimates on epidemiological studies where exposure was only to chrysotile in the most relevant industry. For the remainder of the above uncertainties it is unclear what influence they have on the risk estimates and how they should be accounted for. For example, recently there has been some debate in the literature as to whether a threshold or non-threshold model should be used when predicting risk due to chrysotile exposure. Meldrum (1996) states that based on balance of toxicological evidence, the linear no-threshold model for chrysotile-induced lung cancer may not be appropriate. ... Epidemiological data alone are not able to clearly distinguish between the possibility of a threshold or a non-threshold model due to the relatively high background rate of lung cancer in the human population. There is at present no consensus with respect to a threshold level of exposure for chrysotile below which there is no risk of disease” [pp 70-71].

Table 13 gives an estimate of lifetime mesothelioma risk from exposure to low levels of chrysotile (1.0 f/ml and 0.1 f/ml), based on dose-response data for the Wittenoom cohort, and assuming lower potencies for chrysotile than crocidolite (i.e. 1/12⁰, 1/30⁰ and 1/80⁰).
TABLE 13: ESTIMATES OF LIKELY MESOTHELIOMAS RELATED TO CHRYSOTILE INHALATION AT AIRBORNE CONCENTRATIONS OF 1.0 AND 0.1 f/ml, ASSUMING A CARCINOGENIC POTENCY 1/12TH, 1/30TH OR 1/80TH THAT OF CROCIDOLITE

<table>
<thead>
<tr>
<th>Airborne fibre concentration; Potency</th>
<th>Numbers to age 85 Mesotheliomas/million persons</th>
<th>Duration of exposure* (years)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>1 yr</td>
</tr>
<tr>
<td>1.0 f/ml - 1/12\textsuperscript{th}</td>
<td></td>
<td>252</td>
</tr>
<tr>
<td>1.0 f/ml - 1/30\textsuperscript{th}</td>
<td></td>
<td>113</td>
</tr>
<tr>
<td>0.1 f/ml - 1/12\textsuperscript{th}</td>
<td></td>
<td>28</td>
</tr>
<tr>
<td>0.1 f/ml - 1/30\textsuperscript{th}</td>
<td></td>
<td>11</td>
</tr>
<tr>
<td>0.1 f/ml - 1/80\textsuperscript{th}</td>
<td></td>
<td>4</td>
</tr>
</tbody>
</table>

*Starting at age 20 yrs.

Dr. Infante:

5.206 Interventions on chrysotile cement products can result in extremely high atmospheric fibre concentrations (Rodelsperger et al., 1980) and studies of roofers have demonstrated asbestosis from such exposures (Stauder et al., 1982). A study of workers involved in inside finishing work with concrete asbestos containing 30 per cent chrysotile asbestos has also shown that airflow obstruction among workers can be caused by such exposure (Harless et al. 1978). Open-air cutting such as that involved in roofing and inside finishing operations also will result in exposure to other workers not directly involved in the manipulation of the asbestos, e.g., standby exposure. Such manipulation of asbestos cement will also expose the general population.

5.207 A study of 404 roofers with long term exposure to cement dust indicated that 14 per cent had significantly increased small irregular opacities with profusion in 13 per cent (Stauder et al., 1982). The prevalence of these abnormalities was significantly greater than that observed in the control group. The study by Harless et al. (1978) indicated that approximately 50 per cent of workers exposed for about six months to dust from asbestos cement developed airflow obstruction. The risk of developing lung pathology per unit of asbestos fibre exposure cannot be determined from these studies because of lack of exposure data. The studies do indicate, however, that uncontrolled manipulation of chrysotile cement products can result in a high rate of lung pathology. Lung function can be adversely affected as a result of exposure over a very short period of time.

5.208 A large number of reports indicate mesothelioma related to car mechanics involved in brake repair. General population exposures from such work would be minimal except for those situations where individuals would engage in their own brake repair work.

Dr. Musk:

5.209 Interventions on asbestos-cement products can release fibres therefore a risk of disease exists as in 1(c).

1.(c) Can occasional interventions on high-density chrysotile products, either in occupational circumstances (such as electricians, plumbers, repairers, insulation workers, etc.) or by private individuals (“handyman” type) release fibres, thus presenting a possible risk to the individual making such interventions or to the public in general? Can you quantify this risk?

Dr. de Klerk:

5.210 Yes of course, see (c) and (d).
Dr. Henderson:

5.211 To answer the second question first, I am unable to quantify potential risk, because there are no systematic observational data available for this type of work, to the best of my knowledge (but please see Tables 12 and 13 above in my answer to Question 1(d)).

5.212 The first part of the question has been covered in the preceding answer, with the observation that occasional interventions of this type would predictably produce low cumulative exposures, with a lower risk, for the reasons discussed earlier. Please also refer to AMR 99, for data on mesotheliomas among electricians, carpenters, plumbers, insulation workers and so forth (it is acknowledged that most if not all these mesotheliomas are a consequence of exposure to asbestos-containing materials that included a mixture of asbestos types, including chrysotile and one or more of the amphiboles); in drawing attention to AMR 99, my purpose is simply to use mesothelioma rates as a reflection of past exposures and hence evidence that airborne fibre concentrations were produced by these types of operation, without regard to the fibre types. My own cases of mesothelioma also include a number of individuals whose only exposure to asbestos took the form of maintenance work and renovations carried out on the patient’s home, where there were asbestos-cement building materials. Again, in drawing attention to this type of background, it is not my intention to address fibre type, but simply to indicate that mesothelioma as an outcome of this type of exposure indicates that elevated concentrations of respirable airborne fibres were produced.

5.213 EHC 203 gives the following account (pp 122-123):

“Although the odds ratio for lung cancer associated with exposure to “asbestos” has been estimated in many case-control studies, the studies have not been in general able to distinguish between chrysotile and amphibole exposure, and are therefore less informative for the present evaluation ... In a multisite case-control study from Montreal, Canada, however, exposures to chrysotile and to amphiboles were separated, although exposure to amphiboles was not controlled for in the analysis for exposure to chrysotile (Siemiatycki, 1991). In this study, the occupational history of male cases (age 35-70) of cancer at 20 sites and of 353 population controls was evaluated by a team of industrial hygienists and chemists to assess exposure to 293 agents. Overall, the lifetime prevalence of exposure to chrysotile was 17% and that of exposure to amphiboles 6%. The main occupations involving exposure to chrysotile that were considered were motor vehicle mechanics, welders and flame cutters, and stationary engineers. When lung cancer cases (N = 857) were compared with cases of all other types of cancer, the odds ratio (OR) of any exposure to chrysotile was 1.2 (90% CI = 1.0-1.5; 175 exposed cases), and that of 10 or more years of exposure with at least 5 years of latency (‘substantial exposure’) was 1.9 (90% CI = 1.1-3.2; 30 exposed cases). Corresponding ORs of exposure to amphiboles were 1.0 and 0.9. The OR of exposure to chrysotile was higher for oat cell carcinoma than for other types of lung cancer. Twelve cases of mesothelioma were included in this study. The OR of any exposure to chrysotile was 4.4 (90% CI = 1.6-11.9; 5 exposed cases) and that of substantial exposure was 14.6 (90% CI = 3.5-60.5; 2 cases). Corresponding ORs of exposure to amphiboles were 7.2 (90% CI = 2.6-19.9; 4 cases) and 51.6 (90% CI = 12.3-99.9; 2 cases).”

5.214 Please see also Tables 5, 9, 10, 11, 12, 13, in EHC 203.

5.215 It is my perception that there is no dispute among experts that such interventions release fibres; disagreement is likely over the magnitude of the risk.

Dr. Infante:

5.216 The worker making the intervention would be the most highly exposed and presented with the greatest risk of asbestos related diseases. The extent of exposure to the
worker as well as to those in the surrounding area would depend on the nature of the
intervention, e.g., the circumstances under which the chrysotile asbestos product is
manipulated in terms of work practices, the controls, or lack of controls in place and the type
of personal protective equipment provided to the worker. While data on fibre exposure
levels in these situations are sparse, data on mesothelioma indicates association with workers
who have jobs that result in occasional interventions to asbestos products. Because these
exposures are not routine and the hazard often goes unrecognized, these operations are
unlikely to be well controlled, i.e., they are not anticipated so proper training and educa-
tion about these types of exposures is often lacking.

5.217 It is difficult to quantify this risk because atmospheric measurements are usu-
ally not made during these interventions. However, the identification of cases of mesothel-
ioma associated with these interventions in the literature indicate that they are perhaps
the most detrimental to human health. Mesothelioma has been identified from these expo-
sure situations because it is a marker cancer related to asbestos exposure. What goes uniden-
tified and unmeasured from these situations is the much larger burden of disease and death
from pneumoconiosis and lung cancer. The attributable burden from these latter diseases
will be much greater than that from mesothelioma, but they are not usually recognized
because lung cancer has a high background rate in the general population and asbestosis
may be diagnosed as another type of pneumoconiosis unrelated to asbestos exposure.
Mesothelioma has also been documented among the wives of construction workers, indi-
cating that the family member portion of the public in general is also at risk. These latter
cases of mesothelioma are most likely the result of carry-home exposure from contami-
nated clothing.

5.218 If one considers that the handyman types of exposures are considered as expo-
sures to the general public, then this segment of the general population would also be at an
elevated risk of developing asbestos related diseases. Exposures to family members that
might result from interventions by home owners would depend on the nature and location
of the removal or manipulation of the asbestos. The general public is also exposed through
manipulation of asbestos in residential buildings that is not carried out with appropriate
controls in place and by carry home exposures from contaminated work clothing.

Dr. Musk:

5.219 Occasional interventions on asbestos-cement products by anyone can release
airborne fibres, therefore there is some risk as in Question 1(c).

1.(f) Are chrysotile fibres from chrysotile-cement dust released during interventions
(cutting, sawing, etc.) on chrysotile-cement products as dangerous as pure chrysotile fi-
bres? Is the physical and chemical composition of asbestos-cement dust different from
pure asbestos dust?

Dr. de Klerk:

5.220 Risk from fibres depends on size, shape and durability (and their quantity).
Asbestos cement contains about 10-20 per cent asbestos, so that the dust concentration is
going to be less than if the sheets were pure asbestos. However, cement does not form
tiles, so that any airborne fibre measurements made would only reflect the asbestos con-
centration in the air.

Dr. Henderson:

5.221 To answer the second part first, the physico-chemical composition of asbestos-
cement dust does differ from pure asbestos dust, because the asbestos in asbestos-cement
products is diluted by the cement (asbestos = 10-15 per cent by weight); this being so, one
expects the asbestos fibres to be diluted by cement dust, in comparison to equivalent op-
erations on pure asbestos materials.
5.222 To return to the first part of this question: the claim may be made that chrysotile fibres released from asbestos-cement products by high-speed cutting are altered physically or chemically, with a predominance of short-length fibres not implicated in carcinogenesis. For example, in Canada’s first oral submission, the following comments are made:

“The European Community has also advanced the thesis that the ban is necessary because France has no control over trade persons or the ‘handyman’ who will cut into chrysotile-cement and, in so doing, free some of the chrysotile that was locked into it. Canada is puzzled by France’s assertion that la République Française is unable to regulate its handymen. In any event, there are three technical reasons why France’s concern is misplaced.

First, this thesis is based on the misconception that cutting high-density locked-in non-friable asbestos-containing materials releases substantial amounts of chrysotile. In fact, even if improper tools such as high-speed saws are used to cut chrysotile-cement, the dust released from such an operation contains only a very small amount of pure, respirable-size chrysotile fibres, if any at all.

Second, science tells us that most of the chrysotile fibres released during high-speed cutting have been chemically altered: the resulting entity is chemically and structurally different, and has a biological potential to induce harmful effects, which is different, and less than amphiboles. Similarly, the dust that results from abrading chrysotile-containing resin or plastic reinforced products contains very small amounts of chrysotile fibres. The same is also true of the dust that comes from wear and abrasion of friction materials: analysis of brake shoes shows that almost all of the chrysotile fraction of the finished product is found to be transformed into a totally different, biologically-inactive material called forsterite ....”

5.223 The first paragraph of the Canadian statement is dealt with in later discussion in this report (my answers to Question 5). For the second and third paragraphs of this statement, one can state that in other situations, only a small fraction of airborne asbestos fibres are of respirable size: as one example, about 0.67 per cent of airborne asbestos fibres within indoor air of buildings were longer than 5 µm in length. However, in the Japanese study reported by Kumagai et al. [4] on cutting asbestos-cement pipes with a high-speed disc cutter, where airborne fibre concentrations within the hole used to gain access to the pipes averaged 92 fibres/ml (range 48-170 fibres/ml), the study dealt with fibres longer than 5 µm (i.e. dimensions in the range for which carcinogenicity has been reported). Please see also Table 11 in EHC 203 where Rödelsperger et al. recorded airborne fibre concentrations of 4-5 f/ml and 5-10 f/ml for fibres longer than 5 µm, from blowing off and grinding brake blocks, including truck brakes. (Please see also Table 11 and my answer to Question 2.)

5.224 Clearly, there is disagreement between the parties to the dispute and their respective experts over the issue of whether chrysotile fibres released from high-density products are dangerous. For the reasons outlined above and in later discussion, it is my perception that at least a small proportion of the fibres has dimensions that are associated with carcinogenicity.

Dr. Infante:

5.225 As long as the interventions result in the release of chrysotile fibres, the exposures should be considered as dangerous as pure chrysotile fibres because respirable-sized asbestos fibres will be released. The study by Spurny (1989) indicates that the fibres released from weathered and corroded chrysotile asbestos cement products have the same carcinogenic potency as “standard” chrysotile fibres. Although fibres released by weathering may be somewhat different from fibres released from cutting or drilling asbestos cement products, the former fibres show a potency similar to that of pure chrysotile fibres.
Moreover, because of the potential for cleavage during interventions on asbestos cement products, the dust resulting from cutting, drilling, etc. on asbestos cement may actually contain a greater portion of the asbestos fibres being thinner and more respirable than those that were initially mixed into the cement during the manufacturing process. Therefore, the fibres released from cement during interventions should be considered at least as dangerous as “pure chrysotile fibres.” I can find no data to support an opinion that fibres released from interventions on chrysotile asbestos cement products would be less carcinogenic, or less dangerous. Further, asbestos related pathology has resulted from such situations.

5.226 The asbestos cement dust would be somewhat different in physical and chemical composition from pure chrysotile asbestos because the cement dust would contain respirable asbestos fibres, crystalline silica plus other substances added to the cement.

Dr. Musk:

5.227 It is my opinion that in general airborne fibres released from asbestos-cement products pose a risk. This may differ from other sources of chrysotile depending on the characteristics of the fibres. The area of fibre characteristics and their relationship to different sources is not within my area of expertise.

1.(g) What is the risk to human health associated with demolition and removal of high-density chrysotile products, such as chrysotile-cement products? Can you quantify this risk?

Dr. de Klerk:

5.228 See my answers to Questions 1(c) and (d).

Dr. Henderson:

5.229 I am not aware of any studies that have specifically focussed on either of these situations: therefore, no firm data are available, but one would expect the biohazards to be related to cumulative doses of respirable fibres (i.e. airborne fibre concentrations and the frequency of exposure from these types of work). This being so, one would expect the risks to be equivalent to other operations of like frequency that generated similar airborne fibre levels (Tables 12 and 13).

Dr. Infante:

5.230 Exposure to high density chrysotile products through demolition carries with it the potential risk of lung cancer, asbestosis and mesothelioma. Testimony presented at the OSHA hearings related to its Final Asbestos Standard that was promulgated in 1994 indicated that removal of intact chrysotile asbestos “transite” panels that were held in place by screws can result in airborne fibre concentrations that exceeded 1 f/cc. In this situation, the exposed surfaces were wet prior to removal and the operation was done within a negative pressure enclosure. Many transite panels used in interior wall construction consist of rough inner surfaces from which asbestos fibre is readily released into the air. Other testimony (OSHA, 1994) presented evidence that transite panels can be removed in a manner that results in exposure well below 0.1 f/cc when appropriate work practices are followed. Because of concern for the potential release of asbestos fibres into the air from such demolition, the OSHA standard requires that a “competent person” supervise such activities, e.g., make an assessment and determine that the type of controls being used are appropriate for the removal situation and that the required work practices are being followed. Therefore, the extent of the risk during demolition of chrysotile cement products depends upon the compliance with mandated requirements. (See my answer to Question 5(c) regarding compliance with procedures to reduce risk of disease from asbestos exposure.)
Dr. Musk:

5.231 In so much as demolition activities may result in airborne fibres there is a risk (as above).

1.(h) What is the risk to human health associated with high-density chrysotile wastes, such as chrysotile-cement waste? Can you quantify this risk?

Dr. de Klerk:

5.232 This depends on how the waste is treated and stored, depending of course on the chance of any fibres becoming airborne and hence respirable. Otherwise, see my answers to Questions 1(c) and (d).

Dr. Henderson:

5.233 See my response to Question 1(g).

Dr. Infante:

5.234 I have not researched this issue, but I am inclined to believe that there would not be much potential for fibre exposure from the handling of such waste unless a person at a waste site was hauling asbestos cement and not aware of the product he/she was moving.

Dr. Musk:

5.235 The risk posed by waste products will also be dependent on the chances of fibres becoming airborne as above.

1.(i) Can high-density chrysotile products wastes, such as chrysotile-cement wastes, be dealt with so as to eliminate risks to human health?

Dr. de Klerk:

5.236 They can, by following approved methods for disposal which ensure that fibres are sealed from airborne release. There is of course the chance that subsequent work (for example, waste removal) may disturb the waste and release fibres.

Dr. Henderson:

5.237 In theory, YES — once the asbestos-cement or other high-density product has been removed from its in-place location (though few data are available on exposure levels produced by the actual removal). For example, in Australia, imported chrysotile is delivered to production facilities in sealed plastic bags, so that the same procedure for bagging or encapsulation of high-density wastes should also be applicable, and should prevent release of asbestos fibres once the encapsulation or bagging exercise is complete, unless the bags are ruptured for one reason or another.

5.238 According to NICNAS 99 (p 74), in Australia:

“Waste chrysotile, the polyethylene bags in which it is supplied, and chrysotile containing materials from the manufacturing process, are disposed to landfill by licensed disposal contractors. As chrysotile fibres are unlikely to be mobile in the soil or water table, landfill is not inappropriate from a public health perspective.”

Dr. Infante:

5.239 I have not researched this issue.
Dr. Musk:

5.240 These risks may be eliminated if the fibres could be successfully sealed so that they cannot become airborne.

Question 2:

What is the risk to human health associated with other current applications of chrysotile asbestos (in particular, friction materials and textiles)? In occupational circumstances? In non-occupational circumstances?

Dr. de Klerk:

5.241 While the industries themselves may be well regulated, controlled and compliant with standards, the major problem again could occur in “downstream” users: boiler-makers, plumbers, brake mechanics etc. Fibres released from friction products have a higher proportion of shorter fibres than those from textiles, which release the highest proportion of longer fibres.

Dr. Henderson:

(i) Friction Products (e.g. brake linings)

5.242 Automotive mechanics and garage workers constitute a large population of workers potentially exposed to chrysotile derived from brake linings. For example, brake blocks and linings used in Australia have contained only Canadian chrysotile for many years, and the materials are either imported as pre-formed brake blocks and linings, or chrysotile is imported into Australia for subsequent manufacture of these products. It has been estimated that this group of mechanics amounts to at least 900,000 workers in the US, and the figure may be even higher if one adds in all those who have ever worked in the automotive repair industry but then moved into other employments, and those who have retired.

5.243 For Australia, the number of persons employed as mechanics in 1991 amounted to 85,155 (84,293 males); for 1996, the corresponding figures are 83,647 (82,827 males), out of a total population of 16,852,256 in 1991 (8,363,677 males); for 1996, the total population was 17,892,423 (8,849,224 males). These figures for Australia include all mechanics, including automotive, brake and engine mechanics, together with supervisors and apprentices; the figures for 1996 also include mechanics’ assistants (not included for the 1991 figures).

Taking into account the fact that the Australian population is less than 1/10th that of the United States population, these statistics appear to be roughly comparable.

5.244 The literature contains anecdotal reports of malignant mesothelioma among automotive and brake mechanics. However, the question that arises is whether these anecdotal reports are explicable as the chance occurrence of spontaneous or background mesotheliomas among a large population of mechanics, or whether this group of workers has sustained other significant exposures to asbestos, including one or more amphiboles. In other words, the question is whether there is a general increase in the incidence of mesothelioma among automotive and brake mechanics with no other exposures to asbestos.

5.245 Brake repair workers are potentially exposed to asbestos during a number of procedures, which include removal of dust from bakes by air hoses, and a variety of other manipulations that include bevelling, grinding and drilling. Clearance of dust from brakes by use of an air hose can create a cloud of visible dust, and airborne fibre concentrations of 2.0 to 29.4 f/ml have been recorded in the immediate vicinity [225, 226]. Please see also Table 11 in EHC 203 (pp 42-43).

5.246 In North America, chrysotile has been used almost exclusively in brake linings since the 1940s; chrysotile is also the type of asbestos used in brake linings in Europe (and
also Australia). As I mentioned previously, commercial chrysotile (e.g. Canadian chrysotile) contains on average small quantities of contaminant amphiboles in the form of tremolite (usually = 1%).

5.247 However, the significance of this type of potential exposure among brake repair workers is complicated by a number of factors:

- During moderate braking of automobiles, temperatures as high as = 500°C can be reached within the brakes, and at this temperature a portion of the chrysotile undergoes dehydroxylation and recrystallization to form the mineral forsterite, which is not implicated in mesothelioma induction.

  “Heating of chrysotile at 700°C for an hour converts it to an amorphous, anhydrous magnesium silicate material … Intensive dry grinding also destroys the structure of chrysotile. Analysis of wear debris from brake linings made with asbestos has shown that virtually all of the chrysotile fibre is converted to amorphous material, in association with the mineral forsterite (a recrystallization product). The conversion is explained by localized temperatures above 1000°C at the point of contact between the brake lining and the drum” … [EHC 203, p 14].

- Most chrysotile fibres released from brakes comprise short-length fibres < 0.4 µm in length (> 80 per cent of all chrysotile fibres from brakes). However, some fibres > 5 µm in length and even > 10 µm in length appear to survive (e.g. please see Table 11 in EHC 203). The short-length fibres appear to have only questionable or limited carcinogenicity, and this property is thought to reside primarily in fibres > 5 µm in length. In addition, limited fibre burden studies on brake repair workers have shown a low pulmonary asbestos content.

  “The fibres found in the brake wear debris are predominantly (99%) less than 0.4 µm in length … Rödelsperger et al. (1986) found less than 1% of fibres longer than 5 µm.” [EHC 203, p 14]

- One also needs to remember that assessment of mesothelioma risk among brake repair workers can be confounded by other occupational exposures to asbestos [227].

5.248 In a review of changing risk groups for malignant mesothelioma, Huncharek [228] gives the follow account for brake mechanics:

  “A major problem with epidemiologic studies of this workforce is the difficulty in tracing a large, non-unionized group of workers. Estimation of disease risk has been impeded by lack of quantitative data on exposure levels among individuals with long-term exposure. …

In 1976, investigators at the Mount Sinai School of Medicine studied asbestos exposure among brake repair workers in New York city. Both clinical examinations and fibre counts produced by various operations in brake maintenance workers were analyzed. Samples taken at a distance of 3-5 feet from brake drums during periods of blowing dust showed fibre concentrations of 6.6 to 29.4 fibres/ml, with a mean of 15.9 fibres/ml. In addition, ten samples of brake drum dust were analyzed by phase contrast, optical microscopic examination and transmission electron microscopic examination to determine the percentage of short
fibres (i.e., 25-500 angstroms x 760 to 3750 angstroms\(^\text{198}\)). Eighty-three percent of all chrysotile fibres were in this category, and almost 20% of the total mass of 10 samples was chrysotile (determined by electron diffraction). ‘Throughout the examination by electron microscopy, attention was given to the morphology of the fibres. A majority of fibres showed little alteration in the typical chrysotile fibre’.

In an additional report from Mount Sinai, Rohl et al. analyzed residual dust recovered from brake linings and made direct measurements of the free asbestos fiber content of ‘workroom air’ in areas in which brake lining maintenance and brake shoe installation occurred. Airborne asbestos dust concentrations were similar to those cited by Lorimer et al. (i.e., mean airborne fiber concentrations during compressed air blowing of brake drums ranged from 2.6 fibers/ml at a distance of 10-20 feet to 16.0 fibers/ml at 3-5 feet). Samples of brake drum dust showed that the proportion of chrysotile in this material averaged 3% to 6% (both as free fibre and as particulates in pulverized binder).

Regarding the health effects resulting from the exposures described, Langer and McCaughey published ... a case of mesothelioma in a brake repair worker ... a 55-year-old man who had worked in the used car, tire and car repair business since the age of 19 years. He reported routinely servicing automobiles, including the replacement of brake linings. No other source of asbestos exposure was found.

Analysis of lung tissue showed the presence of chrysotile fibres (no amphibole was found) confirmed by electron diffraction techniques. Ten percent of the fibers found were longer than 10 microns.\(^{199}\) The authors point out that ‘Controversy over the potential of chrysotile to cause mesothelioma has continued despite evidence from asbestos textile fabricators, thought to have used only chrysotile, from workers making brake pads, from chrysotile miners and millers, and from animal studies’. They also state ‘The risk of malignant asbestos disease among these workers seems to be low but mortality data have yet to be thoroughly evaluated.’

The most recent report of mesothelioma in a brake mechanic reviews a pleural mesothelioma occurring in a 47-year-old male automobile mechanic whose only known exposure to asbestos was from clutch and brake repair work during an 11-year period. ...

Another case of mesothelioma in a brake repair worker was recently published. In this report, a 56-year-old male elevator mechanic ... reported working as an elevator mechanic for 30 years. He reported exposure primarily from elevator brake linings that he routinely cut, fitted, and removed during elevator installation and maintenance.

Several recent studies from Scandinavia on this topic also deserve mention. Hansen, from the Institute of Community Medicine in Denmark, completed a historical cohort study examining the mortality of car mechanics from ischaemic heart disease and malignant neoplasms. The study cohort was identified using records of a nation-wide census carried out in Denmark in November 1970. Comparison was made with another cohort of skilled male workers who were not exposed to asbestos or ‘petrochemical substances’. Of 583 observed deaths, one case of pleural mesothelioma was found.

Likewise, Jarvholm and Brisman, in a 1998 report, used the Swedish death register and the census of 1960 to study the occurrence of asbestos-associated tumors in car mechanics. One hundred and eighty-seven deaths attributable to cancer were observed, whereas 154 were expected. Thirty-nine were caused by lung
cancer, whereas only 23 were expected. Again, one death from pleural mesothelioma was found. ... 

It has been estimated that 20,000 deaths from asbestos-related cancer will occur during the next 40 years among automotive maintenance workers in the United States. With the many difficulties faced by epidemiologists studying this workforce, it is unclear how accurate this estimate will prove to be. Clearly, what is needed is better information on duration and intensity of exposure to respirable asbestos fibers in this occupational group. Additional study is needed to accurately determine the incidence of mesothelioma among members of this workforce.” [pp 2704-2705].

5.249 The situation is further complicated by other reports on garage mechanics and workers involved in friction products manufacture [229, 230]. These studies have been reviewed briefly by Wong [231]: three found no increase in the RR for mesothelioma among garage mechanics (RR = 0.9, 0.65 and 1.0 respectively).

5.250 An analysis of > 13,000 workers at a UK friction products factory showed no detectable excess mortality due to lung cancer or other cancers; 13 mesotheliomas were found but 11 had known exposure to crocidolite [229, 232].

5.251 McDonald et al. [230] identified excess mortality from lung cancer among friction products workers, but there were no mesotheliomas:

“A study by McDonald et al. (1984) investigated mortality due to lung cancer, mesothelioma and asbestosis in three US factories manufacturing friction products and packings. The cohort comprised 3641 men employed between 1938-1958. During the 1930s exposures for most processes were 1-5 mpcf (millions of particles per cubic foot) and > 10 mpcf during dry mould mixing. By the 1960s most exposures were < 0.5 mpcf. A significant excess of deaths (reference was to mortality rates for Connecticut) due to respiratory cancer was observed however this was not related to duration of employment. No cases of mesothelioma were reported. There was limited evidence of an increase in risk of lung cancer with increasing exposure. However the SMR for lung cancer was noted in workers with less than one year of service.

A study by Finkelstein, (1989) investigated mortality rates among 1657 employees at two Ontario factories manufacturing chrysotile friction materials. The study population consisted of workers employed for at least 12 months after 1 January 1950. The study showed a significant increase in mortality from laryngeal cancer and lung cancer. No increase in mortality was noted from gastrointestinal cancer or from non-malignant respiratory disease. One or two deaths may have been due to pleural mesothelioma. Case-control analysis demonstrated a lack of association between the risk of death from laryngeal or lung cancer and the duration of employment or employment in departments where chrysotile had been used. The author also noted that cigarette smoking is a risk factor for laryngeal cancer and lung cancer, and therefore, increased risk may be in part attributable to differences in smoking habits.” [NICNAS 99, p 65].

5.252 Similarly, Woitowitz and Rödelsperger [227, 233] found that:

“There is no evidence that car mechanics are exposed to an increased risk of mesothelioma even if they do brake repairs, but asbestos exposure in other employment is an important confounding factor, so that if there is a mesothelioma risk for car mechanics but it was small, it would not be detectable.”

5.253 Nonetheless, the 1999 Report for the Australian Mesothelioma Register299 (AMR 99) records 58 mesotheliomas among brake mechanics with no other exposures to asbestos, during the almost 13-year period between 01 January 1986 and 31 October 1999 (total
cases with a stated history of asbestos exposure = 2585). Mechanics who frequently or consistently work on brake linings and brake blocks represent only a sub-fraction of the total workforce of mechanics in Australia. If one takes the 1996 census figure of 82,827 for male mechanics, this amounts to 58 mesotheliomas in 1,062,946 person-years (= 54.6 mesotheliomas per million person-years). If one rounds off the workforce to 100,000 male mechanics, the figure becomes 45 mesotheliomas per million person-years. If one then doubles the workforce population to take into account retirees and other workers who moved on to other occupations (although a figure of 200,000 is almost certainly an overestimate because it would include all mechanics, whereas brake mechanics constitute a smaller sub-class), the mesothelioma rate becomes 22.6 per million person-years — well under the rate of 337 mesotheliomas per million person-years for the Quebec chrysotile miners and millers but still substantially above the upper limit of the estimated background rate of 1-2 mesotheliomas per million person-years (about 10-fold). One might suspect that mesotheliomas in brake mechanics will cluster in those involved in the grinding, bevelling and other operations on new brake blocks and brake linings (i.e. brake materials unaltered by heat).

Using an earlier set of data for Australia, NICNAS 99 came to a similar conclusion:

"Out of 2119 mesothelioma cases registered (with a response to history) for the period 1986-1995, 46 cases were listed for the category 'brake lining - manufacture/repair', 40 of which were recorded in car mechanics, of which 37 were exposed to asbestos in this occupation only ... Overall the numbers indicate a slight increase of around 1-2 cases per year, which is roughly proportional to the growth rate of all mesothelioma cases in Australia." ... [p 66].

It is apparent that these considerations apply to occupational circumstances.

Evidence indicates that the general population is exposed to only very low levels of asbestos derived from the braking of passing automobiles, and that most of these fibres represent short-length fibres and heat-altered chrysotile. NICNAS 99 has this to say on the subject:

"It is claimed that the amount of asbestos found in the dust arising from braking is rarely more than 1% of the wear product (Asbestos Information Committee, 1975). It is not known what quantity of chrysotile is imported in brake linings and other friction materials, but ABS [Australian Bureau of Statistics] data indicates in excess of 750,000 articles (brake linings, pads and clutch facings) being imported in 1997 containing asbestos and therefore possibly containing chrysotile. Assuming each unit weighs 200 g and contains 50% chrysotile, this equates to around 150 tonnes of chrysotile per annum. Assuming a further 1000 tonnes of chrysotile present in friction products manufactured in Australia, it is estimated that (assuming a worst case scenario of 1% release per annum, i.e. all products are completely worn in one year), around 11.5 tonnes of chrysotile will be released per annum countrywide or 32 kg per day spread all around the country. It is acknowledged that this figure may be an overestimate, as studies have shown that some of the chrysotile is degraded to magnesium silicates and forsterite ... In addition, some of the debris will be retained in the brake system and removed and disposed of under controlled conditions." [p 78].

Table 14 represents a reproduction of Table 7 in EHC 203 for exposure estimates in the South Carolina chrysotile textile plant (1930-1975) before and after controls on exposure levels. As can be seen from this Table, the application of controls on exposure levels produced a significant reduction of exposure, and currently available control technology allows even lower levels to be attained (EHC 203).
EHC 203 refers to a study in Japan which reported a geometric mean concentration of 0.1-0.2 f/ml in the period 1984-1986, for asbestos spinning. From published studies, it seems clear that asbestos textile workers are at greater risk for asbestosis (historically) and lung cancer than mesothelioma. EHC 203 also gives the following comments:

**TABLE 14: EXPOSURE ESTIMATES IN A CHRYSOTILE TEXTILE PLANT (1930-1975)**

<table>
<thead>
<tr>
<th>Operation</th>
<th>Without controls</th>
<th>With controls</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fibre preparation</td>
<td>26.2-78.0</td>
<td>5.8-17.2</td>
</tr>
<tr>
<td>Carding</td>
<td>10.8-22.1</td>
<td>4.3-9.0</td>
</tr>
<tr>
<td>Spinning</td>
<td>4.8-8.2</td>
<td>4.8-6.7</td>
</tr>
<tr>
<td>Twisting</td>
<td>24.6-376.0</td>
<td>5.4-7.9</td>
</tr>
<tr>
<td>Winding</td>
<td>4.1-20.9</td>
<td>4.1-8.4</td>
</tr>
<tr>
<td>Weaving</td>
<td>5.3-30.6</td>
<td>1.4-8.2</td>
</tr>
</tbody>
</table>

From Dement et al. (1983)

"Studies that correlate disease prevalence or symptoms with cumulative exposure can under-estimate disease risk due to progression of disease after employment ceases. Although workers were exposed to both chrysotile and crocidolite (the latter being approximate 5% of all asbestos used), results for 379 men employed at least 10 years in the Rochdale asbestos textile plant are informative... Exposure estimated from work histories range from an average of 2.9 to 14.5 f/ml. Overall, small opacities (> 1/0) were reported in 88/379 (23%) of chest radiographs, with evidence of a gradient seriously confounded by date of first employment and transfer of subjects with suspected asbestosis to less dusty conditions. On the basis of data on incidence, the authors drew conclusions on exposure-response between cumulative exposure and prevalence or incidence of crepitations, possible asbestosis and certified asbestosis - all three depending on clinical opinion and judgement. The authors state that possible asbestosis occurs in no more than 1% of men after 40 years of exposure to concentrations between 0.3 and 1.1 f/ml" [EHC 203, p 105].

On p. 114, EHC 203 goes on to discuss other consequences of exposures sustained during textile manufacture:

"The health of employees has been studied in any detail in only three asbestos textile plants. These comprise a factory at Rochdale, England, originally studied by Doll (1955) and more recently by Peto et al. (1985), another located in Mannheim, Pennsylvania, USA, studied by McDonald et al. (1983b) and a plant in Charleston, South Carolina, USA. Only the study in South Carolina is considered primarily relevant for assessment of the health effects of chrysotile. Although the SMRs for lung cancer in these plants were broadly equivalent, the rates of mesothelioma varied considerably, which may reflect the greater proportions of amphiboles in the Mannheim and Rochdale cohorts.

The textile workers in South Carolina plant have been studied in two separate but overlapping cohorts ... The only amphibole used in this plant was approximately one tonne of imported crocidolite from the early 1950s until 1972, plus a very small quantity of amosite for experimental purposes in the late 1950s. The crocidolite yarn was processed at a single location only, so Charleston can be considered an almost pure chrysotile operation. Exposure levels for workers at this plant were estimated by Dement et al. (1983a) using nearly 6000 exposure measurements covering the period 1930-1975 and taking into account changes in plant processes and engineering controls (Table 7). The conversion of past exposures measured in mpcm (mpcf) to f/ml was based on both paired sample data (100 pairs) and concurrent samples (986 samples) by these two methods collected in plant operations during 1968-1971."
The most recent update of the Charleston study by Dement et al. (1994) demonstrated an overall lung cancer SMR of 1.97 (126 observed) and an overall SMR for non-malignant respiratory diseases... of 3.11 (69 observed). The data for white males, for which data were more complete, demonstrated an overall lung cancer SMR of 2.34 for those achieving at least 15 years of latency. The risk of lung cancer was found to increase rapidly in relation to cumulative exposure. Data for the entire cohort demonstrated an increase in the lung cancer risk of 2-3% for each fibre/ml-year of cumulative chrysotile exposure. Two mesotheliomas were observed among this cohort and an additional mesothelioma was identified among plant workers, occurred after the study follow-up period. Analyses of an overlapping cohort from the same factory... provided similar results.

... the regression line slopes for relative risks of lung cancer in relation to accumulated exposure in the Charleston plant are all some 30 times steeper than those observed in chrysotile mining and cement product manufacture."

5.260 From the foregoing discussion, it is plain that these risks apply to occupational circumstances, and not to non-occupational situations. It is my perception that there is debate among experts over the carcinogenicity of chrysotile released from or associated with friction products and textiles respectively.

Dr. Infante:

5.261 Exposure to chrysotile asbestos through the manufacturing and downstream manipulation of friction products and textiles carries with it the risks associated with exposure to asbestos, most notably, lung cancer, asbestosis and mesothelioma. This is mostly an occupational issue except for consumers, who do their own brake replacement which would put them at risk of developing these diseases.

5.262 Epidemiological studies of workers involved in the manufacturing of friction products demonstrate an elevated risk of lung cancer (McDonald et al. 1994). Other investigators have not observed an excess of lung cancer in the manufacturing of chrysotile and crocidolite containing friction products, but have identified cases of mesothelioma related to both types of asbestos used to produce these products (Berry and Newhouse, 1983). Cases of mesothelioma also have been reported among car mechanics who serviced brakes containing chrysotile fibres only and were exposed to levels estimated to be below 1 f/cc-year cumulative exposure (Woitowitz and Rodelsperger, 1991). Epidemiological study results indicate a high risk of disease related to chrysotile textiles. See my responses to Questions 4(a)-4(c).

Dr. Musk:

5.263 It is my opinion that there is a risk of disease from the release of chrysotile fibres from friction materials (as in brakes and clutches) or textiles (as in asbestos blankets and suits). In general the risk will be dependent on the degree of exposure (as above) and is therefore likely to be greater in people with occupational rather than non-occupational exposure. Fibres released from friction materials may be shorter than from other sources. These fibres may clear more rapidly from the lungs and possibly be related to lower risks but I am not aware of any direct data on this.

Question 3:

The parties disagree as to the relative pathogenicity of amphibole and chrysotile asbestos. Canada argues that, due to chemical and physical differences, a crucial distinction is to be made between amphibole and chrysotile asbestos: the latter is less pathogenic than the former. The European Communities, on the other hand, argues that chrysotile is as dangerous as amphiboles. In answering the four sub-questions below, please specify to what extent your opinion is based on epidemiological data, in vivo or in vitro evidence.
3. (a) **For the purpose of assessing the risk to human health arising from exposure to asbestos fibres, should a distinction be made between chrysotile and amphibole asbestos?**

**Dr. de Klerk:**

5.264 In terms of risks to human health, epidemiological evidence is clear that, for a given quantity (intensity and duration) of exposure, chrysotile imparts less risk than amphibole fibres.

**Dr. Henderson:**

5.265 YES — a clear distinction should be made between chrysotile and the amphibole forms of asbestos. On a fibre-for-fibre basis, amphiboles such as crocidolite and amosite are substantially more carcinogenic for the mesothelium than chrysotile. This potency differential is to some extent confounded by the far greater usage of chrysotile both now and historically (> 95 per cent of world asbestos production). In addition, it is worth reiterating that Canadian chrysotile on average contains trace quantities of fibrous tremolite (an amphibole).

5.266 It is my perception that there is broad agreement among experts that the amphiboles are more potent carcinogens for the mesothelium than chrysotile.

**Dr. Infante:**

5.267 For purposes of assessing disease risk from exposure to asbestos fibres, I see no basis for making a distinction between chrysotile and amphibole asbestos. Several high quality epidemiological studies of workers exposed to chrysotile asbestos demonstrate an elevated risk of death from lung cancer, asbestosis and mesothelioma. The risk of death from lung cancer and asbestosis related to chrysotile exposure appears to be similar to that from exposure to other forms of asbestos. Although epidemiological study suggests that the risk of death from mesothelioma from chrysotile exposure may be less than the mesothelioma risk from amphibole asbestos, it is somewhat difficult to make this comparison. Many of the studies of workers exposed to amphibole asbestos go back further in time and less information is available on the quantitative aspects of exposure. Thus, the role of error in exposure estimation on the perceived difference is difficult to determine. On the other hand, some experimental inhalation studies demonstrate that chrysotile asbestos may be more potent than other forms of asbestos in the induction of mesothelioma (and lung cancer) in relation to the amount of dust deposited in the lungs (Wagner et al., 1974). In any event, the attributable risk of respiratory diseases from exposure to asbestos is going to be heavily weighted by lung cancer and asbestosis. So, even if exposure to chrysotile asbestos would result in a slightly lower relative risk of death from mesothelioma, the overall risk of the asbestos related diseases combined, i.e., lung cancer, asbestosis, mesothelioma, decrements in pulmonary function, will not be perceptively different for chrysotile as compared to the amphiboles. Thus, a distinction should not be made between chrysotile asbestos and amphibole asbestos. In my opinion, exposure to all forms of asbestos results in a significant disease burden to society.

5.268 I believe that there is a need to distinguish chrysotile asbestos from amphiboles based on the epidemiological data at least.

3. (b) **What are the key properties which cause pathogenicity of, respectively, amphiboles and chrysotile fibres for (i) asbestosis, (ii) lung cancer, (iii) mesothelioma, and (iv) other asbestos-related pathologies?**

**Dr. de Klerk:**

5.269 The properties are the same: size, shape and durability in the lung, that is fibres have to be of certain size and shape to be deposited in the lungs and have to stay there long
enough to produce a response. Since most of the body’s responses to fibres appear to be stochastic in nature, the additional features are of course the intensity and duration of exposure as outlined above. All the types of asbestos differ according to these properties, the main difference with chrysotile is that it is less durable in lung tissue than amphibole fibres: it is more soluble and the fibres tend to break more readily into smaller fibrils, it also tends to be more curly rather than dead straight in shape.

Dr. Henderson:

5.270 As discussed already, the pathogenicity of asbestos appears to reside in the physical properties and biopersistence of the fibres, summarized as the 3 Ds — namely dose, fibre dimensions and durability (bio-persistence). All commercial asbestos has the capacity to induce asbestosis, lung cancer, mesothelioma and other pleural abnormalities (e.g., parietal pleural fibrous plaques, benign asbestos pleuritis with effusion, and diffuse pleural fibrosis). Chrysotile and the amphiboles have different potencies in generating these disorders: for example, the amphibole varieties of asbestos appear to be substantially more pathogenic than chrysotile for the induction of asbestosis and mesothelioma, whereas the differential is not sustained for the induction of lung cancer, for which chrysotile is associated with one of the lowest lung cancer rates (in Quebec chrysotile miners and millers) and the highest rate of lung cancer (South Carolina asbestos textile workers, who used Canadian chrysotile).

5.271 Asbestosis: There is good evidence that asbestosis is a dose-dependent disorder with a threshold effect. There is widespread agreement that asbestosis in general is a consequence of high intensity exposure (or lower intensity but more prolonged exposure) than mesothelioma not associated with asbestosis, so that the concentration of asbestos bodies and uncoated asbestos fibres within the lung tissue of asbestotics is considerably higher than the concentrations encountered in mesothelioma patients without asbestosis and in individuals with parietal pleural plaques. In addition, some studies have shown that the severity of asbestosis and its liability to progression are related to the asbestos body and fibre concentration in lung tissue. In the past, asbestosis as a consequence of high-dose exposure was a progressive disorder leading to progressive respiratory insufficiency and death, whereas many cases of asbestosis encountered during the 1980s and 1990s represent milder and static forms of the disease.

5.272 Churg [234] points out that:

"... a considerably greater burden of chrysotile (with its accompanying tremolite) than of amosite or crocidolite is required to produce any particular diseases. ... For example, the mean burden of chrysotile plus tremolite in the lungs of [Quebec chrysotile] miners and millers with asbestosis is 17 times the asbestosis burden in the lungs of shipyard workers with asbestosis.” [p 294].

5.273 To some extent, this differential in potency may reflect fibre dimensions and the generation of oxidants, but a more likely explanation is the faster clearance of chrysotile from lung tissue than any of the amphiboles. However, Churg’s comments seem to apply to Quebec chrysotile miners and millers in particular; in contrast, the study reported by Green et al. [191], which reported histological asbestosis at relatively low cumulative exposures was carried out on South Carolina asbestos textile workers who also worked with Canadian chrysotile (please see following discussion).

5.274 There is also evidence that long fibres are implicated in the development of asbestosis, but this may reflect in part the anatomical distribution of long versus short fibres in lung tissue. For example, some studies have shown that short fibres in the vicinity of 1 um in length have a biological potency similar to that of the longer fibres for initiation of the inflammatory changes implicated in the development of asbestosis, but they do not penetrate the walls of bronchi or bronchioles to the same extent as the longer fibres, to reach the alveolar interstitium. Another study related the development of asbestosis to the total surface area of deposited fibres, rather than to particular lengths of fibres.
5.275 The effect of dose: There is good evidence that the inhaled dose of asbestos affects: (i) the development of the disease itself; (ii) the latency period between exposure and the onset of the disease; and (iii) the severity and progression of disease.

5.276 Fibre burden studies on human lung tissues show that patients with asbestosis in general have higher tissue burdens than patients with asbestos-related diseases other than asbestosis. Accordingly, Mossman and Churg [202] state that:

"... asbestosis is clearly a fiber-dose driven disease but, nonetheless, only a fraction of any cohort exposed to a fibrogenic dose of asbestos develops asbestosis. It has been proposed that person-to-person variations in either fiber deposition or fiber clearance may account for this phenomenon." [p 1671].

5.277 The Ontario Royal Commission [235] noted that asbestos exposures < 25 fibre-years would be unlikely to produce clinical asbestosis (about = 1 per cent of individuals exposed at this level may develop clinical or radiological asbestosis), whereas Browne [236] considered that the minimum dose required to produce clinical asbestosis was in the range of 25-100 fibre-years.

"Chest X-ray changes among textile and friction product workers in China were reported by Huang (1990). A total of 824 workers employed for at least 3 years in a chrysotile products factory from the start-up of the factory in 1958 until 1980, with follow-through to September 1982, were studied. Chest X-ray changes compatible with asbestosis were assessed using the Chinese standard system for interpretation of X-rays. Cases were defined as Grade I asbestosis (approximately equivalent to ILO = 1/1). Overall, 277 workers were diagnosed with asbestosis during the follow-up period, corresponding to a period prevalence of 31%. Exposure-response analysis, based on gravimetric data converted to fibre counts, predicted a 1% prevalence of Grade I asbestosis at a cumulative exposure of 22 f/ml-years." [EHC 203, p 106]

5.278 In an autopsy study of South Carolina asbestos textile workers who used Canadian chrysotile — the same group studied by Dement et al. [171, 172, 237-241] and McDonald et al. [161, 242] — Green et al. [191] showed that histological asbestosis was usually present at = 20 fibre-years of exposure, and a few cases were observed at 10-20 fibre-years. Thimpont and de Vuyst [243] reported that around 50 per cent of specimens of lung tissue removed because of lung cancer showed low-grade airway and interstitial fibrosis with asbestos bodies, when the asbestos body concentration was = 5000 per gram dry lung tissue.

5.279 These different threshold doses are not inconsistent with each other, because they deal with identification of asbestosis by different modalities (i.e. clinical/radiological asbestosis versus histological asbestosis). In this respect, histological examination is generally considered to represent the most sensitive and specific technique for the diagnosis of asbestosis, followed in descending order by high-resolution CT scanning, conventional CT scans and chest X-rays (which will fail to detect asbestosis in about 20 per cent of cases, especially low-grade asbestosis). In other words, early (grade I) asbestosis may be undetectable by clinical investigations.

5.280 Latency interval: there is also evidence that the latency period between first exposure to asbestos and the subsequent diagnosis of asbestosis is roughly inversely proportional to the exposure level, so that short latencies are encountered with high exposures (e.g. the Wittenoom cohort).

5.281 Mossman and Churg [202] also state that:

"Fibre burden studies also indicate that there is a correlation between pathologic severity of asbestosis and increasing burden of asbestos bodies (which are largely markers of amphibole exposure) or uncoated amosite and crocidolite fibres" ... [p 1670].
There are two additional points that are worth emphasis:

- There appears to be considerable variation among individuals in their propensity to develop asbestosis (or variation in the latency intervals for equivalent exposures). For example, I have seen cases where there was a high content of amphibole asbestos in lung tissue (up to 100 million fibres per gram dry lung tissue or more), in the absence of histological asbestosis — at the time when the fibre burden analysis was carried out — whereas other cases had clinical asbestosis with much lower tissue fibre burdens. Of course, one caveat about this situation concerns latency, whereas another relates to the time when the fibre burden analysis was carried out, which is different from the time when the disease was developing.

- In addition to high fibre burdens, the severity and progression of asbestosis can be influenced by other factors, such as cigarette smoke (though tobacco smoke by itself cannot cause asbestosis, of course).

Mesothelioma: Please see discussion in Section C.1.(f) to(h).

Lung cancer: Please see discussion in Section C.1.(i).

Other asbestos-related pathology: The effect of dose is less clear for the induction of parietal pleural fibrous plaques than for other asbestos-related disorders. There is evidence that the frequency and extent of plaques are dose-related, so that plaques tend to be more extensive with higher exposures. However, plaques can also follow trivial exposures to asbestos and asbestos-like minerals, so that the frequency of asbestos-related pleural plaques appears to correlate more closely with duration since exposure than the level of exposure. Plaques are known to be endemic in Finland, apparently as a consequence of very low-level exposure to asbestos-like fibres in the general environment; on the other hand, in societies where plaques are not endemic — e.g. North America, Western Europe and Australia — about 80-90 per cent of radiologically well-defined plaques are a consequence of occupational exposure to asbestos. In such societies, pleural plaques also represent a useful tissue marker for prior asbestos exposure. At the same time, it is also worth emphasising that parietal pleural plaques by themselves do not predispose to any other asbestos-related disorder, the liability being related to inhaled dose and fibre types.

The key properties of the pathogenicity of amphiboles and chrysotile fibres, although not known for certain, are thought to be related to the physical characteristics of the fibres, namely the diameter, length, aspect ratio of length to diameter, surface area and perhaps surface charge of the fibres. Toxicological data suggest that long thin asbestos fibres may be relatively more potent than other asbestos fibres in their ability to induce asbestos related diseases. There is also experimental evidence, however, that shorter fibres can produce asbestos related diseases, though not in as great a magnitude. Any manipulation of these fibres that results in their diameter becoming thinner may provide a relatively greater contribution to the associated pathology and a greater toxic response than if the fibres were not manipulated in any way. The issue of solubility has been raised in terms of asbestos fibres and their capability to cause disease. The role of solubility with asbestos fibres and disease potential, however, is not so clear because chrysotile asbestos seems to have the same overall potency as other forms of asbestos, yet chrysotile fibres appear to be relatively more soluble than amphibole fibres.

Dr. Musk:

It is my opinion that the key characteristics determining the pathogenicity of asbestos for asbestosis, lung cancer, mesothelioma and other diseases are determined by the physical and chemical characteristics of the asbestos. The physical characteristics in-
clude length, diameter and “straightness” of the fibres. The chemical properties determine the durability of the fibres.

3.(c) What is the respective capacity of amphiboles and chrysotile to induce (i) asbestosis, (ii) lung cancer, (iii) mesothelioma, and (iv) other asbestos-related pathologies?

Dr. de Klerk:

5.288 Comparisons between chrysotile and the amphiboles in their capacity to produce mesothelioma and lung cancer have been extensive, for the other diseases, much less so. There is some in vitro and in vivo evidence that amphiboles, particularly crocidolite, are more fibrogenic than chrysotile, but there is no clear epidemiological evidence on this. Pleural plaques appear to be more common among anthophyllite workers than others while crocidolite workers have more diffuse pleural thickening, and benign asbestos pleurisy also seems to be more common after crocidolite exposure. Historically, asbestosis occurred commonly after heavy exposure to all types of asbestos. For mesothelioma, it is thought that for a given cumulative exposure, chrysotile is between one tenth and one hundredth as potent as crocidolite. There is some controversy over the relative capacity of amosite and crocidolite, but amosite seems to carry about one tenth of the risk of crocidolite. For lung cancer, amosite and crocidolite seem to have similar capacities, with chrysotile weighing in around one tenth to one fiftieth of this.

Dr. Henderson:

5.289 See my response to Question 3(b).

Dr. Infante:

5.290 The quantitative risk of dying from lung cancer as a result of exposure to chrysotile asbestos is at least as great as that from exposure to other forms of asbestos. Quantitative risk assessments based on several epidemiological studies indicate a very high risk of lung cancer (potency) among workers exposed to chrysotile asbestos. The cohort study by Dement et al., first published in 1983 and updated in 1994 contains one of the best estimates of worker exposure to chrysotile accompanied by the estimates of relative risk (RR) for lung cancer and asbestosis. In this study, the investigators determined conversion ratios from the available industrial hygiene data by evaluating sampling results from surveys that used the impinger method to measure dust in million particles per cubic foot (MPPCF), and membrane filter samples that allowed for the counting of fibre concentrations. For early exposure periods, the investigators converted one MPPCF to the equivalent of 3 f/cc > 5 um in length for all areas except for preparation where a conversion factor of 8 fibres for every MPPCF was used. In my opinion, this study is the strongest in its methodology for estimating asbestos exposure among cohort members. Quantitative risk assessment based on data for the entire cohort estimates an increase in the RR of lung cancer ranging from 2-3 per cent for each fibre/cc-year of cumulative chrysotile exposure. To my knowledge, this is the highest estimated risk of lung cancer among asbestos exposed workers that has been corroborated by other investigators, i.e., the same population was also studied by McDonald et al., (1983) and the results are remarkably similar. Studies of lung cancer risk among chrysotile asbestos textile workers from two additional occupational cohorts by McDonald et al., (1982) and Peto et al., (1985) also provide similar results. Using data from the McDonald et al., (1983) study and a conversion factor of 6 fibres per MPPCF, Peto et al., (1985) estimated an increase in the RR of lung cancer among chrysotile textile workers to be 1.25 per cent per fibre/cc-year of exposure. From their own study of Rochdale textile workers, Peto et al., (1985) estimated the excess lung cancer risk to range from 0.5-1.5 per cent per f/cc-year of cumulative exposure depending upon whether the estimate was based on the entire cohort or on those employed in 1951 or later, respectively. These estimates are of surprisingly similar magnitude given that they are from epidemiological studies that incorporated retrospective exposure estimation that had to rely upon the conversion of measurements from particles to fibres per cc. Thus, three separate populations of chrysotile asbestos textile workers demonstrate remarkably similar elevated
risks of lung cancer and therefore add to the confidence of the estimates of excess lung cancer risk per unit of fibre exposure that these studies demonstrate.

5.291 Studies of workers exposed to chrysotile in several industries demonstrate a significantly elevated risk of lung cancer. Studies of workers exposed to a combination of chrysotile and crocidolite, or to chrysotile only in cement production (Hughes et al., 1987) indicate a virtually identical excess risk of mortality from lung cancer. In a study of workers exposed to crocidolite in mining, de Klerk et al., (1989) estimate an elevated relative risk of lung cancer of 1 per cent per f/cc-year of exposure. Essentially, the excess relative risk for lung cancer indicated in all of these studies is close to about 1 per cent per fibre per cc-year of exposure (Stayner et al., 1997). The risk of lung cancer from chrysotile asbestos exposure is at least as great as the risk of lung cancer associated with amphibole asbestos exposure.

5.292 Analyses based on the study by McDonald et al. (1993) indicates a much lower dose response for lung cancer in relation to chrysotile asbestos in mining and milling as compared to the risk estimates from other studies, particularly chrysotile textile workers. I suspect, however, that the fibre exposure from this study may be overestimated, particularly among the portion of the cohort that is comprised of miners and that a fair amount of misclassification took place in the amount of fibre exposure estimated for individual cohort members.

5.293 Gibbs and Lachance (1972) who published the initial exposure estimates for this population of workers stated that their cumulative dust concentrations for the cohort members may have been far from their actual experience. Subsequent dose response analysis of this cohort applied only a single conversion factor to estimate fibre exposures from the dust count data for the entire cohort. In the study of Siberian chrysotile miners and millers (Tossavainen et al., 1999), those involved in mining experienced average exposures of 0.08 f/cc, while those involved in two separate mills experienced average exposures of 3.62 f/cc (ranges of average exposures for different milling operations 0.37-6.21 f/cc) and 0.65 f/cc (range of average exposures 0.20-1.26 f/cc). The gravimetric sampling results indicate a 5-fold difference in average exposure for miners versus millers, while the sampling results for fibre exposures (the exposure of concern) indicate a 45-fold difference in average exposure concentrations between miners and millers. Taking into consideration the variation within mining and milling jobs, the difference between exposures to workers involved in these jobs would be even greater. The large difference in the fibre exposures between miners and millers that was observed in the Tossavainen et al. (1999) study adds further support to my concern that applying a single conversion factor from dust samples in estimating chrysotile fibre exposures for miners and millers is the most likely explanation for the slope of the dose response for lung cancer in the McDonald et al. study being so different from the slope for lung cancer based on the studies of chrysotile textile workers. Stayner et al. (1996) provide a summary of data indicating that the lung cancer risk from exposure to chrysotile in either experimental animals or in humans is similar to the risk from exposure to amphibole asbestos in these species.

5.294 With regard to asbestosis, Stayner et al. (1997) published a dose response analysis based upon the updated study of chrysotile textile workers by Dement et al. (1994). Their analysis indicates an excess risk of 2 deaths from asbestosis per 1000 employees exposed to 0.1 f/cc for an occupational lifetime exposure of 45 years, or 0.2 per cent from a cumulative exposure of 4.5f/cc-years. Two additional studies estimate similar risks of death from asbestosis in relation to cumulative chrysotile exposure. The study by Berry et al. (1979) estimates a risk of 1 per cent for those categorized as “probably having asbestosis” among chrysotile textile workers who were exposed to an estimated range of 0.3 f/cc to 1.1 f/cc for 40 years, or a cumulative exposure of 12-44 f/cc-years. For those categorized as “certified asbestosis” a 1 per cent excess risk of death was associated with 63 f/cc-years of exposure. Huang (1990) estimates a risk of asbestosis of 1 per cent associated with 22 f/cc-years of exposure during the manufacturing of chrysotile textile and friction products. Data from Dement et al. (1994) allow for an estimate of 2 per cent asbestosis associated with 22.5 f/cc-years of exposure. Studies of workers exposed to a mixture of chrysotile
and crocidolite asbestos in the manufacture of cement products have demonstrated an
elevated the risk of “certified asbestosis” of 1 per cent associated with 10 f/cc-years of
exposure (Finkelstein 1982). Finkelstein was of the opinion that he identified a 1 per cent
excess risk of asbestosis related to a lower cumulative dose of asbestos exposures than
Berry et al. (1979) because of the longer period of follow-up of the cohort which gave more
time for the asbestosis to clinically manifest itself.

5.295 I am not aware of evidence that similar cumulative exposures to amphibole forms
of asbestos will result in a greater risk of asbestosis. Thus, it is difficult to make any distinc-
tion in potency between the amphiboles and chrysotile asbestos in relation to asbestosis.

5.296 Chrysotile exposures related to numerous jobs and occupations has been associ-
ated with mesothelioma through epidemiological studies and case reports. In some situ-
atations, standby exposure only was associated with mesothelioma. Based on epidemiologi-
cal studies, the potency of chrysotile to induce mesothelioma may be less than that from
other forms of asbestos. However, the rarity of mesothelioma in the general population
and the difficulty in determining asbestos exposure levels experienced by cohort members
decades before measurements were taken, coupled with conversion of dust counts in par-
ticles to fibres per cc, make it difficult to determine differences in potency estimates with
regard to the various forms of asbestos and mesothelioma. Based on toxicological study
results, in terms of the quantity of dust deposited and retained in the lungs, chrysotile may
be more potent than other forms of asbestos in the induction mesothelioma and fibrosis
(Wagner et al., 1974). The population attributable risk of contracting mesothelioma from
chrysotile, however, will be greater than from other forms of asbestos because of the much
greater potential for exposure to chrysotile.

5.297 I have not seen any quantitative data related to decrements in lung function for
either chrysotile and amphibole asbestos. On the basis of mortality from asbestosis, I as-
sume there is little difference in lung function related to the various forms of asbestos.

5.298 In summary to this question, in evaluating the epidemiological evidence, I see
no basis for concluding that the overall disease potential from exposure to the amphiboles
is any different than that from exposure to chrysotile asbestos with the possible exception
that amphiboles may be more potent in causing mesothelioma and chrysotile may be more
potent in causing lung cancer. Studies in experimental animals demonstrate the ability of
chrysotile asbestos as well as the amphiboles to induce fibrosis, lung cancer and mesothe-
lioma. From a public health standpoint, in terms of quantification of disease, it would be
extremely difficult to make any distinction between exposure to the amphiboles and
chrysotile asbestos fibres.

Dr. Musk:

5.299 Broadly it is my understanding that the relative pathogenicity of the different
fibres for the various diseases is different (see Table)

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Question 4:

The parties in this dispute disagree as to the risk to human health associated with chrysotile asbestos fibres at low levels of exposure, i.e. either prolonged exposure to low concentrations of fibres or occasional peaks of exposure. The European Communities considers that, because of a lack of data at low levels of exposure, it is appropriate to endorse the linear relationship model to assess risks associated with such low levels of exposure. On the other hand, Canada is of the view that, at such low levels of exposure, empirical evidence suggests that there is a practical threshold below which chrysotile asbestos fibres present no measurable effects on health.

4.(a) Are epidemiological data available for low levels of exposure to chrysotile fibres and what do they show?

Dr. de Klerk:

5.300 There have been several epidemiological studies that have shown no increased risk for low levels of exposure to chrysotile, particularly in friction products industries.

Dr. Henderson:

5.301 So far as I am aware, there are no exposure-response data for such levels of exposure.

5.302 For example, EHC 203 states the following:

"Few data on concentrations of fibres associated with the installation and use of chrysotile-containing products are available, although this is easily the most likely place for workers to be exposed" [EHC 203, p.3].

"Overall, the available toxicological data provide clear evidence that chrysotile fibres can cause fibrogenic and carcinogenic hazards to humans. The data, however, are not adequate for providing quantitative estimates of the risk to humans. This is because there are inadequate exposure-response data from inhalation studies, and there are uncertainties concerning the sensitivities of the animal studies for predicting human risk" [EHC 203, p.7].

"There is evidence that fibrous tremolite causes mesothelioma in humans. Since commercial chrysotile may contain fibrous tremolite, it has been hypothesized that the latter may contribute to the induction of mesotheliomas in some populations exposed primarily to chrysotile. The extent to which the observed excesses of mesothelioma might be attributed to the fibrous tremolite content has not been resolved" [EHC 203, p.8-9].

"Epidemiological studies that contribute to our understanding of the health effects of chrysotile conducted to date and reviewed in this monograph have been on populations mainly in the mining or manufacturing sectors and not in construction or other user industries. This should be borne in mind when considering potential risks associated with exposure to chrysotile" [EHC 203, p.137].

Dr. Infante:

5.303 One means of determining the risk from low exposure levels for carcinogens is to estimate the risk from studies where exposure information is of reasonably good quality and the estimates of risk were made using sound epidemiological principles and methodology. Once these studies have been identified, an appropriate way to determine quantitative risk from low exposure levels is to use all of the available data in a particular study and to estimate dose response. As mentioned in my response to Question 3(c) above, several
studies of workers exposed to chrysotile asbestos can be used to estimate risk from low levels of exposure. The studies mentioned above, which represent three separate populations of chrysotile textile workers demonstrate an excess relative risk of lung cancer ranging from 0.5-3 per cent for each fibre/cc-year of exposure. Risk assessment based on the study of South Carolina chrysotile textile workers (Stayner et al., 1997) indicates that individuals exposed to 0.1 fibre/cc-year for an occupational lifetime of 45 years, e.g., 4.5 fibre/cc-years cumulative exposure to chrysotile, have an elevated risk of 5 extra deaths from lung cancer and 2 extra deaths from asbestosis per 1000 workers. Dose response for mesothelioma could not be estimated because there were too few deaths from this cause in the study.

5.304 Epidemiological study results and case reports indicate that a large number of jobs which entail occasional peak exposures to chrysotile asbestos have resulted in workers being diagnosed with mesothelioma. Furthermore, mesothelioma has been diagnosed among household contacts of asbestos cement workers (Magnani et al., 1992; Ascoli et al., 1996), among individuals living near chrysotile mining and milling operations (Begum et al., 1992), or living in houses constructed with asbestos cement (Ascoli et al., 1996), or who were exposed to low cumulative amounts of chrysotile asbestos from brake lining work (Woitowitz and Rodelsperger, 1991), or from standby exposure as bakers (Ascoli et al., 1996). This information contributes to the evidence that very low exposure to all forms of asbestos can induce cancer. These observations of mesothelioma should be considered as sentinel events for the pathologies other than mesothelioma that are more difficult to identify among large populations that experience relatively more remote exposure to chrysotile asbestos. In my opinion, these studies constitute high level risk from low levels of exposure to chrysotile asbestos.

Dr. Musk:

5.305 It my understanding there are epidemiological studies which do not show a statistically increased risk of disease from low levels of exposure to chrysotile. However, the absence of demonstrating an increased risk does not mean that there is not some risk as it is not possible to prove a negative and a threshold has not been demonstrated for any carcinogen (nor in my opinion is it biologically likely that one exists).

4.(b) Is there a threshold below which exposure to chrysotile fibres does not induce (i) asbestosis, (ii) lung cancer, (iii) mesothelioma, and (iv) other asbestos-related pathologies, such as pleural plaques? If there is such a threshold, is it a practical one or is it scientifically established?

Dr. de Klerk:

5.306 It is extraordinarily difficult to demonstrate lack of an effect, or a threshold effect, in epidemiological studies because of the ubiquitous problems of bias, confounding and chance. In particular, the smaller effect that needs to be demonstrated, the larger the study needs to be, both in population size and follow-up time and such studies can rarely be done, even with animals.

Dr. Henderson:

(i) Asbestosis

5.307 Please see my answer to Question 3(b).

(ii) Lung Cancer

5.308 Please see discussion in Section C.1.(i). Please see also the statement from EHC 203 below on the question of a threshold for carcinogenesis by asbestos. Some authorities favour a linear no-threshold model, whereas others argue that a threshold probably exists; nonetheless, there is no general agreement on a numerical threshold for asbestos-induced
lung cancer. Dement et al. [171] observed odds ratios of > 2.5 at 2.7-6.8 fibre-years of exposure among South Carolina asbestos textile workers.

(iii) **Mesothelioma**

5.309 As indicated in previous discussion in this report, a linear dose-response model has been identified for mesothelioma induction by the amphiboles, and the dose-response relationship is maintained at low occupational levels of exposure that overlap with environmental exposures: e.g. definite dose-response relationship reported by Rödelsperger et al. [25, 137] at asbestos fibre concentrations in lung tissue of 100,000-200,000 fibres per gram dry lung tissue, with an indication that this relationship is maintained at lower levels of 50,000-100,000 fibres per gram dry lung (the level of 100,000-200,000 fibres corresponds to a cumulative dose of 1-2 fibre-years). Iwatsubo et al. [136] identified an increase in the relative risk at 0.5-0.99 fibre-year. No threshold has been identified for amphibole-related mesothelioma. For chrysotile exposures, a dose-response relationship has also been identified at high exposures, but to the best of my knowledge, there are no dose-response data for low-level exposures to chrysotile.

5.310 On this point, EHC 203 states that “No threshold has been identified for carcinogenic risks” [for chrysotile; p 144]. At the same time, no increase in risk for mesothelioma has been identified at very low levels of exposure, of the type associated with well-maintained asbestos in place, in public buildings. However, it is impossible to ascertain whether or not there is an increase in risk at this order of exposure, because no control or reference group can be assembled where there is no asbestos content in lung tissue. If a threshold exists, it must lie somewhere in this area, between no exposure, low-level environmental exposure, and low-level occupational exposure.

5.311 De Klerk [115] has also commented on the difficulty or impossibility of distinguishing between background versus environmental mesotheliomas:

“There have been increases in the incidence rates of malignant mesothelioma in women, in those without identified exposure to asbestos and, possibly, those younger than 35 years of age in Australia and Western Australia. Although part of the first two increases, at least, may be attributable to specific exposure to asbestos, mathematical modelling of the Western Australia data suggests that there has been about a twofold increase in incidence rates from the 1970s to the 1980s that may be due to increased general environmental exposure to asbestos. ... The excess of 1 per million person years over this presumed ‘background’ rate is also, coincidentally, the amount that was estimated as possibly caused by exposure of school children to 1 fiber per liter ... , a level that might result from use of asbestos-based insulation or other general contamination of the environment with asbestos.

A final consideration in the use of national trend data for estimating environmental effects is the comparison of the likely extrapolated risk from occupational data with the background risk estimated here or from Peto’s Los Angeles data. From the Peto paper ... , the incidence of mesothelioma is related to age in the following way:

\[
\text{Incidence} = 1.7 \times 10^{-12} \times (\text{age})^{13}
\]

which translates to a lifetime risk to age 80 of just over 100 per million people, which is much greater than any of the estimated environmental risks described earlier. An equivalent risk from the adjusted Western Australian data is about 160 per million lifetimes. The question remains as to how these background risks and environmental risks interact. Is the postulated environmental incidence already included in the background incidence, or should the risks be added or even multiplied together? This question is almost certainly unanswerable using epidemiological methods. ...
It is doubtful whether epidemiological methods ... could ever be definitive in deciding whether there is an appreciable hazard from general environmental exposure to asbestos ... or, more importantly, whether the hazard is large enough to justify specific remedial action ...” [pp 29-31].

5.312 Clearly, this issue is one major focus of dispute between experts; from the preceding discussions, it is my perception that — with the exception of asbestosis — no threshold has been delineated, and that even those who claim that a practical threshold must exist cannot delineate such a threshold in precise numerical terms (in this respect, I do not know what the expression “practical threshold” really means).

Dr. Infante:

5.313 Thresholds have not been demonstrated for any substance known to cause cancer and there is no theoretical basis to assume a threshold for the diseases related to chrysotile, or other forms of asbestos, particularly when the mechanisms involved in the pathology are not fully understood. Furthermore, it is not possible to determine thresholds from epidemiological studies because of lack of statistical power to distinguish that the risk is virtually zero. [Note: At times, some investigators state that a single point estimate from the lowest dose evaluated in an epidemiological study that does not demonstrate a significant elevation in cancer risk constitutes a threshold level for the carcinogen. Such a conclusion is scientifically invalid. When estimating dose-response, one has more confidence in the risk related to a particular dose level by using all of the data available in the study. Using only a single point estimate results in more instability in the estimate of risk for that data point in contrast to using all of the available data in the study.]

5.314 Dose response analyses and modeling specifically for chrysotile asbestos exposure and lung cancer and asbestosis have been conducted recently by Stayner et al. (1997) using data from the Dement et al. (1994) study. Alternative exposure-response models were evaluated as part of the study. A model designed to evaluate evidence of a threshold also was fitted for asbestos exposure in relation to lung cancer and asbestosis. There was no significant evidence for a threshold in models pertaining to either lung cancer or asbestosis.

5.315 With regard to mesothelioma, and other asbestos-related diseases, I am not aware of any evidence of a threshold pertaining to either chrysotile, or other forms of asbestos. Furthermore, from a practical standpoint, even if there were a threshold for the chrysotile related diseases, the exposures that workers will routinely encounter in the future through continued use of chrysotile in commerce, will expose them to concentrations of asbestos that have already been related to pathology in humans. In other words, continued use of asbestos will continue to expose individuals to levels and exposure circumstances that have already been related to disease. The threshold question, therefore, seems moot.

Dr. Musk:

5.316 It is my understanding that a threshold for disease has not been scientifically established.

4.(c) Is the linear relationship model an appropriate method for assessing the risk to human health posed by exposure to chrysotile asbestos at low levels of exposure?

Dr. de Klerk:

5.317 The linear relationship model is generally used as a so-called “conservative” estimate, that is, if it is incorrect, it is more likely to err on the side of safety. In some ways, how one extrapolates risk assessment outside the range of available data is more of a societal decision than a scientific one. Biological plausibility could probably be given to any model.
Dr. Henderson:

5.318 In the absence of alternatives because of unavailability of data on exposure-response relationships at low levels of exposure to chrysotile, the linear relationship model is widely employed. Under these circumstances, this may be an appropriate method for risk assessment at low levels of exposure. Whether or not it is a valid method is unknown.

5.319 NICNAS 99 (p. 72) observes that:

“There are many problems associated with low-dose risk extrapolation, such as the assumption of a linear relationship. However, as insufficient data exist to indicate a threshold exposure for effect, the linear extrapolation methodology provides a conservative worst-case scenario estimate of risk. Other confounding factors in estimating risks from epidemiological data are possible contamination by other fibre types and inaccurate estimates of historical exposures.”

5.320 Clearly, this too is one major focus of dispute between experts.

Dr. Infante:

5.321 A linear relationship model is appropriate for determining dose response for chrysotile exposure and lung cancer, and perhaps asbestososis and mesothelioma as well, but the most reasonable model for the latter two diseases are less clear than for lung cancer. With regard to this issue, Stayner et al. (1997) evaluated exposure-response relationships for chrysotile asbestos, and lung cancer and asbestososis by applying several alternate models. The exposure-response relation for asbestos and lung cancer gave the best fit when using a linear model. This observation is consistent with the conclusions of other investigators, who have evaluated dose response for chrysotile asbestos textile workers, or other asbestos workers and mortality from lung cancer (McDonald et al., 1983; Peto et al., 1985; Enterline, Hartley & Henderson, 1987). Furthermore, there appears to be a linear relationship between asbestos exposure and lung cancer over a wide range of exposures where such data are available. Therefore, it seems reasonable to accept a linear relationship for lung cancer when extrapolating risks to exposures below the ranges that have been evaluated in epidemiological studies. Moreover, I am not aware of any literature that convincingly proves that the dose response for asbestos and lung cancer is non-linear. Thus, in my opinion, the linear model is the most appropriate model for estimating dose-response for chrysotile exposure and lung cancer.

5.322 A linear relationship might also be used for chrysotile asbestos exposure and asbestososis although one might make the argument that a non-linear model is also appropriate for asbestososis. Stayner et al. (1997) evaluated this issue using data from the Dement et al. (1994) study and concluded that the association between chrysotile exposure and asbestososis appeared to be non-linear. Stayner et al. (1997) used a non-threshold, non-linear model and the estimates of asbestososis predicted from the model seem to fit very closely with point estimates for asbestososis from other studies of chrysotile exposed populations as mentioned in my responses to Questions 3(c) and 4(a).

5.323 An analysis by Peto et al. (1985) of chrysotile asbestos textile workers shows that a linear model fits the data for mesothelioma with the cube of time since first exposure. In this non-threshold model, the response is linear with dose of asbestos, but exponential with time since initial exposure. The predicted number of mesotheliomas by dose and time since first exposure was in reasonable agreement with the observed number. According to the authors, however, there were too few cases to test the model stringently and they did not attempt to fit other models to their data. Nevertheless, given the consistent observations of the long latency period between initial exposure to various forms of asbestos and the clinical manifestation of mesothelioma, it seems reasonable to use a model that is linear with exposure and exponential with time from initial exposure for chrysotile asbestos and mesothelioma.
Dr. Musk:

5.324 It is my opinion that the linear relationship model is the most appropriate one.

4.(d) Are there scientifically acceptable methods other than the linear relationship model which could be used to assess the risk to human health at low levels of exposure? What results do they suggest?

Dr. de Klerk:

5.325 While a threshold model suggests a lack of risk below a fixed level, it is unlikely that this risk would be completely zero, so that if applied to a much larger population, such a risk could lead to cases of disease.

Dr. Henderson:

5.326 I am not aware of any other methods that have met with broad scientific acceptance or a consensus. It has been suggested that an S-shaped curve might be more appropriate, but I have not seen any data on what the form of the S-curve might be; in other words, the S-shaped model appears to presuppose the existence of a threshold, but no such threshold has been established to the best of my knowledge.

5.327 The problem with arguing that there exists a practical threshold level for lung cancer and mesothelioma induction is that it is impossible to delineate such a threshold in numerical terms, because of a lack of observational data. (Please see also my answer to Q.5(c)).

Dr. Infante:

5.328 From the public health perspective, it has been the convention to use non-threshold linear models for estimating cancer risk to humans. This is particularly the case for substances known to cause cancer in humans. One might deviate from this concept if the mechanism(s) by which the substance causes the cancer were known. This is not the case with chrysotile, or any other form of asbestos. In the particular case of chrysotile asbestos, Stayner et al. (1997) selected several Poisson regression models to explore the shape of the exposure-response relationship between chrysotile asbestos exposure and risk of death from lung cancer. The models were capable of reflecting a wide range of exposure response patterns, including linear, sublinear and supralinear relationships. They also considered a threshold model to determine whether there was evidence that exposures below a certain exposure concentration were equivalent to zero, i.e., that a threshold was present. As mentioned above, for lung cancer, a linear model gave the best fit; for asbestosis, the response preferred was that based on a non-linear, non-threshold model. In both cases, the models did not provide any support for the existence of a threshold. Thus, in my opinion, these models are appropriate to assess risk for these diseases as a result of occupational exposure to chrysotile asbestos. With regard to lung cancer and asbestos, I am not aware of any public health organization, or governmental agency that has ever used a non-linear model to estimate risk. During the hearings held by the U.S. Occupational Safety and Health Administration (OSHA) as part of its rulemaking related to the standard promulgated for asbestos in 1994, numerous scientists were of the opinion that a non-threshold linear model was the preferred model to use for estimating the relationship between asbestos exposure and lung cancer. It is my opinion that a non-linear model is not an acceptable model to use in estimating dose response for asbestos exposure and risk of death from lung cancer. For mesothelioma, I would favor a non-threshold model that incorporates a linear relationship with exposure.

Dr. Musk:

5.329 I believe that there is no reason to discard the linear model as no threshold for any carcinogen is known to exist.
4.(e) To what concentration of chrysotile fibres and for how long must a person be exposed in order to be considered at risk of developing a chrysotile asbestos-related disease (lung cancer, mesothelioma or other asbestos-related pathology)?

Dr. de Klerk:

5.330 A person is “at risk” of developing a chrysotile asbestos-related disease after any exposure to chrysotile asbestos, the lower the amount of exposure, the lower the risk. For example, it could be estimated that there was a 50-50 chance that exposure to 1 fibre of crocidolite could cause 1 case of mesothelioma among the whole population of the world (including all those who have ever lived), i.e. a very small probability, but still greater than zero.

Dr. Henderson:

5.331 This question iterates the issue of a threshold exposure. The answer is essentially the same as for Questions 4(a)-4(d) in the absence of exposure-response data at low levels of exposure.

Dr. Infante:

5.332 The answer to this question depends upon the amount of risk that is considered unacceptable by a particular country. It is a matter of health policy. In the United States, the Environmental Protection Agency (EPA) regulates risk to a level below one extra death in a population of 100,000 people over its entire lifetime. I have already provided estimates for excess risk of death from lung cancer and asbestosis from chrysotile exposure. These risk estimates, however, are average risks to a group of individuals that are based on maximum likelihood estimates (MLEs) and they do not incorporate statistical uncertainty in terms of variability, e.g., they are not based on upper 95 per cent confidence limits as is usually the health policy when estimating adverse health effects to a group of individuals at risk from exposure to an environmental insult. In addition, the risk estimates may only be appropriate to workers’ health risks. They are derived from a group of healthy adults, who were able to pass a physical in order to gain employment. They are not representative of individuals in the general population who may be exposed to asbestos and have a compromised immune system, or be exposed to other conditions which may exacerbate their risk of contracting the various diseases related to chrysotile asbestos as estimated from a healthy worker population. There will always be some risk from exposure to asbestos and the degree of that risk will depend upon the amount of asbestos exposure in relation to the susceptibility of the individuals exposed in terms of their health status and other factors that interact to produce clinical manifestation of disease.

5.333 It is noteworthy, as mentioned in my response to Question 1(a) above, that the population surrounding the Quebec mining and milling operation (Begin et al., 1992) that was exposed to background levels of chrysotile asbestos, developed mesothelioma at an incidence of 62.5 cases per million population per year, or 0.625 per 10,000 per year. This risk level translates to 0.5 per cent of the population developing mesothelioma over an 80-year lifetime from this background exposure. This estimate may well represent an underestimate of risk since the identification of cases was based on a workman’s compensation board review, and additionally, out-migration of inhabitants would also result in loss of some cases. In the same report, it was pointed out that the largest increase in the mesothelioma rate in Quebec was among individuals that had occupations where their exposure would be occasional only, and that 33 per cent of these cases were exposed for less than a 5-year period. When one adds to this information to additional cases of mesothelioma reported in the literature that are associated with standby exposure to chrysotile, it leads one to the conclusion that occasional exposure for a short period of time, or constant low level exposure to chrysotile asbestos leads to death from mesothelioma (and lung cancer and asbestosis). [Note: while an excess of lung cancer was not identified in the Camus et al. (1998) study of women in the same Quebec population, the study had limited power to detect an excess of lung cancer; it did, however, demonstrate an excess of asbestosis and
mesothelioma even though out-migration may have resulted in the loss of all three of these diseases in the study.

Dr. Musk:

5.334 In my opinion any level of exposure to chrysotile (or other form of asbestos) constitutes some risk and that the level of “acceptable risk” is not a scientific issue but an issue for society to debate and determine at different times according to the evidence as they perceive it.

Question 5:

Canada states that, with controlled use, “health risks associated with occupational exposure throughout the life-cycle of chrysotile asbestos can be reduced to acceptable levels already recognized as such by competent international organizations. The European Communities questions this assertion and says that “les données scientifiques disponibles montrent que l'utilisation dite “sécure” de l’amiante chrysotile ne permet pas d’empêcher un grand nombre de cas d’exposition entraînant des pathologies mortelles”.

[“available scientific evidence shows that so-called “controlled” use of chrysotile asbestos does not make it possible to prevent many cases of exposure causing fatal pathologies”]

5.(a) Is there a generally agreed methodology applicable to any use of chrysotile-cement products and other high-density chrysotile products throughout their life-cycle that can be referred to as “controlled use”? Is it embodied in international standards?

Dr. Henderson:

5.336 In principle, regulation and control of chrysotile and high-density chrysotile products is feasible at some points of the life-cycle (manufacture and disposal), but in reality not others (please see following discussion).

5.337 The manufacture of high-density products is usually carried out under closed conditions with dust extraction. As one example, the manufacture of chrysotile friction materials in Australia involves the following processes: following transfer of other ingredients required for the product mix, unopened 50 kg plastic bags of raw chrysotile are placed in the mixer and opened under dust extraction. The empty bag is then delivered into a second plastic bag attached to the mixer. When full, this second bag is sealed and taken to a controlled disposal site. Mixing is a closed process. After mixing, the material is emptied under dust extraction before decanting into smaller buckets for weighing and use in moulding and finishing processes. The moulding is a hot process and when complete, the moulded product undergoes finishing processes that include grinding, grooving and drilling — all carried out under dust extraction. The finished disc pads and commercial vehicle brake blocks and linings are then wrapped and packed into sealed containers.

5.338 The potential for exposure includes opening and emptying chrysotile bags into the mixer, the moulding and finishing processes, and handling of damaged bags containing raw chrysotile. The workforce amounts to a few hundred workers, and the maximum exposures per employee vary from minimal, to the largest group involved in the processing operations. Airborne asbestos fibre levels are assessed by personal monitoring. About 84 per cent of 461 samples between 1992 and 1997 were < 0.1 f/ml; 10 per cent = 0.01= 0.2 f/ml; 6 per cent were = 0.02< 0.5 f/ml, and < 1 per cent were = 0.5 f/ml. The manufacture of compressed asbestos fibre sheeting — most for export and the remainder processed into finished cut gaskets for industrial applications is also a closed process carried out under similar conditions to the manufacture of friction products (please see NICNAS 99, pp 32-34). A total of 232 personal samples between 1991 and 1996 showed similar low airborne
fibre concentration (58 per cent < 0.1 f/ml and only one sample = 0.5 f/ml). Static samples recorded during guillotine and trimming activities were all = 0.05 f/ml.

5.339 Although controlled use of this type is feasible for these manufacturing processes, and for disposal of materials left over (e.g. empty polythene bags), it is my perception that, historically and in reality, it is almost impossible to extend analogous controlled and regulated use to the end-users of asbestos products such as workers involved in building construction and demolition (e.g. builders’ labourers, carpenters, electricians, painters, plasterers and plumbers), or to individuals who carry out maintenance or renovation work on their own homes, or to brake mechanics (please see AMR 99). This is because these groups add up to a large population of disparate and varied workers; many such individuals work for small businesses or are self-employed, so that it is difficult or impossible to extend controlled use or training to all of them.

5.340 Some of the supporting documentation submitted to the WTO refers to ILO recommendations, but it is also worth emphasizing that prohibition or regulation of asbestos-containing products varies from one nation to another, with different upper limits of airborne fibre concentrations (e.g. < 1 f/ml or < 0.1 f/ml). These are summarized in Tables 27 and 28 and Appendix 7 in NICNAS 99.

5.341 From the literature cited throughout this report and the reasons discussed, it is my perception that broad agreement exists among experts that controlled use of chrysotile (or other varieties of asbestos) is not a feasible option in the real world for certain worker groups, notably those involved in construction trades (e.g. see EHC 203).

Dr. Infante:

5.342 I am not familiar with any “agreed upon” methodology applicable to chrysotile-cement products and other high-density chrysotile products related to “controlled use” in the sense that these products could be used without harm to human health. Perhaps the consensus is that when using asbestos, the exposure should be controlled, and various countries have developed programmes or standards that recommended, or require, specific engineering controls, work practices, training and education and personal protective equipment to control exposures to asbestos to the extent feasible. This, to me, is different from the concept of “controlled use” in the dialogue being used by Canada, which seems to imply that using or manipulating asbestos or asbestos containing products can be done in such a manner that people are not exposed, or that the risk from such exposure is de minimus.

5.343 I also am not aware of international standards related to “controlled use” of asbestos products. What you may be referring to here are recommendations from international organizations, or recommendations or regulations (in the context of being enforceable by law) from various countries. These documents, however, should not be considered as international standards that lead to the “controlled use” of asbestos. For example, in 1994, the United States, promulgated a new standard for occupational exposure to asbestos which required the permissible exposure limit (PEL) to be no higher than 0.1 f/cc as an 8-hour time-weighted-average (TWA). In the opinion of OSHA, workers exposed to this PEL over an occupational lifetime (45 years) are still at a significant risk of developing asbestos related diseases. Therefore, the Agency included in the standard several provisions ancillary to the PEL. One could argue that the PEL plus the ancillary provisions constitutes one of the best examples of the concept of “controlled use” of asbestos as I understand it in the concept being brought forward by Canada. Yet, in the United States, the PEL for asbestos as well as the ancillary provisions that include training and education about the health hazards of exposure, work practices, requirements for personal protective equipment, medical surveillance, etc. are not complied with for various reasons in a large number of workplaces. Based on the observations of violations of several provisions of the asbestos standard in the United States, discussions with occupational safety and health personnel from other countries and my review of the literature, it is my opinion that a controlled use concept for chrysotile asbestos is not realistic in workplace situations. It would be much less realistic as applied to non-occupational situations where individuals would make repairs involving the manipulation of asbestos products in their homes.
I am not aware of any international standard that embodies “controlled use” of asbestos in the context that the manipulation of asbestos, or asbestos containing products will result in exposures that will not result in harm to many of those exposed. Furthermore, it should be recognized that programmes to control asbestos in many countries are “agreements” and as such, they are not enforceable by law. Even more disconcerting is the observation that countries like the United States that promulgated stringent requirements to control asbestos find that their standards are often violated. A general downfall with the concept of “controlled use” is that it relies upon human behaviour, which cannot be controlled in too many situations. Hence, it is unreliable.

Dr. Musk:

In my opinion “controlled use” of asbestos is theoretically possible but not practically feasible. The second half of the question is not within my area of expertise.

Dr. de Klerk:

This is definitely outside my area of expertise.

Dr. Henderson:

In theory, training of specific workers (e.g. products manufacture) and some other personnel in the controlled use of chrysotile ought to be feasible (as for other potentially hazardous materials such as radioactive materials used in nuclear reactors). As a matter of common sense, training is most likely to be effective when there is a small and cohesive workforce using materials that are not accessible to most other workers who have limited or no training, and when the workers have a clear understanding of the hazards and risks of the materials handled.

However, training of this type becomes less feasible or impossible in practice when there is a large and non-cohesive workforce and when there is general accessibility to the materials in question (e.g. builders’ labourers, carpenters, electricians, plumbers and so forth at building sites, and brake mechanics).

Even so, I am sceptical about the consistent and universal effectiveness of training programmes in the real world, even when the workforce is small and cohesive, and involved in the handling of materials that present a clear hazard (e.g. radioactive isotopes). For example, following the recent mishap at a nuclear reactor in Japan, a BBC report broadcast by ABC News Radio on Sunday 31 October 1999 pointed out that the workers at the Japanese nuclear plant had been poorly trained, with a poor understanding of the risks.

An analogous report was printed in The Guardian Weekly (October 28-03 November 1999, p. 9):

“Britain’s key nuclear warheads factory this week admitted to more than 100 breaches of safety in the past year but ... the Director of Communications at the Atomic Weapons Establishment in Aldermaston ... branded claims that only luck had prevented an accident worse than that in Japan as ‘irresponsible scaremongering’. ... The Plant’s emphatic denials came after The Observer newspaper published details of a leaked report highlighting more than 100 dangerous incidents since September last year. ... Among these were eight breaches of the ‘criticality’ rules ... and eight instances of environmental contamination outside the site. There were also eight occasions when materials — including plutonium — were incorrectly packaged or labelled, and 19 highly serious health and safety incidents, including all fire-fighting pumping appliances being unfit to
The lead-in of the catalogue of breaches comes as the environment agency prepares to decide whether to prosecute the privatised Aldermarston for dumping tritium, a radioactive substance, in a stream from which Reading’s drinking water is sourced ... it also follows the imposition of a £22,000 fine on the company after two workers breathed in radioactive particles from plutonium that had escaped from a laboratory in August last year. ... The establishment insisted it was ‘regulated up to the eyeballs’ by external agencies and said it was penalised for its ‘openness and transparency’ in reporting breaches ...

Dr. Infante:

5.351 Training is beneficial in reducing worker exposure to toxic substances in some circumstances, but minimizing exposure to asbestos to the extent that a significant amount of disease will not occur, is not one of these circumstances because of the extent of the training provisions necessary to reduce exposures, the lack of ability to reach all of the potentially exposed populations, and the wide-spread use of chrysotile asbestos. Training would be relatively better achieved in the manufacturing sector where the employed population is relatively more stable as compared to the construction sector wherein many workers are transient employees. Because of the transient nature of the workforce and the cost involved to train workers, there is a tendency to not train those who will only be employed for short periods of time.

5.352 In terms of studies related to the feasibility of using chrysotile asbestos in a “controlled manner” OSHA health compliance data may offer some insight. The United States has an asbestos standard that includes training requirements that are enforceable by law, and violations of which are punishable by monetary penalties. Yet, improper work practices (presumably a reflection of lack of training) and violations of the permissible exposure limit continue to be identified. Since 1980, OSHA compliance officers have identified almost 14,000 violations among establishments for failure to comply with provisions of its asbestos standard. During the recent 3-year period of 1996-98, over 4,000 violations have been cited. Because of the small number of OSHA compliance staff in relation to the number of facilities in the United States, it has been estimated that compliance officers are able to visit industrial workplaces an estimated one time in every 84 years. Thus, the non-compliance with provisions of the United States asbestos standard as identified by its compliance officers, represents a “tip of the iceberg” in identifying non-compliance with provisions of the standard that are thought necessary to control asbestos exposure in the workplace. As difficult as this situation is in the occupational setting, I am inclined to believe that training leading to “controlled use” of chrysotile would be even more difficult to achieve outside of the occupational setting.

Dr. Musk:

5.353 As above in my opinion controlled use of chrysotile is theoretically feasible but probably not practically possible. The second question part of the question is not in my area of expertise.

5.(c) Can controlled use, when properly applied, reduce exposure levels to chrysotile fibres to below 0.1 f/ml? Can controlled use provide the assurance that there will be no peaks above this figure for any type of use of high-density chrysotile products? For workers or other persons exposed to chrysotile asbestos at this level, can you quantify the risk?

Dr. de Klerk:

5.354 See Table in paragraph 5.197 above.

Dr. Henderson:

5.355 When properly applied, controlled use in specific situations (e.g. friction products manufacture) can reduce airborne chrysotile concentrations to < 0.1 f/ml for most of
the time: e.g. as mentioned in my answer to Question 5(a), at the Australian friction products factory in Victoria, 84 per cent of 461 samples (1992-1997) showed an airborne fibre concentration < 0.1 f/ml; however, the remainder showed airborne fibre levels above this figure, although the concentrations were still low. This also demonstrates that one cannot always guarantee that fibre concentrations will never exceed 0.1 f/ml, even in highly regulated circumstances such as the manufacture of high-density chrysotile products. For the reasons discussed in preceding sections of this report, it is not possible to quantify the risks (e.g. lung cancer or mesothelioma) from occasional peak exposures > 0.1 f/ml because these risks necessarily depend upon fibre type, the intensity of the exposure, and the frequency and duration of exposure; in addition, quantification of the risk necessarily involves extrapolation from a linear dose-response model, which is the subject of dispute, because there are no observational data on risks from low-level chrysotile exposure.

5.356 Tables 12 and 13 above (see my response to Question 1(d)) give some estimates of lung cancer and mesothelioma risk at various levels of exposure to chrysotile.

5.357 Finally, occasional bursts or peak concentrations of fibres can be released from buildings by catastrophic and uncontrollable events that include, for example, destruction by fire [244-247], earthquake, or explosions [246], including war (e.g. bombing of cities). Although disasters like these are infrequent — and the fall-out from fires is probably of little consequence to nearby residents or the public in general [247] — they pose a potential health risk for some occupational groups likely to encounter asbestos fall-out and debris more often, such as fire fighters and those involved in clean-up operations.

Dr. Infante:

5.358 Since I believe that “controlled use” is a misnomer in relation to potential for exposure to asbestos, I prefer to answer this question in the context of whether a stringent standard for the control of asbestos in the workplace can reduce exposures to below 0.1 f/cc. In many situations, when standards are properly applied, or adhered to, it is possible to maintain exposures below 0.1 f/cc. However, in many situations today, even in the presence of a stringent standard to control exposure, the 0.1 f/cc level for asbestos is exceeded, specified work practices and housekeeping provisions that are required to reduce exposures are not adhered to, communication of hazards is not provided, and appropriate personal protective equipment is not used. As a result, both average exposures and peak exposures above 0.1 f/cc will occur. With regard to high density chrysotile products specifically, over 2,000 violations of the OSHA asbestos standard have been identified in standard industrial classification (SIC) sectors where potential for exposure to chrysotile asbestos would occur, namely: wrecking and demolition work, asbestos products manufacturing, roofing, siding and sheet metal work, manufacturing of gasket, packing and sealing devices and automotive repair shops.

5.359 In addition to the above examples, in my opinion, many work practices are simply not good enough in some situations to reduce exposures below 0.1 f/cc. For example, even though asbestos cement products may be “pre-sized” in manufacturing, cement sheets need to be modified, drilled, cut and ground to fit them into specific areas. Although wetting may be used in some of these situations for cutting asbestos cement, the material dries and the dust becomes airborne resulting in uncontrolled asbestos dust being released into the work environment. Even in such situations where workers are provided with respirators to prevent exposure, the respirators supplied for the situation are often the wrong ones. In addition, respirator effectiveness depends upon how well the masks fit the face, how often they are replaced, cleaned and repaired and how well the employees are trained to select, clean and repair their respirators. In spite of rules requiring good respirator practices, employers often simply do not comply with the regulations.

5.360 In a more general sense, non-compliance with regulations that could assist in the control of asbestos to exposure levels below 0.1 f/cc is often the result of human error, (not recognizing that asbestos is present, not understanding the requirements to protect workers from exposure in specific situations), willful non-compliance, poor judgement, ac-
cidents, inability to adequately reach the potentially exposed population with proper train-
ing and education of the health hazard, etc. Based on my experience in occupational health
and the results of OSHA compliance data and the literature, it is my opinion that it is
impossible to use asbestos and assure that exposures can be kept below 0.1 f/cc in a great
number of situations.

5.361 Regarding health risks from chrysotile exposure to 0.1 f/cc, as mentioned in my
answer to Question 3, the risk assessment conducted by Stayner et al. (1997) based on the
study by Dement et al. (1994) provides a good estimates of disease risks from exposure to
chrysotile asbestos which are not much different from those provided from other studies.
The analyses estimate that exposure to 0.1 f/cc of chrysotile asbestos for an occupational
lifetime of 45 years, e.g., 4.5 f/cc-years of cumulative exposure, results in five extra deaths
from lung cancer and two extra deaths from asbestosis per 1000 workers. The analysis did
not include mesothelioma because there were too few deaths from this cause in the study to
provide estimates of risk. I have not seen risk assessments for mesothelioma based on
chrysotile exposure. Thus, I cannot provide a quantitative estimate of risk for this cause of
death.

5.362 Of these diseases, the risk of developing asbestosis is most likely underestimated.
This underestimation is a reflection of the fact that most risk assessments are based on
mortality studies and individuals diagnosed with asbestosis often die from other causes.
For example, in the study by Finkelstein (1982), of 24 individuals with certified asbestosis,
who had died, 14 (58 per cent) died from lung cancer or mesothelioma and three (12.5 per
cent) died from ischemic heart disease. Thus, 70 per cent of the individuals with asbestosis
who died would not have been identified from a mortality study of this population of
exposed workers. Asbestosis may also be underestimated in a study because it may be
confused clinically with other non-malignant lung diseases. Therefore, the risk of death
from asbestosis based on mortality studies may usually be underestimated. I am not fa-
miliar with quantitative estimates of risk from other pathologies related to chrysotile as-
bestos.

Dr. Musk:

5.363 The question is not within my area of expertise except that risks from exposure
could be calculated if exposure levels are known.

5.(d) Is it possible to control the risks to human health presented by exposure to high-
density chrysotile products, in particular chrysotile-cement, throughout the life-cycle of
the product? Is controlled use a feasible and practicable option in everyday life for work-
ers who are exposed occasionally to potentially high levels (“peaks of exposure”) of
chrysotile asbestos (such as plumbers, electricians, maintenance, repair, insulation, demo-
lition, waste management and “handyman” type persons)?

Dr. de Klerk:

5.364 See my response to Question 5(a).

Dr. Henderson:

5.365 From my perspective, these questions are crucial to the dispute before the WTO.
In my opinion, the answer to both questions is NO. I do not see how asbestos in place — or
existing chrysotile products — can be controlled at every point of end-use. For example,
EHC 203 indicates repeatedly that exposure is most likely to affect workers involved in
building construction or demolition; this is because of the large number of these workers
involved in myriad different tasks; in addition, these various workers represent a non-
cohesive workforce, many self-employed or working as part of “small business”.

5.366 My perception is also based on studies such as that reported by Kumagai et al.
[4] on Japanese workers involved in the repair of asbestos-cement pipes, where fibre con-
centration for fibres > 5 µm in length ranged from 92 f/ml inside the hole where this work took place (range 48-170 f/ml) and up to 15 f/ml outside the hole; the final sentence of the Abstract for this Report indicates that only about 18 per cent of the workers used a protective respiratory device. In addition, a survey in Finland found occasional high fibre concentrations inside personal protectors during asbestos removal work, suggesting that these devices are not always effective.

5.367 Another factor that merits consideration in some societies such as Australia, is poor worker compliance with controls. For example, I am aware of non-compliance in the use of protective equipment (despite penalties), because respiratory protective devices may be cumbersome and uncomfortable — especially in hot climates like Australia — with skin irritation from sweat accumulating within them.

5.368 From the literature cited throughout this report and the reasons discussed, it seems clear that there is a broad consensus among experts that controlled use of chrysotile (or other varieties of asbestos) is not feasible in practice for certain worker groups, notably those involved in construction trades (e.g. see EHC 203).

Dr. Infante:

5.369 As mentioned in my responses to several previous questions, in my opinion, it is not possible to control the risk to human health posed by exposure to chrysotile asbestos throughout its life cycle. “Controlled use” is not a feasible and practical option in everyday life for workers. While “controlled use” is relatively more achievable in the manufacturing sector, violations of regulations enforceable by monetary fines still occur. In the construction sector, control of exposure to asbestos is much more difficult to achieve as compared to manufacturing. Workers who have jobs as plumbers, electricians, maintenance personnel, repairmen, insulators, demolition, waste management and handymen will most likely experience intermittent peak exposures to asbestos. These exposures result from lack of awareness of the hazard, lack of recognition of the hazard, lack of personal protective equipment, lack of training on the maintenance of the protective equipment, etc. as mentioned in responses to other questions above.

Dr. Musk:

5.370 In my opinion controlled use is probably not practically possible but this is not an area of my expertise.

5.371 Is it possible to control the risks to human health presented by exposure to high-density chrysotile products, in particular chrysotile-cement, in non-occupational circumstances, such as intervention on these products by private individuals (cutting, sawing, removal, etc.)? Is controlled use a feasible and practicable option for this category of the population?

Dr. de Klerk:

5.372 As a follow-on to my answer to the preceding question, my answer to both of these questions is also NO. However, the risks from occasional or infrequent interventions on chrysotile-only products (e.g. by home “handymen”) - although not quantifiable because of absence of data - must be very small for lung cancer and mesothelioma, and non-existent for asbestosis.

Dr. Infante:

5.373 As difficult as it is to control exposure to chrysotile asbestos during intervention with cement products in the occupational setting, it is much more so in non-occupational
circumstances because there is no effective means of identifying the potential population at risk. As a result, there is a lack of awareness of the hazard, lack of recognition of the hazard, lack of personal protective equipment, lack of training on the maintenance of the protective equipment, etc. as mentioned in responses to other questions above. Therefore, in my opinion, it is not possible to limit exposure in such circumstances.

Dr. Musk:

5.374 Controlled use is probably not practically possible, in my inexpert opinion.

Question 6:

The parties disagree as to the relative pathogenicity of chrysotile fibres vs. substitute fibres, in particular cellulose fibres, para-aramid fibres, glass fibres and polyvinyl alcohol (PVA) fibres. Canada considers that, overall, substitute fibres have not been demonstrated to be less toxic than chrysotile fibres, and that, by banning chrysotile, France has replaced the “much studied but nonetheless undetectable risk associated with modern uses of chrysotile with the unknown, and perhaps greater risk associated with the use of substitute fibres” [Premier exposé oral du Canada, paragraph 90]. On the other hand, the European Communities argues that none of the substitute products -fibrous or non-fibrous- for chrysotile, and in particular none of the substitutes for chrysotile-cement, has been classified as a proven carcinogen to humans; hence, overall, substitute products present less of a risk to human health than chrysotile asbestos [see Deuxième soumission écrite, pp. 10-15].

6.(a) Is it correct to argue that non-fibrous substitutes are safe or less hazardous than chrysotile and that concern over potential health risks should be focused on fibrous ones? In this context, could you elaborate upon the “effet fibre” (“fibre effect”) of substitute fibres? What general conclusions can be drawn as to the respirability and biopersistence of substitute fibres?

Dr. de Klerk:

5.375 As outlined above, the pathogenicity of fibres is related to their size, shape, durability and quantity. Thus, all the parts to this question can be answered in the same way. The argument here is whether it is safer to stick with the well-studied chrysotile that has a semi-quantifiable and definite carcinogenic risk, than to use other substances which have the potential to increase risk in an unquantifiable way, ie. the “better the devil you know” principle. For example, para-amid fibres have recently been classified by IARC in Group 3, that is, ‘not classifiable as to its carcinogenicity’.

5.376 Substitutes need to be compared to chrysotile in terms of the parameters listed above, namely, size, shape, durability and quantity. These are all properties of fibres and therefore “concern should be focused on fibrous substitutes”. Substitute fibres can then be compared with chrysotile on the four parameters. I am inexpert in commenting on the “extent to which hazardous concentrations can be controlled” but it is my understanding that all four substitutes mentioned involve less dusty operations than equivalent ones involving chrysotile. As far as the other three parameters are concerned: all four substitutes except glass fibre produce a larger proportion of non-respirable fibres than chrysotile does, but respirable fibres are similar for all substances and glass fibre is the least durable; all four except cellulose are less durable than chrysotile, but cellulose is much less dusty and has also been in use for a long while without evidence of ill effect.

5.377 On balance, the substitute fibres appear less likely to cause adverse effects (from their fibres) than chrysotile.

Dr. Henderson:

5.378 Current thinking on this issue indicates that the bio-hazards — specifically the carcinogenic risks - of all fibres are determined by the three Ds: dose, fibre dimensions and
durability (bio-persistence) [248-250]. Therefore, substitute materials that have engendered most concern are fibrous materials as opposed to non-fibrous substances (non-fibrous materials may or may not show different effects in terms of toxicology, but this discussion focuses on carcinogenic risks). For example, refractory ceramic fibres (RCF) are a cause for concern [251] because they may have dimensions similar to those of the amphibole varieties of asbestos and RCF have been reported to induce mesothelioma in experimental animals.

5.379 In a 1995 review, de Vuyst et al. [248] concluded that:

“The group of man-made mineral or vitreous fibres (MMMFs or MMVFs) includes glass wool, rock wool, slag wool, glass filaments and microfibres, and refractory ceramic fibres (RCFs). Experimental observations have provided evidence that some types of MMVF are bioactive under certain conditions. The critical role of size parameters has been demonstrated in cellular and animal experiments, when intact fibres are in direct contact with the target cells. It is, however, difficult to extrapolate the results from these studies to humans since they bypass inhalation, deposition, clearance and translocation mechanisms. Inhalation studies are more realistic, but show differences between animal species regarding their sensibility to tumour induction by fibres. Fibre biopersistence is an important factor, as suggested by recent inhalation studies, which demonstrate positive results with RCF for fibrosis, lung tumours and mesothelioma. There is no firm evidence that exposure to glass-, rock- and slag wool is associated with lung fibrosis, pleural lesions, or nonspecific respiratory disease in humans. Exposure to RCF could enhance the effects of smoking in causing airways obstruction. An elevated standard mortality ratio for lung cancer has been demonstrated in cohorts of workers exposed to MMVF, especially in the early technological phase of mineral (rock slag) wool production. During that period, several carcinogenic agents (arsenic, asbestos, polycyclic aromatic hydrocarbons (PAH)) were also present at the workplace and quantitative data about smoking and fibre levels are lacking. It is not possible from these data to determine whether the risk of lung cancer is due to the MMVFs themselves. No increased risk of mesothelioma has been demonstrated in the cohorts of workers exposed to glass-, slag- or rock wool. There are in fact insufficient epidemiological data available concerning neoplastic diseases in RCF production workers because of the small size of the workforce and the relatively recent industrial production” [abstract].

5.380 In a 1999 review published in French, Boillat et al. [250] came to similar conclusions:

“The group of man-made mineral fibres includes slagwool, glasswool, rockwool, glass filaments and microfibres, as well as refractory ceramic fibres. The toxicity of mineral fibres is determined by several factors such as the diameter (< or = 3-3.5 microns) and the length of the fibres (< 100 microns), their biopersistence, which is much shorter for man-made mineral fibres than for asbestos fibres, their physicochemical structure and surface properties, and the exposure level. The chemical composition of the various types of man-made mineral fibres depends directly on the raw material used to manufacture them. While naturally occurring fibres are crystalline in structure, most man-made mineral fibres are amorphous silicates combined with various metal oxides and additives. Observations using intracavitary administration have provided evidence that some types of man-made mineral fibres are bioactive in cellular and animal experiments and may induce lung tumours and mesothelioma. It is difficult to extrapolate these results to humans since they bypass inhalation, deposition, clearance and translocation mechanisms. Inhalation studies show more realistic results but differences are observed between animal species regarding their sensibility to tumours. There is no firm evidence that exposure to various wools is associated with lung fibrosis, pleural lesions or nonspecific respiratory disease in humans. A possible exception may be mentioned for refractory ceramic fibres. A slightly elevated standard mortality ratio for lung cancer has been documented in large cohorts
of workers (USA, Europe and Canada) exposed to man-made mineral fibres, especially in the early technological phase. It is not possible to determine from these data whether the risk of lung cancer is due to the man-made mineral fibres themselves, in particular due to the lack of data on smoking habits. No increased risk of mesothelioma has been demonstrated in these cohorts. Epidemiological data are insufficient at this time concerning neoplastic diseases in refractory ceramic fibres” [abstract].

5.381 In one study on RCF, Glass et al. [252] reported that:

“In recent inhalation experiments conducted with both rats and hamsters … at the highest dose tested … there was an increased incidence of tumours in both species. Lower doses were only examined in the rat and at these doses there was no significant excess of lung tumours. Epidemiological investigations of workers engaged in the manufacture of ceramic fibres have shown a small excess of pleural plaques. This phenomenon is being further investigated but could be due to confounding exposures. The populations available for study are small and their exposures fairly short, but it is considered prudent that they should remain under surveillance for some time to come. This is despite the fact that present exposures in the ceramic fibre industry are low (< 1 f/ml) and are being reduced” [abstract].

5.382 Okayasu et al. [253] also found that RCF-1 fibres were less cytotoxic and mutagenic than chrysotile:

“Cytotoxicity and mutagenicity of tremolite, erionite and the man-made ceramic (RCF-1) fibre were studied using the human-hamster hybrid A(L) cells. Results from these fibres were compared with those of UICC Rhodesian chrysotile fibres. The A(L) cell mutation assay, based on the S1 gene marker located on human chromosome 11, the only human chromosome contained in the hybrid cell, has been shown to be more sensitive than conventional assays in detecting deletion mutations. Tremolite, erionite and RCF-1 fibres were significantly less cytotoxic to A(L) cells than chrysotile. Mutagenesis studies at the HPRT locus revealed no significant mutant yield with any of these fibres. In contrast, both erionite and tremolite induced dose-dependent S1- mutations in fibre-exposed cells, with the former inducing a significantly higher mutant yield than the latter fibre type. On the other hand, RCF-1 fibres were largely non-mutagenic. At equitoxic doses (cell survival at approximately 0.7), erionite was found to be the most potent mutagen among the three fibres tested and at a level comparable to that of chrysotile fibres. These results indicate that RCF-1 fibres are non-genotoxic under the conditions used in the studies and suggest that the high mesothelioma incidence previously observed in hamster may either be a result of selective sensitivity of hamster pleura to fibre-induced chronic irritation or as a result of prolonged fibre treatment. Furthermore, the relatively high mutagenic potential for erionite is consistent with its documented carcinogenicity” [abstract].

5.383 An important consideration is that fibre dimensions for some substitute materials (e.g. fibreglass) can be varied according to the manufacturing processes employed, so that they can be designed to have fibre characteristics and dimensions different from asbestos, or similar to asbestos: as one example, the dimensions of fibreglass can be varied and when implanted into experimental animals, fibres of the “right” size can induce mesothelioma.

5.384 For this reason, testing of substitute materials with fibre dimensions similar to those of asbestos should be carried out before these materials are used in products available to the general public (e.g. testing for toxicology, clastogenicity, DNA strand breaks, mutagenicity and free radical generation using in vitro systems and/or testing in vivo — such as the intraperitoneal test in rats) [248, 249, 251-256].
5.385 Nonetheless, it is my perception that lumping all substitute fibres together is as erroneous as lumping amphibole and chrysotile fibres into the same category. For example, RCF are the subject of continuing concern, but other substitute fibres such as cellulose fibres, para-aramid fibres and polyvinyl alcohol (PVA) fibres appear to be different from chrysotile, in terms of fibre dimensions and especially bio-persistence.

5.386 NICNAS 99 summarizes these considerations in the following terms:

“Any substitution of chrysotile should be with a less hazardous substance. There has been ongoing debate regarding the health effects of alternatives, such as synthetic mineral fibres (SMF), natural organic fibres and synthetic organic fibres.

In general, less data on health effects of alternative materials (in comparison to asbestiform fibres) are available and because of this, it is difficult to make an assessment of the pathogenicity and potential carcinogenicity of many substitutes.

Although not the only determinant of potential pathogenicity, fibre dimensions (length, width and aspect ratio) are considered to be [some] of the most important factors associated with carcinogenic (lung cancer and mesothelioma) potential ... The commonly accepted ‘peak hazard’ dimensions ... are > 5 µm long (length) and < 3 µm wide (diameter).

The most commonly used alternatives in Australia (and overseas) for friction materials are aramid fibres, attapulgite, fibreglass, refractory ceramic fibres (RCF), semi-metallics, mineral wool, steel wool, cellulose, titanate fibres and wollastonite, and for gaskets are glass fibre, carbon fibre and aramid fibre.

... It should also be noted that ... differences in fibre length, diameter and surface properties may lead to entirely different toxicological profiles.

A recent report by EC concludes that the available data are generally supportive of the conclusion that PVA, cellulose, p-aramid, glass wool and slag wool are likely to be safer in use than chrysotile. However, RCFs are the subject of ongoing concern ...” [p 125].

Dr. Infante:

5.387 I have not seen any information that indicates that non-fibrous substitutes for chrysotile are carcinogenic, or cause non-malignant lung diseases. I would focus attention on the fibrous substitutes in terms of their ability to reach lung tissue (respirability) and their known toxicity. Clearly, if the substitute fibres are not respirable, there is little concern for their “potential” to cause lung diseases. (Attention would then focus on adverse effects from exposure to the skin and eyes.) If the substitute fibres are respirable, then attention needs to focus on their toxicity relative to that of chrysotile in their ability to cause lung cancer, non-malignant lung diseases and mesothelioma.

5.388 The data I have reviewed in this area of investigation appear to indicate that polyvinyl alcohol fibres (PVA) are mostly in the range of 10-16 microns in diameter and hence are too large to respirable and thus cause lung disease. In terms of biopersistence, if they were respirable, they would degrade very slowly. Para-aramid fibres are also generally 10-12 microns in diameter and they also would have little chance of being respirated. These fibres, however, contain fibrils of about 0.2 microns in diameter that can be liberated with high energy input and they would be respirable. P-aramid fibrils greater than 5 microns in length are less biopersistent than chrysotile fibres greater than 5 microns in length (Searl, 1997). Data for dimensions of cellulose fibres show a median length and diameter of about 7.5 and 1.50 microns, respectively, which indicates that they are in the respirable
range (Muhle et al. 1997). In terms of biopersistence, cellulose fibres had a mass half time in the rat lung of 72 days and bioaccumulated in the lungs. Data on the distribution of glass fibres indicates that the majority are in the respirable range, but the fibre size distribution of glass filaments indicates that a small portion are in the respirable range. Glass fibres are less biopersistent than chrysotile fibres. In general, in terms of the combination of respirability and biopersistence, with the exception of cellulose fibres, it appears that the substitute fibres would have less bioaccumulation in the lung than chrysotile fibres because they are either less respirable, or they are not as biopersistent.

5.389 The role of biopersistence in relation to toxicity is complicated. Chrysotile fibres are less biopersistent than amphibole fibres, yet experimental data demonstrate a similar potency for lung cancer, mesothelioma and fibrosis.

Dr. Musk:

5.390 I agree with the Canadian argument philosophically. However there is no evidence that I know of carcinogenicity of substitutes in animal studies and only rockwool has been associated with increased lung cancer risk in epidemiological studies.

6.(b) To what extent do physical characteristics and chemical properties of substitute fibres determine their toxicity? Is it correct to say that man-made fibre substitutes are superior to natural fibre ones in terms of the extent to which exposure to hazardous concentrations can be controlled during the various stages of production? Is your opinion based on one or more of the following evidence: (i) chemical/physical characteristics of the substitute fibres, (ii) epidemiological data, (iii) in vitro evidence, (iv) in vivo evidence?

Dr. de Klerk:

5.391 See my response to Question 6(a).

Dr. Henderson:

5.392 These questions are covered to a large extent in my answer to the preceding question. Again, dose, fibre dimensions (including surface chemistry) and bio-persistence appear to represent the properties that determine the toxicity and carcinogenicity of fibres of any type. The issue of controllability during various stages of production is an engineering and industrial question, and falls outside my expertise.

5.393 My opinions concerning the potential bio-hazards of these fibres are based on the physical characteristics of the substitute fibres, in vivo evidence (tumour induction in experimental animals) and in vitro studies (mutagenicity analogous that reported for other known carcinogens). To the best of my knowledge, there are no large-scale epidemiological studies on cellulose fibres, para-aramid fibres or PVA fibres; two large epidemiological investigations on slag wool fibres in both Europe and the United States did show an increase in the relative risk for lung cancer among the production workers, but this effect may have been explicable by other confounding factors involved in the manufacture of these materials.

Dr. Infante:

5.394 As a matter of general toxicology, I would focus concern on the “potential” for adverse health effects from any fibrous material of dimensions and aerodynamic diameter that will result in its being respirable. This is discussed in my response to Question 6(a) above. The toxicity of the substitute fibres and whether that information was determined on the basis of epidemiological or toxicological evidence is discussed in Question 6(c). The role of the chemical properties of fibres to induce cancer is not clear to me. The nature of the production process makes the substitute fibres more amenable to control than asbestos fibres.
Dr. Musk:

5.395 I do not know: but it is my broad understanding that the physical and chemical properties of the substitute fibres suggests less risk of disease.

6.(c) The parties focus part of their arguments on cellulose fibres, para-aramid fibres, glass fibres and polyvinyl alcohol (PVA) fibres. What evidence exists with respect to the toxicity and health risks of these substitutes? Does the existing evidence suggest that these products are less/equally/more toxic than chrysotile asbestos fibres?

Dr. de Klerk:

5.396 See my response to Question 6(a).

Dr. Henderson:

5.397 In experimental studies on para-aramid fibres in comparison to chrysotile, Warheit et al. [12, 257] found that p-aramid is bio-degradable in the lungs of exposed rats, with faster clearance than long chrysotile fibres which showed greater bio-persistence. In their 1996 study, these authors [12] found that:

"...p-aramid is biodegradable in the lungs of exposed rats; in contrast, the clearance of long chrysotile fibres was slow or insignificant, resulting in a pulmonary retention of long chrysotile asbestos fibres. The dimensional changes of asbestos fibres as well as the pulmonary cell labelling data indicate that chrysotile asbestosis fibres may produce greater long-term pulmonary effects when compared to inhaled para-aramid fibrils" [abstract].

5.398 The present status of knowledge has been summarized by Harrison et al. [19] in a recent review of the comparative hazards of chrysotile and its substitutes:

"There are now practicable substitutes for the major remaining uses of chrysotile. Although lack of a full health and toxicological data set precludes a comprehensive assessment of the safety of substitute fibers, the application of basic principles of fiber toxicology enables a pragmatic decision to be made on the relative safety of potential substitutes. Our judgement is based on relative considerations of the intrinsic properties of fibers, on the pathogenicity of chrysotile in comparison with that of substitute fibers, and on the potential for uncontrollable exposures. The three parameters of dose, dimension (especially diameter), and durability are key to determining the differential hazards. Due consideration of these factors leads us to the following conclusions regarding chrysotile and its main substitutes.

Chrysotile per se can cause lung cancer and asbestosis; it is less clear that chrysotile alone can cause mesothelioma in humans, and indeed it may not, whereas tremolite and other amphiboles certainly can do so. There is no definitive evidence for a threshold exposure level for lung cancer induction, although some studies suggest that a threshold does exist.

The intrinsic hazardous properties of chrysotile can never be ‘engineered out’, and the potential for harm will always remain. Prevention of ill health will thus always rely on the control of exposure, something that history has shown cannot be guaranteed.

Unlike chrysotile, substitute fibres can often be designed or selected to have particular characteristics. Criteria for the substitution of asbestos by other fibers include a) the substitute fibers are not in the respirable range, do not readily
fibrillate, and/or are less durable than chrysotile; b) other materials that must be incorporated into the replacement product do not, in combination with the replacement fiber, produce more harm overall than chrysotile alone; c) the replacement product has an equivalent or acceptable performance; and d) substitution would result in overall lower fiber exposures during manufacture and use and disposal, taking into account likely exposures. The same general principle can be applied to substitute fibers others than those considered here.

We judge that PVA fibers will pose less risk than chrysotile because they are generally too large to be respirable, do not fibrillate, and the parent material causes little or no tissue reaction. Aramid fibers have a reduced potential for exposure when compared to chrysotile because they are generally of high diameter and the production of respirable fibrils is energy intensive. The fibrils are less pathogenic than chrysotile, are less biopersistent, and are biodegradable. Cellulose has the benefit of long experience of use in a variety of industries without having raised significant concern. The potential for the generation of respirable fibers seems to be less than is the case for chrysotile, although fibrillation is possible. Cellulose is durable in the lung, and its biological properties should therefore be investigated further. However, exposure levels for current uses are low, and it is biodegradable in the environment.

We believe that the continued use of chrysotile in asbestos-cement products is not justifiable in the face of available and technically adequate substitutes. Likewise, there seems to be no justification for the continued residual use of chrysotile in friction materials” [pp 610-611].

5.399 From known past uses of asbestos and surveys of current uses in Australia, it is evident that alternatives have replaced chrysotile to a large extent for the following products [NICNAS 99, p 111]:

"Products where chrysotile use has been completely replaced:

- Cement sheeting, tubes and piping.
- Roofing tiles.
- Textiles.
- Fibre insulation.
- Railway brake blocks.
- Brake disc pads in new automotive vehicles (only 1 new vehicle model was identified as being supplied with asbestos pads in Australia).

Products where a major proportion of chrysotile use has been replaced:

- Clutch facings (in automotive vehicles and industrial machinery e.g. tractors, centrifuge drives).
- Brake disc pads (in older taxi and courier vehicles, and industrial machinery).
- Gaskets, such as spiral wound and head gaskets.
- Washers.
- Packing material.
- Rotor blades (e.g. in high vacuum pumps)".

5.400 It is notable that chrysotile is no longer used for brake linings in new passenger cars produced in Australia by most manufacturers, having been replaced by substitute materials: NICNAS 99 comments that:

"Out of 26 companies, 25 stated that they are using non-asbestos original equipment in all current models. One company (Ford Motor Australia) reported that
they are still using asbestos parts in two current models: asbestos head gaskets for the Econovan and asbestos rear brake linings for the Ford utility. Ford Australia introduced non-asbestos components for their most popular models (e.g. Laser, Falcon and Fairlane) between 1989 and 1995. Other current models manufactured by Ford have been asbestos-free since their introduction. ... Asbestos parts are imported by 6 of the 26 companies (BMW, Ford, Mazda, Mitsubishi, Nissan and Toyota) with five companies using asbestos parts for superseded vehicles and one company (Ford Australia) using asbestos parts in superseded and current models ... the majority of the vehicle manufacturing companies stated that they have had policies in place in regard to not using asbestos components in new vehicles for the last 5 to 10 years” [p 22].

5.401 This trend to use of brake linings free of asbestos is shown in the following Table 15 — in comparison to the usage of asbestos brake linings — between 1994 and 1998 (asbestos-containing brake linings appear to be used primarily on older and superseded vehicle models).

<table>
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<tbody>
<tr>
<td>Asbestos brake linings, passenger cars</td>
<td>492,295</td>
<td>47,735</td>
<td>43,087</td>
<td>771,182</td>
<td>(548,692)</td>
</tr>
<tr>
<td>Non-asbestos brake linings, passenger cars</td>
<td>70,109</td>
<td>321,472</td>
<td>485,812</td>
<td>2,084,963</td>
<td>(4,057,143)</td>
</tr>
</tbody>
</table>

Source: NICNAS 99.

Dr. Infante:

5.402 There is no information to my knowledge that cellulose fibres, para-aramid fibres or ployvinyl alcohol (PVA) fibres are carcinogenic. Cellulose fibres have not been studied experimentally for carcinogenicity. It is noteworthy, however, that cellulose has been used in the paper industry for hundreds of years and to date an elevated risk of death from lung cancer and mesothelioma has not been observed. Excess incidences of pharyngeal and/or laryngeal cancers were reported in two studies, but these observations have not been corroborated in other studies (IARC, 1987). Wood dust is associated with sinonasal cancer, but not with lung cancer or mesothelioma. A relatively greater risk appears to be associated with hard woods as compared to soft woods, which suggests that the cellulose may not be the primary factor in the induction of these cancers. Workers exposed to cotton dust also do not demonstrate an excess of lung cancer or mesothelioma even though they develop byssinosis. The debate as to whether this disease is due to cotton dust per se, or to contaminants of the cotton fibre, however, is not resolved.

5.403 Para-aramid fibrils have been studied for carcinogenicity in experimental animals by inhalation and by intra-peritoneal injection. No cancer response was observed. As concluded by IARC (1997), there is inadequate evidence for the carcinogenicity of para-aramid fibrils in experimental animals. The carcinogenicity of para-aramid fibrils has not been evaluated in humans. Likewise, IARC (1987) concluded on the basis of its review of animal cancer tests that there is no evidence for the carcinogenicity of PVA fibres. PVA fibres have not been evaluated for carcinogenicity in humans. It is also my opinion that there is no evidence that these fibres present any risk of cancer to humans.

5.404 With regard to glass fibres, IARC (1988) concluded there was sufficient evidence for the carcinogenicity of glass wool in experimental animals and that there was inadequate evidence for the carcinogenicity of glass wool to humans. Subsequent to the IARC (1988) review, my colleagues and I have reviewed the toxicological and epidemiological
studies related to exposure to glass fibres. In our opinion, there is conclusive evidence from implantation and inhalation studies that glass fibres are carcinogenic in experimental animals (Infante et al. 1994). Studies of workers exposed to glass fibres also demonstrate a significantly elevated risk of death from lung cancer. It is our interpretation of these studies that employment in the manufacturing of glass fibres carries with it an elevated risk of death from lung cancer. Is it proven beyond a doubt through epidemiological study that fibrous glass is a human carcinogen? In my opinion, it is not. However, given the positive animal cancer test results, knowledge that these fibres can be inhaled and retained in the lungs, evidence that workers employed in the manufacturing of these fibres die at a significantly elevated rate of lung cancer, it is my opinion that it is more likely than not that glass fibres are carcinogenic to humans and that employment in this industry carries with it an elevated risk of death from lung cancer.

5.405 It is also my opinion that glass fibres are not as potent as chrysotile asbestos in causing disease. With regard to the capability of glass fibres to cause lung cancer, I have previously published the opinion that on a fibre-per-fibre basis, glass fibres may be as potent or even more potent than asbestos in causing lung cancer. This opinion was based on epidemiological studies which generally demonstrated 10 per cent to 20 per cent elevation in the relative risk of lung cancer (the 1987 study of Canadian workers by Shannon demonstrated a 2-fold risk) as a result of exposures to glass fibres that were reported to be fairly low. Within the past year, however, I have had the opportunity to discuss occupational exposures during glass fibre manufacturing with workers formerly employed at the Canadian facility that manufactured fibrous glass. According to several workers, they were also exposed to crystalline silica many times over the permissible limit through the dumping of sand into the hopper that was used to feed the furnace for melt down. They were exposed to asbestos that lined the furnace when they removed the insulation from the oven doors by hand, or chiselled it away in the absence of respiratory protection; they would then add water to asbestos fibre to make “asbestos mud” that was applied to the oven doors by hand, or by trowel. Workers were so uninformed of the hazard that they sometimes would throw “asbestos mud balls” at each other. The workers at this facility were also exposed to phenol formaldehyde resin that was used as a binder for the glass fibres; they were also exposed to tar that was applied to paper that was then applied to the glass fibre pack. Exposures to glass fibres only is mentioned in the study of these workers that was published by Shannon (1987).

5.406 Furthermore, there is less evidence from epidemiological studies that exposure to glass fibres is associated with a pneumoconiosis as compared to the data for chrysotile asbestos exposure and asbestosis. There is no evidence that exposure to glass fibres is associated with mesothelioma. For the few cases of mesothelioma that have been identified among workers exposed to glass fibres to date, there is claim that they also had been exposed to asbestos fibres. Therefore, it is difficult to attribute these cases of mesothelioma to the glass fibre exposures. Therefore, the totality of disease related to chrysotile asbestos exposure would be greater than that related to a similar amount of exposure to glass fibres.

5.407 In conclusion, only one of the fibres (glass fibres) that may play any significant role in substitution for chrysotile asbestos demonstrates evidence of being carcinogenic. Data for the total toxicity related to these fibres, however, is less than that for chrysotile asbestos fibres. Cellulose fibres have not been tested for carcinogenicity in experimental animals, but epidemiological studies of workers exposed to cellulose in three separate industries, i.e., furniture manufacturing, cotton textile manufacturing and the paper products industry, have not demonstrated an elevated risk of contracting lung cancer or mesothelioma. Regarding non-malignant lung disease among cotton dust exposed workers, it is not known whether the cotton dust per se, or contaminants of the cotton fibre, are responsible for the byssinosis observed in these workers.

Dr. Musk:

5.408 I understand that substitutes have been shown to be less toxic in animals.
5.409 In-place asbestos is widely distributed in industrialized societies and much includes mixtures of chrysotile and amphiboles — although chrysotile has been the predominant type of asbestos used throughout Western Europe for many years (about 94-97%).

5.410 Lung cancer and mesothelioma are the most important bio-hazards from asbestos in place and the continued use of asbestos.

5.411 Because of the prolonged lag-time between exposure and the subsequent development of either lung cancer or mesothelioma, most mesotheliomas in the 1990s and beyond can be attributed to exposures sustained decades before; the mesothelioma “epidemic” predicted for Europe over the next three decades can be attributed to exposures before, during and after the 1960s and 1970s, especially to one or more of the amphibole varieties.

5.412 For the amphibole forms of asbestos and mixtures of asbestos types, a linear dose-response relationship has been found at high levels of exposure; a dose-response relationship with an increase of the relative risk of mesothelioma to > 2.0 has also been observed at low levels of exposure, in the order of 0.5-1.0 fibre-year (which overlaps with non-occupational environmental exposures). No lower threshold dose for mesothelioma induction has been delineated for the amphiboles.

5.413 Chrysotile also has the capacity to induce mesothelioma, although it is less mesotheliomagenic than the amphiboles (my estimate is 1/10th-1/30th).

5.414 Commercial Canadian chrysotile on average contains trace quantities of tremolite, including fibrous tremolite (< 1%).

5.415 Tremolite — a non-commercial amphibole — also has the capacity to induce mesothelioma.

5.416 The carcinogenicity of Canadian chrysotile may be attributable to the trace tremolite content, but it is not possible to separate the dose-response effects for the chrysotile and the tremolite.

5.417 At high levels of exposure to Canadian chrysotile, a linear dose-response relationship has been observed.

5.418 To the best of my knowledge, there are no epidemiological or observational data on dose-response effects of chrysotile only at low levels of exposure.

5.419 No lower threshold dose for the carcinogenic effects of chrysotile has been identified (EHC 203).

5.420 To the best of my knowledge, there are no observational data on the potential carcinogenic effects of inhaled chrysotile when superimposed upon a pre-existing burden of amphiboles ± chrysotile in lung tissue.

5.421 Although the amphiboles are far more potent than chrysotile for mesothelioma induction, this differential in carcinogenicity may be less obvious or absent for lung cancer induction, but this is still the subject of some dispute; chrysotile is associated with a low risk of lung cancer among Canadian chrysotile miners and millers, but the highest risk for lung cancer induction has been observed for South Carolina asbestos textile workers who used Canadian chrysotile almost exclusively.

5.422 A linear dose-response relationship has also been observed for the risk of lung cancer versus cumulative asbestos exposure. Although some authorities favour a linear no-threshold model for lung cancer induction, others suggest that a threshold may exist, but this has not been delineated in numerical terms.
5.423 In contrast, asbestosis is a dose-dependent non-cancerous disorder, with clear evidence of a threshold effect, although the threshold may be lower than previously supposed, at least for histological asbestosis; there is no risk of asbestosis at low levels of chrysotile exposure.

5.424 Although reduction of airborne asbestos fibre concentrations in the mining and manufacturing industries has been achieved, it is too early to evaluate the effects of these reduced exposures, because no epidemiological data are available; however, with reduction of cumulative exposures, a reduction in the incidence of both asbestos-related mesothelioma and asbestos-related lung cancer can be expected.

5.425 The risks from low-level occupational exposure to chrysotile, or from occasional peak concentrations, have not been delineated but are predictably small.

5.426 Carcinogenic hazards from ultra-low levels of atmospheric chrysotile fibres (e.g. simple occupancy of public buildings) appear to be minuscule, negligible or undetectable.

5.427 Therefore, health concerns over chrysotile dust exposure narrow down to a workplace issue.

5.428 There is evidence of an increased incidence of mesothelioma among, say, brake mechanics in Australia exposed to chrysotile derived from brake blocks and lining.

5.429 With the reductions of airborne fibre concentrations in the asbestos mining, milling and manufacturing industries, construction trades workers constitute the group of workers at greatest risk from exposure to asbestos-cement products (e.g. builders, builders’ labourers, carpenters, electricians, plumbers and roofing workers). This group constitutes a large, disparate and non-cohesive workforce for which controlled use of asbestos is not achievable, for the reasons discussed earlier in this report.

5.430 Therefore, chrysotile asbestos should not be used in building materials, because of the hazards imposed by installation, maintenance and removal operations (EHC 203); these risks may be compounded for some groups by catastrophic events affecting buildings — e.g. fires (with a burst of asbestos fibres into the atmosphere and the necessity for clean-up operations), and other disasters.

5.431 Substitutes for chrysotile are available for many applications (e.g. cellulose fibres, para-aramid fibres and polyvinyl alcohol); evidence indicates that these fibres are less bio-persistent than chrysotile and, therefore, national health authorities (EHC 203, NICNAS 99) have recommended phasing out or prohibition of chrysotile whenever safer substitute materials are available.

5.432 Therefore, from a perspective of caution and prudence for occupational health and safety, it follows that chrysotile should either:

(a) Be restricted to only a few and well-defined applications so that it is inaccessible to the great majority of workers and is available for use by only small and cohesive specialized worker groups that can be trained effectively in its controlled use (e.g. analogous to nuclear fuels); in effect, this means that chrysotile should not be used in building products (e.g. high-density fibre-cement materials such as asbestos-cement sheets) or friction products.

OR

(b) It should be made inaccessible to everyone, by prohibition, unless the alternatives pose equal or greater hazards and equal or greater problems with control.
These views are also expressed in EHC 203, wherein it is stated:

“a) Exposure to chrysotile asbestos poses increased risks for asbestosis, lung cancer and mesothelioma in a dose-dependent manner. No threshold has been identified for carcinogenic risks.

b) Where safer substitute materials for chrysotile are available, they should be considered for use.

c) Some asbestos-containing products pose particular concern and chrysotile use in these circumstances is not recommended. These uses include friable products with high exposure potential. Construction materials are of particular concern for several reasons. The construction industry workforce is large and measures to control asbestos are difficult to institute. In-place building materials may also pose risk to those carrying out alterations, maintenance and demolition” … [p 144]

d) The combined effects of chrysotile and other insoluble respirable particles needs further study.

e) More epidemiological data are needed concerning cancer risks for populations exposed to fibre levels below 1 f/ml, as well as continued surveillance of asbestos-populations” … [p 145]

NICNAS 99 sets out a similar set of recommendations:

- “Chrysotile is a known human carcinogen.

- Prudent OHS [occupational health & safety] policy and public health policy favours the elimination of chrysotile wherever possible and practicable.

- The main exposure to Australian workers arises from manufacture, processing and removal of friction products and gaskets. Home mechanics are also exposed during ‘do-it-yourself’ replacement of brake pads/shoes. ... In Australia, chrysotile is no longer used in high density materials such as chrysotile-cement.

- Current overseas experience with the phasing out of chrysotile products indicates that a range of alternatives is available to suit the majority of uses. Good OHS practice dictates that use of chrysotile products should be restricted to those uses where suitable substitutes are not available, and alternatives should continue to be sought for remaining uses”.

Whether the objective of removal of chrysotile from the workplace and the general environment is achievable by enforcement of controlled use for a few restricted applications — or by prohibition — is essentially a societal question and a public health policy issue. For the reasons discussed in this report, a complete ban is more certain to accomplish this objective (paragraph 5.432(b)). Therefore, as a cautious and prudent approach to national occupational health policy, a complete ban is neither unreasoned nor unreasonable, on the balance of prevailing scientific evidence and uncertainties discussed in this report, such a policy seems defensible and, arguably, justifiable as a national health measure. Perhaps it is best to let Bradford Hill have the last words:

“All scientific work is incomplete — whether it be observational or experimental. All scientific work is liable to be upset or modified by advancing knowledge. That does not confer upon us a freedom to ignore the knowledge we already have, or to postpone action that it appears to demand at a given time.”
4. **Endnote by Dr. Henderson**

5.436 Wishing to add two further pertinent references after completing his Report, Dr. Henderson attached the following Endnote. These references deal with the following:

5.437 Clearance of chrysotile fibres from human lung tissue: In the past, the kinetics of chrysotile clearance from lung tissue have been investigated mainly in experimental models using rodents. In an autopsy study published in 1999, Finkelstein and Dufresne [1] investigated clearance of chrysotile from the lung tissue of 72 Quebec chrysotile miners and millers in comparison to 49 control subjects, using regression analyses, with the following findings:

- There was a significant association between the duration of occupational exposure and the tissue burdens of chrysotile and tremolite.

- The concentration of chrysotile decreased with time after exposure ceased but the concentration of tremolite did not.

- The clearance rate varied inversely with the length of chrysotile fibres. For fibres > 10 µm in length - i.e. fibre lengths in the reported range for carcinogenicity - the clearance half-time was estimated to be eight years. In other words, the tissue bio-persistence of chrysotile fibres in this study seems substantially more prolonged than in rodent experiments, and presumably corresponds to persistent high chrysotile fibre concentrations for many years after cessation of occupational exposure in humans, as discussed in paragraphs 5.112 - 5.113. It is also notable that the concentration of 6,250,000 chrysotile fibres mentioned in those paragraphs (for an individual but by no means unusual patient) is probably above the level at which Rogers et al. [2] identified an odds ratio for mesothelioma of > 8.5 (even allowing for differences in fibre size between the two different laboratories), and even the duration of 16 years after exposure stopped (as opposed to its commencement: 24 years) falls into the lag-time range lung cancer induction by asbestosis.

- Studies like this suggest that clearance mechanisms can be overwhelmed and break down at occupational levels of exposure in humans, with the existence of a long-term sequestered fraction of chrysotile fibres.

5.438 Mesothelioma rates in men and women in Sweden: attached to this Endnote is a recent paper by Jarvholm et al. [3] on trends in mesothelioma incidence in Sweden, which re-emphasizes some of the points made earlier in this report.

D. **COMMENTS BY THE PARTIES ON THE RESPONSES FROM THE EXPERTS**

1. **Canada**

5.439 Canada is pleased that the experts agree with Canada on certain crucial aspects of the debate in this case. Most importantly they opine that:

- Chrysotile is significantly safer than amphibole asbestos (three of the four experts agree);

- there is no risk to the public from low-level environmental exposure to chrysotile or from exposure in buildings that contain chrysotile (all of the experts agree);

- there is no risk to workers in mines or factories where use of chrysotile is controlled (three of the four experts agree); and
there is no risk to “handymen” or “do-it-yourselfers” who disturb chrysotile products, because their exposure is intermittent and thus inconsequential (three of the four experts appear to agree).

5.440 In short, although the experts agree on the inadequacy of a data (statistical limitation to support a threshold), their findings are consistent with the view that low levels of exposure to chrysotile asbestos create no detectable health risks. Indeed, the only population that the experts view as having problematic exposure is tradesmen, e.g., plumbers, electricians and mechanics, who disturb or modify chrysotile cement and friction products. At this point, the experts and Canada diverge; also, at this point, the experts stray beyond their specialities (as several admit). Canada maintains that adequate controls for these exposures can be developed and applied and has set forth such controls in Comments to Experts’ Answers to Question 5.

5.441 Several other aspects of the experts’ answers require comment. Some of the responses of the experts appear not to distinguish between chrysotile and amphibole exposure, and between modern uses (e.g., chrysotile friction and cement products) and historical uses (e.g., insulation containing amphiboles). In many places, for example, the experts appear to draw conclusions regarding chrysotile based in part or in whole on data from individuals exposed to amphiboles and/or amphiboles and chrysotile. This is of greatest concern to Canada regarding the experts’ conclusions on tradesmen; as the experts no doubt agree, the greatest risk to tradesmen is not exposure to modern chrysotile products, but the disturbance of flocking or insulation containing amphiboles. Similarly, the experts do not always distinguish peak and cumulative exposures. For the purposes of defining health risks, the cumulative measure, not the peak measure, is key.

5.442 It is crucial that this proceeding forms on the pertinent issues. The key issue is whether exposure to modern uses of chrysotile can be controlled to ensure worker safety or if a total ban is required to achieve an equivalent level of safety. The experts’ answers help to focus the proceeding on tradesmen exposure.

Question 1(a)

5.443 Canada believes that the Panel should take note of the clarifications to this question proposed by Drs. de Klerk and Musk. The former writes that “the more relevant question here is: who is likely to receive the most exposure and therefore have the greatest risk of disease”. Dr. Musk rephrases the question in almost identical terms, taking the expression “risk of exposure” to mean “who is most likely to receive the most exposure and therefore be at the greatest risk of developing asbestos-related disease”. In Canada’s view, because the evidence suggests different risk per unit fibre exposure in different sectors, it is the combination of level of exposure, duration of exposure and risk per unit fibre exposure that is important.

5.444 The experts have confirmed that any risk from exposure to chrysotile will depend on the nature of an individual’s specific occupational setting and the risk per unit fibre exposure in that setting, certain sectors being the subject of more stringent controls than others. For example, the experts echo the Parties’ agreement to the effect that the mining and manufacturing sectors have successfully controlled the risks to which their workers had previously been exposed. Certain settings pose lower risk per unit fibre exposure than others do.

5.445 Canada does not disagree with the statement that the so-called secondary user sector is the most diverse. Canada nevertheless understands that the experts do not believe that the diversity of this particular workforce is to be considered the only factor that may contribute to a greater likelihood of exposure; rather, as Dr. Musk puts it, “the risk of developing asbestos-related disease [...] (also) depend(s) on [...] the type of asbestos being produced or used or otherwise encountered. It would also depend on the conditions of work such as indoors versus outdoor etc.” As the Panel also knows, the specific uses or products also entail more or less risk.
Canada does not believe that the diversity of this workforce precludes effective control. The diversity of a specific workforce is not indicative of the quality of the work practices actually observed by the members of that workforce. A typical construction site offers numerous examples of sound safety practices: from hard hats to proper footwear, from the use of common sense to following trade-specific work practices, measures are taken to insure safety and avoid trauma.

Canada notes that the experts have not commented on the assertion by the European Communities that there is a correlation between the amount of chrysotile used by France and the incidence of asbestos-related disease. Clearly no such correlation can be made, in logic or in fact. The logic on which the European Communities purports to base this assertion is a sophism, and should be dismissed accordingly. From a factual point of view, the following factors suggest that the correlation is false: the relative difference in potency and in biopersistence of amphiboles and chrysotile, the historical uses of each fibre type, and the differences in risk per unit fibre exposure in different sectors.

Canada notes that Dr. Infante assimilates friable amphibole or mixed fibre type exposure circumstances to those of high-density chrysotile products, thereby answering the wrong question. He correctly identifies worker contact with insulation as being the “typical scenario” in which exposure to asbestos will occur. But most insulation is friable, as opposed to high-density, and most friable insulation products contained amphiboles or mixed fibre types. It is not clear how this answer based on friable mixed asbestos products responds to a question relating solely to safety of high-density chrysotile products.

Question 1(b)

Canada takes note that the experts have indicated that the risk of human health associated with the various uses of chrysotile throughout its life cycle is overwhelmingly a workplace issue, and therefore not related to the “handyman”.204

Question 1(c)

The answers given by the experts indicate that, on their own, chrysotile cement products do not pose a health risk because of their normal weathering, erosion or general degradation, and that “there is little or no dispute among experts on this issue”.205

Canada wishes to draw the attention of the Panel to the results of the investigation carried out by the Western Australia Advisory Committee on Hazardous Substances (WAACHS), cited by Dr. Henderson.206 This report contains different sections describing asbestos cement products, their production and use and their health effects, as well as surveys of schools and other relevant measurements of asbestos concentrations. In addition to pertinent recommendations, the report contains several appendices, including one on the Effects of Asbestos Cement Products – A Review of the Literature and another on Acceptable Air Concentrations of Asbestos Fibres in the General Environment, both prepared by one of the experts to this Panel, Dr. de Klerk.

On low level air concentrations, Dr. de Klerk writes: “[M]ost of these estimates are on or below the level of what the Royal Society would consider acceptable [...] The 1986 IPCS report did not even bother to estimate such risks and summarised the risk exposure unrelated to occupation as being undetectably low”.207 Indeed, the executive summary of the WAACHS report indicates: “[...] [T]he level of risk is low enough to be considered to be negligible relative to these other risks in our society”.208 Similarly, in his report to the Panel, Dr. Henderson underlines that compared to the fibre concentrations observed in the vicinity of asbestos-cement roofing, “a greater risk to health would arise from [workers] falling from or through the roofs”.208

High-density chrysotile on buildings has been extensively studied. Indeed Teichert found the following: “the study of emission conducted on coated and uncoated roofing materials revealed low asbestos fibre concentrations, even though severe corrosion
was observed on uncoated asbestos cement roofs and a considerable quantity of material containing asbestos could be removed by blowing or suction. The asbestos fibre concentrations that were measured in populated areas are well below the level considered acceptable by the Health Authorities of the Federal Republic of Germany, i.e. clearly below 1000 fibres/m³ (or 0.001 f/ml). Felbermayer and Ussar, for their part, write: "a comparison of the asbestos fibre concentrations in those areas with and without asbestos-cement roofing (...) lead to the conclusion that there is no statistical significant connection between the use of asbestos-cement materials and the asbestos fibre concentrations found in the various measurement areas."

5.454 Finally, Canada would like to bring to the Panel’s attention the following recommendation of the WAACHS report, which is: "[A]n asbestos cement roof, which has not deteriorated to an extent where physical safety or structural integrity is of concern, should not be replaced. In addition, an asbestos cement roof should not be treated with a coating on the basis of risk to health. Other asbestos cement products are generally less prone to deterioration and do not require attention for health purposes". Nonetheless, many chrysotile-cement products are coated with protective sealant agents.

Question 1(d)

5.455 The experts agree that the degree of risk to the health of workers intervening on high-density chrysotile cement products will depend on the manner in which an intervention is carried out. As noted by Dr. Infante in his response to question 1(e), "the extent of the exposure to the worker (...) would depend on the nature of the intervention, e.g., the circumstances under which the chrysotile asbestos product is manipulated in terms of work practices, the controls, or lack of controls in place and the type of personal protective equipment provided to the worker". Dr. Henderson illustrates this proposition when he writes that "cutting (chrysotile-cement) with hand saws produced lower concentrations."

5.456 Canada accepts that abrasion and cutting of high-density chrysotile products can release materials. However, the degree of exposure, if any, will depend on the methods and controls used. Canada notes that the experts disagree as to the exact composition of the materials that would be released by such interventions (see question 1(f)), although there is apparent agreement that cutting chrysotile cement releases crystalline silica, an IARC Class 1 carcinogen. Cutting chrysotile cement using simple work practices such as those outlined in ISO Standard 7337 will therefore provide protection from any potentially harmful material contained in such a product. Wetting the product before cutting and/or using commonly found suction attachments when sawing are techniques that can be used as added, but perhaps unnecessary, precautions. A final safety barrier would be for the worker to wear a facemask: this step would render it virtually impossible for the worker to inhale dust.

5.457 Neither the European Communities nor the experts have demonstrated that such practices would subject workers to cumulative exposures presenting health risks. An American survey estimated that a worker would spend less than 1/16th of his work time on tasks that would involve aggressive interventions on chrysotile-cement of the type susceptible of releasing any substantial amount of dust. Canada submits that the European Communities have not identified any population of workers that would be subject to a detectable risk because of professional contact with high-density chrysotile cement. The European Communities’ contentions vis-à-vis the “handyman” are therefore even less convincing (see next answer).

Question 1(e)

5.458 Canada agrees with Dr. Henderson’s conclusion that “occasional interventions (...) would predictably produce low cumulative exposures, with a lower risk (...)”. Dr. Henderson also affirms that for “electricians, carpenters, plumbers, insulation workers and so forth”, “it is acknowledged that most if not all these mesotheliomas are a consequence
of exposure to (...) a mixture of asbestos types, including chrysotile and one or more of the amphiboles.”

5.459 The Panel has not been presented with evidence that contradicts Canada’s assertion that occasional interventions do not pose a risk that is significantly different from zero (statistically). Therefore, the experts have not validated the EC’s claim that an alleged risk for workers or the “handyman” is something more than undetectable.

5.460 Nor has the Panel been presented with evidence or expert opinion that supports the European Communities’ claim with respect to the “handyman”. Given that cohorts exposed to relatively high concentrations of chrysotile over entire occupational lifetimes show no increase of disease, it is unlikely that occasional interventions by a “handyman” would produce more than an equally undetectable risk. Obviously the “handyman” or bricoleur du dimanche will not encounter high-density chrysotile-cement products on a daily basis, nor devote his “handiwork” exclusively or principally to cutting such products. Rather, the typical “handyman” will rarely, if ever, come into contact with chrysotile cement products, let alone be sawing them.

5.461 The Panel should note that no evidence has been presented that shows any fatality in workers, let alone in “handymen”, who would have been subject to any form of exposure, high or low, from contact with chrysotile cement products; the argument presented by the European Communities has been based entirely on hypothetical scenarios.215

Question 1(f)

5.462 There is debate in the scientific community and among the experts appointed by the Panel as to the exact physical and chemical composition of what is contained in dust from certain interventions on chrysotile cement products. Dr. Infante writes, however, that this dust (indeed, all cement dust) will contain “crystalline silica”, a known IARC Class 1 carcinogen found in all cement.

5.463 A 1992 IARC publication determined that “in asbestos-cement products, the asbestos fibres usually represent 10-15 per cent of the total weight and are embedded in the cement. Therefore, it is not certain a priori that dust generated from asbestos-cement products will have the same effect as dust from pure chrysotile. [...] In asbestos-cement dust most of the asbestos fibres form aggregates with cement particles ... those which do not form aggregates ... appear to be coated with a calcium-containing layer. In absorption experiments, the asbestos-cement dust behaves more like cement dust than like asbestos dust.”216 Because the surface properties of asbestos fibres are altered by certain heating, pH, and abrasion conditions217, it can be deduced that the composition and effect of the final aerosol would be different than that suggested by studies of concentrations of fibres alone. And again, controlled use procedures limit release, and proper breathing equipment precludes exposure.

Question 1(g)

5.464 Canada believes that the Panel was not presented with any quantification of this risk, or indeed its existence. Dr. Infante describes how the removal of chrysotile cement panels can be accomplished with negligible release of respirable fibres. Most other chrysotile cement products are found in the form of underground water pipes. Studies show that these products remain intact for decades after installation.218 Hence, very little of this product will need to be disturbed. Moreover, the excavation and removal of pipes is not executed by manual labour, the bulk of any removal being done by heavy machinery.

5.465 The Panel should also note that the removal of chrysotile cement products does not generally entail crushing. Rather, if and when necessary, chrysotile cement products can be removed, transported, and disposed of by means that do not constitute a detectable risk to human health. The French Circulaire 97-15 accomplishes this goal for the high-density products at issue in this proceeding.219 Also, if France is ensuring the safe removal
and disposal of friable asbestos materials known to contain amphiboles or mixed fibre types, the Panel should conclude that the removal and disposal of high-density chrysotile cement products can be accomplished even more safely, since high-density materials are indisputably recognised, even by France, as much easier to manage than anything in friable form.

**Question 1(h)**

5.466 See comments on previous question.

**Question 1(i)**

5.467 Canada wishes to add the following comments on the answers to this question. Once removed from a building, a chrysotile cement panel, even if broken into several pieces, remains as intact as when it formed part of that building. Studies referred to above indicate that chrysotile cement roofing does not contribute (< 0.001 f/ml) to the levels of chrysotile occurring naturally in the environment. Likewise, chrysotile cement piping is generally found below ground, and therefore does not contribute to the levels of chrysotile naturally occurring in the atmosphere. If removed from roofing, or if excavated and removed from a water system, chrysotile cement products are transported to a landfill and buried anew beneath a layer of earth. Consequently, Canada is of the view that used chrysotile cement products can be eliminated safely.

5.468 Canada also notes that recent technology has enabled safe (in some cases, on-site) disposal of chrysotile products. For example, chrysotile can be treated with chemicals and/or subjected to high temperatures so as to render the end product entirely harmless and, in fact, suitable for enhancing the quality of soils. For example, in the United States, a foam has been developed that eliminates the risk associated with removing asbestos from buildings; when this product is sprayed onto asbestos fireproofing, the fibres turn into harmless globs of magnesium silica. A U.S. building contractor recycles asbestos by subjecting it to a chemical bath and high temperatures resulting in a totally inert end-product suitable for soil improvement. A Japanese company, responding to a government law mandating pollution-free disposal of asbestos, melts asbestos into harmless glass.

**Question 2**

5.469 Canada has advocated the use of chrysotile in high-density products only; textiles are not of that category, and had been banned in France prior to the adoption of the measure that is the subject of this dispute. Friction materials using chrysotile have not been shown to constitute a risk to human health. Indeed, the contrary is probably true: lesser braking action of linings manufactured without chrysotile is cited by France as the safety concern for which it exempted certain military vehicles from the purview of the Decree.

**Question 3(a)**

5.470 Three of the four experts concur with the position of Canada and the WHO that a clear distinction must be made between chrysotile and amphiboles. Dr. Musk believes that “there is a need to distinguish chrysotile asbestos from amphiboles based on the epidemiological data at least” and that the relative pathogenicity of some amphiboles to chrysotile may, in some cases such as mesothelioma, be 100 to 1. Dr. de Klerk affirms that the “epidemiological evidence is clear that, for a given quantity (intensity and duration) of exposure, chrysotile imparts less risk than amphibole fibres.” The difference in pathogenicity is, according to Dr. de Klerk, up to 50-fold in the case of lung cancer and up to 100-fold for mesothelioma. Dr. Henderson concludes that: “a clear distinction should be made between chrysotile and the amphibole forms of asbestos.”

5.471 Domestic legislation and international standards have long recognized the relative pathogenicity of different asbestos fibre types by permitting higher exposures to chrysotile than to amphiboles. In the European Communities in 1998, for example, the maximum exposure level for amphiboles was 0.3 f/ml, whereas it was 0.6 f/ml for chrysotile.
In Canada (Quebec), it is 0.2 f/ml for crocidolite and 1 f/ml for chrysotile. Similarly, international instruments such as the ILO’s Convention 162 and Recommendation 172 advocate an outright ban on crocidolite, while recommending replacing chrysotile if and only if safer substitutes exist.

5.472 Dr. Infante acknowledges epidemiological data to the effect that chrysotile is less dangerous than amphiboles, but sees no basis for distinguishing between asbestos fibre types. Dr. Infante’s dissonant view to the question of relative pathogenicity between asbestos fibres – one which echoes the European Communities’ argument but simply begs the question – is that because amphiboles and chrysotile are both classified as carcinogens, no distinction should be made.

5.473 In 1998, the WHO affirmed that a distinction should be made between chrysotile and amphiboles because using data from exposures to amphiboles “contribute[s] less to our understanding of the effects of chrysotile, due to concomitant exposure to amphiboles.” The distinction between chrysotile and amphiboles is crucial in this instance since the current problem of asbestos in France is due to past uses of friable materials, high-level exposures, and the use of amphibole fibres. The distinction between chrysotile and amphibole asbestos is also important because the extrapolations made by INSERM to assess the risks associated with chrysotile are based on exposures to amphibole fibres in proportions of up to 100 per cent in circumstances which have nothing to do with the current uses of chrysotile.

**Question 4(b)**

5.474 Physical properties, as well as chemical properties that determine biopersistence, are identified as relevant factors of pathogenicity by Drs. Musk, Henderson and de Klerk and by the WHO.

5.475 Dr. de Klerk, for example, has written that:

> “[T]he important carcinogenic properties of asbestos are related to the physical properties of size and shape of the fibers, and to their quantity. To cause any harm, fibers must be able to reach the target organs [...]” [...]

> “[I]n all occupationally exposed series of mesotheliomas, none have occurred in cohorts where amphibole asbestos has never been used or detected. Chrysotile asbestos has not been directly implicated in any case of peritoneal mesothelioma. [...] The main differences between the effects of chrysotile and amphibole fibers are:

1. Industries using a mixture of asbestos types have higher rates of disease than similar industries using only chrysotile.
2. Chrysotile fibers are eliminated more readily from the lungs than are amphibole fibers.
3. Much smaller doses of amphibole fibers than chrysotile fibers can induce mesothelioma.”

5.476 All four experts recognize the lower biopersistence of chrysotile. INSERM, citing numerous studies, also acknowledges the lower biopersistence of chrysotile:

> “Les études expérimentales ont montré que la biopersistante des fibres de chrysotile était inférieure à celle des amphiboles (Wagner et al., 1974; Davis et al.; Davis and Jones, 1988; Churg et al., 1989; Churg, 1994).”

5.477 Dr. Infante identifies the physical characteristics as also relevant to the relative pathogenicity of asbestos fibre types, but, unlike the three other experts and the WHO, believes that the role of biopersistence, through the element of solubility, “is not so clear.”
5.478 Chrysotile fibres are “curly” and downy while amphibole fibres are straight and rigid like needles. Drs. de Klerk and Musk both specifically address the “straightness” element. The WHO has observed that:

“Inhalation of respirable straight fibres [amphiboles] is reported to be associated with greater penetration to the terminal bronchioles than in the case of ‘curly’ fibres [chrysotile].”

5.479 Once they have entered the respiratory tract, chrysotile asbestos fibres, because of their curly shape, are more easily cleared by the mucociliary process than are straight and rigid amphibole fibres. Dr. Henderson writes: “[I]t is well known that chrysotile fibres are cleared more rapidly than amphiboles, especially in long-term studies (Churg, 1994).” This is confirmed by a 1994 European study by Dr. Albin: “[A]dverse effects are associated rather with the fibres retained (amphiboles), than with the ones being cleared (largely chrysotile).”

5.480 For chrysotile fibres that do nonetheless manage to become lodged in the lungs, the solubility of the fibres and the action of macrophages come into play to make chrysotile a much less potent fibre. First, as the WHO recognizes, chrysotile has a lower resistance than amphiboles in acidic environments such as the lungs. Second, the macrophages responsible for eliminating fibres from the lungs are able to deal more easily with chrysotile fibres than with amphibole fibres. A 1997 report of the French Government (G2SAT) referred to by the European Communities, recognizes that as a result of the chemical dissolution process that takes place in the lungs, carcinogen activity is subsequently practically nil:

“Il a été démontré que le chrysotile est nettement plus facilement éliminé du poumon humain que les autres formes [amphiboles]. Par ailleurs, il ne présente pratiquement plus d’activité cancérigène (par injection intra-cavitaire) après attaque acide, laquelle dissout la majorité du magnésium.”

5.481 Dr. Wagner, in his 1988 study of asbestos-related diseases, concluded: “Chrysotile is the least harmful form of asbestos in every respect and […] more emphasis should be laid on the different biological effects of amphibole and serpentine asbestos fibre.”

5.482 It should also be noted that gravimetric comparisons between amphiboles and chrysotile – widely used in the past in experimental work – tend to grossly misrepresent the relative pathogenicity of the fibres. According to the WHO, chrysotile “may contain more than 10 times more fibres per unit weight.” Recent studies that use both the fibre mass and the number of fibres as dose units confirm that, on a per fibre basis, amphiboles are far more pathogenic than chrysotile.

**Question 3(c)**

(i) **Asbestosis**

5.483 Dr. Henderson asserts that: “[T]he amphibole varieties of asbestos appear to be substantially more pathogenic than chrysotile for the induction of asbestosis and mesothelioma.” According to Dr. Henderson, “[A]sbestosis is a dose-dependent disorder with a threshold effect […] There is widespread agreement that asbestosis in general is a consequence of high intensity exposure (or lower intensity but more prolonged exposure).”

5.484 INSERM also supports the existence of a threshold for asbestosis, and according to INSERM, current low-level exposures to chrysotile pose no threat of asbestosis: “les expositions actuellement relevées dans les industries directement utilisatrices d’amiante devraient conduire à la disparition des cas d’asbestose confirmée (Doll et Peto, 1985)” It is clear, therefore, that asbestosis is not relevant to this dispute.
(ii) Lung Cancer

5.485 Dr. Musk believes that lung cancer risks are more than ten times greater in the case of amphiboles than in the case of chrysotile asbestos. Dr. de Klerk suggests the difference may be up to 50-fold.

5.486 Dr. Henderson states that the “greater carcinogenicity of the amphiboles [...] appears not to extend to the induction of lung cancer” but he admits that “chrysotile is implicated in one of the lowest rates of asbestos-associated lung cancer (in Quebec chrysotile miners and millers).” Dr. Henderson’s reluctance to conclude the greater carcinogenicity of amphiboles seems to be caused by the results of Dr. Dement’s study of the Charleston, South Carolina asbestos textile industry.

5.487 The Charleston data has recently been revisited by Bruce Case, André Dufresne, A.D. McDonald, J.C. McDonald and Patrick Sébastien in a study released in Maastricht in October 1999 at the VIIth International Symposium on Inhaled Particles, a symposium attended by some of the world’s leading experts. This study shows that a significant amount of crocidolite and amosite fibres was found in the textile workers’ lungs. This analysis sheds new light on the issue and explains the extreme results of the original study by Dr. Dement and the subsequent study by Dr. Stayner. These studies of textile workers exposed to crocidolite and amosite can thereby no longer be used to demonstrate the risks associated with chrysotile fibres.

5.488 The seminal findings of Case et al. may cause Dr. Infante to reconsider his view – based principally on the studies by Dement and by Stayner – that “chrysotile may be more potent in causing lung cancer.”

(iii) Mesothelioma

5.489 On the relative risks of mesothelioma, Dr. Henderson observes that: “[T]here is general though not universal agreement of a differential potency between the amphiboles versus [chrysotile] for mesothelioma induction.” He believes amphiboles may be greater than 60 times more likely than chrysotile to induce mesothelioma. Drs. Musk and de Klerk estimate that the potency of amphiboles may be 100 times greater. And although Dr. Infante also concedes that “amphiboles may be more potent in causing mesothelioma”, he fails to conclude from this that a distinction exists between chrysotile and amphibole fibres.

5.490 This distinction is also emphasized in pathology medical reference books: “It is important to make the distinction between various forms of amphiboles and serpentines, because amphiboles, even though less prevalent, are more pathogenic than the serpentine chrysotile, particularly with respect to induction of malignant pleural tumors (mesotheliomas). Indeed, some studies have shown the link is almost invariably to amphibole exposure.”

(iv) Other Diseases

5.491 Dr. de Klerk links other asbestos-related diseases such as pleural plaques and pleural thickening more with amphiboles than with chrysotile: “[P]leural plaques appear to be more common among anthophyllite workers than others while crocidolite workers have more diffuse pleural thickening, and benign asbestos pleurisy also seems to be more common after crocidolite exposure.” Dr. Henderson also raises the issue of types of fibres in dealing with parietal pleural plaques.
Question 4(a)

5.492 Drs. de Klerk and Musk agree that the existing epidemiological data show no excess health risks at low-level chrysotile exposures. Dr. Henderson is not aware of exposure-response data for low-level exposures. Dr. Infante again relies heavily on Stayner’s study, a study on one single cohort of textile workers now known to be based on textile workers exposed to amphiboles as well as to chrysotile. Newhouse and Sullivan studied exposures to chrysotile in the manufacturing setting: “[I]t is concluded that with good environmental control, chrysotile asbestos may be used in manufacture without excess mortality.”

5.493 Thomas et al. concluded similarly for an asbestos cement factory: “[T]hus the general results of this mortality survey suggest that the population of the chrysotile-cement factory studied are not at any excess risk in terms of total mortality, all cancer mortality, cancers of the lung and bronchus, or gastrointestinal cancers.”

5.494 There is clearly no increased risk of lung cancer in the friction products manufacturing industry at levels below 356 f/ml-years. This means that there was no chrysotile-related increase in lung cancer risk for persons exposed to the equivalent of up to 8.9 f/ml for 40 years. Even if we allowed a 10-fold protection factor this would be 0.9 f/ml for 40 years for lung cancer. More recently in 1997, McDonald et al. concluded from the analysis of a cohort of 10,000 asbestos workers with average exposures to 45 f/ml over 20 years that: “[...] from the point of view of mortality [...] exposure in this industry to less than 300 mpcf.years [approximately 45 f/ml over 20 years] has been essentially innocuous.” This unequivocal data comes from the longest term study of the largest group of chrysotile workers ever conducted. A review of eight studies of cohorts exposed to chrysotile only led its authors to conclude: “[T]he evidence for chrysotile shows that for lung cancer and mesothelioma there exist levels of exposure below which risks are for practical purposes zero.”

Question 4(b)

5.495 According to Dr. Henderson, whether a threshold exists generally is a much-debated issue. For the case at hand, i.e. low-level exposure to chrysotile, Dr. Henderson states that: “[I]f a threshold exists, it must lie somewhere in this area, between no exposure, low-level environmental exposure, and low-level occupational exposure.” He also points out that, although no threshold has been identified, “[a]t the same time, no increase in risk of mesothelioma has been identified at very low-levels of exposures.” Drs. Musk and de Klerk agree that the epidemiological data show an absence of risk at low exposure levels, but are unwilling to commit to the existence of a threshold. If there is agreement that low level exposures show no increased health risk, admitting the existence of a threshold is academic.

5.496 The extreme difficulty of proving a threshold scientifically is echoed by the European Communities’ DG XXIV Report:

“In fact, a threshold implies the demonstration that an effect does not occur at or under a given dose level. The unequivocal demonstration (i.e. identification) of a ‘negative’ is tantamount to impossible.”

5.497 The corollary to the proof of a threshold is the proof of the absence of a threshold. The proof that no threshold exists would need to explain the absence of an excess risk of lung cancer or mesothelioma in chrysotile-only cohorts, as well as the lack of any chrysotile-related increase in lung cancer mortality in workers exposed to less than 900 f/ml-years in the 10,000 miners and millers studied in Quebec. Dr. Henderson does acknowledge the existence of a threshold for asbestosis in his answer to Question 3: “Asbestosis is a lung dependent disorder with a threshold effect [...] There is widespread agreement that asbestosis in general is a consequence of high intensity exposure (or lower inten-
sity but more prolonged exposure).” INSERM also supports the existence of a threshold for asbestosis:

“La plupart des données épidémiologiques recueillies dans des populations professionnelles exposées suggèrent que l’asbestose cliniquement et/ou radiologiquement caractérisée n’apparaît qu’à partir d’expositions suffisamment élevées […] un seuil minimal de 25 f/ml-années a ainsi été avancé (Doll et Peto, 1985).”

5.498 Why could there not be a threshold for other asbestos-related diseases? Dr. de Klerk asserts that:

“[I]t is now widely believed that the risk for chrysotile workers in fibrous cement and friction product manufacturing is so slight as to be undetectable. It is widely held that this kind of negligible risk level ‘threshold’ exists at different levels for all types of asbestos for all relevant diseases.”

5.499 Some experts advising the EC believe there is a threshold for diseases other than asbestosis:

“It is very likely that there is a practical level of exposure below which it will be impossible to detect any excess mortality or morbidity due to asbestos. […] Thus, it is possible that there is a level of exposure (perhaps already achieved in the general public) where the risk is negligibly small.”

5.500 This links to Dr. de Klerk’s observation that: “[T]he smaller the effect that needs to be demonstrated, the larger the study needs to be.” Dr. Infante, who dismisses the Panel’s question as “moot”, points out that “it is not possible to determine thresholds from epidemiological studies because of the lack of statistical power to distinguish that the risk is virtually zero.” Canada argues – epidemiological data in hand – just that low-level exposures to chrysotile pose a risk that is “virtually zero”: “un risque indétectable.” Dr. Infante uses Stayner’s data once again to claim that the chrysotile data fit with a linear no-threshold model. With the new analysis on the Charleston cohort data discussed above, this argument does not hold.

Question 4(c)

5.501 Drs. de Klerk and Musk agree that there is epidemiological data indicating no increased risk at low-level exposures, but the experts believe the linear model may be appropriate. However, “[W]hether or not it is a valid method is unknown.” According to international experts from the Health Effects Institute-Asbestos Review (HEI-AR), such as Julian Peto, David G. Hoel and W. Nicholson, the linear model is not used for its validity, but precisely because it tends to overestimate risk. Dr. de Klerk shares this view and states that the model provides a “conservative estimate.”

5.502 The limits of the linear model and the conditions under which extrapolations are made must be clearly set out. Extrapolations from high-level exposures and exposures to amphiboles should not be taken at face value to ban chrysotile in today’s context of low-level chrysotile-only exposures. Canada’s critical view of the linear model is supported by a 1999 report by the Australian National Industrial Chemicals Notifications and Assessments Scheme (NICNAS) cited by Dr. Henderson:

“There are many problems associated with low-dose risk extrapolation, such as the assumption of a linear relationship. However, as insufficient data exist to indicate threshold exposure for effect, the linear extrapolation methodology provides a conservative worst-case scenario estimate of risk. Other confounding factors in estimating risks from epidemiological data are possible contamination by other fibre types and inaccurate estimates of historical exposures.”
5.503 Not only does the linear model provide a worst-case scenario, it provides a grossly exaggerated estimate of risk when "confounding factors", as Dr. Henderson calls them, are so clearly present. INSERM made extrapolations from high-level amphibole exposures to mixed fibre type exposures, as well as from exposures in the textile industry and during the installation of low-density products such as flocking. Amphiboles are much more potent than chrysotile, and the risks in the textile industry cannot be compared with the risks in the high-density chrysotile products, as Dr. Henderson points out in citing Boffetta: "[I]n general, the risk of lung cancer ... is highest in studies of asbestos textile workers." 265

5.504 Another important consideration is the human biological defence mechanisms that are naturally much more effective at low-levels of exposure, i.e. clearance, biopersistence and DNA repair mechanisms. 266 Given these mechanisms, the reasoning behind the threshold model is both intuitively and scientifically sound, as well as epidemiologically validated. To illustrate this, consider the following illustration: the effect of 50 fibres in the lungs will be more than five times the effect of ten fibres.

5.505 According to Sir Richard Doll, who first demonstrated the link between asbestos and lung cancer (as well as between smoking and lung cancer), "[W]e have no real ground for postulating that a linear relationship for lung cancer can be extrapolated back to the levels of dose with which we are concerned in non-occupational settings." 267 Ames and Gold are of the same view: "[L]inear extrapolation from the maximum tolerated dose in rodents to low-level exposure in humans has led to grossly exaggerated forecasts in mortality." 268 Fournier and Efthymiou are even more categorical: "[L]inear extrapolation to zero is an unscientific methodology whose social consequences are so immense that it warrants unconditional elimination." 269 INSERM acknowledges the limits of the linear model's application when it states that it provides nothing more than food for thought: "cette extrapolation ne crée pas une information scientifiquement certaine, elle représente une aide à la réflexion en matière de maîtrise de risque." 270

5.506 As Dr. de Klerk points out, "how one extrapolates risk assessment outside the range of available data is more of a societal decision than a scientific one."

**Question 4(d)**

5.507 Situations where there is no increased risk at low levels of exposure have been used by Stayner et al. to establish NOAELs [i.e. no observable adverse effect levels] for silica. A similar model is used for asbestosis. Canada believes that the use of such a model is warranted for other asbestos-related diseases, particularly since it has been acknowledged by Dr. Musk and Dr. de Klerk that epidemiological data exists to justify such an approach.

**Question 4(e)**

5.508 We concur with Dr. Henderson’s view that "[T]his question iterates the issue of a threshold exposure." Canada nonetheless notes the use by Dr. Infante of a 1992 study by Bégin et al. to demonstrate the risks related to “background levels” is erroneous. As has been pointed out by Canada in its factual arguments, 271 this study is based on exposures to a mix of chrysotile and amphiboles in the manufacturing and construction industry, and therefore is not relevant to exposures from the current uses of chrysotile.

**Question 5(a)**

5.509 Clearly the answers given by the four experts are based on their concept of what is meant by controlled use. It is also evident that the controlled use concept as espoused by Canada was not the approach that resulted in their answers. We must therefore respectfully disagree with the answers given by the experts in respect of controlled use of chrysotile and high-density chrysotile containing products. The fact that they agreed that controlled use of chrysotile and high-density chrysotile products is feasible at some points of the life cycle, but not in others, suggests that they are not far from the view of Canada. The only
difference is that Canada believes that the experts misunderstand the controlled use principle and that, as properly understood and implemented, use can be controlled throughout the full life cycle of high-density chrysotile containing products. The basis for our view, with supporting evidence, is set out below.\textsuperscript{272}

(i) \textit{Canada’s understanding of the “Controlled use” principle}

5.510 The Canadian government’s review of the experts’ reports and answers to the questions posed by the Panel reveals that there is one crucial issue, which seems to override all other issues. This is the question of whether the application of the controlled use principle is feasible and credible in all stages in the life cycle of a product. While there is a reasonably high degree of agreement among experts that controlled use can be a reality in the mining and manufacturing sectors, serious doubts are expressed that controlled use can be applied in a few sectors of use - installation, maintenance and demolition. However, the basis for this view is not documented, except by Dr. Infante and Dr. Henderson.

5.511 By “controlled use”, the Canadian government means “stewardship” based on the total life cycle. This is outlined in the document \textit{The Mineral and Metals Policy of the Government of Canada: Partnerships for Sustainable Development}.\textsuperscript{273} With regard to asbestos, this “controlled use” is based on the following general principles:

- Only the chrysotile variety is used;
- only a limited number of well-defined product applications, where it has been demonstrated that they can be handled safely, are allowed (i.e. where the fibres are encapsulated in a matrix such as cement, bitumen, plastic, resin, etc.);\textsuperscript{274} and
- new product applications may be introduced only after a strict evaluation to ensure that a certain level of fibre release is not exceeded during its life cycle.

5.512 With regard to the downstream use sectors, “controlled use” implies that all distributors/manufacturers of asbestos will be required to have an import permit. This permit will be withdrawn if the company does not meet the following commitments:

- To distribute its products only to companies (users) licensed to purchase these products. Those companies must have workers trained and licensed to install products, and must be in compliance with regulations. Approved users shall not resell to third parties, and any unused materials must be returned to the manufacturer;
- to provide a list of users of products to the responsible government agency;
- to provide products cut to specification and to establish centres equipped to cut the products to size, and where persons cutting the products are trained and are licensed to work with asbestos; and
- to police the downstream users in co-operation with the government. The product manufacturer visits, monitors and reports on the performance of the downstream users at regular intervals. There are penalties for failing to provide this product stewardship.

5.513 While high-density products in most countries are not considered to pose any occupational or environmental health risk, disposal should only be undertaken by approved and appropriately trained persons.

5.514 Dr. Infante’s description of the permissible exposure limit for chrysotile asbestos, as well as programmes or standards that recommend or require specific engineering
control, work practices, training and education and personal protective equipment to control exposures to asbestos corresponds, to some extent, to Canada’s approach. Dr. Infante seems to suggest that because some workers do not comply with standards and regulations on controlled use in the United States, controlled use is not feasible. As explained in Appendix A on friction material and Appendix B on asbestos cement, the controlled use approach can minimize, if not eliminate, workers’ non-compliance. 275

5.515 Canada does not propose that any chrysotile products produced, sold or used without the implementation and enforcement of very stringent control procedures. Taking into account the types of products being manufactured and used in France at the time of the ban, Canada does not advocate re-introducing any product that cannot be handled according to the safety criteria outlined above. Canada is not advocating the introduction anywhere in the world of manufacturing facilities of products for which the technology does not exist to protect workers from exposure to chrysotile at levels where risks would be above epidemiologically based practical thresholds.

5.516 The experts have indicated that the level of exposure is such that they are not concerned about asbestos-related disease for persons living in buildings containing chrysotile asbestos products, including friable insulation. As none of the chrysotile products that will be used in the future are friable, this conclusion would be further reinforced. If the procedures envisaged under the “controlled use” policy are followed by licensed practitioners, the public will not be placed under any practically determinable increased risk of disease as a result of the manufacture and use of chrysotile containing products. Unlike friable insulation products where janitorial staff, electricians, carpenters, and others may be required to work regularly in an environment where exposures to asbestos would occur, the nature of the high-density products will ensure that exposures are a much rarer event.

5.517 Canada recognizes that the clock cannot be turned back. Friable mixed products produced in the past are now in place, and trades such as electricians or telephone engineers face situations where the potential health risk from exposure is considerably greater than any additional risk that new high-density chrysotile products would present. It is evident that the protection of workers who come into contact with friable products must be assured by the responsible jurisdictions through training in trade schools, appropriate information programmes by unions, and by governments and employers ensuring that the appropriate equipment and tools are made available to workers. 276

5.518 Regarding high-density products, Canada believes that no less stringent measures should be required, even though the evidence shows that the risk from exposure to high-density chrysotile products is minuscule compared to the risk from friable products, in many cases containing mixtures of chrysotile and amphibole fibres. Furthermore, in the absence of sound scientific data to the contrary, the same criteria should be applied to the handling of all products in which respirable fibres, including asbestos substitutes, may be released.

(ii) International Standards

5.519 None of the experts acknowledges that controlled-use of chrysotile asbestos cement products and other high-density chrysotile products stems from international standards. Dr. Infante even denies the existence of international standards on controlled-use of high-density chrysotile products. Canada wishes to remind the Panel that international standards, as the term is defined in the Agreement on Technical Barriers to Trade, do exist. Regulatory developments on asbestos fibres have been guided by ILO Convention 162 concerning Safety in the Use of Asbestos. 277 ILO convention 162 provides for: (i) the prescription of adequate engineering controls and work practices; (ii) the prescription of special rules and procedures for the use of asbestos or certain types of asbestos or products containing asbestos or for certain work processes; (iii) where necessary to protect the health of workers and technically practicable, the replacement of asbestos or of certain types of asbestos by other materials or the use of alternative technology scientifically evaluated by
the competent authorities as harmless or less harmful; and (iv) total or partial prohibition of the use of asbestos or of certain types of asbestos in certain work processes.276

5.520 The Code of Practice on Safety in the Use of Asbestos of the International Labour Office referred to by Canada in all its submissions is another international standard on controlled-use.277 The objects of the Code are: (i) to prevent the risk of exposure to asbestos dust at work; (ii) to prevent harmful effects on the health of workers arising from exposure to asbestos dust; and (iii) to provide reasonably practicable control procedures and practices for minimising occupational exposure to asbestos dust. To do so, the Code gives detailed guidance on the limitation of exposure in respect of asbestos cement and friction materials. Finally, Canada has referred the Panel to International Standard ISO 7337: Asbestos Reinforced Cement Products - Guidelines for On-Site Work Practices.280 This international standard gives guidelines for tools and working methods to be used on site with a view to maintaining the dust emission at the lowest practicable level. It applies to asbestos-cement products.

5.521 The ILO Convention 162 and the Code of Practice on Safety in the use of chrysotile should be supplemented by a national policy on responsible use based on the recognition and acceptance of the principles that both international standards set forth.281 As explained above, the objective of responsible use is to limit the handling of chrysotile to companies that comply with the national regulations or that have submitted action plans and formal commitments in writing with a view to bringing their activities into line with these regulations.

Question 5(b)

5.522 The experts recognize that training could be achieved in the manufacturing sector, where there is a small and cohesive workforce, but assert without support that it cannot be achieved in the construction sector, where there is a large and non-cohesive workforce. Dr. Infante wrongly equates non-compliance with regulated training requirements to non-feasibility of training for controlled-use of chrysotile asbestos.282

5.523 In Europe, as in other countries, there are now requirements for training workers. In Canada, both levels of government require training at all workplaces. It is possible for training to be made available by industry. In fact, information and training is one of the most important elements of a company’s preventive control programme. In line with the controls suggested at paragraphs 5.511 and 5.512, France could require through legislation that all construction workers handling asbestos products attend training sessions. France could also require that only designated, properly trained workers be allowed to work with those asbestos products that need to fall under a controlled regime.

5.524 During manufacture, controls such as wet processes and exhaust ventilation, essentially eliminate all exposure. On the work site, process changes are reduced by the industry manufacturing products requiring no, or virtually no, modifications on site. The controlled use approach includes the use of pre-cut and pre-drilled asbestos cement products, and provides for designated locations where chrysotile asbestos cement sheets or pipes are cut and drilled and where the appropriate controls are in place. The monitoring process is similar to that for other workplaces: all complaints are submitted to government inspectors for evaluation. The supplier has the responsibility for ensuring that all companies to which they supply have in place the proper equipment and training to ensure safe use of the product throughout its life cycle. Finally, the removal of high-density chrysotile products is carried out in accordance with government codes.

Question 5(c)

5.525 Both Dr. Henderson and Dr. Infante agree that, in many situations, when standards are properly applied, it is possible to maintain exposure below 0.1 f/ml. Also, as explained in Appendix A on the friction industry and Appendix B on the asbestos cement industry, experience shows that a level below 0.1 f/ml can be achieved because the
technology and work practices exist to control exposure during manufacture. No guarantee can be offered that there would never be a situation in which 0.1 f/ml might be exceeded as a peak exposure. However, there is no evidence that occasional peak exposures increase the risk of lung cancer or mesothelioma in chrysotile exposed workers. For example, the health experience of brake mechanics, i.e. no evidence of an increased risk of mesothelioma or lung cancer, is based on exposures that involved peak exposures, such as occurred during the blowing out of brake wear debris and the occasional grinding of brake linings. These operations involved short exposures above 0.1 f/ml. The actual concentrations associated with various tasks have been reported by Kauppinen and Korhonen, and by Rödelsperger. In spite of these short-term peak exposures, the average exposure of auto mechanics was less than 0.05 f/ml.

5.526 A person repairing their own brakes periodically nowadays (using disc brake pads mainly) would have extremely low cumulative exposures compared to full time auto mechanics and there is no reason for them to have any, even short term, exposures exceeding 0.1 f/ml. The risks associated with cumulative exposure to chrysotile at these levels would not be epidemiologically detectable for handymen handling friction or asbestos cement products.

5.527 Rödelsperger made dust measurements on about 40 buildings sites in Germany. He reported peak exposures of more than 100 f/ml in the vicinity of a grinding machine used to cut asbestos cement sheets. However, when he used the standardized work histories of 61 roofers, who had a mean duration of exposure of 16 years, he found that their mean cumulative exposure was 1.6 fibre-years/ml. These measurements were made 20 or more years ago, with the products and technology available then and for regular construction workers. It is evident that even under these circumstances, lifetime cumulative exposures were low. Thus, a handyman, even if he did not take proper precautions would still have a low cumulative fibre exposure because peak exposures are of short duration and he would be at a very low, undetectable risk of health effects.

5.528 It is generally agreed that at the levels of exposure associated with the use of the modern high-density products, they would not even put a full-time worker at increased risk of asbestosis and, therefore, this would not be of concern for a handyman working occasionally with the product. It has been amply demonstrated that the risk of lung cancer increases with increasing cumulative lifetime exposure that combines duration and level of exposure. A person exposed at 0.1 f/ml for 40 years has a cumulative lifetime exposure of 4 f/ml-years. If that person worked on a project only once each week for four hours for 40 years, he would not achieve the same lifetime exposure unless he was exposed to 1 f/ml continuously for the four hours of exposure every time he was exposed for 40 years. Thus occasional peak exposures of a few minutes contribute very little to cumulative lifetime exposure which is important in evaluating the risk of chronic diseases such as lung cancer or mesothelioma.

5.529 Gardner found no increased risk of lung cancer or other asbestos-related disease in a chrysotile asbestos cement plant where exposures were less than 1 f/ml. This was in a cohort of workers employed between 1941 and 1983. It is evident that any risk would have been well below the detection limit at 0.1 f/ml. A study of chrysotile cement production workers by Thomas and Neuberger & Kundi identified no chrysotile-related increased risk of lung cancer and Weill, while reporting an increased risk of lung cancer in asbestos cement workers, found the increased risk only in those with asbestosis. In this study, there was little evidence of asbestosis below 30-40 f/ml-years of exposure. This is about 0.75-1 f/ml continuous exposure for 40 years. Thus, there is little evidence to support a detectable increase in risk of lung cancer in workers with a 40 years cumulative lifetime exposure at 4 f/ml-years.

5.530 Any risk estimates obtained by linear extrapolation from high exposures to such low exposures are somewhat hypothetical and both Lash and Camus have shown that the risk estimates made by the U.S. Government have overestimated lung cancer risks.
5.531 Canada disagrees with the views of Dr. Henderson\textsuperscript{290} and Dr. Infante that controlled use of chrysotile asbestos is not feasible for workers involved in the construction trade and that service and maintenance workers such as carpenters, plumbers, and electricians will experience peaks of exposures to asbestos that place them at risk. The nature of high-density chrysotile asbestos products is such that few of the trades listed above will ever need to work on the products, with the possible exception of demolition workers. Again, there is evidence that during demolition exposure concentrations associated with chrysotile asbestos cement products is very low.\textsuperscript{294} Today, with chrysotile cement products and controlled use procedures, health risks become insignificant.

5.532 Recommended installation methods can eliminate the need to cut or drill into chrysotile-based products at construction sites, since those products are distributed in a variety of pre-cut and pre-drilled sizes, according to buyers’ specifications. In fact, many asbestos cement products are pre-formed ready for use. They are factory-made to the correct size and shape including holes so that a minimum of on-site preparations is needed. Once installed, chrysotile asbestos cement pipes are below ground and pose no risk to workers. Even if dug up, they pose no risks unless comminuted, ground or sawed, and, when this is necessary, the use of appropriate tools and controls will keep the release of dust and exposure well within the level considered safe by the WHO. Chrysotile asbestos cement sheets are used for roofing and exterior building walls. Once installed, there is no need to modify the roof until the life of the product is over. Similarly, there is no need to modify chrysotile asbestos sheets used as walls once they have been installed. The product can be painted without fibre release.

5.533 Chrysotile cement products are unlikely to release fibres into the environment or breathing zones of workers such as janitors, plumbers, electricians, repair men, etc., unless these workers have to actually cut or drill the product. Unlike insulation products, there will rarely be a need for anyone to perforate, saw, or grind installed chrysotile cement products. Where cutting or drilling is required, hand tools and low speed power tools are recommended in combination with wetting to keep dust levels to a minimum. Dust levels for various types of on-site working have been measured both in laboratories and in the field and these facts showed that risks could be maintained below detection limit.

5.534 Dr. de Klerk and Dr. Musk wrote that efficiency of controlled-use in the case of home handymen is outside their area of expertise. However, both Dr. Henderson and Dr. Infante have concluded that it is not possible to control exposure to chrysotile asbestos high-density products in non-occupational circumstances (occasional interventions by home handymen). Neither bases his conclusion on data. Dr. Henderson adds to his answer that although such risks are not quantifiable because of absence of data, these risks must be very small for lung cancer and mesothelioma, and non-existent for asbestosis.

5.535 Controlled use will reduce and even eliminate risks. The risk of chrysotile-related health effects is tied to cumulative exposure, that is, duration and level of exposure. Rarely will an individual under non-occupational circumstances achieve the exposure of a full-time worker. Occasional uncontrolled exposures for a handyman would not result in appreciable cumulative exposure. Data published by Brown\textsuperscript{295} showed time-weighted average (TWA) levels during demolition of weathered asbestos-cement roofing between 0.3 and 0.6 f/ml. One can likely guess that a handyman would not practice such an activity more than 40 hours in 25 years. This would average out to a TWA of 0.015 f/ml for the year of this activity, and a TWA of 0.0006 f/ml each year of the worker’s adult life. This is 1 million times less than past asbestos workers are. It is equivalent to exposure levels in schools containing ACM.\textsuperscript{296}

5.536 Based on INSERM\textsuperscript{297} and HEI-AR risk tables, which are based on mixed asbestos exposures, the resulting lifetime cancer risk would be between 10 and 20 in a million
depending on the time occurrence of this exposure scenario. More accurately however, the lifetime risk would be near zero per million, based on chrysotile friction workers who were exposed to similar fibres (species and dimension wise), and about 1 in a million, based on the risks of past chrysotile miners and millers. The casual user of a high density product, even if the product were weathered, is not likely to be at any increased risk of an asbestos related disease. If the supplier follows through on the requirements of controlled use, the casual purchase of chrysotile asbestos-cement products by the handyman will not be possible. However, there is probably no way of stopping any individual from doing something to any product if they can obtain it. This is a problem that exists for any product many of which pose serious health risks if abused.

Question 6(a)

5.537 Canada respectfully disputes the conclusions of the experts regarding the risk from substitute fibres, and with respect to one expert, the ability of substitutes to serve as suitable replacements to chrysotile. Canada notes that the treatment of the issue by two of the experts is terse, comprising only several sentences. To their credit, Drs. de Klerk and Musk indicate that the use and control of substitute fibres is not within their areas of special expertise. They offer some responses nonetheless. Canada is concerned, in particular, by their lack of familiarity with the relevant studies and actual modes of production, use and disposal of substitute fibres. For example, they apparently are unaware of research conclusively demonstrating the significant health risks from exposure to refractory ceramic fibres (RCF), which are discussed below.

5.538 This concern applies to Dr. Infante as well. Dr. Infante further appears unaware of (or ignores) recent research demonstrating that chrysotile is less biopersistent than many substitute fibres. Dr. Infante also ignores the population the experts agree are most at risk from exposure to any fibre – tradesmen – when he concludes (without support or, even, explanation, Canada notes) that the “nature of the production process makes substitutes more amenable to control than asbestos fibres.” Assuming he is not again conflating chrysotile with amphiboles when referring to “asbestos,” his point, were it true, would be irrelevant. The experts agree that chrysotile and chrysotile products can be safely mined. The key is exposure to tradesmen. And, for that exposure, no rationale exists suggesting that the ability to impose effective controls differs based on the type of the fibre.

5.539 Dr. Henderson, for his part, recognises that, as with all fibres, the pathogenicity of substitutes is defined by the “3Ds” (dimension, dose, durability). He seems also to understand that, due to the (lack of) historical use of substitutes, we cannot fully know the risks of using them. However, he then seems to ignore the importance of these facts.

5.540 All of the experts fail to take into account several very important factors. First, the chrysotile products at issue in this proceeding are quite few. Second, the exposure levels during the manufacture, use and disposal of these products are extremely low. Third, the data demonstrate that these few products have been and can be used without detectable health effects in humans. Moreover, in order to assess whether a substitute is safer to use than chrysotile in a product: (i) it is fundamental that the characteristics of the fibres being compared be those of the fibres as they are used in the product or as they are released from the product throughout the product’s lifecycle; (ii) it is essential that data on at least the key parameters (exposure, biopersistence and dimensions) be available to make this assessment. Unfortunately, the experts have not addressed these topics. In short, the experts have based their opinions on very limited, if any, data. While the experts reach conclusions that various substitutes (PVA fibres, glass fibres, cellulose and para-aramid fibres) are safer to use than chrysotile, they provide no systematic comparison of risks and very limited, questionable scientific data in support of their opinions.

5.541 Canada presents below a survey of the studies and concepts that the experts ignored. These studies give a picture of risks from substitute fibres starkly different from that suggested by the Panel’s experts. As demonstrated below, the situation concerning risk from substitute materials is as Canada set out in its factual arguments.
(i) The Fibres to Compare

5.542 Experimental data for a wide range of fibres have shown that the physical characteristics (diameters, lengths, density) of fibres are important in determining their respirability, when they are deposited in the respiratory system, and their capacity to induce fibrosis and cancers. Further, the risk of effects also depends on dose (exposure). Thus, differences in the risk of disease in various industrial sectors would be expected to occur because of differences in these, as well as other factors. As the characteristics of chrysotile and any substitute fibres are likely to be dictated by the product in which they are used, it is not appropriate to assess the risks associated with friction products or asbestos cement products using data from other industrial sectors. The data that should be available and used for the purpose of comparing risks should be those for the fibres as used in the specific products under review. Canada’s presentation proceeds on this basis.

5.543 Davis pointed out that while materials like wool, cellulose and other fibres have in some cases been used for many years, they are now being used in quite different applications, about which knowledge is very limited. As a consequence, the characteristics of the fibres used in the newer applications may not be the same as those in the conventional products manufactured in the past. Such changes can modify the respirability and biological activity of the materials. There is a further complication for substitutes that is not addressed by the experts. This is the fact that substitution does not always involve replacement of chrysotile by a single fibre, but often by several different materials or substitute fibres. For example, cocktails of fibres are needed to meet technical requirements in friction products. In addition, when substituting for chrysotile, other materials such as silica or other fibres, fire retardants or biocides must often be added. These agents may themselves be toxic or carcinogenic, and may act synergistically.

(ii) The Outcomes to be Measured

5.544 While it is reasonable to compare the risks of lung cancer and mesothelioma between the various fibre types, it must be remembered that different sized fibres may lead to fibre deposition at different locations in the respiratory system. For example, if more fibres of one material than another are likely to be deposited in the nasal passages, one should consider the possibility of an increased frequency of nasal cancer in evaluating the substitute. Dr. Infante mentioned the increased risk of nasal cancer in woodworkers, which has been well established. This might raise a question concerning the sources of the cellulose used as a substitute and whether controls are in place to avoid exposure to cellulose from woods that have caused such cancers. Also, some materials may cause dangerous allergic responses. Certain glass fibres cause skin irritation. Harrison notes that there are indications of an accumulation of oligomers in the kidney in some circumstances, so that attention should be given to the molecular weight of PVA used “especially if a smaller diameter material were to be produced.”

5.545 In considering risks, the composition of the dusts and fibres to which workers are exposed when handling the “raw substitute” materials, manufacturing the product, cutting, grinding, manipulating or disposing of the product also must be considered. For example, it is important to know whether the fibres of para-aramid, PVA or cellulose are opened (fibrillised) or comminuted during preparation or manufacture of the product? Does manipulation, sawing or drilling of the product give rise to narrower diameter respirable “fibres” such as result with polyester fibres during weaving? Do these fibre fragments have biological significance? What are actual use concentrations? It must be remembered throughout that there exists a substantial body of information concerning chrysotile. Unfortunately, in the case of substitutes, there are rarely any human epidemiological data available and even experimental data are limited. Perhaps this fact led Dr. de Klerk to conclude that “[G]iven the comparative lack of knowledge about the health effects of substitute materials, the continued use of chrysotile under [controlled] circumstances seems sensible.”

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The data that need to be compared in an evaluation of the relative safety of chrysotile and substitutes include the following:

- Epidemiological data that provide direct evidence of the risks associated with the products.
- Experimental data by inoculation of fibres or by inhalation experiments in experimental animals.
- Dimensions of fibres in the respirable airborne dust during the manufacture of the product.
- Dimensions of fibres in the respirable airborne dust during the use of products containing the fibre.
- Dimensions of fibres in the lungs of workers engaged in the manufacture of products containing the fibre.
- Dimensions of fibres in the lungs of persons exposed during the use of products containing the fibre.
- Dimensions of fibres in the respirable airborne dust and in the lungs following exposure during the disposal of the fibre or products.
- The biopersistence of the fibres in humans and animals.
- The cumulative exposure (i.e.: concentration x time) of workers engaged in all phases of manufacture, use and disposal of the product.
- Data on alterations or modification of the fibres chemically, physically and biologically during their life cycle that might affect their potential to cause health effects.

Even if one were to narrow the requirements to a smaller number of key parameters such as fibre dimensions, biopersistence and exposure-response, the available data are still inadequate to provide a credible basis for an adequate comparison. Thus, the unqualified wholesale affirmation that “substitutes are safer than chrysotile” is not well founded and potentially very dangerous. For example, prior to the finding of a very high risk of mesothelioma for persons exposed to very low concentrations of fibrous zeolite erionite in Turkey, there had been no indications world-wide that such fibres might after 30 years from first exposure at such very low levels produce such a high rate of mesotheliomas in humans. In South Africa, crocidolite had been used for about 60 years before Wagner reported that mesotheliomas were associated with crocidolite exposure. In humans, mesotheliomas do not occur until 40-60 years after first exposure. Thus, caution is needed in the absence of data regarding substitute fibres. As one expert stated: “better the devil you know” than the devil you do not know.

The experts present no data that show that the dimensions of all fibrous substitutes are outside the respirable size range during the substitute product’s lifecycle. This is because no such data exist.
(ii) PVA & Aramid Fibres

5.549 Views from the experts appear to be mixed. Dr. Henderson quotes Harrison’s review stating that PVA and aramid fibres are too large to be respirable. Dr. de Klerk states that all substitutes except glass (cellulose, aramid, PVA) produce a larger proportion of non-respirable fibres than chrysotile but respirable fibres are similar for all substances. Dr. Musk offers no opinion. Dr. Infante states that PVA fibres are “mostly” in the size range 10-16 µm and aramid fibres 10-12 µm. However, he notes, quite correctly, that, as mentioned below, the aramid fibres can and do split into fibrils of about 0.2 µm in diameter.

5.550 In assessing fibre respirability, none of the experts accounts for the fact that respirability depends on density as well as fibre diameter. The densities of PVA and paraaramid fibres are both considerably less that that of chrysotile. This means that much larger diameter substitute fibres would be respirable. In fact, the upper limits of diameters that are respirable for these fibres, as reported by Harrison306, are approximately 7 µm and 6-7 µm respectively. The equivalent upper level diameter for chrysotile is about 3-3.5 µm. Thus, fibres of much greater diameter can penetrate into the alveolar region of the lung. A review of the available information in the literature is that there is a general opinion without data that the respirable fraction of PVA fibres is small. However, there do not appear to be any data on the dimensions of airborne fibres during mixing with cement or other materials, or as released from the products during processing and use.

(iii) Glass and Cellulose Fibres

5.551 As far as cellulose and glass fibres are concerned, none of the experts provided any actual measurement data on the sizes or respirability of the fibres. Also, the dimensions of fibres at various stages of the processing, use and disposal of cellulose have not been reported. The actual dimensions of fibres in the airborne dust will depend on the specific glass fibres used and how they were prepared.

(iv) Biopersistence

5.552 It is well known that biopersistence is a key parameter. Indeed, the human evidence for chrysotile indicates that it is likely to be one of the main reasons why chrysotile is less dangerous than the amphiboles in respect to mesothelioma risk. This is clearly recognized by three of the four experts, as well as by INSERM.307

(v) Cellulose

5.553 Drs. Infante, Henderson and de Klerk recognise that cellulose is durable in the lung. In fact, the data show that some cellulose fibres have half lives of about 1000 days in the lung, which are many times longer than even those published data for amphibole fibres, much less chrysotile fibres.308

(vi) PVA

5.554 Dr. Musk and Dr. Henderson had no comment on PVA durability. Dr. de Klerk presented no data, but expressed the view that PVA was less durable than chrysotile. Davis, in a review in 1998, found no published data on the biopersistence of PVA fibres. Data was not published until 1999, when Harrison [1999] reported that PVA “will degrade very slowly, if at all in the lung.”309 There do not appear to have been any systematic studies of the biopersistence of PVA fibres, a crucial parameter in assessing the hazard associated with PVA fibres.

(vii) Para-aramid Fibres

5.555 Based on a study by Searl,310 who compared chrysotile and para-aramid fibres, the general view of the experts is that para-aramid fibres are less biopersistent. However,
Searl failed to check the lung tissue to confirm that the retained fibres were chrysotile. Based on studies using a standard protocol, Dr. David Bernstein has found that the biopersistence of chrysotile is in fact less than that of para-aramid fibres.311

(viii) Glass Fibres

Dr. Musk, de Klerk and Henderson presented no data on the biopersistence of glass fibres. Dr. Infante, without identifying the specific glass fibres, reports that glass fibres are less biopersistent than chrysotile. In fact, the recent work by Dr. Bernstein, in which the same protocol was used as for synthetic fibres, found that long [i.e.: > 20um] pure chrysotile fibres are removed from the lung faster than most, if not all, of the glass fibres reported in the published literature.312

(ix) Chrysotile

As far as chrysotile is concerned, it is well accepted that chrysotile is readily removed from the lung. This is why the lungs of chrysotile millers and miners, exposed to chrysotile, have been found at autopsy to contain more tremolite (an amphibole asbestos mineral) than chrysotile.313 The chrysotile cleared, but the tremolite fibres remained in the lung because of their much greater biopersistence. There are various estimates of the half time for chrysotile clearance. Oberdöerster314 studied baboons and estimated a 90-110 day half time for chrysotile fibres. The study by Searl was mentioned above. The estimates by Dr. Bernstein are even shorter (< 10 days).315 For direct comparison purposes, the clearance rates for fibres in the same ranges of dimensions must be studied. In addition, it is crucial that the fibres be tested using the same methodology. The studies by Bernstein best fit these criteria and show that, size for size, chrysotile has a very short half-life.

Question 6(c)

(i) Exposure-Response

In the absence of exposure-response data, it is not possible to quantify the risks associated with the various fibres. The question to be addressed is not: is one material more dusty than another? Nor is it: is the concentration higher when working with one material compared to another? Rather, the question that must be asked is: what is the risk for workers manufacturing or using the product? The decision on which fibre is safer has to be made on the basis of an assessment of the risk of disease for workers when manufacturing and using a product containing chrysotile compared to that when manufacturing and using the same product containing the substitute when subjected to the same or equivalent handling.

Three sources of data might be considered: experimental animal studies; human (epidemiological) studies; and in vitro studies. The latter (in vitro) are of little value for estimating risk as they only involve tests of, for example, biological activity in cells isolated from the processes which occur in a complete organism. Thus, they are an inadequate basis of comparison to the effects of inhalation on animals, much less humans.

(ii) Animal Studies

The first approach involves the exposure of animals to fibres of well-defined characteristics and concentrations by inhalation and following them for their lifetimes. Such studies have been done for a wide range of synthetic mineral fibres. Problems with this approach are many, as has been demonstrated in the considerable work done in recent years on synthetic mineral fibres. First, the animal species may have a limit on the size of fibre that it can inspire. Second, there are marked differences in the sensitivity of different animal species. For example, a refractory ceramic fibre (RCF) which produced one or two mesotheliomas in rats, produced mesotheliomas in 40 per cent of the hamsters exposed. Third, the lifetime of rats is about two years. In order to produce an effect within the lifetime of the animals they are subject to enormous exposures. Such exposures can pro-
duce the abnormal situation of lung overload so that the real reason for any biological effect is not clear. Fourth, an animal must produce an effect within two years (before it dies of natural causes). If biopersistence is important, fibres that are readily removed in humans over the course of a human’s life are not removed from the lung of an experimental animal because of the high exposures and shorter life. Fifth, the interpretation of much of the experimental work must be done with caution, because until recently, fibre exposures were reported on mass not number basis. As the materials tested can have quite different dimensions, the same mass can lead to exposures involving considerably different concentrations of fibres on a number basis.

(iii) PVA

5.561 There are no studies relating the long-term effects of exposure to PVA fibres.

(iv) Cellulose

5.562 Studies that have been done with cellulose have shown that it initiates a severe inflammatory response and fibrosis. Unfortunately, no chronic exposure data have been published.

(v) Glass Fibres

5.563 While there have been many studies of glass fibres, the only study in which the same methodology was applied to the study of synthetic mineral fibres and chrysotile asbestos is that of Hesterberg. He found that while there is an increased risk of lung cancer identified at high concentrations (as for glass and other fibres) the animal data suggest that at low level exposure, the risk associated with the chrysotile exposure is considerably less than that associated with the synthetic mineral fibres tested.

(vi) Para-aramid Fibres

5.564 While in recent years the information base concerning aramid fibres has increased greatly, there remain several issues. The only data are those derived from experimental studies in animals. While studies of biopersistence suggest that long fibres are shortened by enzymes in the lungs of animal experiments (Searl) and hence removed from the lung, the situation in humans is not known. Two researchers (Davis and Pott) have produced mesotheliomas by intra-peritoneal injection of these fibres, so their potential to produce mesotheliomas cannot be dismissed. The interpretation of “proliferative keratin cysts” observed during inhalation experiments remains unclear. Minty et al., in a criteria document for an occupational exposure limit (OEL) in the UK, summarized what was known about the para-aramid fibres at that time and drew several parallels with chrysotile. For example they state that “[T]he balance of evidence suggests that respirable aramid fibres possess a low potential to produce mesothelioma which is likely to be at least as low as with chrysotile.”

5.565 Referring to chrysotile, they conclude that mesothelioma “would only be detectable following very heavy and prolonged exposures.” The recent evidence that the mesothelioma risk for chrysotile miners and millers is associated with tremolite, will render the threshold of mesothelioma for downstream workers even more remote. These authors considered a clear no-effect level of 2.5 f/ml for pulmonary toxicity and a recommended OEL of 0.5 f/ml to allow for “uncertainties in interspecies differences.”

(vii) Epidemiological Data

(i) PVA

5.566 Drs. Musk, de Klerk, Henderson and Infante did not identify any epidemiological studies of PVA fibre workers. In fact, there is one study involving a small number of PVA fibre production workers (about 400 exposed employees). Even though the length
of exposure thus far is quite short, already two lung cancer deaths have occurred in the
cohort to date. Clearly a much longer follow-up is needed. Regarding mesothelioma, it
must be noted that with such a small population, even if half were dead and there were
one mesothelioma, the risk would be 0.5 per cent which is more than the risk of mesothel-
lioma found in Quebec miners and millers exposed to tremolite contaminated chrysotile.
(Also, the Panel should note that there were no mesotheliomas among 1267 deaths in
chrysotile exposed friction product manufacturing workers exposed to chrysotile.) Thus,
this study cannot detect either mesothelioma or lung cancer risks as low as at that already
known for chrysotile. Clearly, there are no human data on which to assess risk to conclude
that the risk is less than it is for chrysotile either per f/ml of exposure or globally from
work with products manufactured using PVA fibres.

(ii) Cellulose

5.567 Dr. Infante states that there are three studies in which cellulose exposures have
been investigated, but he does not identify them. The other two experts do not suggest
any epidemiological data. Studies in which there is no overall increase in mortality from
lung cancer are not adequate to investigate the risk of exposure. To assess this risk, the
relationship between lung cancer and cellulose fibre exposures on a per fibre basis must
be, but has not been, examined.

(iii) Para-aramid Fibres

5.568 None of the experts reported any epidemiological data. Clearly, para-aramid
fibres can be inhaled, as experimental animals inhaled them. However, because para-aramid
fibres have been used for such a short time, there are no data on the relationship between
levels of fibre exposure and the risks of lung cancer, mesothelioma or other adverse effects
for persons working with this substitute or products manufactured using it.

(iv) Glass Fibres

5.569 There have been several studies of workers exposed to glass fibres during fibre
manufacture. Studies have also included rock [stone] and slag wool exposures. The latter
were associated with an increased risk of lung cancer even at very low levels of exposure.
Doll324 concluded that the risks from such exposures were greater than those associated
with chrysotile asbestos. Doll summarized the situation as follows: "an occupational haz-
ard of lung cancer has been demonstrated in the rock and slag wool section of the industry
and possibly the glass wool section." The human evidence since that time has not dis-
pelled concern about the risks associated with these fibres. This question is still not re-
solved.

5.570 Dr. Infante325 and co-workers once reached the same conclusion for glass fibre
(although he has changed his opinion in his current report). In his report, Dr. Infante men-
tions that after speaking with workers, he now thinks that there was asbestos exposure at
the plant studied by Shannon in Ontario, Canada, where a high level of risk of lung cancer
was found in glass fibre workers. A recent discussion with Dr. Harry Shannon about his
study of glass fibre workers reveals that to his recollection, no one had raised the question
of asbestos as a potential confounder in his study. He noted that, as the study was pub-
lished many years ago, it seems unlikely that this issue – if it actually existed – would not
have been raised and studied, especially by the glass fibre industry.326 Clearly, no new
analyses have been done, so the impact of supposed asbestos exposure, if it took place, is
not known. Dr. Infante’s reversal of opinion does not seem justified, as no new data are
presented. For example, it is not known whether the “asbestos exposed workers” had
high or low glass fibre exposure. If they had low glass fibre exposure, then the risk associ-
ated with the glass fibre exposures might increase. Thus, without additional analyses, the
best estimates at present are the original analyses by Shannon.327

5.571 It was noted earlier that exposure levels during production may not be the same
as those during product use. While it has not been possible to find data on the use of glass
fibres in chrysotile cement or friction products, there has been an estimate of the risk associated with glass fibres as installed in homes. In this study, Wilson\textsuperscript{328} used animal data to derive lung cancer risk estimates for glass fibre exposure. They assumed an exposure of 1f/ml for one year based on available data and estimated that the lung cancer risk in smokers associated with blown glass wool without binder in a smoker without a respirator would be $2.4 \times 10^{-4}$. If one uses the same methodology as applied by them to derive a chrysotile estimate (based on epidemiological data), but for friction product manufacture, the risk would be very much lower: $0.12 \times 0.00058 = 0.00007$ or $7 \times 10^{-5}$. This is a lower risk than calculated for glass fibres. In fact, there is no demonstrated increased risk of lung cancer in the friction industry, so even this chrysotile risk is hypothetical and certainly an overestimate. Wilson acknowledges this in their paper.

5.572 In this light, it is safer to work with chrysotile in friction products than to work with glass fibres. While it might be argued that there has been no report of an increased risk of mesothelioma in humans as a result of manufacturing glass fibres, in the case of chrysotile, there is greater confidence concerning this lack of risk because there is no evidence of an increased risk of mesothelioma associated with friction products throughout their lifecycle, and the studies are far more voluminous and varied in approach. There are no systematically gathered data available concerning downstream risks for glass fibres as used as a substitute in cement or friction products. Similarly, with regard to the asbestos cement industry, Harrison\textsuperscript{329} reports that most studies have not found an increase in mesothelioma: certainly this is true for chrysotile asbestos cement plants. Thus, it is evident that there are clear no epidemiological or experimental data to conclude that “glass fibres” are safer than chrysotile, indeed, there is evidence to suggest the contrary.

5.573 In summary, the experts have based their opinions on very limited, if any, data. The data that do exist suggest that the conclusions of the Panel’s experts concerning the relative safety of the substitutes and chrysotile at low concentrations are incorrect.

2. The European Communities

(i) Introduction

5.574 Each of the four scientific experts appointed by the Panel has recently responded to the points which the Panel wished to clarify. The European Communities note that the four experts consulted unanimously and unambiguously corroborate the analysis that led France to adopt the Decree 96-1133 banning asbestos. This analysis was communicated to the Panel in the two written submissions of the European Communities of 21 May and 30 June 1999 and is based on the following points:

(a) All forms of asbestos, including chrysotile asbestos, are carcinogens, and there is no scientifically established threshold below which exposure to asbestos would be without risk for humans;

(b) exposure to asbestos, including chrysotile asbestos, is the cause of many cancers, the vast majority of which affect secondary users, particularly workers coming into contact with materials containing asbestos, including asbestos cement;

(c) so-called “controlled” use of asbestos is in fact impossible in practice;

(d) there are asbestos substitutes which are far less dangerous for human health.

5.575 In this document, the European Communities do not wish to make systematic and detailed comments on all the replies by the four experts consulted, but will simply refer to the main conclusions and give a summary of their replies in the annex.\textsuperscript{330}
The four experts consulted agree that all types of asbestos, including chrysotile, are carcinogens and that there is no established threshold under which exposure to asbestos is without risk for humans.

5.576. The four scientific experts unanimously consider that chrysotile asbestos, as well as amphiboles, can cause mesothelioma and lung cancer *inter alia*.

5.577. The four experts also unanimously agree that there is no scientifically established threshold below which exposure would not pose any risk of cancer for humans. All the experts state that the risk of cancer is proportional to the cumulative level of exposure and all consider that the non-threshold linear model is the most scientifically appropriate model for guaranteeing the level of health protection decided upon by France in this particular case. This explains and confirms that the non-threshold linear model has always been used, without exception, by the authorities in all those countries that have so far carried out scientific assessment of the cancer risk.

The four experts consider that exposure to asbestos, including chrysotile asbestos, is the cause of many cancers that mainly affect secondary users, particularly workers in contact with materials containing asbestos, including asbestos cement.

5.578. The four experts consider that the vast majority of the risks concern so-called "secondary" users, in other words, workers making interventions (building workers, electricians, plumbers, maintenance workers, handymen, etc.) because of their large number and the nature of their activities, even if the individual risks are sometimes lower.

5.579. For example, most cases of mesothelioma now affect this category of workers in all the industrialized countries, including Canada (Quebec) and Australia, countries which produce asbestos. The four experts point out that the levels of exposure in the course of occasional contacts with asbestos cement products are very high, much higher than the levels at which a risk of cancer has been definitely and scientifically established.

The four experts consider that the so-called "controlled" use of asbestos is not practically possible.

5.580. The four experts unanimously agree that so-called "controlled" use aimed at ensuring a constantly low level of release of the fibres into the atmosphere is absolutely impracticable in the vast majority of work situations where workers have to deal with friable or non-friable materials containing asbestos.

5.581. The four experts consider that it might be possible in very special situations where a small number of workers carry out a very precise task. They also indicate that interventions on materials such as asbestos cement can release very large quantities of asbestos fibres; that protective equipment is not or not always effective and not always used; that the recommended procedures are rarely or incorrectly followed in small enterprises such as those in the building sector; that it is quite impossible to apply them to non-professionals (for example, handymen, etc.).

**E. Supplementary Remarks from Dr. Henderson**

1. **Concerning the comments from the European Communities**

5.582. The comments from the European Communities are very brief, occupying only four pages in English translation, so that only a short comment is needed. The tabular summary of the four experts' reports appears to represent a fair précis of my conclusions and opinions, if an oversimplification. In para. 5.579, the European response refers to Canada (Quebec) and Australia as countries that produce asbestos. As pointed out in para 5.27, Australia is no longer an asbestos producer.
Concerning the comments from Canada

5.583 At 62 pages and with over 50 Annexes, the comments from Canada are far lengthier than the response from the European Communities; the Canadian documents include new information, necessitating more extensive discussion. Some general comments follow; other issues are discussed later under specific sub-headings.

5.584 In para. 5.441, the comment is made that some of the answers from the experts “appear not to distinguish between chrysotile and amphibole exposure” or between “modern uses ... and historical uses ... .” Throughout my own Report, I tried to make this distinction wherever appropriate, and my answers to the Panel’s questions deal almost exclusively with chrysotile (like EHC 203 [1]) - e.g. my discussion of the risks to brake mechanics and the tabulation of risk estimates for lung cancer and mesothelioma (Tables 12 and 13 in paras. 5.203 and 5.205. At the same time, it seems worth reiterating that commercial chrysotile from Canada on average contains variable trace amounts (about <1 per cent) of tremolite (fibrous tremolite is a non-commercial amphibole; e.g. please see EHC 203). In relation to Canada’s concerns about the “experts’ conclusions on tradesmen” (e.g. building construction workers), my perspective seems to concur with the IPCS/WHO monograph on chrysotile (EHC 203):

“... (c) Some asbestos-containing products pose particular concern and chrysotile use in these circumstances is not recommended. These uses include friable products with high exposure potential. Construction materials are of particular concern for several reasons. The construction industry workforce is large and measures to control asbestos are difficult to institute. In-place building materials may also pose risk to those carrying out alterations, maintenance and demolition” ... [p 144].

5.585 My recognition that chrysotile is substantially less potent than the amphiboles on a fibre-for-fibre basis for mesothelioma induction - and that present exposures overall are substantially less than in the past - explains why my Report dwelt mainly on workplace exposures (e.g. to building materials and friction products). Non-workplace exposures (e.g. non-occupational exposures including household contact or neighbourhood exposures [2-4]) - and exposures to friable insulation materials - received less attention and were included mainly to put the present situation into historical perspective.

5.586 In para. 5.489, the Canadian comments state that: “... He [Henderson] believes amphiboles may be greater than 60 times more likely than chrysotile to induce mesothelioma ... .” In fact, in para. 5.103, I had stated that:

“There is general though not universal agreement of a differential potency between the amphiboles versus chrysotile for mesothelioma induction; in this respect, the amphiboles are substantially more potent, with estimates ranging from 2-4X, to 10X, to 12X on a fibre-for-fibre basis, to 30X, to a 30-60X greater potency, or more (e.g. please see EHC 203).”

5.587 Later in my Report (paras. 5.141 and 5.413), I gave my estimate that chrysotile has a potency 1/10th-1/30th the carcinogenicity of crocidolite for the mesothelium. This estimate has not changed.

5.588 Some clarification of my answer to Question 1(b)334 from the Panel seems necessary, taking into account Canada’s quotation of my view and the comment that: “Canada takes note that ... [this is] ... overwhelmingly a workplace issue, and therefore not related to the ‘handyman’.” In my answer, I took “workplace” to refer to any situation where work of any type is carried out (e.g. cutting, sawing, drilling, grinding, rasping or sanding of asbestos-cement building products), whereas I interpreted the expression “a larger part of the population” to refer to general environmental exposure to asbestos (e.g. simple occupancy of buildings, or potential exposure of urban dwellers in general to asbestos derived from the brakes of passing vehicles). Obviously, any risks to handymen who carry out
maintenance on homes only occasionally will be much less than the risks to professional tradesmen such as carpenters, who work day in, day out at building construction sites - because the frequency and duration of exposures for the handyman will be less (with lower cumulative exposures), assuming the types of asbestos to be the same. However, this may not always be so. For example, I know that some “handymen” in Australia specialize in buying and living in dilapidated houses, to carry out extensive renovations on these dwellings (e.g. throughout each weekend or more often) before selling them a year or more later; because the house qualifies as a principal place of residence, the profit is not taxable. The handyman then repeats this exercise on another “handyman special” house, and so on. Many such handymen also regularly carry out maintenance and renovation work on other homes, so that their exposures may approach those of professional tradesmen, but they style themselves as “home handymen”. The activities of such handymen are virtually unregulated because they are self-employed and a number work on a “strictly cash” basis.

5.589 In commenting on the experts’ responses to Questions 1(b), Canada reiterates the proposition that “chrysotile is readily removed from the lung”, and estimates of the half-life of chrysotile are given as 90-110 days, and even a shorter estimate of < 10 days from Dr. David Bernstein. Canada goes on to state that “size for size, chrysotile has a very short half-life.” I again draw attention to the 1999 study by Finkelstein and Dufresne [5], who found a lung tissue half-life of eight years for chrysotile fibres > 10 µm in length; this investigation was discussed briefly in the Endnote to my report (see Section V.C.4):

“... in the past, the kinetics of chrysotile clearance from lung tissue have been investigated mainly in experimental models using rodents. In an autopsy study published in 1999, Finkelstein and Dufresne [5] ... investigated clearance of chrysotile from the lung tissue of 72 Quebec chrysotile miners and millers in comparison to 49 control subjects, using regression analyses, with the following findings:

There was a significant association between the duration of occupational exposure and the tissue burdens of chrysotile and tremolite.

The concentration of chrysotile decreased with time after exposure ceased but the concentration of tremolite did not.

The clearance rate varied inversely with the length of chrysotile fibres. For fibres > 10 µm in length - i.e. fibre lengths in the reported range for carcinogenicity - the clearance half-time was estimated to be eight years. In other words, the tissue bio-persistence of chrysotile fibres in this study seems substantially more prolonged than in rodent experiments, and presumably corresponds to persistent high chrysotile fibre concentrations for many years after cessation of occupational exposure in humans, as discussed on p 31. It is also notable that the concentration of 6,250,000 chrysotile fibres mentioned on p 31 (for an individual but by no means unusual patient) is probably above the level at which Rogers et al. [6] identified an odds ratio for mesothelioma of > 8.5 (even allowing for differences in fibre size in the counts by the two different laboratories), and even the duration of 16 years after exposure stopped (as opposed to its commencement: 24 years) falls into the lag-time range for lung cancer induction by asbestos.

Studies like this suggest that clearance mechanisms can be overwhelmed and break down at occupational levels of exposure in humans, with the existence of a long-term sequestered fraction of chrysotile fibres.”

5.590 This study seems to be of particular significance for the tissue bio-persistence of chrysotile fibres in comparison to substitute materials (please see below, paras. 5.642 to 5.652).

5.591 I also emphasize that some of the estimates given in my Report were conservative, with potential under-estimation of effects. For example, after discussing the inci-
It is mentioned that the incidence rate for spontaneous mesothelioma unrelated to asbestos is in the range of 1-2 mesotheliomas per million person-years - whereas the likely true figure is probably less than one [4] - I nonetheless used the upper figure of two cases/million for comparison with mesothelioma incidence in some occupational groups (e.g. the incidence of mesothelioma among male automobile/brake mechanics in Australia; please see para. 5.253). In a similar way, I referred to a 30-fold differential rate for lung cancer among the South Carolina (Charleston) chrysotile textile workers in comparison to the Quebec chrysotile miners and millers, whereas others give the differential as up to "about 50 times higher in Charleston" [7].

5.592 I also draw attention to the occurrence of mesothelioma among various cohorts and studies other than the Quebec chrysotile miners/millers, as set out in paragraphs 5.124-5.141, and to the incidence of mesothelioma among mechanics in Australia as shown in the 1999 Report for the Australian Mesothelioma Register [AMR 99] and in NICNAS 99 (see my answer to Question 2).

5.593 In my Report, I discussed the limitations or deficiencies of those studies which reported an increased risk of lung cancer only among workers with pre-existing asbestosis (e.g. the Hughes-Weill study [8]) - and the uncertainties of the study by Camus et al. [9] on lung cancer risk from non-occupational exposure to chrysotile among females in Quebec [50] - but the comments from Canada (para. 5.529) reiterate the Hughes-Weill conclusion that an increased risk of lung cancer occurs "only in those with asbestosis." (Please see paras. 5.73-5.74 and 5.152-5.162 above; this subject was reviewed extensively by Henderson et al. [13] in 1997. Lung cancer risk among the South Carolina chrysotile textile workers versus the Quebec chrysotile miners/millers is also discussed in paras. 5.596 to 5.620 below).

5.594 At various points (paragraphs 5.475, 5.498 and 5.545), the comments from Canada quote de Klerk and Armstrong [14], in a chapter on The Epidemiology of Asbestos and Mesothelioma, in the book Malignant Mesothelioma, for which I was the senior editor and a co-author. I shall leave it to Dr. de Klerk to respond.

5.595 In passing, I point out that Malignant Mesothelioma was published in 1992; the text for those chapters which I wrote was current up to September 1990, and the manuscript was sent to the publisher shortly thereafter. Much new information on asbestos-related diseases has accumulated since that time (e.g. references 15, 16, 113, 125, 126, 131-133, 140, 141, 170-172, 177-179, 181, 185-187, and 190-194 in my Report, to list but a few). My views on many aspects of asbestos-related disorders have changed very substantially since Malignant Mesothelioma was published (e.g. my views on asbestos and lung cancer - please see references [13, 15-18] in these Supplementary Remarks).

(a) Lung cancer rate among South Carolina (Charleston) chrysotile textile workers versus the Quebec chrysotile miners/millers

5.596 With respect to this question, Canada states (see paras. 5.485-5.486): "Dr Henderson states that the "greater carcinogenicity of the amphiboles [...] appears not to extend to the induction of lung cancer [p 40], but he admits that 'chrysotile is implicated in one of the lowest rates of asbestos-associated lung cancer (in Quebec chrysotile miners and millers)’ [where I also stated that chrysotile is also implicated in the highest lung cancer rate]. Dr Henderson's reluctance to conclude the greater carcinogenicity of amphiboles seems to be caused by the results of Dr Dement's study of the Charleston, South Carolina asbestos textile industry [...]".

"The Charleston data has [sic] recently been revisited by Bruce Case, André Dufresne, A.D. McDonald, J.C. McDonald and Patrick Sébastien in a study released in Maastricht in October 1999 at the VIIIth International Symposium on Inhaled Particles, a symposium attended by some of the world’s leading experts. This study shows that a significant amount of crocidolite and amosite fibres was..."
found in the textile workers' lungs. This analysis sheds new light on the issue and explains the extreme results of the original study by Dr Dement [...] and the subsequent study by Dr Stayner [...]. These studies of textile workers exposed to crocidolite and amosite can thereby no longer be used to demonstrate the risks associated with chrysotile fibres."

5.597 Subsequently, the manuscript for a paper by Case et al. [19] entitled *Asbestos Fibre Type and Length in Lungs of Chrysotile Textile and Production Workers: A Preliminary Report* arrived by facsimile transmission. I offer the following comments on this document (and, later, on the Abstract for the corresponding presentation at the Maastricht meeting [20]):

5.598 A disclaimer beneath the title [19] indicates that this is a "DRAFT DOCUMENT: SUBJECT TO REVIEW - NOT TO BE CITED". It is cited nonetheless. There is no indication that this document has gone through a process of peer review and been accepted for publication.

5.599 This study revisits the study reported in 1989 by Sébastien et al. [7], and the draft manuscript indicates that the same grids were examined (but fewer cases). The main difference between this investigation and the earlier study by Sébastien et al. [7] is that Case et al. [19, 20] analysed long fibres > 18 µm in length, whereas Sébastien et al. [7] studied fibres > 5 µm in length, with an aspect ratio > 3:1. (It is common practice for fibre burden analyses to focus on fibres = 5 µm in length and there is no evidence that the carcinogenicity of asbestos fibres - in terms of lung cancer induction - is restricted only to fibres about 20 µm in length or more.)

5.600 Another study on the lung fibre content of the Charleston chrysotile textile workers was reported in 1997 by Green et al. [21]; this investigation studied all fibres resolvable by electron microscopy and with an aspect ratio > 3:1. For this study, lung tissue was analysed from 39 textile workers versus 31 comparable controls matched closely for age (median age at death for the asbestos workers was 56.0 years, versus 59.0 years for the controls).

5.601 In the Green et al. [21] study, the Charleston chrysotile workers had a higher lung content of chrysotile in comparison to the controls (geometric mean = 33,450,000 versus 6,710,000 fibres/g dry lung), with a higher content of tremolite (3,560,000 vs 0.260,000); the asbestos workers also had a slightly elevated mean amosite/crocidolite content of 470,000 fibres/g. 210,000 for the controls (please see Table 1).

**TABLE 1: MINERAL FIBRE CONTENT OF LUNG TISSUE, SOUTH CAROLINA ASBESTOS TEXTILE WORKERS VS. CONTROLS (ALL COUNTS = FIBRES X 10^6 / G DRY LUNG)**

<table>
<thead>
<tr>
<th></th>
<th>Textile workers</th>
<th>Controls</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Age at death (median; years)</strong></td>
<td>56.0 (M); 57.0 (F)</td>
<td>59.0 (M); 62.5 (F)</td>
</tr>
<tr>
<td><strong>Year of death (median)</strong></td>
<td>1971 (M and F)</td>
<td>1972 (M); 1971 (F)</td>
</tr>
<tr>
<td>Chrysotile (fibres x 10^6 / g dry lung)</td>
<td>33.45</td>
<td>6.71</td>
</tr>
<tr>
<td>Tremolite</td>
<td>3.56</td>
<td>0.26</td>
</tr>
<tr>
<td>Amosite/crocidolite</td>
<td>0.47</td>
<td>0.21</td>
</tr>
<tr>
<td>Anthophyllite</td>
<td>0.16</td>
<td>0.13</td>
</tr>
<tr>
<td>Mullite</td>
<td>1.63</td>
<td>4.01</td>
</tr>
<tr>
<td>Other</td>
<td>1.02</td>
<td>1.9</td>
</tr>
<tr>
<td><strong>All fibres</strong></td>
<td>52.46</td>
<td>16.02</td>
</tr>
</tbody>
</table>

*Modified from Tables 1 and 3 in Green et al. [21]; M = men; F = females.
In the discussion section, Green et al. [21] commented that:

“The population was exposed almost exclusively to chrysotile asbestos from Quebec. The native ore contained about 1% tremolite asbestos. The high concentrations of chrysotile and tremolite asbestos found in the lungs of the asbestos textile workers are also consistent with their exposure histories. Our finding on enrichment of tremolite relative to chrysotile in the lungs of the asbestos workers is in keeping with the use of small quantities of crocidolite between 1950 and 1975, but the values were only slightly greater than those found in the control population. The increased risk of lung cancer in the asbestos textile workers is also unlikely to be due to differences in exposure to tremolite asbestos, as Sebastien et al. have shown that the textile workers had less tremolite asbestos in their lungs than miners and millers of the original ore after matching for exposure intensity. Differences in exposure to other commercial amphiboles (crocidolite and amosite) may have played a small part based on our own data ... and on the data of Sebastien et al., which showed a small excess of these amphiboles in the lungs of the textile workers compared with the miners; however, it is very unlikely that this is the whole explanation as commercial amphiboles formed a very small proportion of the total amphiboles in both studies. Moreover, review of the 10 cases with lung cancer in this study on whom lung fibre analyses were made, showed only one case with substantially increased (> 1 x 10^6 fibre/g dry lung) crocidolite or amosite”.

In this study, it is also notable that the lung cancer cases on which fibre burden analysis was carried out were not representative of the cohort as a whole: e.g. autopsies were carried out on only about 10 per cent of all deaths in the cohort, and the mean lifetime cumulative exposure for the ten lung cancer cases was 94.6 fibre-years in comparison to 67 fibre-years for male lung cancer cases across the whole cohort [21, 22].

There are even greater concerns about the representativeness of the cases on which fibre burden analysis was carried out by Sébastien et al. [7]. For example, this study was confined to tissue from 72 autopsies among 857 deaths (8.4 per cent) among the Charleston cohort, and there were only seven lung cancer cases out of 66, whereas Case et al. [19] list 126 lung cancers, so that the fibre burden data reported by Case et al. [19] appear to deal with no more than 5.56 per cent of the Charleston lung cancers. It is also notable that the mean age at death in the Charleston group was about a decade younger than the age at death for the Thetford group which formed the basis for comparison in the 1989 study by Sébastien et al. [7]a

In addition, as reported by Sébastien et al. (see Table 3 in reference [7]), those cases from the Thetford group that came to autopsy showed an over-representation of asbestos-related diseases (lung cancer, mesothelioma and pneumoconiosis) than the Thetford cohort overall - so that cases of lung cancer + mesothelioma + pneumoconiosis added up to 37 out of 89 autopsies (42 per cent), in comparison to 306 out of 4463 deaths across the whole cohort (7 per cent) [7]. For the Charleston cohort, the figures were more comparable, so that lung cancer + mesothelioma + pneumoconiosis cases added up to 13 out of 72 autopsies (18 per cent) in comparison to 10 per cent across the cohort [7].

In the more recent study from Case et al. [19], there is a further point on which the two study groups (Thetford versus Charleston) are not comparable: the time following cessation of exposure was a median of eight years for the Thetford group, in comparison to 20 years for the Charleston cohort (please see Table 2). Therefore, it is clear that those lung cancer cases on which fibre burden analysis was carried out from each cohort were not representative of each cohort, and that there were also substantial differences between the two cohorts for the same types of case. Finally, the manuscript from Case et al. [19] does not include a control group against which the two cohorts can be compared (the only one of the three investigations that does is the Green study [21]).
TABLE 2: COMPARISON OF THE SOUTH CAROLINA AND QUEBEC CHRYSOTILE WORKER COHORTS*  

<table>
<thead>
<tr>
<th>South Carolina textile workers</th>
<th>Quebec miners/millers</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cohort number</td>
<td>3022</td>
</tr>
<tr>
<td>Cohort deaths</td>
<td>1258</td>
</tr>
<tr>
<td>Age at death (years)</td>
<td>67 ± 10 (??)**</td>
</tr>
<tr>
<td>Lung cancers in cohort</td>
<td>126 [SMR 197]</td>
</tr>
<tr>
<td>Mesotheliomas in cohort</td>
<td>2</td>
</tr>
<tr>
<td>Years since cessation of exposure (median)</td>
<td>20</td>
</tr>
<tr>
<td>Geometric mean exposure (mpcfy)*****</td>
<td>3.63</td>
</tr>
<tr>
<td>Subjects studied</td>
<td>64</td>
</tr>
<tr>
<td>Lung cancer cases studied</td>
<td>7</td>
</tr>
<tr>
<td>Chrysotile (fibres x 10⁶ / g dry lung)</td>
<td>0.054</td>
</tr>
<tr>
<td>Tremolite</td>
<td>0.027</td>
</tr>
<tr>
<td>Amosite/crocidolite</td>
<td>0.037</td>
</tr>
<tr>
<td>Total amphibole (tremolite + amosite/crocidolite)</td>
<td>0.064</td>
</tr>
</tbody>
</table>

*Modified from Case et al. [19]. Fibre counts represent geometric means, all expressed as fibres x 10⁶/g dry lung; **see footnote 160; ***mpcfy = millions of particles per cubic foot-years.

5.607 From the above Table, it is evident that the amosite/crocidolite content of lung tissue from the textile workers is slightly (< 2-fold) higher than the amosite/crocidolite in the lung tissue from the Quebec miners and millers (37,000 fibres > 18 µm in length versus 24,000). This difference in concentration seems to be insufficient to explain the "huge" [19] risk difference (about 30-fold) in the slope of the lung cancer dose-response line between the two groups. In addition, it is noteworthy that the tremolite content of lung tissue was higher in the Quebec miners and millers than the Charleston textile workers (325,000 versus 27,000 for fibres with a mean fibre length of 21.7 versus 21.9 µm). The point is that the total amphibole content (tremolite + amosite + crocidolite) is higher in the Quebec miners and millers at 349,000 f/g dry lung in comparison to a total amphibole content of 64,000 among the Charleston textile workers. In this respect, there is no evidence that tremolite is substantially less potent than the other amphiboles for lung cancer induction, as shown by the high lung cancer incidence (SMR = 285) among Montana vermiculite miners exposed only to tremolite/actinolite (please see paragraph 5.107-5.111).

5.608 From these studies, it appears that the amosite/crocidolite content of lung tissue among the Charleston textile workers may in part be a reflection of low level exposure to the small amount of crocidolite (< 1000 kg total) used in the plant from 1950-1975 to make an asbestos tape or braided packing. The material was received at the plant as a yarn ready for weaving, and no fibre preparation, carding, spinning or twisting was done using crocidolite. Packing workers were not included among the textile worker cohort, and analysis of lung cancer risk by operation in the plant shows all operations to be at about the same lung cancer risk after controlling for chrysotile exposure in a logistic model (Dement, personal communication, 1999).

5.609 A portion of the amosite/crocidolite content may also be explicable by general environmental (non-occupational) exposure, taking into account the small differences between the amphibole content in the textile workers versus the controls in the study reported by Green et al. [21]. In this respect, amphibole concentrations of up to 100,000-200,000 fibres per gram (f/g) dry lung tissue can be expected for about 5 per cent of the population in Germany [23]. Therefore, it seems that the amosite/crocidolite cannot explain the risk of lung cancer in the Charleston cohort in comparison to either matched controls (also matched for smoking) versus the Thetford miners and millers.
5.610 If major significance is to be assigned to the small difference in amosite/crocidolite content of lung tissue between the Charleston workers versus the Thetford miners/millers for lung cancer induction, a question that immediately arises is: *where are the mesotheliomas among the Charleston workers?* Case et al. [19] suggest that misclassification of mesotheliomas as lung cancers among the Charleston workers could have produced under-estimation of the true number of mesotheliomas “while having virtually no effect on the lung cancer excess or lung cancer exposure-disease slope of risk.” No evidence in support of this proposition is adduced, and Case et al. [19] state that this is “speculation.” The larger number of mesotheliomas in the Quebec cohort may be explicable in part by the higher mean total amphibole content for this group, but this still leaves unexplained the disproportionately larger numbers of lung cancers in the Charleston group (e.g. the ratio of lung cancers to mesotheliomas in the Thetford group is $657/38 = \approx 17:1$, whereas the ratio for the Charleston group is $126/2 = 63:1$).

5.611 Case et al. [19] are also rather more cautious in their interpretation than the propositions put forward in Canada’s responses to the reports from the experts. For example, on the last page of text they state:

“... comparison of groups of individuals using this technique is valid only insofar as those studied are representative of the larger groups ... from which they are derived. We cannot be certain to what degree our groups of chrysotile miners/millers and textile workers are representative of the cohorts from which they are derived... the two groups are not directly comparable in some ways: not only was exposure much higher in the miners/millers, but the interval between cessation of employment and death was shorter ... . Our results closely parallel those reported by Sébastien et al. ... Any other result would be surprising since the subjects were drawn from the latter study. ... Caution remains in interpretation. ... One continuing mystery, given the apparent non-trivial long-fibre commercial amphibole exposures is the low level of reported mesotheliomas in this cohort ...”

5.612 Given the data on fibre lengths across the cohorts, in comparison to the data in Sébastien et al. [7], the difference in lung cancer rates between the two groups cannot be explained by differences in fibre length. This is stated explicitly by Case et al. [19].

5.613 However, on looking at the data, it seems that the differences between the two cohorts might be explicable in part by the exposure estimates. Differences in exposure assessment are not refuted by the “new” study reported in draft form by Case et al. [19] or by the earlier study reported by Sébastien et al. [7]: e.g. the difference between 20 years (Charleston) and eight years (Quebec) for clearance after exposure ceased could have a large effect. One can calculate the final exposure (end of exposure) ($N_0$) from the final fibre content in lung tissue at death ($N$), from the equation

$$N/N_0 = e^{-\hat{e}t}$$

where $\hat{e}$ represents a clearance coefficient ($\hat{e} = 0.693 / T_{1/2}$) and $t =$ half-life in tissue ($T_{1/2}$). For $T_{1/2} = 8$ years [5], $\hat{e} = 0.693/8$, so that for the chrysotile miners/millers, where $N = 0.231$, $N_0 = 0.462$. For the Charleston textile workers, where $N = 0.054$, $N_0 = 0.306$.

5.614 If $T_{1/2}$ is shorter (e.g. one year), then $N_0$ for the miners/millers = 59.2 and the corresponding $N_0$ for the textile workers = 56456.

5.615 Therefore, for a half-life of eight years, one would expect the ratios of exposure (exposure miners/millers ÷ exposure textile workers) to be 0.462/0.306 = 1.5. For a half-life of one year the ratio becomes (exposure miners/millers + exposure textile workers) 59.2/56456 = 0.001. (For tissue half-lives of 90-110 days or < 10 days, the differences would be even more drastic.) However, the ratio of the estimated exposures (mpcfyQuebec/
mpcfy_{Charleston}) is \( \frac{186}{3.63} = 50 \), suggesting that one or other particle count estimate is incorrect.

5.616 In this respect, it might be argued that the exposure estimates for the Charleston cohort represented an under-estimation of exposure, but this suggestion is not supported by the low tremolite content in the lung tissue of the Charleston workers, and is explicitly rejected by Sébastien et al. [7], who state (p. 187):

“The hypothesis of a systematic underestimation of exposures to asbestos in Charleston, which would have accounted for the difference in risk, must therefore be rejected and other explanations sought.”

5.617 Given that contamination of the Charleston chrysotile by mineral oils has now been excluded, one possibility that remains is over-estimation of the exposures for the Quebec chrysotile miners/millers (with under-estimation of risk). If this explanation is unsustainable, it follows that the paradox remains, it remains unexplained, and seems likely to remain so.

5.618 Finally, I draw to the attention of the Panel the following comment by Case and Dufresne [20] in the Abstract for their presentation at the Maastricht meeting:

“Risk assessment for asbestos exposure is based on lung cancer risk for textile workers, rather than miners/millers.”

5.619 In the draft manuscript, Case et al. [19] state only that:

“... suggestions that the textile worker mortality data [are] suitable for chrysotile risk assessment [for lung cancer] should be re-evaluated ... .”

5.620 Therefore, even if one accepts this proposition for the moment, the claim that the South Carolina cohort can “thereby no longer be used to demonstrate the risks associated with chrysotile fibres” goes beyond the data in this study. For the reasons discussed in this section, I conclude that the data in Sébastien et al. [7] and in Case et al. [19] do not detract from the conclusions drawn by myself and other authorities from the investigations carried out the South Carolina cohort by Dr. Dement and his colleagues [22, 24].

(b) The question of a threshold for the carcinogenicity of chrysotile (lung cancer and mesothelioma)

5.621 On this question, I simply reiterate EHC 203:

“Exposure to chrysotile asbestos poses increased risks for asbestosis, lung cancer and mesothelioma in a dose-dependent manner. No threshold has been identified for carcinogenic risks” [p. 144].

5.622 In the absence of a threshold or an agreed alternative (non-linear) exposure-response model, the linear relationship model is widely employed for risk assessment at low levels of exposure.

5.623 As indicated, the precision or validity of this model is not known at low levels of exposure and, as stated by Dr. de Klerk, the model provides a “conservative estimate”. This is the point: in the absence of direct observational data or credible alternative models, the linear model errs - if it does err - on the side of safety, which is appropriate for risk assessment as a prelude to the formulation of occupational health and safety and public health policy. The principle is: if there is doubt, play safe (i.e. first, do no harm; primum non nocere).

5.624 In relation to prudent approaches to occupational and public health policy, The Minerals and Metals Policy of the Government of Canada™ states the following (p 7):

828
“The precautionary principle is an important factor when the Government needs to make a decision in the face of scientific uncertainties about cause and effect, and when the potential environmental consequences are generally considered to be serious or irreversible. This principle was enunciated clearly as Principle 15 in the 1992 *Rio Declaration on Environment and Development* (the Rio Declaration) of the United Nations Conference on Environment and Development (UNCED), to which Canada is a signatory:

‘Where there are threats of serious or irreversible damage, lack of full scientific certainty shall not be used as a reason for postponing cost-effective measures to prevent environmental degradation.’

The principle complements science-based approaches for the management of risks. Its use is premised on the recognition that our scientific understanding of the potential magnitude and consequence of impacts on human health and the environment of the production and uses of some minerals and metals may be incomplete. While there is a need to work toward closing such gaps in our understanding, there is also a requirement, where potential impacts are ‘serious or irreversible’, to consider a cost-effective precautionary approach.”

5.625 Later, on page 12, the same *Minerals and Metals Policy* document states:

“... It is generally accepted that, in some cases, the risks associated with certain products or product uses cannot be properly controlled or managed. Consequently, where such a situation exists, the Government [of Canada] will either discontinue or prohibit the specific product or product use.”

5.626 Three additional points are worth iteration:

- Exposure to commercial Canadian chrysotile is not “chrysotile-only”, but usually chrysotile + trace tremolite exposure, although evidence indicates that chrysotile when uncontaminated by tremolite also has the capacity for the induction of lung cancer and mesothelioma.

- Risk estimates for lung cancer and mesothelioma for low levels of chrysotile exposure were set out in Tables 12 and 13 (see my response to Question 1(d)).

- As stated in section C.1(f)(viii) and in paragraph 5.95, there are no observational data on the interactive effects of inhaled commercial chrysotile fibres when these are superimposed separately and later upon a pre-existing amphibole + chrysotile burden within lung tissue (?superimpositional additive or multiplicative carcinogenic effect). In my Report, I emphasized that it has been estimated that up to 15-20 per cent of men in industrialized societies may have sustained occupational exposures to asbestos (chrysotile/amphiboles), and Rödelsperger et al. [23] indicate that fibre concentrations of 100,000-200,000 amphibole f/g dry lung tissue may be expected for about 5 per cent of the population in Germany. Rödelsperger et al. [23] have also identified a dose-response relationship for mesothelioma induction at these low fibre concentrations. We do not know what the effect of subsequent chrysotile fibre inhalation superimposed upon an existing amphibole burden of this order might be, but NICNAS 99 states the following (p. 61):

  “... multivariate analysis of cases found a dose-response relationship for lung fibre content of crocidolite, amosite and chrysotile and the development of mesothelioma. Either a multiplicative or additive model could be used to fit the
relative risk/dose coefficients for the various asbestos types. A progressive increase in relative risk with increasing fibre content was reported for all fibres ...

5.627 Because the risks of both lung cancer and mesothelioma show a dose-response effect related to total cumulative exposure levels, it may be expected that later superimpositional inhalation of chrysotile±tremolite fibres would aggravate the overall consequences of a pre-existing asbestos burden (i.e. increase the risk further).

(c) The feasibility in practice of “controlled use” of chrysotile asbestos

5.628 In para. 5.510, Canada identifies the feasibility in practice of “controlled use” of chrysotile asbestos as “one crucial issue, which seems to override all other issues” (i.e. the question of whether the application of the “controlled use principle” is feasible and credible at all stages in the life cycle of chrysotile asbestos).

5.629 As already indicated (see my reply to Question 5), I agree that this proposition is crucial to the dispute before the WTO. However, for the reasons discussed in my Report, I see no requirement to resile from my perception that - although regulation and control of chrysotile and high-density chrysotile products may be achievable at some points of the life-cycle (e.g. the manufacture of friction and high-density products) - “controlled use” of this type is not feasible in reality or in practice at others (e.g. in building construction and other points of end-use).

5.630 No airborne fibre measurements are available for the overwhelming majority of asbestos-related diseases encountered in my everyday practice, even for exposures throughout the 1970s and in some cases extending into the late 1980s. Among my series of asbestos-associated lung cancers and mesotheliomas, I cannot recollect ever seeing actual airborne dust and fibre measurements at points of end-use (e.g. at building construction sites or shipyards).

5.631 The concentration of total asbestos fibres in lung tissue from one of my cases of asbestos-associated lung cancer was up to 125,000,000 f/g dry lung (up to 108,000,000 amosite + crocidolite fibres), for a worker who had been employed at a major asbestos-cement manufacturing facility for about 2-3 years (lag-time = 28 years) [15]. I also note Dr. de Klerk’s comment about demolition of an old asbestos-cement factory in Sydney in the latter half of 1999 (probably the same factory) “where no observable precautions of any kind were being taken”339 (Dr. de Klerk, response to Question 1(a)).

5.632 Other interventions on high-density asbestos-cement materials that can lead to high fibre concentrations are discussed in my first Report (e.g. Kumagai et al. [25]; my answer to question 1(d); please see also the 1980 report by Rödelsperger et al. [26] on exposure to asbestos-cement dust at building sites, which refers to a daily mean airborne fibre concentration of 0.6 f/ml for fibres > 5µm in length, and “peak concentrations of more than 100 fibres/ml”).

5.633 In para. 5.532, the comments from Canada include the statement that chrysotile “can be painted without fibre release” (presumably including building products). However, painting of such products can cover warning notices and disguise the true nature of the product, so that workers who later carry out maintenance or renovation work on the same product - and those who recycle the same material - may be unaware of its true nature. In my own series of mesotheliomas, it is not uncommon to encounter cases for which the patient was unaware or unsure that he (or less often she) had worked in the past with an asbestos-containing product.

5.634 In one recent case, the patient worked (1973-1988) at a factory where tins and pails were produced. In about 1979 she had worked for several months at a conveyor belt that carried the tins and pails into a fan-forced oven, which appears in retrospect to have been lined by asbestos-containing insulation. The patient was present when maintenance work on the oven was carried out, and she recalled hot air continually blowing from the
oven into her face as she worked on the conveyor. After diagnosis of her mesothelioma and its treatment in the late 1990s by radical pneumonectomy, an asbestos body and fibre analysis on her lung tissue revealed a count of 1,640 asbestos bodies per gram dry lung, and a total asbestos fibre count of 34,120,000 f/g dry lung (30,770,000 chrysotile fibres + 3,350,000 crocidolite fibres). This was the only history of exposure that was obtainable on exhaustive questioning.

5.635 A similar history was also obtained in another mesothelioma case seen on referral in 1999, where a radio assembly worker had used asbestos-containing cloths used to clean soldering irons, together with a history of about one fibre-hour of exposure to four asbestos-cement building sheets used for maintenance work on his home; only later did I discover that during his work at the radio factory, he often entered a walk-in fan-forced oven apparently lined by insulation bricks.

5.636 Again, please see the spread of occupations in AMR 99 attached to my Report; a similar spread of occupations is listed by Hodgson et al. [27] in a 1997 report on mesothelioma mortality in Britain - e.g. see Table 1 and Fig 1 in the original reference. In footnote 96 to the comments from Canada, it is stated that:

“The ‘controlled use’ approach has been endorsed by the WHO in its 1998 Environmental Health Criteria 203: Chrysotile Asbestos, p. 144. ‘Control measures, including engineering controls and work practices, should be used in circumstances where the occupational exposure to chrysotile can occur. Data from industries where control technologies have been applied have demonstrated the feasibility of controlling exposure to levels generally below 0.5 fibres/ml. Personal protective equipment can further reduce individual exposure where engineering controls and work practices prove insufficient.’”

5.637 I interpret this passage from EHC 203 differently when it is taken in the context of the preceding paragraphs; apart from the heading, the complete text on page 144 of EHC 203 is:

a) Exposure to chrysotile asbestos poses increased risks for asbestosis, lung cancer and mesothelioma in a dose-dependent manner. No threshold has been identified for carcinogenic risks.

b) Where safer substitute materials for chrysotile are available, they should be considered for use.

c) Some asbestos-containing products pose particular concern and chrysotile use in these circumstances is not recommended. These uses include friable products with high exposure potential. Construction materials are of particular concern for several reasons. The construction industry workforce is large and measures to control asbestos are difficult to institute. In-place building materials may also pose risk to those carrying out alterations, maintenance and demolition. Minerals in place have the potential to deteriorate and create exposures.

d) Control measures, including engineering controls and work practices, should be used in circumstances where occupational exposure to chrysotile can occur. Data from industries where control technologies have been applied have demonstrated the feasibility of controlling exposure to levels generally below 0.5 fibres/ml. Personal protective equipment can further reduce individual exposure where engineering controls and work practices prove insufficient.

e) Asbestos exposure and cigarette smoking have been shown to interact to increase greatly the risk of lung cancer. Those who have been ex-
posed to asbestos can substantially reduce their lung cancer risk by avoiding smoking.”

5.638 When seen in the context of para. c), I take d) to mean that in those situations where exposure is likely or unavoidable, exposure can be reduced or minimized by certain procedures appropriate to the circumstances (e.g., engineering controls in manufacture/production or best work practices), but EHC 203 has already identified friable products and building products as materials of “particular concern” and their use is “not recommended”, in part because of difficulties of control in the construction industry. I do not see paragraph d) as an endorsement of ongoing “controlled use”.

5.639 A similar sentiment is expressed in NICNAS 99:

“Prudent OHS [occupational health and safety] policy and public health policy favours the elimination of chrysotile wherever possible and practicable [p 139]

... Best practice must be implemented to minimise occupational and public exposure, and to minimise environmental impact, over the remaining period(s) of use [p 140].

A risk reduction strategy using all available and appropriate measures is required to ensure that the risks posed by chrysotile are continually reduced and eliminated wherever possible” [p 140].

5.640 NICNAS 99 also goes on to state (p 140):

“In achieving this it is further recommended that:

a) Specific phase-out periods should be set, with stages (over the shortest possible period of time) to encourage and reflect the availability and suitability of alternatives [to chrysotile].

b) Action is taken in the immediate future to prohibit the replacement of worn non-chrysotile original equipment with chrysotile products, as alternatives are now available.

c) No new uses of chrysotile or chrysotile products should be introduced (i.e., an immediate prohibition on new uses).

d) Occupational health and safety authorities take the lead role in considering this recommendation and specific strategies to implement it as worker health is identified as the major concern.”

5.641 As stated in the paper by Jarvholm et al. [28] attached to the Endnote (Section V.C.4) for my report:

“... The first regulation of asbestos [in Sweden] was introduced in the early 1960s and subjects who started their occupational career in the 1960s should have been exposed to lower doses on average, than those who started earlier. On the other hand, by the 1960s asbestos was being used more extensively so the number of people exposed to asbestos may have increased. ... More stringent regulations of asbestos were introduced in the mid-1970s, which led to the sharp decrease in its use. People who have only worked under such conditions were born from 1955 onwards. They have not yet reached a sufficient latency time for possible mesotheliomas to have developed so the number of cases [is] few. However, the first indication is that they may have a decreased risk compared with earlier birth cohorts. A more certain conclusion can probably not be drawn for another ten years. Thus, the preventive measures of the mid-1970s can probably not be evaluated with reasonable precision until around 2005, 30 years later.
The present situation in Sweden, that mortality from mesothelioma due to early use of asbestos is of a similar size to the total number of fatal occupational accidents, is caused by a situation in which at least 90% of the asbestos used was chrysotile. However, we have no information about the type of exposure to asbestos among the cases of mesothelioma - whether they had an exposure to crocidolite or amosite. There is some pressure from the asbestos industry worldwide to change the asbestos regulations to allow the use of chrysotile. To evaluate such an experiment would take at least another 20 years. Even if the major cause of mesothelioma in Sweden was from types of asbestos other than chrysotile, it is difficult to see how the benefits from an increased use of asbestos in Sweden could outweigh the uncertainty of the risks. A similar prudent approach would also be appropriate in other European countries ...

(d) Are substitute fibres safer than chrysotile?

5.642 In para. 5.539, Canada states:

“Dr. Henderson, for his part, recognises that, as with all fibres, the pathogenicity of substitutes is defined by the “3 Ds” (dimension, dose, durability). He seems also to understand that, due to the (lack of) historical use of substitutes, we cannot fully know the risks of using them. However, he then seems to ignore the importance of these facts.”

5.643 My comments on the safety or potential biohazards of substitute fibres were based on the following:

- The dimensions and respirability of substitute fibres. For example, it appears that synthetic fibres can be engineered to be either shorter than the lengths of asbestos fibres that have been associated particularly with carcinogenicity, or to be predominantly non-respirable. In contrast, according to Harrison et al. [29]:

“The intrinsic hazardous properties of chrysotile can never be “engineered out”, and the potential for harm will always remain. Prevention of ill-health will thus always rely on the control of exposure, something that history has shown cannot be guaranteed. ... Unlike chrysotile, substitute fibres can often be designed or selected to have particular characteristics.”

- Dose: reported airborne fibre concentrations from the manufacture or use of substitute (e.g. synthetic) fibres are low - comparable to or lower than the airborne fibre concentrations produced by the manufacture or subsequent use of chrysotile-containing materials. This being so, my conclusions about the relative safety of chrysotile versus substitute fibres are based primarily on fibre dimensions (discussed above) and biopersistence (discussed below).

- Durability (biopersistence): in para. 5.552, Canada states the following:

“It is well known that biopersistence is a key parameter. Indeed, the human evidence for chrysotile indicates that it is likely to be one of the main reasons why chrysotile is less dangerous than the amphiboles in respect to mesothelioma risk. This is clearly recognised by three of the four experts, as well as by INSERM.”

Canada then emphasizes the rapidity of clearance of chrysotile from lung tissue, with reference to a 90-110 day half-life for chrysotile in lung tissue, and an even shorter estimate of < 10 days. Again, I draw attention to the recent study from Finkelstein and Dufresne [5] who estimated a lung tissue half-life of eight years.
for chrysotile fibres > 10 µm in length in Quebec chrysotile miners and millers. Accordingly, in my survey of the literature, I placed particular emphasis on the biopersistence of substitute fibres in comparison to chrysotile.

- The relative potency of substitute fibres or chrysotile fibres to produce pathological changes (e.g. genotoxicity/mutagenicity and the capacity for tumour induction).

5.644 Warheit et al. [30] claim that p-aramid fibres are biodegradable in the lungs of exposed rats, with faster clearance times than long chrysotile fibres, which showed greater biopersistence.

"... p-aramid is biodegradable in the lungs of exposed rats; in contrast, the clearance of long chrysotile fibres was slow or insignificant, resulting in a pulmonary retention of long chrysotile asbestos fibres. The dimensional changes of asbestos fibres as well as the pulmonary cell labelling data indicate that chrysotile asbestos fibres may produce greater long-term pulmonary effects when compared to inhaled para-aramid fibrils" [Abstract].

5.645 In 1993, Hesterberg et al. [31] compared the effects of size-separated respirable fractions of fibrous glass (FG) with refractory ceramic fibres (RCF) and chrysotile fibres. They found that:

"Exposure to chrysotile asbestos (10 mg/m³) and to a lesser extent RCF (30 mg/m³) resulted in pulmonary fibrosis as well as mesothelioma and significant increases in lung tumours. FG (fibreglass designated MMVF10 and MMVD11) exposure was associated with a non-specific inflammatory response (macrophage response) in the lungs that did not appear to progress after 6-12 months of exposure. The cellular changes are reversible and are similar to the effects observed after inhalation of an inert dust. No lung fibrosis was observed in the FG-exposed animals. Further, FG exposure resulted in no mesotheliomas and no statistically significant increase in lung tumour incidence when compared to that of the negative control group. These findings, along with previous inhalation studies, suggest that respirable fibrous glass does not represent a significant hazard for fibrotic or neoplastic lung disease in humans" [Abstract].

5.646 In a later (1995) study, Hesterberg et al. [32] found that exposure of rats to crocidolite and chrysotile asbestos and to RCF by inhalation induced pulmonary fibrosis, lung tumours and mesotheliomas (41 per cent of hamsters exposed to RCF developed mesothelioma343); fibreglasses MMVF10 and MMVF11, slagwool (MMVF22) and stonewool (MMVF21) did not produce a significant increase in lung tumours or mesotheliomas344.

5.647 In a further study published in 1998, Hesterberg et al. [33] investigated the biopersistence of synthetic vitreous fibres and amosite in the rat lung, together with refractory ceramic fibres (RCF1A). They found that "the very biopersistent fibres were carcinogenic" (amosite, crocidolite, RCF1 and two relatively durable special application fibreglasses designated MMVF32 and MMVF33), whereas "the more rapidly clearing fibres were not" (including rock [stone] wool designated MMVF21, HT stonewool designated MMVF34, slag wool, and insulation fibreglasses designated MMVF10 and MMVF11).345

5.648 An Annex from Canada346 also includes a 1995 document on p-aramid fibres from the Health & Safety Executive (HSE) in the United Kingdom. In a summary statement (p. 22) the HSE document states that:

"The balance of evidence suggests that aramid fibres possess a low potential to produce mesothelioma, which is likely to be at least as low as for chrysotile asbestos. While chrysotile is thought to present a hazard with respect to mesothelioma development, current knowledge indicates that the risks for human expo-
sure are low, and would only be detectable following very heavy and prolonged exposure. Thus, if in terms of mesothelioma production, aramid fibres are equally, or less hazardous than chrysotile, it can be concluded that the risks at occupationally relevant levels of exposure would be extremely low.”

5.649 The HSE then set an exposure limit of 2.5 f/ml, but in a subsequent document on Substitutes for Chrysotile (White) Asbestos, the HSE commented that:

“There are many long-established alternatives to chrysotile which do not rely on fibre technology. For example, corrugated polyvinylchloride (PVC) and steel sheeting can be used instead of asbestos cement sheets.

Several types of non-asbestos fibres can also be substituted for asbestos; they have been developed for use in a wide range of products. The main non-asbestos fibres in current use are polyvinyl alcohol (PVA), aramid and cellulose. A considered scientific view on their safety has recently become available. In July 1998, the UK’s Department of Health Committee on Carcinogenicity (CoC) concluded that these three asbestos substitutes (PVA, cellulose and aramid) are safer than chrysotile. This view was endorsed by the European Commission Scientific Committee on Toxicity, Ecotoxicity and the Environment in September 1998.”

5.650 More recently, a press release from the UK Health and Safety Commission (HSC/HSE) announced a prohibition on the importation, supply or use of chrysotile in Great Britain, effective from 24 November 1999.

5.651 I also re-emphasize the comments in the reviews quoted in my original Report (answer to Question 6), including the review by Harrison et al. [29] who comment along the following lines:

“The diameter of PVA [polyvinyl alcohol] fibres, as manufactured, is well above the respirable limit and most of them are not inhalable. ... the fibres are mostly in the range of 10-16 µm diameter. There is evidence that they do not fibrillate (split lengthwise). Many of the particles seen in the atmosphere are non-fibrous. ... Although the published toxicologic information on PVA is relatively sparse, the parent material has been used extensively in surgery and has food contact clearance, presumably based on unpublished studies. Indications of an accumulation of oligomers in the kidney in some circumstance ... mean that the spectrum of molecular weight of material in the fibres as used should be considered, especially if a smaller diameter material were to be produced. The material will degrade only slowly, if at all, in the lungs. ... Thus, substitution of PVA for asbestos fibers in products such as asbestos-cement should result in reduced exposures. This prediction has been confirmed in industrial applications where very low fibre counts have been experienced. Misuse of installed material would not result in significant exposure.

... On balance, the use of aramid fibers should result in reduced levels of fiber exposure as compared to chrysotile asbestos and the fibrils released will be no more toxic and will be less biopersistent. The predicted reduction in absolute exposure levels has been achieved in industrial practice. Misuse of installed material would not be expected to give significant exposures.

... On balance, the coarse fiber structure and the long experience in use indicate that substitution of cellulose fiber for chrysotile asbestos should result in reduced occupational exposures to fiber and lower levels of deposition in the lung. The apparent biopersistence of cellulose in the lung would be a possible cause for concern if the potential for limited lung damage is confirmed.
... We believe that the continued use of chrysotile in asbestos-cement products is not justifiable in the face of available and technically and adequate substitutes. Likewise, there seems to be no justification for the continued residual use of chrysotile in friction materials.”

5.652 These comments also coincide with one of the recommendations in NICNAS 99: “... Current overseas experience with the phasing out of chrysotile products indicates that a range of alternatives is available to suit the majority of uses. Good OHS practice dictates that use of chrysotile should be restricted to those uses where suitable substitutes are not available, and alternatives should continue to be sought for remaining uses” [p 139].

(e) Summary

5.653 It is my perception that the conclusions in my Report submitted already to the WTO concur with mainstream thinking and approaches to occupational and public health policy from national and international health authorities; these include, inter alia:

- The National Occupational Health & Safety Commission in Australia (WorkSafe Australia). (Please see NICNAS 99.)
- The World Health Organization (EHC 203).
- INSERM (France).
- The National Health & Safety Commission/Health & Safety Executive (HSC/HSE) in Great Britain.
- Medical Research Council (MRC) Institute for Environment and Health at the University of Leicester (UK).
- National health authorities in other European Nations.
- The Collegium Ramazzini.

5.654 This being so, it is my perception that the dispute before the WTO is, to some extent, focused upon inappropriate issues. There has been on-going argument among scientists on the health hazards of chrysotile asbestos (the chrysoilohiles versus the chrysoilophobes). Given the extent and complexity of the scientific literature - with contradictory observations on some important issues and with uncertainties related to gaps in observational data - it is almost inconceivable that this controversy can be resolved by the WTO Panel, or, indeed, that it will be resolved in the foreseeable future (partly because no control group free from asbestos exposure can be assembled to ascertain the true spontaneous mesothelioma rate).

5.655 The point to be emphasized is that there exists a substantial body of independent scientific and medical opinion - embodied in national and international health authorities - that chrysotile is carcinogenic with no delineated threshold; that it cannot be controlled at all points of end use; and that existing scientific evidence indicates that safer substitute materials are available.

5.656 To me, this body of opinion is no tendentious artifice designed only to secure a commercial advantage. From my perspective, this is perhaps the crucial issue, from the so-called precautionary principle, given that neither side is likely to concede that the other has proven its case at a high order of scientific probability. In other words, the question is not so much whether there exists a proven health risk or virtually no risk from the continued use of chrysotile, but whether there exists a body of independent and reputable opinion that the possible risks or uncertainties about risk justify a policy of highly restricted use or non-use.
5.657 From this perspective, restriction of chrysotile to only a very few special applications - or its prohibition - is a reasonable and defensible measure designed as a cautious and prudent approach to public and occupational health policy.

5.658 Therefore, I re-affirm the conclusions set out in my original Report (paragraph 5.431) that chrysotile should either:

(a) Be restricted to only a few and well-defined applications so that it is inaccessible to the great majority of workers and is available for use by only small and cohesive specialized worker groups that can be trained effectively in its controlled use (e.g. analogous to nuclear fuels); this means that chrysotile should not be used in building products (e.g. high-density fibro-cement materials such as asbestos-cement sheets) or friction products.

OR

(b) It should be made inaccessible to everyone, by prohibition, unless the alternatives pose equal or greater hazards and equal or greater problems with control.

5.659 In this latter circumstance, the principle is that minimization of exposure is more certain when no new chrysotile-containing products are introduced into the workplace or the general environment, so that the total amount of asbestos in-place will diminish over time; the problem then becomes primarily one of minimization of exposure to existing asbestos products during maintenance, repair, removal, demolition and disposal.

VI. SUBMISSIONS FROM NON-GOVERNMENTAL ORGANIZATIONS

6.1 The Panel received four amicus briefs from the following non-governmental organizations:

- Collegium Ramazzini, dated 7 May 1999
- Ban Asbestos Network, dated 22 July 1999
- Instituto Mexicano de Fibro-Industrias A.C., dated 26 July 1999
- American Federation of Labor and Congress of Industrial Organizations, dated 28 July 1999

6.2 These amicus briefs were transmitted to the parties for their information. In their written rebuttals of 30 June 1999, the EC incorporated by reference the submission of the Collegium Ramazzini. In a letter dated 18 August 1999, Canada notified the Panel that, bearing in mind the general nature of the opinions expressed by the non-governmental organizations in those submissions, they would not be useful to the Panel at this advanced stage of the proceedings. Should the Panel nonetheless accept the submissions as amicus briefs, Canada believed that the parties should be given the possibility to respond to the factual and legal arguments set out in them. In a letter dated 3 November 1999, the EC informed the Panel that it was incorporating by reference the amicus brief submitted by the American Federation of Labor and Congress of Industrial Organizations, as that body supported the EC’s scientific and legal arguments in this dispute. The EC also proposed to the Panel that it reject the submissions from the Ban Asbestos Network and the Instituto Mexicano de Fibro-Industrias A.C., as those documents contained no information of relevance to the dispute. In a letter dated 10 November 1999, Canada again urged the Panel to reject the four amicus briefs as it was inappropriate to admit them at this stage in the proceedings. Should the Panel nevertheless consider these submissions, Canada considered that, for the sake of procedural fairness, the parties should have an opportunity to comment on their content.

6.3 In a letter dated 12 November 1999, the Panel informed the parties that, in the light of the EC’s decision to incorporate into its own submissions the amicus briefs submit-
ted by the Collegium Ramazzini and the American Federation of Labor and Congress of Industrial Organizations, the Panel would consider these two documents on the same basis as the other documents furnished by the EC in this dispute. It was also on that basis that the Panel submitted those two submissions to the scientific experts for their information. At the second substantive meeting of the Panel with the parties, the Panel gave Canada the opportunity to reply, in writing or orally, to the arguments set forth in these two amicus briefs. At that same meeting, the Panel also informed the parties that it had decided not to take into consideration the amicus briefs submitted by the Ban Asbestos Network and by the Instituto Mexicano de Fibro-Industrias A.C.

6.4 On 27 June 2000, the Panel received a written brief from the non-governmental organization ONE ("Only Nature Endures") situated in Mumbai, India. The Panel considered that this brief had been submitted at a stage in the procedure when it could no longer be taken into account. It therefore decided not to accept the request of ONE and informed the organization accordingly. The Panel transmitted a copy of the documents received from ONE to the parties for information and notified them of the decision it had taken. At the same time, it also informed the parties that the same decision would apply to any briefs received from non-governmental organizations between that point and the end of the procedure.
NOTES

2David S. Bernstein, Summary of the Final Reports on the Chrysotile Bio-Persistence Study (Geneva Switzerland; 2 October 1998).
4See para. 4.30 below.
5Official Journal of the European Communities, C 135/108 (14 May 1999) (30 September 1998 answer of Mr. Bangemann to Written Question E-2736/98 of Christine Oddy (PSE)). See also Official Journal of the European Communities, C 13/123 (18 January 1999) (24 July 1998 answer of Mr. Bangemann to Written Question E-1950/98 of Anita Pollack (PSE)) (“[I]t is important to mention that a new ban would not lead to a lower risk of exposure to existing asbestos for workers, nor would it reduce the number of deaths from past exposure to asbestos. Possible contamination from asbestos in existing buildings (e.g. in relation to maintenance activities and asbestos removal operations) will remain an important cause of exposure to workers for many years.”).
6Brazil notes that in the chrysotile-cement industry, the largest present-day use of chrysotile, the manufacturing process uses a water slurry mixture of chrysotile and cement. No dust or pollution is created during this process. See also American Lung Association, Asbestos, pp. 2 and 3 (http://www.lungusa.org/air/en/asbestos.html) (“Asbestos is rarely used alone, and it is generally safe when combined with other materials with strong bonding agents. As long as the material remains bonded so that fibers are not released, it poses no health risk.”); National Cancer Institute, (1996), p. 3 (http://www.nci.nih.gov/clinicalpdq/risk/Questions_and_Answers_About_Asbestos_Extposure.html) (“Asbestos that is bonded into finished products such as walls, tiles, and pipes poses no risk to health as long as it is not damaged or disturbed (for example, by sawing or drilling) in such a way as to release fibers into the air. . . . [N]o fiber type can be considered harmless, and proper safety precautions should always be taken by people working with asbestos.”).

In developing countries such as Brazil, the availability of low-cost, high-quality building and piping materials, such as chrysotile-cement products, is crucial. Substitute products are more expensive and thus less available to those who need them most.

8Id.

Brazil concurs with Canada that the weight of all available scientific evidence, including the INSERM Report, leads to the conclusion that the ban serves no purpose other than restricting trade.

Brazil notes that, because it is only the INSERM Report which preceded the ban, the ban must be supported by the Report alone. Brazil has discussed both the Synthesis and the Report because the former underscores some of the defects of the latter.


IIbid., p. 409 (“France used asbestos much later and to a much lesser degree than other countries, and doubtlessly the asbestos used contained a lower proportion of amphibole-type fibres. Because of these differences, it is not possible to simply transpose to France the results of projections concerning mesothelioma [and cancer] cases prepared recently for Great Britain.”).

Brazil notes that the Cara Brava Mine, in Brazil, for example, has an exceedingly complex and effective air filtration system. The mine is the first and only asbestos mine in the world to have been certified as complying with ISO 14001. It was certified by Det Norske Veritas of Rotterdam, the Netherlands.

sis. Brazil notes that these studies demonstrate that substitute fibres, both when manufactured and used, are likely to present health risks similar to those from chrysotile.

1See Corrosion Proof, 947 F.2d pp. 1226-27 (even while banning asbestos, the EPA conceded that ductile iron pipes and PVC pipes present health (cancer) risks “similar” to those presented by asbestos-cement pipes).

11INSERM, Effets sur la santé des fibres de substitution à l’amiante-synthèse, Paris, 1998, pp. 376 and 428. Brazil notes that the European Commission has also recognized this as an important issue: “There is a key scientific issue which Member States and the Commission agree still needs to be clarified. This is an assessment of the relative risk posed by the substitutes in comparison to the risk posed by chrysotile.” Official Journal of the European Communities, C 13/35 (18 January 1999) (11 June 1998, Answer of Mr. Bangemann to Written Question P-1451/98 of Peter Skinner (PSE)).

18INSERM Synthesis, p. 2.

19Ibid., p. 33.

20According to Brazil, the assumption is contrary to logic because a threshold must exist given that asbestos is ubiquitous in water and air. Only those who have suffered intensive, prolonged, exposure have contracted asbestos-related diseases.


23INSERM Report, pp. 239 and 414.


26Ibid., p. 4.

27Ibid., p. 10.

29Brazilian Law No. 9055 of 1 July 1995.

30Brazilian Decree No. 2350 of 15 October 1997.

31Convention 162, Article 12.

32Article 12 of Recommendation 172 states:

(1) The competent authority, wherever necessary for the protection of the workers, should require the replacement of asbestos by substitute materials, wherever possible.

(2) Before being accepted for use in any process, all potential substitute materials should be thoroughly evaluated for their possible harmful effects on health. The health of workers exposed to such materials should be continuously supervised, if judged necessary. (Emphasis added.)


35Ibid., p. 1215.

41According to Brazil, general rules of pleading, but also Article 2.5 of the TBT Agreement confirm that France has the burden of justifying its trade restrictive measure. According to
Article 2.5, a standard shall be “rebuttably presumed not to create an unnecessary obstacle to trade” when it pursues a legitimate objective and is “in accordance with relevant international standards.” France cannot take advantage of this exception to normal rules of pleading because, as demonstrated below, the ban is contrary to relevant international standards.

Brazil notes that, while no WTO panel or Appellate Body reports have addressed this issue under the TBT Agreement, relevant precedents under the SPS Agreement exist. In Japan - Apples, the Appellate Body found that an SPS measure was justified only if the Member imposing the measure demonstrated a “rational relationship” between the SPS measure and available scientific information. Japan - Measures Affecting Agricultural Products (22 February 1999), WT/DS76/AB/R, para. 84; similarly, in EC - Hormones, the Appellate Body required the EC to establish “an objective relationship between two elements, that is to say, an objective situation that persists and is observable between an SPS measure and a risk assessment.” EC - Measures Concerning Meat and Meat Products (Hormones) (16 January 1998), WT/DS26/AB/R, para. 189; the Appellate Body has also held that a finding that an SPS measure is not based on an actual assessment of health risks is “a strong indication” that the measure does not really protect health but is instead “a trade-restrictive measure in the guise of an SPS measure.” Australia - Measures Affecting Importation of Salmon (20 October 1998), WT/DS18/AB/R, para. 166. This is precisely the case with the ban.


ISO 7337, §§ 4 and 5 (pp. 2-9): Brazil notes that the cutting of plates or tiles for roofing is not a source of emission if ISO-7337 is followed. ISO-7337 addresses the use of chains to break pipes through pressure, low-speed saws, saws equipped with a vacuum dust extractor, and, also, proper wetting of the materials prior to any action. The cutting or grinding of all cement pipe (even that which does not contain chrysotile) emits silica in the air, in the absence of proper controls. The International Association for Research on Cancer (IARC) rates silica as a Type 1 carcinogen (for man), like asbestos. The worker who cuts any cement pipe therefore has an interest in following ISO-7337.


INSEERM Report, p. 434.

Brazil notes that, similarly, under the SPS Agreement, in determining whether an SPS measure is more trade-restrictive than required, the authorities must evaluate whether an alternative, less trade-restrictive, SPS measure would achieve the importing country’s appropriate level of protection. See Australia - Measures Affecting Importation of Salmon (20 October 1998), WT/DS18/AB/R, paras. 208-210.

See paragraph 4.29 above, regarding the conclusions of the American Health Effects Institute, the Royal Commission and the U. S. Court of Appeals for the Fifth Circuit.

Brazil notes that interpretations that render a treaty provision null or void, or consign it to “inutility” are to be avoided whenever possible. See United States – Standards for Reformulated and Conventional Gasoline (20 May 1996), WT/DS2/AB/R, p. 23.

The fact that Article XI applies to the ban is further confirmed by Article XI:2(b) which applies to “import […] prohibitions,” among other restrictions. See also Japan – Trade in Semi-Conductors, L/6309, adopted 4 May 1988, BISD 35S/116, para. 102; United States – Import Prohibition of Certain Shrimp and Shrimp Products, WT/DS58/R (15 May 1998), paras. 7.11 to 7.17.

Brazil notes that a GATT Panel has held that a ban (which, of course, precludes marketing) is not “related” to marketing under Article XI:2(b). See Canada - Measures Affecting Exports of Unprocessed Herring and Salmon, L/6268, adopted 22 March 1988, BISD 35S/98, para. 4.112, paras. 4.2-4.3 (rejecting Canadian argument that a ban on exports of certain unprocessed fish was related to marketing, and finding that, to fall under Exception Two, the regulation in question must apply to “marketing as such,” and that Exception Two does not apply to just any regulation facilitating foreign sales).

According to Brazil, the absence of imports because of the imposition of a ban does not provide a valid basis for asserting that GATT Article III:4 (and TBT Article 2.1) cannot be applied. Interpretations that render a treaty provision null or void, or consign it to “inutility” are to be avoided whenever possible. See United States – Standards for Reformulated and Conventional Gasoline (20 May 1996), WT/DS2/AB/R, p. 23.

Brazil notes that the EC acknowledges this when explaining that the French ban does not include chrysotile diaphragms for use in chlorine environments because substitutes cannot safely be used.


Brazil notes that this criterion was first cited in EEC – Measures on Animal Feed Proteins, L/4599, adopted 14 March 1978, BISD 25S/49, 63, para. 4.2.

Brazil recognizes that Canada has not alleged a violation of GATT Article 1. However, as demonstrated, the French ban violates the most-favoured-nation obligations of both that Article and of Article 2.1 of the TBT Agreement.

The United States notes that its arguments focus on chrysotile asbestos, as that is the subject of the Canadian challenge.


Airborne Asbestos Health Assessment Update, p. 118 (EPA, June 1986) (concluding that “while differences in pleural mesothelioma risk attributable to fibre type may exist, they are much less than differences attributable to other factors”).


The United States notes that the “amphibole hypothesis” postulates that the mesotheliomas among the workers exposed to chrysotile may be explained by confounding exposures to amphiboles, and that chrysotile may have a lower carcinogenic potency than amphiboles.


51 Federal Register 33992, 33993, col.3 (24 September 1986).

The United States notes that, as the 1986 EPA carcinogen risk assessment guidelines point out: “It should be recognized that epidemiological studies are inherently capable of detecting only comparatively large increases in the relative risk of cancer. Negative results from such studies cannot prove the absence of carcinogenic action ... ”. (51 Federal Register 33992 (24 September 1986), pp. 33995-96). Canada’s statement that “no epidemiological study to date has detected a higher health risk [than the linear risk model] resulting from low-level exposures” must be viewed in this light.


51 Federal Register 33992 (24 September 1986), p. 33997. See also EPA Proposed guidelines for carcinogen risk assessment, 61 Federal Register 17960, 17962 (23 April 1996). Although these most recent guidelines are not yet final, they demonstrate that EPA’s reassessment of the issues is similar to the approach taken previously.


Airborne Asbestos Health Assessment Update, EPA, June 1986, p. 23.


Ibid., p. 1207.

Airborne Asbestos Health Assessment Update (EPA, June 1986).


The United States notes that it is not entirely clear what Canada means by the term “undetectable risk”. The presence of asbestos fibres in the air or other media can be detectable or undetectable. A risk can be significant, insignificant, or non-existent. It appears that Canada uses the term “undetectable risk” to refer to a risk that Canada deems insignificant. Significance, however, is a judgment call that can only be made by the regulatory authority responsible for public health and safety. It is up to France to determine what level of risk to the French people from asbestos (or any other hazard) is significant.


The United States notes that, for example, a study of mortality among long-term employees of an Ontario asbestos-cement factory found a substantially increased risk of death from lung cancer and mesothelioma. Finkelstein, M. M., Mortality Among Long-Term Employees of an Ontario [Canada] Asbestos-Cement Factory, 40 Br J. Ind. Med. 138-44, 1983.

The United States notes that this has been recognized by EPA’s asbestos National Emission Standard for Hazardous Air Pollutants (NESHAP), promulgated under §112 of the
Clean Air Act, 42 U.S.C. 7412. 55 Federal Register 48406, 48408-09 (Nov. 20, 1990), codified at 40 CFR part 61, subpart M.


10Ibid., pp. 4-32 and 4-33.

10Ibid., pp. 4-33.

10Project Summary: Airborne Asbestos Concentrations During Buffing of Resilient Floor Tile, EPA, October 1993, p. 4.

10Ibid., pp. 4-32 and 4-33.


10BISD 18S/102, para. 18.


10Ibid. pp. 97, 98.


10Article 5.6 provides in relevant part “when establishing or maintaining sanitary or phytosanitary measures to achieve the appropriate level of sanitary or phytosanitary pro-
tection, Members shall ensure that such measures are not more trade-restrictive than required to achieve their appropriate level of sanitary or phytosanitary protection, taking into account technical and economic feasibility”.

133 This information is based on a report by Reuters News Agency, dated 6 May 1999.
135 Ibid., p. 144.
136 Ibid., pp. 7 and 144. Zimbabwe notes that, in this regard, it is misleading for the EC to cite the 1998 Task Group report for the proposition that there is an international consensus that no threshold of exposure can be identified below which no risk to humans exists. The Task Group in fact merely stated that it could not identify any such threshold on the basis of the available data. See WHO, IPCS Environmental Health Criteria 203 - Chrysotile Asbestos, Geneva, 1998, pp. 7 and 144.
137 Ibid., p. 145.
138 Zimbabwe notes that this is especially true of applications of chrysotile-containing products in industries such as construction, on which the EC has placed great emphasis in its Submission, because “studies have not been in general able to distinguish between chrysotile and amphibole exposure”. See WHO, IPCS Environmental Health Criteria 203 - Chrysotile Asbestos, Geneva, 1998, pp. 122 and 112.
139 According to Zimbabwe, the EC confirms this when stating: “[L]es principales données qui ont été présentées illustrent le caractère ubiquitaire de l’amiante en milieu de travail qui peut, à des niveaux d’exposition suffisamment élevés, entraîner de nombreux cas de maladies mortelles”.
141 For a more detailed discussion of this point, see Zimbabwe’s arguments with respect to GATT Article XX.
145 Ibid., p. 144.
147 In this connection, Zimbabwe notes that it is a truism that it is often easier to define objects negatively than to come up with an exact and exhaustive positive definition.
149 A treaty interpreter is not entitled to assume that [the use of different words in different places] was merely inadvertent on the part of the Members who negotiated and wrote that Agreement”. See EEC - Measures Concerning Meat and Meat Products (Hormones), Appellate Body Report, adopted on 13 February 1998, WT/DS26/AB/R, WT/DS48/AB/R, para. 164.
151 See Section III.B of this Report.
152 Ibid.
153 See Section III.C of this Report.
155 Zimbabwe believes that its interpretation of the TBT Agreement also finds support in Article 1.6 of the TBT Agreement. This Article states that “[a]ll references in this Agreement to technical regulations […] shall be construed to include any amendments thereto and any additions to the rules or the product coverage thereof, except amendments and additions thereof, except amendments and additions of an insignificant nature”. This provision clearly establishes that the term “technical regulation” as used in the TBT Agreement is not to be given a narrow reading, but one which promotes the Agreement’s effectiveness.
157Zimbabwe notes that, with regard to glass fibres, this follows from the definition of HS tariff position 68.11.

158In Zimbabwe’s view it does not matter for the purposes of an inquiry under Article III:4 whether domestic production of the “like product” is substantial or small. Nowhere does Article III:4 lay down a requirement that domestic production needs to be substantial.

159Zimbabwe considers that the Decree falls obviously within the scope of Article III:4 inasmuch as it is a regulation which affects the internal sale of asbestos fibres.


161Ibid., p. 20.


163This is precisely why, in the view of Zimbabwe, the different tariff classification of chrysotile asbestos fibres, on the one hand, and of cellulose and aramid fibres, on the other hand, cannot provide any useful guidance for purposes of determining “likeness” in this case. The Appellate Body has in fact confirmed in its report on Japan – Taxes on Alcoholic Beverages, adopted on 1 November 1996, WT/DS/8/AB/R, WT/DS10/AB/R, WT/DS11/AB/R, p. 22, that the value of tariff classification as a criterion for establishing “likeness” must be assessed on a case-by-case basis.

164According to Zimbabwe, it should be borne in mind in this context that diameter and fibrillosity are in any event relevant only to the extent that these characteristics correlate with health risks to humans.

165Zimbabwe notes that this is all the more true in view of the fact that the kind of diseases at issue here involve long latency periods.

166See Section III.B of this Report.


168According to Zimbabwe, the same is true for glass fibres.

169Zimbabwe notes that the fact that there is an additional and special temporary exemption in Article 7 of the Decree for certain used and agricultural vehicles precisely suggests that no equivalent and affordable substitutes existed at the time the Decree was signed into law and that the sectors concerned successfully lobbied the Government to provide for a temporary exemption.

170Zimbabwe notes that the question to be answered by the Panel here is whether it was necessary for France to discriminate between asbestos fibres and “like” domestic fibres in order to protect human health.


172See Section III.C of this Report.

173Zimbabwe notes that, by way of analogy, it might be added here that many people think that the wearing of seatbelts in cars makes driving more “complicated” and awkward. Yet many countries made the wearing of seatbelt a legal requirement.

174In this connection, Zimbabwe recalls that a tenant or house owner who wants to drill a hole in a wall to hang up a painting, for instance, also needs to know exactly where the electrical wiring and installations are, lest he/she wants to put his/her life at risk.

175Zimbabwe notes that, after all, in most countries, drugs cannot be bought in supermarkets, but only in pharmacies upon production of a doctor’s prescription.

176To give another analogous example, Zimbabwe notes that in many countries the installation of ceiling lamps and other electrical appliances may only be carried out by certified electricians.


178WT/DSB/RC/1, of 11 December 1996.


180According to the European Communities, confirmation can also be found in the chapeau to Appendix 4 to the Dispute Settlement Understanding which provides that “[the following rules and procedures] shall apply to expert review groups established in accordance
with the provisions of paragraph 13”, that is, regardless of whether it is the first or the second sentence of this Article that is being used by the Panel.

181 That also explains the Appellate Body’s reasons for its finding on that matter in the Hormones case. See the report AB/1997-4, para. 147.

182 As the Appellate Body found in Hormones (para. 164), “a treaty interpreter is not entitled to assume that such usage was merely inadvertent on the part of the Members who negotiated and wrote that Agreement”.

183 The third sentence of Article 1:2 of the Dispute Settlement Understanding is not applicable in this case, as the GATT 1994 does not contain contradictory rules and procedures on this matter.

184 The Panel’s interpretation is also contrary to one of the corollaries of the general rule of interpretation set out in the 1969 Vienna Convention, which is that the interpretation must give meaning and effect to all the terms of a treaty. As held by the Appellate Body in the Gasoline case “an interpreter is not free to adopt a reading that would result in reducing whole clauses or paragraphs of a treaty to redundancy or inutility” (AB-1996-1, page 22). Specifically, the Panel has so far refrained from providing explicit substantive reasons for its choice of consultation with individual experts over the establishment of an expert review group.

185 As far as the European Communities are concerned, additional support for the proposition that the term conflict of “interest” should be interpreted as broadly as possible may be drawn from Article III.1 of the Rules of Conduct mentioned above, and in a systematic interpretation (by analogy) of the following provisions: Articles 8:2, 8:3 and 17.3 of the DSU, paras. 2 and 3 of Appendix 4 to the DSU, as well as paras. 2 and 3 of Annex 2 to the TBT Agreement.

186 See the Reports of the Appellate Body in European Communities – Measures Concerning Meat Products (Hormones) (WT/DS26/26-DS48/AB/R), para. 147 (“… in disputes involving scientific or technical issues, neither Article 11.2 of the SPS Agreement, nor Article 13 of the DSU prevents panels from consulting with individual experts. Rather, both the SPS Agreement and the DSU leave to the sound discretion of a panel the determination of whether the establishment of an expert review group is necessary or appropriate”) and Argentina – Measures Affecting Imports of Footwear, Textiles, Apparel and Other Items (WT/DS56/AB/R) para. 84 (“Article 13 of the DSU enables a panel to seek information and technical advice as it deems appropriate in a particular case, and (…) the DSU leaves ‘to the sound discretion of a panel the determination of whether the establishment of an expert review group is necessary or appropriate’.”).

187 See the Report of the Appellate Body in Guatemala – Anti-Dumping Investigation Regarding Portland Cement from Mexico (WT/DS60/AB/R), paras. 65 and 66.


189 For complete references, see Annex III to the Panel Report.

190 The National Cancer Institute’s Surveillance Epidemiology and End Results program.

191 Mark and Yokoi [118] have called into question the existence of mesothelioma in the absence of asbestos exposure, pointing out that the early descriptions of pleural tumours may have dealt with localized fibrous tumours of the pleura (four of the five tumours reported by Klemperer and Rabin [119]) or secondary carcinoma. Thus, mesothelioma might represent a new disease consequent upon the industrial use of asbestos (analogous to AIDS) and it may disappear upon withdrawal of the causative asbestos from the environment (analogous to smallpox). In support of this proposition, these authors cited the records of the Massachusetts General Hospital, where no examples of mesothelioma were diagnosed before 1946, in contrast to 100 autopsy cases thereafter, in a total of 47,000 autopsies. They also referred to the Henke-Lubarsch Handbuch der speziellen pathologischen Anatomie und Histologie, wherein the four pages devoted to tumours of the pleura did not specifically acknowledge the existence of mesothelioma; the Henke-Lubarsch authors concluded that many cases described in the literature as primary pleural neoplasms were cases of lung cancer with spread to the pleura. I find the evidence for Mark and Yokoi’s proposition to be underwhelming and unconvincing. The case reported in the 1920 paper by Du Bray and Rosson [120] is, I believe, a clear example of a mesothelioma, as is the fifth case of Klemperer and Rabin [119]. Failure to diagnose a tumour is hardly synonymous with its non-existence, and the pathological features of many tumours have been delineated in quite recent times. Pathological diagnoses follow prevailing evidence and fashions, and
because the groundwork for modern concepts of mesothelioma was laid down in 1931 by Klemperer and Rabin, it is hardly surprising that the diagnosis became more widespread only after this time.

191Given in various publications as fibres/ml, fb/ml, f/ml, f/mL and fibres/cm³.
192The philosopher of science, Sir Karl Popper ... coined the term 'falsification' to express the concept that scientific theories are not proven by repetition of results but rather survive because they successfully withstand refutation (falsification). His example of the black swan makes this point clearly. Suppose you have a hypothesis that all swans are white ... you observe, say, 10,000 swans and they are all white. Another scientist repeats your efforts and observes another 10,000 swans: they too are all white. So far the theory is standing up well. The repetition helped to strengthen it — but if only a single black swan is sighted, this falsifies the theory: it is no longer tenable. Popper asserted that scientific statements have to be formulated in a manner that subjects them to the possibility of falsification. One of the important demarcating criteria between science and nonscience, according to Popper, is this formulation of statements in a manner permitting falsification” [pp 18-19] [44].
193This figure is inconsistent with the former limit of 0.1 f/ml in France, and contradicts Case’s claim that the Quebec women were exposed at up to 1 f/ml [192]: at a level of 0.0107 f/ml (a figure two orders of magnitude less than 1 f/ml), a cumulative exposure of 5 fibre-years would require residence of > 150 years (adjusted for equivalence to an 8-hour working day) and > 750 years to reach 25.0 fibre-years (using the same adjustment).
194Estimates calculated at my request by Dr. N.H de Klerk.
19610,000 Ångstroms = 1.0 µm.
197This relatively high proportion (10 per cent) in comparison to the smaller fraction of airborne fibres of the same size is presumably explicable by preferential clearance of short fibres from lung tissue, with a proportional increase of long fibres over time.
198The Register is a compilation of all and unselected mesotheliomas throughout Australia.
199This over-estimates the number of brake mechanics, because the figure includes all automotive mechanics, engine mechanics, apprentices, and supervisors: Australian Bureau of Statistics, 12 October 1999.
200Maximum exposure limit which was authorized in France before the ban.
201Henderson, p. 54, citing Multiple Authors, Asbestos Cement Products. Report by the Western Australia Advisory Committee on Hazardous Substances, Perth, 1990, hereinafter the WAACHS Report.
203Henderson, answer to Question 1(d).
204Infante, answer to Question 1(f).
205CONSAD Research Corporation, 1990, No. 8282.
207Henderson, answer to Question 2.

Circulaire no. 97-15 du 9 janvier 1997 relative à l’élimination des déchets d’amiante-ciment générés lors des travaux de réhabilitation et de démolition du bâtiment et des travaux publics, des produits d’amiante-ciment retirés de la vente et provenant des industries de fabrication d’amiante-ciment et des points de vente ainsi que tous autres stock.

Circulaire no. 96-60 du 19 juillet 1996 relative à l’élimination des déchets générés lors des travaux relatifs aux flocages et aux calorifugeages contenant de l’amiante dans le bâtiment.

Canada notes that French regulations indeed recognize a difference in disposal proscriptions between “les matériaux friables” and “l’amiante liée” – see Note DPPR/SDPD/BGTD/LT/LT no. 97-320 du 12 mars 1997 relative aux conséquences de l’interdiction de l’amiante et à l’élimination des déchets, which is as follows:

“III. – Quelles sont les filières d’élimination des déchets contenant de l’amiante?


“Les filières d’élimination des déchets contenant de l’amiante autres que ceux qui ont fait l’objet des deux circulaires précitées peuvent être déterminées par analogie aux prescriptions de ces deux circulaires:

- Les matériaux friables, c’est-à-dire les matériaux susceptibles d’émettre des fibres sous l’effet de chocs, de vibrations ou de mouvements d’air, sont assimilables aux flocages et aux calorifugeages. Ils devront être éliminés dans des installations de stockage de déchets industriels spéciaux ou dans l’unité de vitrification;

- pour les déchets contenant de l’amiante liée, trois cas sont envisageables:

- Si les déchets sont composés d’amiante associée uniquement avec des matériaux inertes, ceux-ci pourront être éliminés conformément à la circulaire du 9 janvier 1997 relative à l’élimination des déchets d’amiante-ciment;

- si l’amiante est associée avec des matériaux, qui lorsqu’ils deviennent des déchets, sont classés déchets ménagers et assimilés, c’est par exemple le cas des dalles vinyle-amiante, ils pourront être éliminés dans des installations de stockage de déchets ménagers et assimilés;

- si l’amiante est associée avec des matériaux, qui lorsqu’ils deviennent des déchets, sont classés déchets industriels spéciaux, ils devront être éliminés soit dans des installations de stockage de déchets industriels spéciaux, soit dans l’unité de vitrification.

“Dans tous les cas, l’industriel ou l’entreprise devra fournir des éléments permettant de caractériser les déchets afin de déterminer les filières d’élimination adaptées.”


Appendix A on Control Use in the Friction Industry, Canada’s Comments to Question 5(a), contained in Annex IV to this Report.


See notably INSERM Report, p. 213.

WHO, IPCS Health Criteria 203 on Chrysotile, WHO, Geneva, 1998, p. 51: “[I]t is considered that the potential respiratory health effects related to [...] airborne concentrations, patterns of exposure, fibre shape, diameter and length (which affect lung deposition and clearance) and biopersistence.”

De Klerk, N.H. and Armstrong, B.K., The Epidemiology of Asbestos and Mesothelioma in


25Henderson, see above para. 5.112.


28INRS, Rapport du Groupe scientifique pour la surveillance des atmosphères de travail (G2SAT), 1997, p. 47.


32Henderson, see above paragraph 5.146.

33Ibid.


36Henderson, see above para.5.103.


38See Canada’s Comments to Question 3.


46INSERM Report, p. 327.

850


260 See Canada’s comments to Question 3.

261 Henderson, answer to question 4(c).


263 Australia National Industrial Chemicals Notifications and Assessments Scheme (NICNAS), Chrysotile Asbestos: Priority Existing Chemical No. 9 (Full Public Report), February 1999 at p. 72, cited by Henderson in his answer to question 4(c).

264 INSERM Report, p. 213.


270 INSERM Report, p. 239 and 414.

271 See above Section III.A.5.

272 Canada notes that the “controlled-use” approach has been endorsed by the WHO in its 1998 Environmental Health Criteria 203: Chrysotile Asbestos, p. 144. “Control measures, including engineering controls and work practices, should be used in circumstances where occupational exposure to chrysotile can occur. Data from industries where control technologies have been applied have demonstrated the feasibility of controlling exposure to levels generally below 0.5 fibres/ml. Personal protective equipment can further reduce individual exposure where engineering controls and work practices prove insufficient.”

273 NRCAN, The Minerals and Metals Policy of the Government of Canada: Partnership for Sustainable Development, Public Works Canada, 1996. Canada notes that the “controlled-use” approach to regulating chrysotile asbestos is well researched as evidenced in the studies and conclusions referred to by Canada in its factual arguments (see above Section III.A.6).

274 To illustrate this point, examples of “controlled-use” of friction products and asbestos-cement are detailed in Appendices A and B respectively to these comments. (These Appendices can be found in Annex IV to this Report).

275 See Section III.A.5 of this Report and Camus M., L’amiante et les risques pour la santé, April 1999.


277 According to Canada, the emphasis of ILO Convention 162 is on controlled-use and not
on product prohibitions. The Convention calls for two specific prohibitions: crocidolite and all products containing crocidolite, and sprayed-on applications of asbestos.


30Canada notes that it should be recalled that his basis for risk assessment is based on the textile industry.

31See Annex IV to this Report.


41Henderson, answers to Questions 5(a) to (d).


45INESM Report.

46Henderson, response to Question 6(b).

47See above Section III.A.6.


49Infante, answer to Question 6(c).


53de Klerk, answer to Question 6(a).

Perspective, 107.
307See Canada’s comments on Question 3.
311Bernstein, D.M., Graph on Biopersistence of p-Aramid Fibres.
314Hesterberg, T.W., Miller, W.C., Theveney, Ph. and Anderson, R., Comparative Inhalation Studies of Man-Made Vitreous Fibres: Characterization of Fibres in the Exposure Aerosol and Lungs, (1995) 39 Ann. Occup. Hyg., pp. 637-653. Hesterberg exposed rats to a concentration of 10,000 WHO fibres/ml of chrysotile, which resulted in 18.9 per cent lung tumours. Rats exposed to 232 f/ml of one type of glass fibre resulted in 5.9 per cent lung tumours, with 4.4 per cent lung tumours reported with other man-made vitreous fibres and 13 per cent with a RCF sample. The air control resulted in 1-3 per cent tumours (At 1000 f/ml, the risk of lung tumours would be just under 2 per cent which is well within the rate of tumours in the control animals).
317Gibbs, G., Phone Communication.
See Annex V to this Report.

See factual arguments by the EC, Section III.A.4.

For complete references of the documents quoted in this Section, see Annex III to this Panel Report.

For decades, brake blocks and brake linings used in Australia have contained Canadian chrysotile asbestos only, with no added amphiboles.

Please see also my answers to Questions 1(e) and 5(a).

Canada’s comments also refer to the meta-analysis of lung cancer risk reported by Lash et al., [10], which identified a low risk. Meta-analysis is a field that lies outside my expertise, but I understand that there are various models for meta-analysis and problems with this approach (e.g. see Blettner et al. [11] who state that “...Meta-analyses using published data are in general insufficient to calculate a pooled estimate since published estimates are based on heterogeneous populations, different study designs and mainly different statistical models [Abstract] ... Meta-analyses using published data are, therefore, restricted and seldom useful to produce a valid quantitative estimate or to investigate exposure relations such as dose-response [p 8] ...”). In a meta-analysis of 69 asbestos-exposed occupational cohorts, Goodman et al. [12] identified “... meta-SMRs of 163 and 148 [for lung cancer] with and without latency, with significant heterogeneity of results ...”

There is a discrepancy between the ages at death in the original paper by Sébastien et al. [7] (i.e. mean = 55.8 ± 9.7 for the Charleston cohort vs 67.5 ± 9.7 for the Thetford group) and the follow-up study by Case et al. [19] (Table 1A, where the ages are reversed: 67 ± 10 for the Charleston group vs 56 ± 6 for Thetford). Clearly, one or the other must be wrong.

Clearly, from the data in Table 2 and the discussion in paras. 6095.604 to 6145.609, they are not representative.

The potential for misuse of asbestos-containing materials remains, as shown by some prosecutions (e.g. please see the UK Health & Safety Executive (HSE) press releases E198:98 and E079:99, http://www.hse.gov.uk/press/e98198.htm and http://www.hse.gov.uk/press/e99079.htm), but cases that come before the courts almost certainly represent only a small fraction of the misuses, most passing unnoticed by regulatory agencies.

Please note the high chrysotile count almost a decade after the patient’s employment ended.


Omitted from my original Report because I did not - and do not - take this to be an endorsement of “controlled use”, and also because the figure of up to 0.5 f/ml is up to five times higher than the level of 0.1 f/ml mentioned in Question 5(c) from the WTO Panel.

The hamster seems to show a propensity for mesothelioma induction in some circumstances (e.g. SV40 inoculation) but not others; in some studies (Research and Consulting Company) chrysotile did not induce mesothelioma or lung in hamsters but in rats it produced pulmonary fibrosis, lung tumours and mesotheliomas, so that the rat has been advocated as the most appropriate model for assessment of the human risk from fibre inhalation [32].

Wilson et al. [34] estimate that fibreglass is 5-10 times less “risky” than chrysotile and they state that “… no one has found any cancer attributable to the manufacture or installation of glass wool fibres … “. In their estimates of lung cancer risk from chrysotile, they use the unit carcinogenicity factor of 0.01 (K, used before them by the US EPA), and they calculate an absolute excess lung cancer risk of 1.2 x 10⁻⁵ for smokers, and 1.4 x 10⁻⁶ for non-smokers; for 40 yrs exposure at 1.0 f/ml, these estimates equate to 4.8 x 10⁻² (smokers) and 5.6 x10⁻³ (non-smokers) - i.e. about 5 per cent and 0.5 per cent respectively, both of which can be considered quite “high”. (Wilson, R., Langer, A.M. and Nolan, R.O., A Risk Assessment for Exposure to Glass Wool, (1999) 30 Regulatory Toxicology and Pharmacology, pp. 96-109.

In the UK HSC Press Release C054:99 announcing implementation of a policy of prohibition of chrysotile from 24 November 1999, the following specific uses are allowable until 2001-2005:

- The use of compressed asbestos fibre (CAF) in gaskets for use with saturated and superheated steam, and with certain flammable, toxic and corrosive chemicals until 1 January 2001;
- The use of CAF in gaskets for use with chlorine until 1 January 2003;
- The use of any sheet which, when in a dry state, has a density greater than 1900 kilograms per cubic metre and is used at temperatures at or above 500°C until 1 January 2003;
- The use of asbestos components in aeroplanes and helicopters where this is crucial for their safe operation until 1 January 2004;
- The use of any product consisting of a mixture of asbestos with a phenol formaldehyde or with a cresylic formaldehyde resin in vanes for rotary vacuum pumps, vanes for rotary compressors, any bearing or its housing or for split-face seals used to prevent water leakage from hydro-electric power generation turbines or from cooling water pumps in power stations until 1 January 2004;
- The use of asbestos in pre-formed joints made from proofed asbestos cloth for sealing the doors of steam boilers until 1 January 2004;
- The use of asbestos in personal protective clothing when used in very high temperatures (500°C or more) until 1 January 2005.
VII. INTERIM REVIEW

A. INTRODUCTION

7.1 The Panel's interim report was given to the parties on 13 June 2000, pursuant to Article 15.2 of the Understanding. On 27 June 2000, Canada and the European Communities sent a request to the Panel in writing asking it to review certain aspects of the interim report. Neither of the parties requested the Panel to hold a further meeting with them. When sending the interim report to the parties, the Panel gave each party an opportunity to transmit in writing its comments on the initial request for review submitted by the other party, if no meeting was requested, provided that the comments were strictly confined to the questions identified by the other party in its initial request for review. The parties submitted their comments on 4 July 2000.

7.2 As a preliminary remark, the Panel notes that the parties requested the Panel to improve upon some of its summaries of their arguments in the findings. The Panel indicates, however, that neither party claimed that the Panel had made any substantive error in its assessment of the facts.

7.3 The Panel also notes that the two Parties have put forward corrections of form and of substance. Wherever appropriate, the Panel accepted the former. The following observations therefore only concern the comments which the Panel considered to be remarks concerning the substance.

B. COMMENTS BY CANADA

7.4 Regarding Canada's comments on substance, the Panel amended footnote 1 to take into account both Canada's remark and the objections of the European Communities. The Panel considers that this summary adequately reflects Canada's arguments in the descriptive part. It did not therefore deem it necessary to replace the text as suggested by Canada. The next comment concerns para. 8.28 (para. 8.29 of the final Report). In response to Canada's remark that the Panel referred to the object and purpose of the WTO Agreement without examining it subsequently, in para. 8.48 of the final Report we have made it clear that we did not find any particularly relevant elements in the preamble to the WTO Agreement relating to the terms we had to interpret. As regards the comment on footnote 37 in the interim report (footnote 41 in the final Report), we did not consider it necessary to delete it because it is not a question of turning to the preparatory work within the meaning of Article 32 of the Vienna Convention on the Law of Treaties (1969), but of a reference to a document showing that the Agreement on Technical Barriers to Trade of the Tokyo Round was already seen as a development of certain rules of the GATT. We have nevertheless edited the footnote in order to make our intention clearer. We have also reworded paras. 8.61 and 8.62 of the final Report and the footnote concerning para. 8.61 in order to make it clear that, in the light of our previous findings, it was not necessary to examine whether the Decree met the five requirements which, in Canada's opinion, are contained in the definition of "technical regulation" in the TBT Agreement. Likewise, we did not wish to give the impression that Canada confirmed the initial view of the Panel that it would be appropriate to analyse the two aspects of the Decree separately. We have also clarified this point. The same applies to para. 8.72 of the final Report. Having reached the conclusion that the legal characterization of the exceptions in the Decree did not affect the legal characterization of the ban, we have simply found that Canada did not make any claims concerning the exceptions and, consequently, we could not reach any findings thereon according to our terms of reference.

7.6 With regard to the section of our findings on application of Article III and/or Article XI of the GATT 1994, we have made our summary of Canada's arguments (para.
8.84 of the final Report) clearer and have revised the rest of the section (paras. 8.87-8.100 of the final Report) in order to respond to the comments concerning paras. 8.85 and 8.95 of the interim report.

7.7 In connection with the discussion on violation of Article III:4 of the GATT 1994, we have also made paras. 8.109, 8.117, 8.121 and 8.145 of the final Report clearer, taking into account Canada’s pertinent remarks.

7.8 Concerning application of Article XX(b) of the GATT 1994, we have amended footnote 128 in the interim report (footnote 142 in the final Report) in order to include the EC’s arguments, while maintaining the reference to Dr. Henderson’s remarks. The Panel has also revised para. 8.187 of the final Report because this was not a reference by Canada but by a Canadian scientist cited by one of the experts consulted pursuant to Article 13 of the Understanding. The Panel did not deem the EC’s objection relating to para. 8.187 to be relevant. In addition, while maintaining the order of paras. 8.196 and 8.197 in the final Report, because the EC referred to Article XX of the GATT 1994, the Panel completed the summary of Canada’s arguments in para. 8.197 by including the factual elements as well, as was done in the summary of the EC’s arguments. The Panel notes that the factual elements mentioned by Canada have already been summarized in para. 8.164 of the final Report, but more concisely.

7.9 With regard to para. 8.233 (para. 8.238 of the final Report), the Panel does not consider that the fall in French imports of asbestos from 1996 onwards (see para. 3.20 above) is proof of protectionist intentions underlying the adoption of the Decree. In this connection, the Panel notes that Canada devotes a long passage (paras. 3.26 and 3.27 above) to showing that the Decree was adopted hastily under the pressure of public opinion. Regarding the industrial and commercial aspects, Canada’s description in para. 3.28 above makes this a secondary element. The French Government’s call for research proposals in July 1996, specifically to develop substitute fibres, does not appear to be the consequence of the choice already made for reasons that bear no direct relation to industrial or commercial considerations.

7.10 Lastly, in our considerations on Article XXIII:1(b) of the GATT 1994, we have harmonized our terminology in paras. 8.293, 8.294 and 8.303 of the final Report.

C. COMMENTS BY THE EUROPEAN COMMUNITIES

7.11 Like Canada, the EC have made a number of comments with a view to amending the Panel’s summary of their arguments in its findings. As was the case for Canada, we have generally taken these comments into account provided that we were able to identify them in the submissions made to the Panel in the course of the procedure.

7.12 This is the case for para. 8.7 in relation to para. 3.6. Regarding the description of the EC’s arguments in para. 8.25 of the final Report, we have taken the relevant parts of paras. 3.264 and 3.265 above, which contain the EC’s arguments on these points. We did not accept Canada’s comments of 4 July 2000 because they essentially repeated arguments already made and taken into account during the procedure.

7.13 We have made clearer the last sentence of para. 8.32 and the first sentence of para. 8.41 of the final Report. We have also amended the content of footnotes 55 and 58 of the final Report and have revised the text of paras. 8.119, 8.149, 8.162 and 8.185 of the final Report, taking into account Canada’s comments of 4 July 2000 and provided that the EC’s suggestions were in line with their submissions to the Panel or the terms of the Decree. We have also made the first sentence of para. 8.149 of the final Report clearer. On the other hand, we did not consider it appropriate to replace the words “Canada’s position” in para. 8.56 of the final Report by the words “Canada’s contextual interpretation”, particularly since Canada opposed this in its comments of 4 July 2000. Likewise, we did not consider that the complementary nature of the EC’s argument on the transitional nature of the exceptions in the Decree warranted the deletion or amendment of para. 8.65 of the final Re-
port. In this connection, the Panel notes that the EC did not clearly state what they wished the Panel to do with this paragraph.

7.14 The EC also requested the Panel to amend paras. 8.66-8.70 and 9.1(a) of the interim report to take account of the fact that Canada did not include the implementing Decree referred to in Article 2 of the Decree in the terms of reference of the Panel and that, consequently, no finding could be made in this respect. Although we agree with the EC that we cannot reach any findings regarding the part of the Decree on exceptions, we consider that this is primarily due to the absence of allegations by Canada regarding these exceptions. In this sense, our assessment is fully in compliance with the practice of the Appellate Body on interpretation of the terms of reference for panels. On the other hand, we do not consider that the fact that the implementing Decree was not specifically mentioned in the Panel’s terms of the reference prevents us from taking it into account when evaluating the legal characterization of the part of the Decree on exceptions. If this were the case, the legal characterization of a measure could depend on the scope of the terms of reference defined by the complaining party, which is unacceptable from a legal point of view. It thus appears appropriate to us to consider the Decree in the light of the implementing measures it provides. For us, the implementing decree is a fact. In addition, we can infer its existence simply by reading the text of the Decree. Consequently, we do not consider that we have exceeded our terms of reference by evaluating the legal characterization of one part of the Decree in the light of the facts before us. We are aware that our efforts to define the legal characterization of the exceptions in the Decree might, at first sight and in the absence of claims from Canada, appear to be an obiter dictum. Nevertheless, this procedure was necessary for two reasons: (a) it provides positive confirmation of our analysis of the scope of the TBT Agreement; and (b) it permits a comprehensive analysis of the elements of the Decree, fulfilling our obligation to make an objective examination of the facts. We did not therefore deem it necessary to amend the paragraphs in question, except for para. 9.1(a), in order to make the meaning of our conclusions clearer in the light of the comments above.

7.15 In response to the EC’s comments and also those of Canada in its communication of 4 July 2000, we have explained in para. 8.273 of the final Report why we considered that the risk of an effective increase in the cost of health measures following application of Article XXIII:1(b) to measures justified under Article XX(b) would be marginal. We have also made the last sentence of para. 8.296 of the final Report clearer.

7.16 Finally, the Panel considered it necessary to add a paragraph concerning the request by a non-governmental organization, received on 27 June 2000, to be allowed to make a written submission and to be heard by the Panel. Moreover, bearing in mind the nature of the comments by the parties, the Panel considered it appropriate to add a footnote to para. 8.35, to revise footnotes 44 and 50, and to amend the first sentence of para. 8.183 of the final Report. The Panel has also clarified its comments in paras. 8.211 and 8.214. Other changes of form have been made in order to take into account the change in the numbering of certain paragraphs. Lastly, the Panel adopted the word “exceptions” to describe the regime applicable under Articles 2 et seq. of the Decree, in conformity with the term used in the Decree itself.

VIII. FINDINGS

A. SUMMARY OF THE FACTS AT THE ORIGIN OF THIS DISPUTE AND CLAIMS BY THE PARTIES

1. Measure at the origin of the dispute

8.1 The measure at the origin of this dispute is Decree No. 96-1133 of 24 December 1996, issued by the Prime Minister of the Government of the French Republic, banning asbestos, implemented pursuant to the Labour Code and the Consumer Code (hereinafter the “Decree”). The relevant provisions of the Decree are set out below:
"Article 1

I. For the purpose of protecting workers, and pursuant to Article L. 231-7 of the Labour Code, the manufacture, processing, sale, import, placing on the domestic market and transfer under any title whatsoever of all varieties of asbestos fibres shall be prohibited, regardless of whether these substances have been incorporated into materials, products or devices.

II. For the purpose of protecting consumers, and pursuant to Article L. 221.3 of the Consumer Code, the manufacture, import, domestic marketing, exportation, possession for sale, offer, sale and transfer under any title whatsoever of all varieties of asbestos fibres or any product containing asbestos fibres shall be prohibited.

III. The bans instituted under Articles I and II shall not prevent fulfilment of the obligations arising from legislation on the elimination of wastes.

Article 2

I. On an exceptional and temporary basis, the bans instituted under Article 1 shall not apply to certain existing materials, products or devices containing chrysotile fibre when, to perform an equivalent function, no substitute for that fibre is available which:

- On the one hand, in the present state of scientific knowledge, poses a lesser occupational health risk than chrysotile fibre to workers handling those materials, products or devices;

- on the other, provides all technical guarantees of safety corresponding to the ultimate purpose of the use thereof.

II. The scope of application of paragraph I of this Article shall cover only the materials, products or devices falling within the categories shown in an exhaustive list decreed by the Ministers for Labour, Consumption, the Environment, Industry, Agriculture and Transport. To ascertain the justification for maintaining these exceptions, the list shall be re-examined on an annual basis, after which the Senior Council for the Prevention of Occupational Hazards and the National Commission for Occupational Health and Safety in Agriculture shall be consulted.

Article 3

I. The manufacture, processing, importation and domestic marketing of any of the materials, products or devices falling into one of the categories mentioned on the list envisaged under Article 2 shall be subject to a statement, signed, as appropriate, by the head of the business establishment, the importer or the party responsible for domestic marketing, which should be addressed to the Minister for Labour. This statement shall be filed in January of each year or, as appropriate, three months before the start of a new activity or the alteration of an existing production activity, by means of a form decreed by the Ministers for Labour, Consumption, Industry and Agriculture.

The statement shall be accompanied by all the supporting documents in the possession of the declaring party making it possible, considering the state of scientific and technological progress, to determine that as of the date of signature of the statement, the activity covered by the statement meets the conditions set forth in Article 2.I.
II. Activities that have not been the subject of a full statement submitted within the set time-frame may not benefit from the exception granted under Article 2.

III. The Minister for Labour may at all times convey to the author of the statement such information as may seem to him to establish that the material, product or device in question, although falling into one of the categories on the list mentioned in Article 2, does not meet the conditions laid down in paragraph I of that same Article. After requesting comments from the declaring party, he may serve notice to said party to cease manufacture, processing, importation or domestic marketing and to observe the ban instituted under Article 1. He may make such notification public.

Article 4

The manufacture and processing of the materials, products and devices falling into the categories on the list mentioned in Article 2 of this Decree must conform with the rules laid down under Chapters I and II and Chapter III, Section 1 of the aforementioned Decree dated 7 February 1999.

Labelling and marking shall conform with the requirements of Article L. 231-6 of the Labour Code and the rules established by the aforementioned Decree dated 28 April 1998.

Article 5

Without prejudice to the application of the penalties envisaged under Article L. 263-2 of the Labour Code in the event of violation of the provisions of Article I.I of this Decree, the act of manufacturing, importing, introducing into the domestic market, exporting, offering, selling, transferring under any title or possessing for sale all varieties of asbestos fibres or any product containing asbestos fibres, in contravention of the provisions of Article I.II shall be punishable by the fine prescribed for fifth class offences.

Article 6

I. Articles 1, 2, 3 and Article 6.I of the above-mentioned Decree No. 88-466 of 28 April 1998 are hereby repealed.

II. In the first subparagraph of Article 4 of the same Decree, the words: ‘bans envisaged in Article 2 above’ shall be replaced by the word: ‘bans’.

III. In Article 6.II of the same Decree, the words: ‘other than those envisaged under Article 2’ shall be replaced by the words: ‘which are not subject to bans’.

Article 7

Until 31 December 2001 and on a transitional basis, the ban on possession for sale, offering for sale and transfer under any title shall not apply to the used vehicles nor to the agricultural or forestry machinery put into circulation before the effective date of this Decree, and covered by Article R.138 of the Traffic Law. [...]

8.2 The Decree entered into force on 1 January 1997.

2. Main claims by the parties

(a) Main claims by Canada
8.3 Canada claims, firstly that the Decree is a technical regulation covered by the Agreement on Technical Barriers to Trade. As such, it is incompatible with paras. 1, 2, 4 and 8 of Article 2 of the TBT Agreement.

8.4 Secondly, Canada claims that the Decree is incompatible with Articles XI and III:4 of the GATT 1994.

8.5 Lastly, Canada requests that, in the event that the Panel is unable to find a violation of Article XXIII:1(a) of the GATT 1994, it nevertheless finds that the provisions of Article XXIII:1(b) of the GATT 1994 apply.

8.6 Main claims by the European Communities

8.7 With regard to the GATT 1994, the EC request the Panel to confirm that either the Decree does not establish less favourable treatment for imported products than for like domestic products within the meaning of Article III:4 or that the Decree is necessary to protect human health within the meaning of Article XX(b). Lastly, the EC ask the Panel to find that Article XXIII:1(b) of the GATT 1994 does not apply.

8.8 Before going further into the claims by the parties, we must first recall certain decisions taken in the course of the procedure.

B. ISSUES ON WHICH THE PANEL HAD TO TAKE A POSITION DURING THE PROCEDURE

1. Introduction

8.9 In the course of the procedure, the Panel took two decisions which it considers should be reproduced as an integral part of these findings. These were (a) its decision to consult scientific experts on an individual basis taking into account Article 13.1 of the Understanding on Rules and Procedures Governing the Settlement of Disputes; and (b) its decision on whether or not to take into account "amicus curiae briefs" received during the procedure.

2. Consultation of experts

8.10 At the first substantive meeting, the Panel informed the parties of its intention to seek the opinion of individual scientific experts except where, in the light of the parties' written rebuttals, it deemed that such a procedure was not necessary. Having consulted the parties and received their comments, the Panel decided to consult experts individually, in accordance with para. 1 and the first sentence of para. 2 of Article 13 of the Understanding. The Panel's decision was reaffirmed on 2 August 1999. The following are the relevant parts of its letter:

"[…] The Panel carefully examined the arguments put forward by the parties concerning the method of consulting experts, in particular the argument of the European Communities that Article 13.2 of the Understanding requires that, when consulting scientific experts, there should be an expert review group according to the terms of Appendix 4 to the Understanding.

We draw attention to Article 13 of the Understanding which states inter alia that:

"each panel shall have the right to seek information and technical advice from any individual or body which it deems appropriate" and "panels may seek information from any relevant source and may consult experts to obtain their opinion on certain aspects of the matter."
We also note that Article 13.2 provides that panels ‘may’ request an advisory report in writing from an expert review group, in particular but not exclusively in order to consider a factual issue concerning a scientific matter. We consider that this text allows the establishment of such an expert review group but does not prevent experts being consulted on an individual basis with respect to scientific matters or any ‘other technical matter’. In our view, this interpretation of Article 13.2 of the Understanding fully complies with the text of this provision interpreted according to Article 31 of the Vienna Convention on the Law of Treaties and the interpretation of the Appellate Body whereby Article 13 of the Understanding does not prevent a panel from consulting experts individually and leaves the panel free to determine whether it is necessary or appropriate to establish an expert review group.

The Panel also took account of the argument of the European Communities that, if the measure in question should be considered as coming under the TBT Agreement, which it contests in any case, Article 14.2 of the Agreement would mean that a technical expert group would have to be consulted for any scientific or technical question, as well as the position of the European Communities that this provision, pursuant to Article 1.2 of the Understanding, would prevail over the provisions in Article 13 of the Understanding.

Article 14.2 of the TBT Agreement is one of the provisions mentioned in Appendix 2 to the Understanding and which, according to Article 1.2 of the Understanding, prevails over the latter if there is any difference between the two. We note, however, that it is only ‘if there is any difference’ between the rules and procedures in the Understanding and a special or additional rule or procedure covered by Appendix 2 to the Understanding that the latter would prevail. As the Appellate Body has already pointed out, it is only when the provisions in the Understanding and the special or additional rules and procedures in Appendix 2 cannot be construed as complementary that the special or additional rules will prevail over those in the Understanding, in other words, in a situation where the two provisions would be mutually incompatible. In this particular case, Article 14.2 of the TBT Agreement provides that a panel ‘may’ establish a technical expert group. Like Article 13.2 of the Understanding, this text allows the possibility of establishing an expert group and determines the procedures applicable to it, where applicable. It does not however make the establishment of such a group mandatory and, in our view, this possibility is not incompatible with the overall possibility given in Article 13 of the Understanding of consulting experts individually. The two provisions can be seen as complementary.

In this case, it appeared to us that consultation of experts on an individual basis was the most appropriate method inasmuch as it would be the most appropriate way of allowing the Panel to obtain useful information and opinions on the scientific and technical issues raised by this case. In view of the wide variety of fields of competence involved, in particular, it appeared to us to be preferable to obtain different information and opinions individually rather than requesting a joint report on the various scientific or technical issues raised.

[...] In the light of the foregoing, we emphasize that our decision to consult experts individually is without prejudice to the question of the applicability of the TBT Agreement to the measure in question, on which the parties disagree.”

8.11 The Panel reaffirms the decision and the reasoning in this letter.

3. **Amicus curiae briefs**

8.12 In the course of the procedure, the Panel received written submissions or “amicus curiae” briefs from four sources other than Members of the WTO. Referring to the position taken by the Appellate Body in United States – Import Prohibition of Certain Shrimp
and Shrimp Products on the interpretation of Article 13 of the Understanding concerning *amicus curiae* briefs, the Panel informed the parties accordingly and transmitted the submissions to them. The EC included two of these submissions in their own submission. Having examined each of the *amicus curiae* briefs, the Panel decided to take into account the submissions by the *Collegium Ramazzini* and the *American Federation of Labor and Congress of Industrial Organizations*, as they had been included by the EC in their own submissions on an equal footing. At the second meeting with the parties, Canada was given an opportunity to respond in writing and orally to the arguments in the two *amicus curiae* briefs.

8.13 On the other hand, the Panel decided not to take into account the *amicus curiae* briefs submitted respectively by the *Ban Asbestos Network* and by the *Instituto Mexicano de Fibro-Industrias A.C.* and informed Canada and the EC accordingly at the second meeting with the parties held on 21 January 2000.

8.14 On 27 June 2000, the Panel received a written brief from the non-governmental organization ONE ("Only Nature Endures") situated in Mumbai, India. In view of the provisions in the Understanding on the interim review, the Panel considered that this brief had been submitted at a stage in the procedure when it could no longer be taken into account. It therefore decided not to accept the request of ONE and informed the organization accordingly. The Panel transmitted a copy of the documents received from ONE to the parties for information and notified them of the decision it had taken. At the same time, it also informed the parties that the same decision would apply to any briefs received from non-governmental organizations between that point and the end of the procedure.

C. ORDER IN WHICH THE PANEL EXAMINED THE CLAIMS

8.15 The Panel notes that Canada considers that the TBT Agreement is applicable to the Decree because the Decree is a "technical regulation" covered by the Agreement. The EC, on the other hand, consider that the Decree is not a technical regulation, that the TBT Agreement does not apply to it and that it should be considered in the context of the GATT 1994 alone.

8.16 We note that the Marrakesh Agreement Establishing the World Trade Organization constitutes a single treaty instrument that was accepted by the WTO Members as a single undertaking within whose framework all the applicable provisions must be given meaning. Both the GATT 1994 and the TBT Agreement form part of Annex 1A to the WTO Agreement and may apply to the measures in question. Consequently, although we do not in principle exclude application of the TBT Agreement and/or the GATT 1994 to the Decree, we have to determine the order in which we should consider this case. According to the Appellate Body in *European Communities – Regime for the Importation, Sale and Distribution of Bananas*, when the GATT 1994 and another Agreement in Annex 1A appear *a priori* to apply to the measure in question, the latter should be examined on the basis of the Agreement that deals "specifically, and in detail," with such measures. In this particular case, as the parties do not agree on the legal nature of the measure itself (technical regulation covered by the TBT Agreement or general ban coming under the scope of the GATT 1994 alone), it is difficult at this stage to determine which Agreement, either the GATT 1994 or the TBT Agreement, deals with the measure in question most specifically and in the most detailed manner without undertaking an in-depth examination of the measure in the light of each Agreement.

8.17 In order to decide upon the order in which our consideration should proceed, in the way suggested by the Appellate Body, the hypothesis should be that, if the Decree is a "technical regulation" within the meaning of the TBT Agreement, then the latter would deal with the measure in the most specific and most detailed manner. Consequently, in our view it must first be determined whether the Decree is a technical regulation within the meaning of the TBT Agreement. If this is the case, we shall start considering this case by examining the ways in which the Decree violates the TBT Agreement. If we find that the Decree is not a "technical regulation", we shall then immediately start to consider it in the context of the GATT 1994.
D. APPLICABILITY OF THE TBT AGREEMENT TO THE DECREE

1. Arguments of the parties and approach adopted by the Panel

(a) Arguments of the parties concerning the applicability of the TBT Agreement to the Decree

8.18 Canada considers that the Decree is a “technical regulation” within the meaning of the definition given in Annex 1 to the TBT Agreement. The EC consider that the Decree does not come under the definition of a technical regulation in the TBT Agreement.

8.19 The following is the definition of “technical regulation” in Annex 1.1 to the TBT Agreement:

“1. Technical regulation

Document which lays down product characteristics or the related processes and production methods, including the applicable administrative provisions, with which compliance is mandatory. It may also include or deal exclusively with terminology, symbols, packaging, marking or labelling requirements as they apply to a product, process or production method.”

8.20 Canada contends that the Decree considers that all asbestos fibres and all materials, products or devices containing such fibres pose risks for people’s health. The Decree is a “technical regulation” in particular because it lays down a characteristic of a product, a process and a production method, as well as administrative provisions applicable to a product.

8.21 Referring to the definition of “technical regulation” in Annex 1 to the TBT Agreement, Canada considers that the ordinary meaning of the word “characteristic” is “that which constitutes a recognizable distinctive feature”. The Decree describes a recognizable distinctive feature because it bans asbestos fibre, particularly chrysotile, in the manufacturing and processing of materials, products or devices placed on the French market. The characteristic of these materials, products and devices is the absence of asbestos fibres. The Decree indicates precisely or describes the features of the materials, products or devices in whose manufacture or processing the incorporation of asbestos is prohibited. The Decree refers to the processing of all asbestos fibres, whether or not included in products. By so doing, it imposes restrictions on the processes and production methods for asbestos fibres. Moreover, the Decree stipulates that manufacturing activities are subject to the standards governing exposure to asbestos dust in places of business and this imposes a manufacturing process. It includes administrative provisions and a declaration mechanism applicable to products which, exceptionally, contain chrysotile fibre. Lastly, the Decree also deals with labelling requirements and its principal provisions are binding.

8.22 The EC claim that the Decree cannot be construed as a “technical regulation” within the meaning of the TBT Agreement because the latter does not cover general prohibitions on the use of a product for reasons to do with the protection of human health, which come under the GATT 1994. It follows from the preamble, from the background to the TBT Agreement and from the actual wording of several of its provisions that the fundamental objective of the Agreement is to monitor the adoption and application of the “standards” and “technical regulations” that relate to the detailed characteristics of products or their methods of production (e.g. the minimum resistance level for seat-belts). The definition in Annex 1 means that the TBT Agreement does not apply to the Decree because, according to this definition, a technical regulation is a document which lays down the characteristics or processes or production methods with which a specific/identified product must comply, in particular if it is to be released for free circulation on a given market.

8.23 For the EC, any other approach would be equivalent to nullifying the effect of certain provisions of the GATT 1994, for example, Articles I and III, which apply to general
prohibitions. The TBT Agreement must be considered as the specific application of the principles of the GATT 1994 to technical regulations. The Decree prohibits asbestos fibres at all stages. It does not specify the characteristics, the processes or production methods for asbestos fibres and asbestos-containing products or the products exempt from the prohibition measure. The EC do not consider that the fact of not containing asbestos serves to characterize products placed on the French market. For the Decree to be able to lay down the “characteristics” of a product, it would have had to identify the product(s) to which the said characteristics relate. The Decree, however, does not designate any product. It lays down the principle of prohibition. As regards the “processes and methods”, the EC emphasize that the Decree does not lay down any means or set of rules governing the production of asbestos fibres or products containing them.

8.24 In Canada’s view, including a general prohibition within the scope of the TBT Agreement is contrary neither to its object nor its purpose. The distinction made by the EC between prohibitions that apply to all products without distinction and measures aimed particularly at a specific product is not supported by the TBT Agreement. The EC’s interpretation is contrary to the principle of effectiveness: it would suffice to give a measure the form of a general prohibition in order to allow it to evade the disciplines of the TBT Agreement. Canada also considers that, in order to determine whether the Decree satisfies the criteria for the definition of a “technical regulation”, its provisions on both asbestos and exceptions must be examined.

8.25 For the EC, the fact that the definition of “technical regulation” is narrow is not a matter of chance, but signifies that the authors intended to limit the scope of the Agreement. The object and purpose of the TBT Agreement is to deal with technical regulations and standards, not to resolve market access problems associated with general prohibitions. This does not result in a legal vacuum because the general prohibitions continue to be covered by other provisions, in particular Articles I and III of the GATT 1994. The general prohibition eliminates these products from the market. A technical regulation, on the other hand, means that the products affected by the regulation can still be placed on the market. Even in the case of limited and transitional exceptions, the sole preoccupation of the French authorities has been to protect human health. The Decree does not define either the processes and production methods for the products that may be exempted from the general prohibition adopted. The provisions of the Decree concerning exceptions, therefore, do not fall within the scope of the TBT Agreement either.

8.26 Canada claims that the fact that the Decree qualifies as a “technical regulation” is confirmed by France’s notification to the Committee on Technical Barriers to Trade. According to Canada, France thereby recognized the applicability of the TBT Agreement in to the Decree. The EC also recognized the applicability of the TBT Agreement in a document from the European Commission and during the consultations relating to this case.

8.27 According to the EC, the fact that the Decree was notified to the Committee on Technical Barriers to Trade in no way prejudices the applicability of the Agreement. The notification was made in good faith for the sake of transparency. Any other interpretation would create additional obligations for Members and would induce them to discontinue, or at least reduce, notifications.

(b) Approach followed by the Panel

(i) Criteria for the application of the TBT Agreement

8.28 The Panel considers that, for the TBT Agreement to apply to the Decree, the measures imposed under the Decree must come within the definition of “technical regulation” in Annex 1.1 to the TBT Agreement. On the basis of the arguments put forward by Canada and the EC, it appears that the parties share this view. The Panel will therefore examine this definition.
According to Article 3.2 of the Understanding, the Panel must clarify the terms of the WTO Agreement in accordance with customary rules of interpretation of public international law, more particularly those set out in Articles 31, 32 and 33 of the 1969 Vienna Convention on the Law of Treaties. The Panel will therefore first consider the ordinary meaning to be given in good faith to the wording of the definition of “technical regulation” in Annex 1 to the TBT Agreement, taken in context and in the light of the object and purpose of the TBT Agreement and the WTO Agreement in general. If necessary, additional ways of interpreting Article 32 will be taken into account. As the WTO Agreement is a treaty with authentic texts in three languages, it is also important to bear in mind the spirit underlying the provisions of Article 33.

(ii) Differentiation among the prohibitions as such and the exceptions

We note that Canada claims that the exceptions to the ban on asbestos in the Decree confirm that the latter is a technical regulation. The EC state that neither the provisions on exceptions nor the prohibition itself are covered by the TBT Agreement.

Notwithstanding the fact that the main purpose of the Decree is to ban the use of asbestos in France, we note that the text is essentially composed of a general prohibition on the one hand (Articles 1 and 5) and, on the other, exceptions, including the administrative regime applicable to them (Articles 2-4). It must therefore be determined whether we should examine the prohibitions and exceptions as one single measure or as separate measures that might come under different Agreements.

We note first of all that it is possible for several components of the same administrative act to be covered by a single Agreement or by different Agreements. While it is the responsibility of the parties to prove the facts they claim, it is within the competence and is one of the duties of the Panel to determine the law applicable to the facts, with the help of the parties’ arguments. Nothing therefore prevents the Panel from examining the Decree as a single and unique measure, if it believes there are legal grounds for this. Likewise, if the Panel considers that certain provisions of the Decree are covered by provisions of the WTO Agreement other than those applicable to the rest of the Decree, nothing prevents it from considering these provisions in the light of those other provisions of the WTO Agreement.

Consequently, without taking any decision at this stage on the ultimate need for a separate study of the two parts of the Decree – we might in fact reach the conclusion that both parts have the same legal characterization or that one should be considered as supplementing the other – we consider that there should be a separate examination of the provisions specifically concerning the ban on asbestos and the provisions on exceptions in order to make our analysis clearer. We shall therefore first consider the applicability of the definition of “technical regulation” in Annex 1 to the TBT Agreement to the provisions of the Decree on the prohibition. On the basis of these conclusions, we shall then consider the legal nature of the exceptions and any effect they may have on the legal status of the Decree as a whole.

2. Is the Decree a technical regulation within the meaning of the TBT Agreement?

(a) Examination of the part of the Decree prohibiting the marketing of asbestos and asbestos-containing products

(i) Preliminary remarks

Canada puts forward its arguments in general in relation to chrysotile fibres and products containing chrysotile fibres (mainly chrysotile-cement products). Although Canada claims that chrysotile fibre is simply an input in other products and is of no use per se, we consider that it is a product. In our examination of the applicability of the TBT Agreement to the Decree, there is no reason to treat chrysotile fibre differently to products containing chrysotile.
8.35 The Panel notes that the measure at issue in this case is a general ban excluding a given product from the French market as such or when it is incorporated in other products not specified in the Decree. Consequently, the Panel wishes to emphasize that its findings regarding the scope of the concept of technical regulation in the context of the TBT Agreement are linked to the specificities of the measure in question and in no way pre-judge the conclusions that any other panel might reach concerning the same provisions of the TBT Agreement in other factual circumstances.

(ii) Analysis

Ordinary meaning of the terms in the definition in Annex 1 to the TBT Agreement

8.36 The Panel notes first of all that the definition of "technical regulation" relates to the characteristics of a product or its processes or production methods. The Panel notes that the definition uses the word *product*. Applying the principle of effectiveness, we must assume that there was a specific purpose underlying the use of the word "product" in the definition in Annex 1.1 to the TBT Agreement and that it does not appear by chance.

8.37 An initial explanation might be that the authors wished to indicate that this definition related to products rather than services, for example. However, as the TBT Agreement is in Annex 1A to the WTO Agreement, which deals with trade in goods, specifying that the definition applies to products was not necessary. Moreover, it is possible to define a technical regulation applicable to products without specifically using the word "product". As an example, we note that paragraph 3.5.1 of the ISO Guide/IEC 2:1991 contains a definition of the words "technical regulation" that does not employ the word "product". We therefore consider that the authors when using the word "product" were pursuing another objective.

8.38 Another interpretation would be that the purpose sought by introducing the word "product" was to create a link between the technical characteristics and one or more given products. In other words, the product(s) to which the characteristics refer must be clearly identifiable in the document in question. If, in the document, the characteristics described do not refer to an identifiable *product*, the document does not meet the criteria in Annex 1.1 to the TBT Agreement. We consider that this interpretation is better able to give a proper meaning to the word "product" in the definition of "technical regulation" in Annex 1.1 to the TBT Agreement.

8.39 We therefore conclude that a technical regulation is a regulation which sets out the specific characteristics of one or more identifiable products in comparison with general characteristics that may be shared by several unspecified products.

8.40 In this connection, the Panel notes that the ban introduced by the Decree is generally applicable both to asbestos and products containing it, in other words, a very large number of products which the Decree does not identify by name nor even by function or category.

8.41 The Panel also notes that the EC contend that the Decree does not describe the characteristics of a product and that its purpose cannot be to describe the characteristics of an imported product because placing asbestos and asbestos-containing products on the French market is prohibited. In our view, according to the applicable customary rules of interpretation, it is also necessary to examine the meaning of the word "characteristics". The ordinary meaning of "characteristic" is "that which constitutes the distinctive feature, is typical of a person or thing" or "a recognizable distinctive feature". There is thus a link between the characteristics of a product and the product itself. Nevertheless, the "characteristics" must be differentiated from the identification of the product itself. It is indeed possible to describe technical characteristics without specifically identifying the product(s) to which they relate. Likewise, the identification of a product does not suffice to show its characteristics. The Panel notes that the reference to "characteristics" is the special feature of the definition of "technical regulation" in Annex 1.1 to the TBT Agreement. The measure
must not only relate to one or more given products, but its purpose must be to define the characteristics, i.e. the criteria or elements which the product(s) concerned must satisfy in order to be introduced into the territory of the Member that adopted the measure.

8.42 By adding the word “product” before “characteristics”, the authors of Annex 1.1 therefore wished to specify the circumstances in which the TBT Agreement applied. In order purely and simply to ban the import of a product, it is not absolutely necessary to define its characteristics. In the same way, if it is desired to exclude certain raw materials as such, it is not necessary to specify the products in which they can be incorporated. On the other hand, if the characteristics of a given product are identified – even those which mean that import is not allowed – at the same time the characteristics of products which can be introduced into the territory of the country applying the measure are identified.

8.43 The Panel therefore concludes that, taking into account the ordinary meaning of the words “characteristics” and “product”, the definition of “technical regulation” in Annex 1.1 to the TBT Agreement applies to the measures which define the technical specifications that one or more given products must meet in order to be authorized for marketing in a Member.

8.44 It may, however, be considered, as Canada does, that the prohibitions in the Decree concern a given product: asbestos, because of its specific characteristics. Nevertheless, as far as the ban on asbestos as such is concerned, the Decree does not concern itself with the characteristics of this product, in any event, not specifically. What is banned is not asbestos which possesses certain characteristics but all types of asbestos. As regards asbestos-containing products, the Panel notes Canada’s argument that the characteristic of the products authorized for import is the absence of asbestos. We note, however, that the definition in Annex 1.1 to the TBT Agreement refers to the technical characteristics of a given product. In this case, the products whose characteristic would be that they contained asbestos are not identified in the Decree. As we have already found, the identification of a product appears to be one of the criteria in the definition in Annex 1.1.

8.45 Lastly, the Panel notes that the ban applicable to asbestos-containing products is in fact one aspect of or complementary to the general ban on asbestos. It can therefore only be considered together with the ban on asbestos as such. It is in fact there in order to ensure that asbestos is not imported into France in any form whatsoever. In such a context, not defining the products concerned is not simply a choice of wording but is the result of the logic of the objective sought (a ban on all forms of asbestos). To specify each product affected would be pointless.

8.46 To the extent that Article 31 of the Vienna Convention contains a single rule of interpretation and not a number of alternative rules, the various criteria in the Article should be considered as forming part of a whole. Although the Panel subdivided its analysis for reasons of clarity, this does not mean that there has to be a hierarchy among the various elements in the rule of interpretation. In this particular case, we deem it preferable to analyse the context after examining the object and purpose because the latter help to clarify the understanding of the context of “technical regulation” in Annex 1 to the TBT Agreement.

8.47 We note that the object and purpose of a treaty can also be found in its preamble. Applying the practice of the Appellate Body in this respect, it is relevant to look not only at the preamble to the WTO Agreement but also at the preamble to the TBT Agreement itself, which provides certain indications.

8.48 In this connection, we note first of all that, although the preamble to the WTO Agreement refers in particular to expanding trade in goods, it does not give sufficiently precise information on the terms which we must interpret. We therefore turn to the preamble to the TBT Agreement. This refers to standards for products and their role in improving
8.49 It might be argued that the fact that the TBT Agreement only refers to elements relating to the marketing criteria for a product does not in itself mean that it does not apply to import bans. Strict application of standards can in fact lead to a ban on the import of certain products. The Panel notes, however, that there is no specific reference in the TBT Agreement to the type of measure involved in this particular case (i.e. a pure and simple import ban without any reference to the characteristics of given products), even though a ban is by its very nature the most restrictive market access measure. Consequently, if the Members had agreed that the TBT Agreement also applied to general bans, they would undoubtedly have mentioned it. It would appear that the purpose of the TBT Agreement is to prevent much more complex situations than a straightforward unconditional ban on a product, which is covered by the very strict provisions in Article X:1 of the GATT 1994. In the Panel’s view the purpose of adopting the TBT Agreement was to control the development and application of standards - situations in which protectionist aims can be better disguised and for which the existing disciplines within the GATT appeared to be inadequate. A general prohibition by nature does not usually involve any technical specifications. It thus appears that the drafters were concerned at the development and application of technical characteristics for protectionist purposes or leading to protectionist effects, as can be seen from the preamble to the TBT Agreement:

"Desiring however to ensure that technical regulations and standards, including packaging, marking and labelling requirements [...], do not create unnecessary obstacles to international trade." (Emphasis added).

8.50 It would therefore seem that in principle the object and purpose of the TBT Agreement were that it should apply to potentially protectionist aspects of the application of technical regulations and standards which, even if their effect might be purely and simply a ban, could circumvent the GATT disciplines more easily for a number of reasons.

8.51 The Panel nevertheless notes that Canada points out that the imposition of technical characteristics applicable to the import of a product implies a ban on products which do not meet these characteristics. Consequently, any technical regulation is also a general prohibition on products that are not in conformity. The Panel does not disagree. It notes however that, in this particular case, none of the products covered by the Decree can be imported, with the exception of those given a temporary exemption. This is a major difference compared with a technical regulation that defines the characteristics of one or more given products. Even if the number of products that can in fact be imported is very restricted, import remains possible, in theory or in practice.

8.52 This does not mean either that a technical regulation which defines the characteristics of a product negatively does not come under the TBT Agreement. It is always possible to define the characteristics of a product negatively as long as these technical characteristics are specified in such a way as to allow the product to be marketed in the importing countries. As we emphasized in para. 8.40 above, the Decree contains a general prohibition that does not define the characteristics of specific products.

Context

8.53 The Panel notes that the provisions of the TBT Agreement are situated within the broader context of Annex 1A to the WTO Agreement. The EC emphasize in this connection that an interpretation reaffirming the applicability of the TBT Agreement to a general prohibition measure would be equivalent to nullifying the effect of Articles I, III, and XI of the GATT. Canada considers that, if the ban in the Decree is not deemed to be a "technical regulation", there is a risk of circumventing the disciplines of the TBT Agreement by intro-
ducing “horizontal” prohibitions instead of technical regulations in the strict sense. We consider that these matters should be examined more specifically in the light of the context of the TBT Agreement because it is by taking into account the content of the obligations in the provisions that form part of the context of the TBT Agreement that we will be able to determine the scope of the definition of “technical regulation” in the TBT Agreement.

8.54 The Panel considers that the concept of circumvention implies that a given attitude makes it possible to avoid an obligation that would otherwise apply. An obligation must be generally applicable before circumvention can occur. Firstly, nothing proves that it is in a Member’s interest to try to evade the obligations in the TBT Agreement because its provisions represent a combination of rights and obligations whose balance was carefully negotiated. Nevertheless, assuming that it was in the Member’s interest, if the object and purpose of the TBT Agreement are to regulate the marketing conditions, a measure that quite simply banned market access would not come under the Agreement. This does not, however, mean that such measures would not be penalized if they created “unnecessary obstacles to international trade”. To the extent that the GATT 1994 applies, especially Articles I, III or XI whose scope is particularly broad, there would not be any legal vacuum.

8.55 We also note that the criteria on the preparation, adoption or application of technical regulations in Article 2.2 of the TBT Agreement are very similar to those in Article XX of the GATT 1994. The preamble to the TBT Agreement in fact repeats some of the wording of Article XX of the GATT. In the Panel’s view the TBT Agreement is a development of the GATT. As mentioned in para. 8.49 above, the TBT Agreement appears to have been adopted in order to strengthen the disciplines applicable to the specific area of manufacturing standards, where they appeared to be insufficient to prevent certain forms of protectionism.

8.56 We also note that the EC claim that Canada’s interpretation of the TBT Agreement would be equivalent to nullifying the effect of Articles I and III of the GATT 1994. We note in this connection that the Appellate Body has on many occasions, most recently in the Argentina – Safeguards case, recalled that “a treaty interpreter must read all applicable provisions of the treaty in a way that gives meaning to all of them, harmoniously”. It is therefore important that our interpretation of the provisions of the TBT Agreement in this case should not result “in reducing whole clauses or paragraphs of a treaty to redundancy or inutility”. Canada is concerned that a particular interpretation might allow Members to circumvent the obligations imposed by the TBT Agreement. As already indicated above, in our view the main reason for circumventing a provision is to avoid certain obligations. If a general prohibition is not covered by the TBT Agreement, this does not mean that it is not subject to any other provision in the WTO Agreement. At the very least, it will come within the scope of Articles I, and/or III, and/or XI of the GATT 1994 and may have to meet the criteria in Article XX in order to be justified under this Agreement. There may be grounds for Canada’s position if it is established that the provisions of the TBT Agreement impose stricter disciplines than those in the GATT 1994 and that excluding general prohibitions from their scope would nullify their effect. The elements before us, however, do not allow us to conclude that this is the case.

8.57 The Panel therefore concludes that, taking into account the terms of the definition considered within their context and in the light of the object and purpose of the TBT Agreement, a measure constitutes a “technical regulation” if:

(a) the measure affects one or more given products;

(b) the measure specifies the technical characteristics of the product(s) which allow them to be marketed in the Member that took the measure;

(c) compliance is mandatory.
In the light of the above, we provisionally conclude that the part of the Decree dealing with the general prohibition on marketing asbestos and asbestos-containing products does not constitute a “technical regulation” within the meaning of the definition in Annex 1.1 to the TBT Agreement.

(iii) Additional arguments by Canada

The Panel notes that Canada puts forward two additional arguments that must be examined before coming to a definite conclusion on this matter. Firstly, in Canada’s view, the EC recognized that the TBT Agreement applied by notifying the Decree to the Committee on Technical Barriers to Trade and during the consultations relating to this dispute.

From a legal point of view, the question seems to be whether there is estoppel on the part of the EC because they notified the Decree or because of their statements, including those during the consultations. This would be the case if it was determined that Canada had legitimately relied on the notification of the Decree and was now suffering the negative consequences resulting from a change in the EC’s position. In this case, however, it does not appear that Canada was able legitimately to rely on a notification to the Committee on Technical Barriers to Trade or on a statement made during the consultations. We consider that notifications under the TBT Agreement are made for reasons of transparency. It has been recognized that such notifications do not have any recognized legal effects. Furthermore, notification under the TBT Agreement is one of the few ways of notifying this type of measure for a Member who wishes to show transparency in good faith. Lastly we consider that both the notification and the comments made by the EC during the consultations or in another context constitute observations on the legal characterization of the Decree. Claims regarding the legal characterization of a fact by the parties, however, cannot bind the Panel.

Lastly, we note that Canada asserts that the Decree contains five of the requirements in the definition in Annex 1.1 to the TBT Agreement, namely, it indicates the characteristics of a product, the processes and production methods relating to a product, the administrative provisions applicable to a product, and packaging or labelling requirements for a product. Finally, compliance is mandatory. We do not in principle rule out that the Decree may include some or even all the requirements mentioned by Canada. We note, however, that our consideration of the terms of the definition of “technical regulation” in Annex 1.1 in their context and in light of the object and purpose of the TBT Agreement in relation to the part of the Decree prohibiting the marketing of asbestos has led us to conclude that this part of the Decree is not covered by the TBT Agreement. Concerning Canada’s arguments, we have found that the part of the Decree prohibiting asbestos and for the production of asbestos-containing products does not define the characteristics of the products whose import is banned. We have also found that this part of the Decree does not cover processes or methods for the production of asbestos or asbestos-containing products. Even if we adopt the approach suggested by Canada for the identification of the five requirements, in our view, the absence of the first two requirements suffices to conclude that the Decree does not meet the criteria for the definition of “technical regulation” in the TBT Agreement. We do not therefore think it necessary to consider whether the three other requirements referred to by Canada are to be found in this part of the Decree.

We nevertheless note that, in its arguments, Canada does not distinguish, as the Panel does, between the part of the Decree banning asbestos and the part relating to exceptions to the ban. It is thus possible that the elements to which Canada refers are to be found in the part of the Decree on exceptions. If this was the case, this would be even greater justification for our decision to analyse the two aspects of the Decree separately.

We therefore conclude that the part of the Decree dealing with the general ban on the marketing of asbestos and asbestos-containing products does not constitute a “technical regulation” within the meaning of the definition in Annex 1.1 to the TBT Agreement.
(b) Analysis of exceptions and the effect of the type of exceptions on the findings concerning prohibitions

(i) The exceptions in the Decree constitute technical regulations

8.64 Bearing in mind the discussion in the preceding section, we point out that the following aspects of a measure are likely to mean that the measure is covered by the definition of “technical regulation” in Annex 1.1 to the TBT Agreement.57

(a) The measure affects one or more given products;

(b) the measure specifies the technical characteristics of the product(s) which allow them to be marketed in the Member that took the measure58;

(c) compliance is mandatory.

8.65 First of all, we note that the EC contend that exceptions are transitional measures limited by their purpose and temporary by nature. Consequently, they should not be covered by the TBT Agreement. We do not consider that the “transitional” nature of a measure suffices to exclude it from the scope of the TBT Agreement. The Agreement does not distinguish between measures according to whether or not they are transitional or temporary.

8.66 We also note the EC’s arguments on the ancillary nature of the exceptions.59 We have already indicated above that, in theory, there is no reason why two parts of the same text could not have a different legal characterization. We note that, pursuant to the accessory theory (accessorium sequitur principale), the legal regime applicable to a good or an action may be related or subject to the regime for a principal good or action.60 Assuming that this theory applies to a public international law situation, we consider that this particular case does not come within its scope. The measures before us do not relate to application of the ban in the strict sense but to an exception which in practice allows chrysotile asbestos to enter the French market. In such a context, we hesitate to consider the exceptions in the Decree as ancillary to the ban on asbestos.

8.67 We note that the part of the Decree concerning exceptions to the general ban on importing asbestos or asbestos-containing products (Articles 2-4) does not define the products benefiting from such exceptions. Article 2.II, however, covers the identification of the materials, products or devices shown in an exhaustive list drawn up by decree by the Ministers for Labour, Consumption, the Environment, Industry, Agriculture and Transport. We therefore consider that the applicable French regulation (i.e. the Decree as implemented by the above-mentioned Decree) identifies the products benefiting from an exception.

8.68 We also note that Article 2 of the Decree sets out the criteria for marketing the products identified in the Decree and not solely the criteria for excluding products from the market. The second sentence in Article 3.I of the Decree completes these criteria.

8.69 In our view, the marketing criteria in Article 2.I of the Decree relate to the characteristics of one or more given products or processes or production methods relating to them. This is particularly true of the second subparagraph on the technical guarantees of safety appropriate to the use, which has to be read in conjunction with the second paragraph of Article 3.I of the Decree. We also note that Article 2.II and Article 3 in particular cover the administrative provisions applicable to the technical regulations. The second paragraph of Article 4 contains provisions on the labelling and marking of products benefiting from an exception. Lastly, compliance with these provisions is mandatory and there is provision for penalties (see Article 3.III).
8.70 We thus conclude that the part of the Decree concerning exceptions to the ban on asbestos comes within the scope of the definition of technical regulation in Annex 1.1 to the TBT Agreement.

(ii) Effect of the legal characterization of the exceptions on the legal characterization of the prohibitions

8.71 The Panel notes that Canada considers that the regime applicable to exceptions proves that the Decree as a whole is a technical regulation. We are of the opinion that there is no legal reason why certain parts of the Decree should not come under one of the WTO Agreements and other parts under another Agreement. We have already referred above to application of the accessory concept in this particular case. Even assuming that the accessory concept might apply to this particular case, the fact that it is the exceptions to the general ban which come under the TBT Agreement and not the principal element of the Decree (namely the general ban on asbestos) makes it difficult to conclude that the ban must follow the exceptions regime. Consequently, the fact that the exceptions in the Decree are covered by the TBT Agreement in our view should not affect the legal characterization of the general ban and especially the fact of whether or not the ban comes within the scope of the TBT Agreement.

8.72 In any event, as the legal characterization of the exceptions does not affect that of the general ban and even though Canada contests the legality of the Decree as a whole, the Panel should not go further than the question of the legal characterization of the exceptions. The Panel considers that Canada did not make any specific claims concerning the exceptions in the terms of reference drawn up under Article 6.2 of the Understanding. The Panel therefore concludes that it does not have to reach any findings concerning the exceptions.

3. Conclusion

(a) The TBT Agreement does not apply to the part of the Decree relating to the ban on imports of asbestos and asbestos-containing products because that part does not constitute a "technical regulation" within the meaning of Annex 1.1 to the TBT Agreement.

(b) The TBT Agreement applies to the part of the Decree relating to exceptions to the ban on imports of asbestos and asbestos-containing products because that part constitutes a "technical regulation" within the meaning of Annex 1.1 to the TBT Agreement. This legal characterization, however, does not affect the legal characterization of the part of the Decree banning asbestos nor our consideration of the rest of this case because Canada did not make any specific claims regarding the exceptions to the general ban.

8.73 Having concluded that the part of the Decree banning asbestos and asbestos-containing products does not come within the scope of the TBT Agreement, we shall now consider Canada’s claims relating to the GATT 1994.

E. APPLICATION OF THE GATT 1994 TO THE DECREE

1. Preliminary questions

(a) The effect of practice within the GATT 1947 and the WTO

8.74 The Panel notes that, in support of their arguments, the parties refer to panel and Appellate Body reports adopted in the WTO. They also refer at length to panel reports under the GATT 1947, some of which were adopted and others not. As certain issues raised by the parties have been dealt with in detail or even exclusively under the GATT 1947, we deem it useful to explain how we intend to take into account the panel reports issued within the framework of the GATT 1947. For this purpose, we shall base ourselves on the approach followed by the Appellate Body in Japan – Taxes on Alcoholic Beverages.
8.75 In this particular case, the Appellate Body emphasized that, with regard to the reports adopted by panels under the GATT 1947, Article XVI:1 of the Agreement Establishing the WTO and paragraph 1(b)(iv) of Annex 1A incorporating the GATT 1994 into the WTO Agreement allow the “legal history and experience under the GATT 1947” to be brought into the new WTO. Adopted panel reports “are an important part of the GATT acquis”. They create legitimate expectations among WTO Members and should thus be taken into account when they are relevant to any dispute.

8.76 Turning to unadopted reports, the Appellate Body stated that unadopted panel reports “have no legal status in the GATT or WTO system since they have not been endorsed through decisions by the CONTRACTING PARTIES to GATT or WTO Members”. [footnote omitted]. Likewise, “a panel could nevertheless find useful guidance in the reasoning of an unadopted panel report that it considered to be relevant”. [footnote omitted].

8.77 Following this approach, we shall duly take into account the relevant adopted panel reports and, to the extent necessary, we shall take into account the reasoning set out in unadopted panel reports when we deem it relevant to the issues raised in our examination.

8.78 Canada claims violation of Articles III:4 and XI:1 of the GATT 1994. The EC considers that, even if there is a violation of these provisions, the measure is justified pursuant to Article XX(b). We note that the Appellate Body reaffirmed the general rules applicable to the burden of proof in the case of United States – Shirts and Blouses.

8.79 Applying these rules, it is our opinion that Canada, as the complaining party, should normally provide sufficient evidence to establish a presumption that there are grounds for each of its claims. If it does so, it will then be up to the EC to adduce sufficient evidence to rebut the presumption. When the EC puts forward a particular method of defence in the affirmative, it is up to them to furnish sufficient evidence, just as Canada must do for its own claims. If both parties furnish evidence that meets these requirements, it is the responsibility of the Panel to assess these elements as a whole. Where the evidence concerning a claim or a particular form of defence is, in general, equally balanced, a finding has to be made against the party on which the burden of proof relating to this claim or this form of defence is incumbent.

8.80 The Panel also notes the extreme factual and scientific complexity of this case, which led to the consultation of experts under Article 13 of the Understanding. It notes that, in the context of the Agreement on the Application of Sanitary and Phytosanitary Measures, where scientific facts and data play a very important role, the Appellate Body has taken a special approach. As it is necessary to examine the application of Articles III:4 and XI:1 of the GATT 1994 before even considering the EC’s arguments based on Article XX, we simply take note of this approach and, if necessary, will return to it when examining the issues raised in connection with Article XX.

8.81 Bearing in mind the above, we consider that it is important for each party to state clearly its arguments and factual evidence in support of its claims. If not, the Panel will only be able to conclude that the party concerned did not make the prima facie case needed to establish its claim. The Panel considers that it can only reach findings concerning a claim or part of a claim, even when it is clearly defined in the terms of reference set by the DSB, if the party making the claim spontaneously makes the prima facie case needed to establish its claim. Information provided by the experts consulted by the Panel pursuant to Article 13 of the Understanding “to help it to understand and evaluate the evidence submitted and the arguments made by the parties”, even where it has been requested by the Panel, can under no circumstances be used by a panel to rule in favour of a party which has not established a prima facie case based on specific legal claims or pleas asserted by it. Put another way, the Panel cannot utilize the information furnished by the experts in
order to establish the validity of a claim by one of the parties if that party has not already established a prima facie case.

8.82 On any legal matter, notably the interpretation of the Agreements concerned, the Panel will be assisted by the arguments of the parties but will not be bound by them. Pursuant to Article 3:2 of the Understanding, our decision on these matters must be in accordance with the customary rules of interpretation of public international law applicable to the WTO Agreement.

(c) Application of Article III:4 and/or Article XI of the GATT 1994

(i) Question before the Panel

8.83 The Panel notes that the parties have differing views concerning the applicability to the Decree of Article III:4 and Article XI:1 of the GATT 1994.

8.84 Canada considers that the Decree should be examined in the light of these two Articles because it comprises two distinct aspects. On the one hand, it prohibits imports of asbestos in a manner incompatible with Article XI:1. On the other, it includes internal regulations prohibiting the sale and use of asbestos in a manner incompatible with Article III:4. Canada considers that two different aspects of the same measure can be examined in the light of two different articles of the same Agreement. Canada considers that the interpretative Note Ad Article III does not apply in this case. If the Panel considers that the Decree cannot be examined under the two Articles, it should be considered as a measure affecting imports and incompatible with Article XI:1. If it cannot be considered as a measure affecting imports, it should be considered as a measure affecting sales and other transactions on the French market and so incompatible with Article III:4.

(ii) Analysis

8.85 The EC assert that either the measure is an internal regulation, in which case it is covered by Article III:4, or it only concerns the import of products, in which case it must be assessed in the light of Article XI:1. The distinction made by Canada fails to take account of the relationship between the two Articles, which means that a single measure that applies both to domestic and imported products must necessarily be covered as a whole by Article III:4 if it is imposed on an imported product at the time or place of importation. Previous practice in the GATT confirms that there can be no cumulative application with Article XI. The practical application of the Decree means that the same measure, namely a general ban on asbestos and asbestos-containing products, applies to all products irrespective of their origin. As one and the same measure is applied to both domestic and imported products – application is at the border for the latter – only Article III:4 is applicable in this case.

Note Ad Article III in the Notes and Supplementary Provisions in Annex I to the GATT 1994 states the following:

“Any internal tax or other internal charge, or any law, regulation or requirement of the kind referred to in paragraph 1 which applies to an imported product and to the like domestic product and is collected or enforced in the case of the imported product at the time or point of importation, is nevertheless to be regarded as an internal tax or other internal charge, or a law, regulation or requirement of the kind referred to in paragraph 1, and is accordingly subject to the provisions of Article III.”
Article XI:1 of the GATT 1994 states the following:

“No prohibitions or restrictions other than duties, taxes or other charges, whether made effective through quotas, import or export licences or other measures, shall be instituted or maintained by any contracting party on the importation of any product of the territory of any other contracting party or on the exportation or sale for export of any product destined for the territory of any other contracting party.”

8.87 The Panel notes first of all that the parties agree that Article III:4 applies to that aspect of the Decree which bans in particular the sale, domestic marketing and transfer under any title of all varieties of asbestos fibres and any product containing them. This aspect concerns the “treatment accorded to products […] in respect of all laws, regulations and requirements affecting their internal sale, offering for sale, purchase, transportation, distribution or use” within the meaning of Article III:4. Canada, on the other hand, considers that Article XI:1 applies to the ban on imports affecting products from Canada.71

8.88 The Panel draws attention to Note Ad Article III, which specifically covers a situation in which a law, regulation or requirement applies both to an imported product and to the like domestic product and is enforced in the case of the imported product at the time or point of importation. The latter is in fact the case. Consequently, the Panel considers it proper to commence its analysis by determining whether the Note Ad Article III applies to this case. As neither of the parties contests the fact that the measure applicable to the imported product is imposed at the time or point of importation, it is necessary to examine whether the Decree “applies to an imported product and to the like domestic product”.

8.89 In Canada’s view, interpretative Note Ad Article III only applies if the measure is applicable to the imported product and to the domestic product. The explicit import ban does not, however, apply to the domestic product because the domestic product is obviously not imported. Moreover, as France neither produces nor mines asbestos fibres on its territory, the ban on manufacturing, processing, selling and domestic marketing is, in practical terms, equivalent to a ban on importing chrysotile asbestos fibre.

8.90 For the EC, the import ban is merely the logical corollary of the general prohibition on the use of asbestos and asbestos-containing products. Article III:4 must be assessed in the light of the interpretative Note relating to it. When a domestic measure applies to both domestic and imported products, Article III must apply.

8.91 The Panel notes that the word “comme” in the French text of Note Ad Article III [“and” in the English text] implies in the first place that the measure applies to the imported product and to the like domestic product.72 The Panel notes in this connection that the fact that France no longer produces asbestos or asbestos-containing products does not suffice to make the Decree a measure falling under Article XI:1. It is in fact because the Decree prohibits the manufacture and processing of asbestos fibres that there is no longer any French production. The cessation of French production is the consequence of the Decree and not the reverse. Consequently, the Decree is a measure which “applies to an imported product and to the like domestic product” within the meaning of Note Ad Article III.

8.92 Secondly, the Panel notes that the words “any law, regulation or requirement […] which applies to an imported product and [“comme” in the French text] to the like domestic product” in the Note Ad Article III could also mean that the same regime must apply to the imported product and the domestic product.72 In this case, under the Decree, the domestic product may not be sold, placed on the domestic market or transferred under any title, possessed for sale, offered or exported. If we follow Canada’s reasoning, products from third countries are subject to a different regime because, as they cannot be imported, they cannot be sold, placed on the domestic market, transferred under any title, possessed for sale or offered. Firstly, the regulations applicable to domestic products and foreign products lead to the same result: the halting of the spread of asbestos and asbestos-containing products on French territory. In practice, in one case (domestic products), they
cannot be placed on the domestic market because they cannot be transferred under any title. In the other (imported products), the import ban also prevents their marketing.

8.93 For the following reasons, we also consider that the wording of Note Ad Article III and practice in the GATT 1947 in this respect do not support Canada’s approach that an identical measure must be applied to the domestic product and the like imported product if the measure applicable to the imported product is to fall under Article III.

8.94 We note that the relevant part of the English text of Note Ad Article III reads as follows: “Any [...] law, regulation or requirement [...] which applies to an imported product and to the like domestic product”. The word “and” does not have the same meaning as “in the same way as”, which can be another meaning for the word “comme” in the French text. We therefore consider that the word “comme” cannot be interpreted as requiring an identical measure to be applied to imported products and domestic products if Article III is to apply.

8.95 We note that our interpretation is confirmed by practice under the GATT 1947. In United States – Section 337 of the Tariff Act of 1930, the Panel had to examine measures specifically applicable to imported products suspected of violating an American patent right. In this case, referring to Note Ad Article III, the Panel considered that the provisions of Article III:4 did apply to the special procedures prescribed for imported products suspected of violating a patent protected in the United States because these procedures were considered to be “laws, regulations and requirements” affecting the internal sale of the imported products, within the meaning of Article III of the GATT. It should be noted that in this case the procedures examined were not the same as the equivalent procedures applicable to domestic products.

8.96 Canada also claims that the import ban is not an internal measure imposed at the border, for administrative reasons. We consider that an internal charge applied to a domestic product must also be imposed on an imported product. Nevertheless, if it is deemed appropriate to impose the charge at the border rather than waiting until the imported product is actually marketed, the same logic applies in the case of a regulatory measure prescribing a ban on marketing. Is it not equally preferable from the administrative point of view and in the interests of the importers themselves to prevent the entry of the like product into the country applying the measure rather than waiting until it is placed in a warehouse before banning its sale?

8.97 The Panel also notes that Canada makes reference to paras. 4.24 and 4.26 of the Report of the Panel in Canada – Import, Distribution and Sale of Certain Alcoholic Drinks by Provincial Marketing Agencies to support its claim that the Decree should be examined under Article XI:1 of the GATT 1994. We note that in paragraph 4.24 of the Report, the Panel considered that according to the Note Ad Articles XI, XII, XIII, XIV and XVIII, restrictions made effective through state-trading operations were “import restrictions” or “export restrictions”. It considered that, in the case of enterprises enjoying a monopoly of both importation and distribution in the domestic market, the distinction normally made between restrictions affecting the importation of products and restrictions affecting imported products lost much of its significance since both types of restriction could be made effective through decision by the monopoly. In this case, the Decree did not institute a monopoly on the import or distribution of asbestos and like products, so the Note Ad Articles XI, XII, XIII, XIV and XVIII is not relevant to settlement of this matter.

8.98 As regards Canada’s reference to paragraph 4.26 of the aforementioned report, we consider that it does not substantiate Canada’s position in this case either. In this paragraph, the Panel refrains from ruling on a violation of Article III:4. It appears to do so, however, for reasons of legal economy because it simultaneously recognizes that Article III:4 could apply to state-trading transactions. Contrary to Canada’s assertion, this paragraph does not confirm the non-applicability of Article III:4 to the part of an internal measure dealing with the treatment of imported products. At the most, it could confirm the application of both provisions. Nevertheless, as explained in the preceding paragraph, the
Panel found that Article XI:1 applied, referring to the Note Ad Articles XI, XII, XIII, XIV and XVIII. This Note only applies to state-trading transactions. In the present case, however, there is no question of a measure applied in the context of state-trading activities.

8.99 For the foregoing reasons, we consider that Article III:4 of the GATT 1994 applies to the ban on importing asbestos and asbestos-containing products imposed by the Decree. On the basis of the grounds for this conclusion, we do not consider it necessary to examine further Canada’s arguments on the exclusive application of Article XI:1.

8.100 Looking at Canada’s arguments, it is difficult to see whether Canada is claiming that, even if the import ban falls under Article III:4, it is also covered by Article XI:1. If Canada does in fact make such a claim for the cumulative application of Article III:4 and Article XI:1 to the part of the Decree banning imports, we do not consider that this forms part of the terms of reference given to the Panel by the DSB and, even if that were the case, Canada’s arguments do not make a prima facie case in the sense given to this concept by the Appellate Body. Consequently, we do not consider it necessary to examine this point any further.

2. Violation of Article III of the GATT 1994

(a) Arguments of the parties

8.101 According to Canada the likeness of products should be assessed on a case-by-case basis, considering, in particular, the end-use of the product, consumers’ tastes and habits, and the properties, nature and quality of the product. To these should be added tariff classification. Precedents under the GATT 1947 and the GATT 1994 do not, however, require that all the criteria be applied when evaluating the likeness of given products. Moreover, “like” does not mean “identical”, it is a matter of showing that the products compared share many similar features. For Canada, applying the criteria in the precedents confirms the likeness of polyvinyl alcohol (hereinafter “PVA”), cellulose and glass fibres and chrysotile fibre, on the one hand, and fibro-cement and chrysotile-cement products on the other.

8.102 The EC contend that asbestos and asbestos-containing products, on the one hand, and substitute products, on the other, are not like products within the meaning of Article III:4 of the GATT 1994. Four criteria in particular can be used to assess the likeness of products: (a) their properties, nature and quality; (b) their tariff classification; (c) their end-use; and (d) consumers’ tastes and habits. In this case, three criteria are relevant: the properties, nature and quality; the tariff classification and the end-use of the product. In the EC’s view Canada is confusing the concept of “like” product in Article III:4 with that of “competitive” or “directly substitutable” product in Article III:2, read in conjunction with the relevant interpretative Note. In this case, although certain fibrous products are indeed “substitutable” for chrysotile asbestos and products containing it, they are nevertheless not “like” products. Asbestos has unique physical characteristics and properties that make it difficult to replace for certain industrial purposes. This is why the Decree envisages exceptions. As asbestos has so many uses, there is no single natural or synthetic product which, alone, could replace it in all the products and materials that contain asbestos.

(b) Issues raised in connection with the arguments of the parties relating to Article III:4 of the GATT 1994

8.103 The Panel considers that the arguments of the parties raise the following issues:

(a) The first issue is the criteria to be applied in order to determine the likeness of the products in question. The Panel notes that the parties do not in fact agree on the scope of the concept of like product under Article III:4. Nevertheless, with regard to the criteria to be taken into consideration when assessing likeness, the parties referred to the criteria contained inter alia in the Report of the Working Party on Border
Tax Adjustments\textsuperscript{52} and in the Reports of the Panel and the Appellate Body in Japan – Taxes on Alcoholic Beverages.\textsuperscript{53} The problem can therefore be summarized as whether the criteria developed in these cases must be applied as such or in a particular way in the context of Article III:4, or whether additional criteria should be taken into consideration. Inasmuch as these are the only criteria developed to date in connection with Article III, we consider it appropriate to commence our analysis by applying them to the facts in this case. If, when applying these criteria, we encounter difficulties specific to Article III:4, we shall deal with them at that stage. Moreover, we note that in the above-mentioned case the Appellate Body specified that each criterion should be assessed on a case-by-case basis.\textsuperscript{84} It is thus not a question of systematically determining the criteria applicable in the context of Article III:4 but rather of deciding which are relevant in this case.

(b) The second issue is related to the fact that the parties often refer in general to substitute products for chrysotile, products containing asbestos in general, chrysotile-cement products or fibro-cement products. Taking into account the practice of the Appellate Body regarding determination of the likeness of products and the burden of proof in this connection, we consider that the identification of the products which have to be compared pursuant to Article III:4 is of importance in our findings.\textsuperscript{85} We also have to decide how to deal with the fibres as such compared to products containing the fibres.\textsuperscript{86}

8.104 In view of the particular implications of the second question for our findings, it appears to us necessary first to devote a special section of the report to this.

(c) The Panel’s approach to product-by-product analysis and certain specific aspects of the burden of proof.

8.105 The Panel notes that the EC have indicated that 90 per cent of imported chrysotile fibre was used for making asbestos-cement products prior to the application of the Decree.\textsuperscript{87} The Panel also notes that Canada “focuses” its arguments on chrysotile fibres and substitute fibres for chrysotile, as well as on chrysotile-cement and fibro-cement products. With regard to substitute fibres for chrysotile, Canada points out that there are more than 150 substitute fibres for chrysotile fibre. Nevertheless, Canada indicates that, in view of Article III:4 of the GATT 1994, it considers that the Panel’s findings and conclusions should take into account the impact of the Decree in relation to three substitute fibres: PVA, cellulose fibres and glass fibres, as well as the fibro-cement products incorporating these. Canada also indicates that it does not invoke the argument of likeness with respect to non-fibrous substitutes for chrysotile fibres (e.g. PVC or ductile iron), nor to non-fibrous products used as substitutes for chrysotile-cement products.

8.106 The EC indicate that in its request for consultations and its request for the establishment of a panel, Canada never claimed that this dispute concerned high-density cement products containing chrysotile. The Panels terms of reference do not refer to these either. Moreover, the EC point out that Canada only exported chrysotile asbestos as a raw material and not in the form of high-density cement products containing chrysotile.

8.107 Taking into account the rules governing the burden of proof\textsuperscript{88}, the Panel in the first place considers that it should restrict its examination of the likeness of products in the context of Article III:4 to those products identified by Canada for which the parties provided evidence or made an adequate prima facie case to establish their likeness or their absence of likeness to chrysotile and products containing it.\textsuperscript{89}

8.108 We note, that although friction products are mentioned, Canada does not include these in its comparisons relating to Article III:4. We therefore conclude that Canada does not intend the Panel to rule on the likeness of friction products containing chrysotile.
and other products. We shall therefore not examine these products in our findings relating to the provisions of the GATT 1994.

8.109 Secondly, although Canada appears sometimes to consider that chrysotile fibre is simply an input and not a product per se, we note that the Decree refers to “varieties of asbestos fibres, … regardless of whether these substances have been incorporated into materials, products or devices”. We also note that the two parties discussed the likeness of asbestos fibres as such, irrespective of the products into which they were incorporated.

8.110 Concerning the EC’s argument that neither in the request for consultations nor in the request for the establishment of a panel nor in the Panel’s terms of reference is it specified that the dispute concerns high-density cement products containing chrysotile, the Panel wishes to recall that it is not necessary for the request for the establishment of a panel to list all the products concerned in detail. The Panel also considers that the fact that Canada did not export high-density cement products containing chrysotile cannot constitute an obstacle to claims relating to these products.

8.111 In the light of the foregoing, it would appear appropriate to structure our analysis as follows:

(a) First of all, to examine whether PVA, cellulose and glass fibres, taken separately (i.e. not incorporated in a product), are products like to chrysotile fibre. Even if they are not a finished product, the fibres themselves are products which are exported and marketed and we believe it is relevant to compare them.

(b) Secondly, with regard to products containing asbestos or substitute fibres, we note that Canada confines its comparison to products combining cement and chrysotile (“chrysotile-cement”) or one or the other of the three aforementioned fibres (“fibro-cement”). We shall therefore limit our findings to these categories.

(c) Lastly, we note that, although Canada makes a specific reference to and cites as an example certain products: pipes, tiles, insulating tiles or boards, it does not furnish an exhaustive list of the products actually affected by the measure and its claims are not arranged on a “product-by-product” basis. Canada refers in generic terms to two major categories of product: (a) chrysotile, PVA, cellulose and glass fibres; and (b) products combining chrysotile and cement or other fibres and cement. We also note that the EC does not distinguish between various fibro-cement or chrysotile-cement products. Even though the different products composed of chrysotile-cement or fibro-cement have different uses (water pipes, thermal insulation, roofing, etc.), we do not consider it necessary to identify each product manufactured in chrysotile-cement or fibro-cement because they are all composed of either fibro-cement or chrysotile-cement and the matters discussed by the parties concerning their likeness are essentially related to whether or not they contain chrysotile.

(d) Analysis of likeness

(i) Introductory remarks

8.112 We note that both Canada and the EC refer to the Report of the Working Party on Border Tax Adjustments, which, using the terms of the Appellate Body in Japan – Alcoholic Beverages, lays down the fundamental principle for interpreting the words “like products” in general in the various provisions of the GATT 1947. This Report states the following:

“ … the interpretation of the term [‘like products’] should be examined on a case-by-case basis. This would allow a fair assessment in each case of the differ-
ent elements that constitute a ‘similar’ product. Some criteria were suggested for determining, on a case-by-case basis, whether a product is ‘similar’: the product’s end-uses in a given market; consumers’ tastes and habits, which change from country to country; the product’s properties, nature and quality”.

8.113 The Panel and the Appellate Body in Japan – Alcoholic Beverages recognized the relevance of this list and added tariff classification in the Harmonized System (“HS”). The Appellate Body also pointed out that the principle elaborated by the Working Party in Border Tax Adjustments had been followed in almost all the reports of subsequent panels. We note in this connection that the panel in United States – Gasoline, in the context of the GATT 1994, applied this principle when examining likeness in relation to Article III:4.

8.114 Finally we note that in Japan – Alcoholic Beverages the Appellate Body reaffirmed that panels must use their best judgement when determining whether products are in fact like products, and this would always inevitably involve a degree of discretionary judgement. The Appellate Body also confirmed that, when making an assessment, no single approach would be appropriate to every single case. The circumstances peculiar to each case must be taken into account. The criteria outlined in the report on Border Tax Adjustments should be examined, but there could be no precise and absolute definition of like.

8.115 We would add that even though, for reasons of clarity, each criterion has to be examined separately, it is more than likely that they are largely interdependent. In other words, it does not appear appropriate to examine each of the criteria in isolation. Our examination should be based on an assessment of each criterion in its context, that is to say in the light of the other criteria deemed relevant in this case.

8.116 Having defined the approach to be used, we commence our analysis by examining the likeness of fibres.

(ii) Likeness of asbestos fibres and substitute fibres

8.117 We note that, in their arguments, the parties have not always distinguished between fibres as such and products containing such fibres. In this section, therefore, we shall examine the relevant criteria provided that they can be related specifically to the fibres.

Properties, nature and quality of the products

8.118 Canada considers that the nature of chrysotile and substitute fibres is the same because they are all fibres. Even if the length, diameter and width-diameter ratio have an effect on pathogenicity, this does not mean that fibres of different dimensions cannot be like fibres. The dimensional parameters set by the WHO do not constitute the criterion of the nature, quality and properties according to which the likeness of fibrous products is determined. Too much importance should not be attached to the special nature of asbestos fibres claimed by the EC. Even if substitute fibres are more costly than chrysotile fibres and have other uses, chrysotile-cement or fibro-cement manufacturers use them for the same purposes and the likeness of the manufacturing processes for chrysotile-cement and fibro-cement shows the similarities of the properties and the nature of these fibres. In order to offer the same technical guarantees as chrysotile-cement products (one of the conditions for substitutability in the Decree), fibro-cement products must indubitably have the same properties, quality and nature. Similarly, the “lower” pathogenicity of substitute fibres should not preclude the conclusion that they are like asbestos. Products may be considered like despite their differing impact on health. There is no contradiction between distinguishing two types of fibre in scientific terms and according to their pathogenicity, on the one hand, and, on the other, applying the criteria derived from WTO and GATT practice for determining whether products are like. The toxicity of a product is not recognized as a criterion for the evaluation of likeness.
8.119 The EC consider that the properties, nature and quality of products are important when assessing likeness within the meaning of Article III:4. Unlike other criteria, this criterion has always been used by panels in connection with Article III:4. In the light of this criterion, the products are in any case different. Asbestos fibres have a very particular fibrous texture (bundles of fibrils that can easily be separated lengthways and have a very small diameter). The physical and chemical characteristics of substitute fibres are not the same as those of asbestos fibres (for example, their diameter is much bigger and their fibrillation capacity is more limited). No single natural or synthetic substitute product is able to combine, or combines, all the properties of asbestos, bearing in mind the unique nature of the characteristics of asbestos fibres. These characteristics also make asbestos fibres particularly dangerous for health. Since 1977, the WHO has classified asbestos fibres in category 1 of proven carcinogens. The EC point out that, in contrast, none of the substitute products for chrysotile asbestos is classified as a proven carcinogen for humans. The nature, composition, physical properties and proven effects on human health of chrysotile make it radically different from substitute products. In such a situation, the health risk posed by the product must necessarily be taken into account. A dangerous product should be regarded as being different in nature and quality from a harmless or less dangerous product.

8.120 The Panel considers that, in addition to the factual elements, the parties’ arguments raise a first issue, namely, in what context should the criterion of the properties, nature and quality of the fibres be taken into account for the likeness test within the meaning of Article III:4? The parties in fact basically take up the question of the way in which the properties, nature and quality of the fibres should be taken into account by the Panel.

8.121 The Panel notes that no party contests that the structure of chrysotile fibres is unique by nature and in comparison with artificial fibres that can replace chrysotile asbestos. The parties agree that none of the substitute fibres mentioned by Canada in connection with Article III:4 has the same structure, either in terms of its form, its diameter, its length or its potential to release particles that possess certain characteristics. Moreover, they do not have the same chemical composition, which means that, in purely physical terms, none of them has the same nature or quality. It could therefore be concluded that they are not like products.

8.122 It should be recalled, nevertheless, that the context for the application of Article III:4 is not a scientific classification exercise. The objective of Article III concerns market access for products. Its purpose is to prevent internal measures from being applied in such a way as to protect domestic production. Article III:4 upholds this objective in respect of laws, regulations and requirements affecting the sale, marketing, purchase, transportation, distribution and use of products on the domestic market. We also note that the criterion includes the concept of the “quality” of a product, which is indicative of a commercial approach, otherwise the word “quality” would no doubt have been used in the plural, in which case it would have been the same as “properties” in the sense of a particular quality of a product. It is thus with a view to market access that the properties, nature and quality of imported and domestic products have to be evaluated.

8.123 Although we share Canada’s view that all the products are “fibres” and thus like products, we do not consider that the examination of the physical structure and chemical composition (which in our view relate to the nature of the product) should be taken to the other extreme, even though it has been argued that other panels followed a narrower approach in this respect. If such an approach was adopted, many products would never be like in respect of their nature, even if they had a similar use. On the other hand, products which bore no relation to each other in terms of their use in everyday life could be considered as like products because of their chemical composition. As mentioned in para. 8.114 above, we note the need to evaluate these criteria on a case-by-case basis, in other words, bearing in mind the factual circumstances. In this particular case, because of its physical and chemical characteristics, asbestos is a unique product. We note, nevertheless, that for many industrial uses other products have the same applications as asbestos. If the chemical and physical characteristics were to be recognized as decisive in this case, we would
have to disregard all the other criteria and this does not appear to us to be consistent with the flexibility given to panels by the Appellate Body when examining the principle of likeness.

8.124 As regards properties, we note that no substitute fibre alone combines all the properties and qualities of chrysotile fibre itself. Article 2, paragraph 1, second subparagraph of the Decree recognizes this by basing the criteria for substitution inter alia on “all technical guarantees of safety corresponding to the ultimate purpose of the use thereof”. A narrow interpretation of the concept of like product might perhaps lead us to exclude the likeness of products which do not always show the same properties in all circumstances. In the context of market access, it is not necessary for domestic products to possess all the properties of the imported product in order to be a like product. It suffices that, for a given utilization, the properties are the same to the extent that one product can replace the other. If the properties of products always had to be the same, the category of like products would be very small, sometimes even just one product. We note in this connection that the Panel in Japan – Customs Duties, Taxes and Labelling Practices on Imported Wines and Alcoholic Beverages considered that gin, vodka, whisky, brandy, liqueurs, still wine and sparkling wine should be considered as like products within the meaning of Article III:2 in view of their similar properties and end-uses we consider that this Report upholds our approach. If products whose nature is not exactly the same and whose properties are not always identical, for example wine on the one hand and whisky on the other, can be considered like products, the same approach can be followed pursuant to Article III:4 for chrysotile fibres on the one hand and PVA, cellulose and glass fibres on the other.

8.125 In this case, even if the end-uses of chrysotile fibres on the one hand and PVA, cellulose and glass fibres on the other are only the same for a small number of their respective applications, in some cases the applications are similar. Their properties are then equivalent, if not identical. This is the juncture of interest to us, the moment when the products are used for the same purpose. As we have already mentioned above, the criteria proposed for determining likeness should not be examined in isolation. In this particular case, we consider that the end-use of the products should affect the way in which we examine the properties of the fibres compared, inasmuch as none of the fibres mentioned by Canada always fulfills the same functions.

8.126 We therefore conclude that, taking into account the properties criterion, chrysotile fibres are like PVA, cellulose and glass fibres. With regard to nature and quality, we consider that these criteria should not be applied narrowly in the factual circumstances of the present case. Consequently, the fact that chrysotile fibres do not have the same structure or chemical composition as PVA, cellulose or glass fibres cannot be decisive for the evaluation of the likeness of these products.

8.127 The second question that must be answered in relation to the application of the properties, nature and quality criterion is that of the relevance of the risk of the product raised by the EC.

8.128 The Panel has noted the EC’s argument that the capacity of chrysotile fibres to break up into extremely fine particles that can penetrate the pulmonary alveoli gave these fibres a property which meant that they were not like because this property was the basis for chrysotile’s potential to cause diseases of the lung and the pleura, mainly lung cancers and mesotheliomas.

8.129 We note first of all that the risk of a product for human or animal health has never been used as a factor of comparison by panels entrusted with applying the concept of “likeness” within the meaning of Article III. In addition to the fact that no other panel has probably ever been called upon to examine a question similar to the one before us, in our view the reason is to be found in the economy of the GATT 1994. Its primordial role is to ensure that a certain number of disciplines are applied to domestic trade regulations. Article XX of the GATT, however, recognizes that certain interests may take precedence over the rules governing international trade and authorizes the adoption of trade measures aimed at preserving these interests while at the same time observing certain criteria.
8.130 We consider that introducing a criterion on the risk of a product into the analysis of likeness within the meaning of Article III would largely nullify the effect of Article XX(b). The protection of human health and life is specifically covered by this Article. Article III, on the other hand, does not refer to this. The burden of proof would not of course be greatly modified because the EC would still have to prove the risk of the product, applying the principle of probatio incumebit ejus qui dixit. We nevertheless consider that other aspects that form part of the rights and obligations negotiated by the Members would be affected. Introducing the protection of human health and life into the likeness criteria would allow the Member concerned to avoid the obligations in Article XX, particularly the test of necessity for the measure under paragraph (b) and the control exerted by the introductory clause to Article XX concerning any abuse of Article XX(b) when applying the measure. As the Appellate Body has emphasized on a number of occasions, all these provisions in the WTO Agreement must be given meaning. Introducing a risk criterion into the examination of likeness under Article III would be contrary to this basic principle of interpretation.

8.131 Finally, if such a criterion was applied, it would make all the other criteria mentioned by the Working Party on Border Tax Adjustments totally redundant because it would become decisive when assessing the likeness of products in every case in which it was invoked, irrespective of the other criteria applied.

8.132 We therefore conclude that, bearing in mind the overall economy of the WTO Agreement, in particular the relationship between Article III and Article XX(b), it is not appropriate to apply the “risk” criterion proposed by the EC, neither in the criterion relating to the properties, nature and quality of the product, nor in the other likeness criteria invoked by the parties.

End-use

8.133 Canada considers that, given the nature of chrysotile fibre (a raw mineral resource), special importance must be attached to the criterion of the product’s end-use. Chrysotile fibre has no use in its raw form. After incorporation into the cement, chrysotile fibre and PVA, cellulose and glass fibres are used for the manufacture of chrysotile-cement and fibro-cement products respectively. At the end-use stage, these products constitute one single product to be used for the same purpose. According to Canada, applying the end-use criterion suffices to conclude that chrysotile fibres and the aforementioned substitute fibres are like products. If end-use is a key criterion for determining whether two products are directly competitive or substitutable pursuant to Article III:2, it is equally important for likeness. It is the market which determines the end-use of a product.

8.134 According to the EC, the end-uses of these fibres are different and this criterion is not decisive when determining “likeness” within the meaning of Article III:4, which is essentially “technical”. Even where products may have some end-uses in common, these uses are not sufficient to classify the products as like products when each of them also has many other end-uses. The criterion of end-use could not, by itself, invalidate the conclusion based on other criteria that these products are not like products within the meaning of Article III:4 of the GATT 1994.

8.135 The Panel notes that Canada limited its arguments to the use of chrysotile in cement products and, at the same time, to the use of PVA, cellulose and glass fibres in cement products.

8.136 We have already found above that the respective properties of chrysotile fibres on the one hand and PVA, cellulose or glass fibres on the other allowed certain identical or at least similar end-uses. We do not therefore deem it necessary to elaborate this further, except to recall that, in our view, the fact that all the end-uses of these fibres are not like uses does not mean that the products are not like products.
Consumers’ tastes and habits

8.137 As regards consumers’ tastes and habits, Canada is of the view that manufacturers of chrysotile-cement or fibro-cement products are the consumers. The drop in chrysotile asbestos imports in 1996 and 1997 is a result of the Decree and not of a change in consumers’ tastes and habits. It is thus inappropriate to consider this criterion. Canada agrees with the practice of panels cited by the EC, and asks the Panel not to take into account consumers’ tastes and habits when determining the likeness of products in this case.

8.138 In the view of the EC, the consumers’ tastes and habits criterion is not a priori relevant because the products concerned are not everyday consumer goods. It might nevertheless be interesting to analyse the consumers’ perception of these products. Informed users will not choose asbestos after the competent international organizations have decided that it is a proven carcinogen.

8.139 We note first of all that the parties do not consider it useful or necessary for us to take this criterion into account. In our view, this is not in itself sufficient reason for us not to take it into account. We do not agree either, that, in view of Article III:4, there are reasons for excluding consideration of this criterion a priori. We consider that it is up to the Panel to decide whether one of the criteria applicable to determining the “likeness” of the products concerned is relevant or not. What is important is to ensure that our analysis takes into account all the relevant elements. In this particular case, we note first of all that, when determining the tastes and habits of consumers, it is necessary to place oneself at the time prior to the entry into force of the ban in the Decree. Even if we do place ourselves prior to that date, however, it would be difficult to determine precisely what were the tastes and habits of consumers at that time. The groups of consumers to be taken into account are very varied and their tastes and habits based on an equally wide variety of considerations. Because this criterion would not provide clear results, the Panel considers that it is not relevant to take it into account in the special circumstances of this case.

8.140 We shall therefore refrain from taking a position on the impact of this criterion on the likeness of the products considered.

Tariff classification

8.141 As regards tariff classification, Canada recalls that the 107 six or eight-digit codes for chrysotile-cement products in the Harmonized Commodity Description and Coding System of the Customs Cooperation Council are identical to the 107 codes for fibro-cement products.

8.142 The EC consider that the tariff classifications are different, whether for asbestos fibres and substitute fibres or for products containing asbestos and substitutes for asbestos. For example, in the Harmonized System, asbestos fibres are classified in their own heading.

8.143 We do not consider that the fact that asbestos fibres are classified in their own heading is decisive in this case. PVA, cellulose and glass fibres respectively are also classified in different tariff headings. We note, however, that such a classification reflects the difference in their nature, whether they are mineral or vegetable, artificial or natural. We have already found that this factor did not affect the fact that their properties and end-use are the same under certain circumstances.

Conclusion

8.144 Above we concluded that chrysotile fibres, on the one hand, and PVA, cellulose and glass fibres, on the other, are, in certain circumstances, similar in properties, nature and quality. We also concluded that these products have similar end-uses. From this it follows that chrysotile fibres, on the one hand, and PVA, cellulose and glass fibres, on the other, are like products within the meaning of Article III:4 of the GATT 1994.
(iii) Likeness between products containing asbestos and certain other products

8.145 The Panel considers that many of the arguments put forward in relation to asbestos, PVA, cellulose and glass fibres are applicable *mutatis mutandis* to products containing those fibres. Thus, if a fibro-cement product, for example, a tile, is compared with a similar tile of chrysotile-cement, the only difference between the products concerned is the presence of either chrysotile or a substitute fibre in the tile, the product itself being a tile and the other component of the material being in both cases cement. It is the presence of chrysotile or some other fibre that gives the cement product its specific function: mechanical strength, resistance to heat, compression, etc.

8.146 Moreover, we note that, in Canada’s opinion, chrysotile-cement and fibro-cement products are both industrial products which cannot be distinguished on the basis of their external appearance. From the standpoint of the tastes and habits of the French consumer, they are interchangeable. According to the EC, informed users would not use products containing asbestos after the international organizations had decided that asbestos was a proven carcinogen.

8.147 Consequently, we consider that in fact the likeness between a chrysotile-cement product and a fibro-cement product depends on two factors: (a) the nature of the product itself and (b) the presence of chrysotile fibres or of PVA, cellulose or glass fibres in the product.

8.148 Using the criteria adopted by the Panel and the Appellate Body in *Japan – Alcoholic Beverages*, we note, first of all, that the HS tariff classification (heading 68.11) is the same for articles of asbestos-cement, of cellulose fibro-cement or the like. This heading covers hardened articles consisting essentially of an intimate mixture of fibres (for example, asbestos, cellulose or other vegetable fibres, synthetic polymer or glass fibres) and cement or other hydraulic binders, the fibres acting as strengthening agents. Other products may fall into different tariff headings. However, many of the products mentioned by Canada appear to fall within this subdivision.115

8.149 We consider, moreover, that one of the EC’s main arguments against the likeness between fibro-cement and chrysotile-cement products with respect to their properties, nature and quality and their end-use is based on the risk associated with the chrysotile which they contain. To the extent that in paragraph 8.132 we concluded that the “risk” criterion cannot be included among the criteria applicable to the determination of likeness under Article III:4, we do not consider it necessary to discuss the applicability of this criterion with respect to chrysotile-cement products. The same applies with respect to consumers’ tastes and habits.

8.150 We therefore conclude that chrysotile-fibre products and fibro-cement products are like products within the meaning of Article III:4 of the GATT 1994.

(e) Less favourable treatment of Canadian products116

8.151 With respect to the existence of less favourable treatment, Canada argues that the Decree alters the conditions of competition between, on the one hand, substitute fibres and products containing them of French origin and, on the other hand, chrysotile fibre and products containing it from Canada. The Decree does not afford chrysotile fibre imported from Canada and products containing it effective equality of opportunities for imported products in respect of the application of laws, regulations and requirements affecting the internal sale, offering for sale, purchase, transportation, distribution or use of products. Not only does the ban apply only to asbestos fibre and products containing it but it is applicable only if there are products like to chrysotile fibre or to the products containing it. Thus, the Decree imposes less favourable treatment in all cases where like products exist. Accordingly, the Decree constitutes *de jure* discrimination between chrysotile fibres and
products containing them, on the one hand, and like products (PVA, cellulose or glass fibres and fibro-cement products containing them), on the other.

8.152 Canada also alleges de facto discrimination. The French PVA fibres industry is in better shape than ever. Moreover, it is not because France has imported a marginal additional quantity of Canadian cellulose fibres that the French domestic industry has not benefited from the ban. The Decree does indeed impose a choice on the French consumer who is now prevented from using chrysotile fibre or products containing it.

8.153 The EC maintain that the contested measure accords with the fundamental purpose of Article III, which is to prevent protectionism, and is not discriminatory, neither de jure nor de facto, inasmuch as it guarantees effective equality of opportunities for domestic and imported products. The context in which the Decree was adopted and its provisions show that the intention of the French authorities was in no way to protect domestic substitute products but to protect human health against the risks associated with asbestos. They also show that the Decree makes no distinction between imported and domestic products, whether it be a question of substitute or asbestos products, and that neither its object nor its effect is to protect domestic production. The Decree does not create any de facto discrimination, since in France most substitute products are imported from various third countries. Moreover, France has a negative trade balance in substitute products. The ban on the use of asbestos on public health grounds has required a painful changeover, in human and financial terms, including the loss of external outlets for French industry. Finally, the Decree is “neutral” in respect of the choices that businesses can make concerning replacement products.

8.154 We note that with regard to the establishment of the existence of less favourable treatment, it is first necessary to determine, as we have done, whether there is a likeness between the imported and the domestic products. Above, both with regard to chrysotile fibres, on the one hand, and PVA, cellulose and glass fibres, on the other117, and with regard to products made of chrysotile-cement, on the one hand, and fibro-cement, on the other, we concluded that they were “like” within the meaning of Article III:4. With respect to the treatment of these products as compared with the like domestic products, we note, first of all, that France does produce substitutes for chrysotile fibres and chrysotile-cement products. We next note that the terms of the Decree in themselves establish less favourable treatment for asbestos and products containing asbestos as compared with substitute fibres and products containing substitute fibres. Thus, paragraphs I and II of Article 1 of the Decree read as follows:

“I. For the purpose of protecting workers, and pursuant to Article L. 231-7 of the Labour Code, the manufacture, processing, sale, import, placing on the domestic market and transfer under any title whatsoever of all varieties of asbestos fibres shall be prohibited, regardless of whether these substances have been incorporated into materials, products or devices.

II. For the purpose of protecting consumers, and pursuant to Article L. 221.3 of the Consumer Code, the manufacture, import, domestic marketing, exportation, possession for sale, offer, sale and transfer under any title whatsoever of all varieties of asbestos fibres or any product containing asbestos fibres shall be prohibited.”

8.155 Inasmuch as the Decree does not place an identical ban on PVA, cellulose or glass fibre and fibro-cement products containing PVA, cellulose or glass fibres, we must conclude that de jure it treats imported chrysotile fibres and chrysotile-cement products less favourably than domestic PVA, cellulose or glass fibre and fibro-cement products.

8.156 Having established de jure discrimination on the basis of the Decree and, moreover, the European Communities not having submitted any evidence that might lead us to believe that the Decree is applied in such a way as not to introduce less favourable treat-
ment for chrysotile fibres and chrysotile-cement products as compared with PVA, cellulose and glass fibres and fibro-cement products containing PVA, cellulose or glass fibres, we do not consider it necessary to determine whether there is any de facto discrimination between these products.

8.157 For these reasons, we conclude that the Decree applies to chrysotile and chrysotile-cement products a treatment less favourable than that which it applies to PVA, cellulose and glass fibres and products containing them, within the meaning of Article III:4.

(f) Conclusion

8.158 On the basis of the above, we find that the provisions of the Decree relating to the prohibiting of the marketing of chrysotile fibres and chrysotile-cement products violate Article III:4 of the GATT 1994.

3. Violation of Article XI of the GATT 1994

8.159 In the light of our observations on the applicability of Article III:4 of the GATT 1994 to the border measures imposed on chrysotile fibres and chrysotile-cement products and considering our findings with regard to the violation of Article III:4 by the Decree, we conclude that it is not necessary to examine the Canadian argument concerning the violation of Article XI:1 of the GATT 1994.

4. Applicability of Article XX of the GATT 1994

(a) Arguments of the parties

8.160 The European Communities contend that the Decree falls under the exception provided in Article XX of the GATT 1994. Thus, the Decree is necessary to achieve the French Government’s public health goals under Article XX(b) and is not applied in such a manner as to constitute a means of arbitrary or unjustifiable discrimination between countries where the same conditions prevail or a disguised restriction on international trade, within the meaning of the introductory clause of Article XX.

8.161 Canada notes that, in accordance with panel and Appellate Body practice, Article XX permits a limited and conditional exception from the obligations set out in the other Articles of the GATT and must be interpreted narrowly. Although the European Communities assert that they have the right to establish the level of protection they desire, in so doing they must comply with their obligations. In this connection, the Appellate Body has condemned abuse of rights under Article XX. Moreover, it is up to the European Communities to demonstrate that the Decree falls under Article XX(b).

8.162 The European Communities argue that asbestos fibres and products containing them are a proven hazard for human health. The risks linked to the use of these fibres are recognized both by scientists and international organizations. By prohibiting the marketing and use of asbestos and asbestos-containing products, the Decree seeks to halt the spread of these risks, in particular for people occasionally and often unwittingly exposed to asbestos, and thereby reduce the number of deaths among the French population. It is the only measure capable of preventing the spread of the risks due to asbestos exposure. According to the EC, the review in the light of Article XX cannot be allowed to undermine the health protection goal set by the Member concerned. Its sole purpose must be to assess whether the trade measure adopted is necessary to attain that goal. This test concerns the trade measure and not the level of protection set by the Member.

8.163 Canada considers that the only exposures that could be affected by the Decree are exposures to chrysotile encapsulated in high-density products. In order to determine whether there is an equally effective alternative that is less restrictive for international trade and that is capable of protecting human life or health just as effectively, the health risk can and must be examined. Otherwise any country could cite the risk – real or not – in support of a prohibition measure.
8.164 Canada is of the opinion that the current uses of chrysotile do not constitute a detectable risk to human health. According to Canada, the European Communities do not explain that the exposures of secondary users are essentially to friable materials, very often containing amphiboles, which are no longer marketed and have been in place for a long time. Contrary to what the EC appear to believe, controlled use substantially reduces exposure. As a result of pre-fabrication, pre-machining, the use of fittings and compliance with work standards, workers are not subjected to high levels of exposure, as the EC claim.

8.165 Canada considers that, inasmuch as the only risk associated with asbestos is that of the past use of amphiboles and the use of friable materials, the Decree, which prohibits the contemporary uses of chrysotile, is not “necessary” to protect human life or health from the risks associated with past uses of asbestos. A ban is not “necessary” because high-density chrysotile products do not pose any detectable risk. On the other hand, controlled use indisputably constitutes an alternative to a total ban that is significantly less restrictive for international trade.

(b) Approach adopted by the Panel and burden of proof

(i) Introductory remarks concerning the approach adopted by the Panel

8.166 At this stage, the Panel wishes to point out that, inasmuch as Article XX provides for a certain number of exceptions to the rules of the GATT 1994, it is necessary and sufficient to examine the applicability of these exceptions solely with regard to measures and in relation to products for which a violation of another provision of the GATT has been identified. In the present case we have found that the Decree violates Article III:4 of the GATT 1994 as regards the treatment accorded to imported chrysotile asbestos fibres as compared with PVA, cellulose and glass fibres. We have also found a violation of Article III:4 with respect to the treatment of chrysotile-cement products as compared with fibrocement products containing PVA, cellulose or glass fibres.

8.167 In accordance with the approach noted by the Panel in United States – Gasoline and the Appellate Body in United States – Import Prohibition of Certain Shrimp and Shrimp Products, we will first examine whether the measure falls within the scope of paragraph (b) of Article XX, the provision expressly invoked by the European Communities. If we decide that it does, we will consider whether, in its application, the Decree satisfies the conditions of the introductory clause of Article XX.

8.168 First of all, as regards Article XX(b), we find that the provisions of this paragraph relevant to the present case require that, subject to fulfilment of the conditions of the introductory clause of Article XX, nothing in the GATT 1994 shall be construed to prevent the adoption or enforcement by any Member of measures

"(b) necessary to protect human ... life or health"

8.169 The Panel notes that in United States – Gasoline, the Panel stipulated that with respect to Article XX(b), the party invoking that provision must prove:

(a) That the policy in respect of the measures for which Article XX is invoked falls within the range of policies designed to protect human life or health; and

(b) the inconsistent measures for which the exception is invoked are necessary to fulfil the policy objective.

8.170 As regards subparagraph (a), we consider that, inasmuch as they include the notion of “protection”, the words “policies designed to protect human life or health” imply the existence of a health risk. We must therefore determine, on the basis of the relevant rules of evidence, whether chrysotile-asbestos, in the various forms we have considered so far, poses a risk to human life or health.
8.171 This said, we note that the panel in United States – Gasoline also made clear that it did not have to examine the necessity of the policy goal. In other words, we do not have to assess the choice made by France to protect its population against certain risks, nor the level of protection of public health that France wishes to achieve. We must simply determine if the French policy of prohibiting the use of chrysotile-asbestos falls within the range of policies designed to protect human life or health.

8.172 As regards the criterion of the necessity of the measure (subparagraph (b) of paragraph 8.169), we note that previous panels that had to assess the “necessity” of a measure under Article XX(b) appear to have done so solely in relation to the existence of other measures consistent or less inconsistent with the GATT in the light of the health objective pursued. Thus, in Thailand – Cigarettes, the Panel ruled that:

“The import restrictions imposed by Thailand could be considered to be “necessary” in terms of Article XX(b) only if there were no alternative measure consistent with the General Agreement, or less inconsistent with it, which Thailand could reasonably be expected to employ to achieve its health policy objectives”.

8.173 In this particular case, the public health objectives pursued by Thailand were confirmed by a WHO expert and accepted by the parties to the dispute. In the present case, Canada does not deny that France’s goal, namely to protect the health of workers and consumers, is a public health objective, but it disputes the existence of a public health problem in relation to chrysotile. Thus, Canada denies that chrysotile fibre poses a public health risk in its applications because once in place chrysotile-cement does not release any fibres and because it is possible to work with chrysotile-cement products in a safe or controlled fashion.

8.174 For the Panel, the assessment of the necessity of a measure could be focused on the existence of other measures consistent or less inconsistent with the GATT 1994 if, as in previous cases, the parties were agreed on the existence and extent of the health problem associated with chrysotile. In this respect, we note that Canada does not dispute that chrysotile fibres in themselves pose health risks. However, Canada does not consider that any risk is posed by chrysotile enclosed in a matrix of high-density cement.

8.175 Consequently, inasmuch as the parties disagree on the extent of the health risk posed by chrysotile-cement products, we consider that our examination of the existence of a general health objective in the light of subparagraph (a) of paragraph 8.169 should include the question of the existence of a health problem in relation (i) to chrysotile fibres as such and (ii) to chrysotile-cement products.

8.176 This being so, we shall have to take expressly into account the extent of the health problem in assessing the necessity of the measure. Thus, if we were to conclude that the health hazard represented by chrysotile or chrysotile-cement was less than the EC allege, less vigorous measures might then be justified.

(ii) Burden of proof

General considerations

8.177 With regard to the burden of proof, we refer, first of all, to our previous remarks concerning those cases in which a party invokes a defence. We consider that the reasoning of the Appellate Body in United States – Shirts and Blouses from India is applicable to Article XX, inasmuch as the invocation of that Article constitutes a “defence” in the sense in which that word is used in the above-mentioned report. It is therefore for the European Communities to submit in respect of this defence a prima facie case showing that the measure is justified. Of course, as the Appellate Body pointed out in United States – Gasoline, the burden on the European Communities could vary according to what has to be proved. It will then be for Canada to rebut that prima facie case, if established.
8.178 If we mention this working rule at this stage, it is because it could play a part in our assessment of the evidence submitted by the parties. Thus, the fact that a party invokes Article XX does not mean that it does not need to supply the evidence necessary to support its allegation. Similarly, it does not release the complaining party from having to supply sufficient arguments and evidence in response to the claims of the defending party. Moreover, we are of the opinion that it is not for the party invoking Article XX to prove that the arguments put forward in rebuttal by the complaining party are incorrect until the latter has backed them up with sufficient evidence.

Considerations specific to the burden of proof as regards the scientific aspects

8.179 As pointed out above, three essential elements must be considered by the Panel when examining the justification of a measure in the light of Article XX(b): (a) the existence of a risk for human health; (b) the level of protection which the Member concerned wishes to achieve; and (c) the existence of other measures consistent or less inconsistent with the GATT 1994 and enabling the same objective of protecting public health to be obtained. The Panel considers that its examination of the scientific data should be exclusively concerned with points (a) and (c), inasmuch as it has long been established that Members are free to set the level of protection of their choice for their populations.

8.180 The Panel has therefore had to determine how it should assess the existence of a health risk in relation to chrysotile and, more particularly, chrysotile-cement and the necessity of the measures in question. The Panel has examined the practice in relation to Article XX of the GATT 1994, but also in the context of other WTO Agreements in which scientific studies are invoked, namely the Agreement on Sanitary and Phytosanitary Measures. The Panel noted that the SPS Agreement contains more detailed provisions than Article XX with respect to the scientific justification of a sanitary or phytosanitary measure and that these provisions have been the subject of clarifications by panels and by the Appellate Body. However, it also noted that in the first dispute settlement proceedings initiated under the WTO Agreement concerning Article XX of the GATT 1994, the Appellate Body had not sought to extend the principles of the SPS Agreement to the examination of the measures for which Article XX(b) had been invoked or even to base itself on them, although the SPS Agreement was already in force. The Panel preferred to confine itself to the provisions of the GATT 1994 and to the criteria defined by the practice relating to the application of Article XX.

8.181 In this context, in relation to the scientific information submitted by the parties and the experts, the Panel feels bound to point out that it is not its function to settle a scientific debate, not being composed of experts in the field of the possible human health risks posed by asbestos. Consequently, the Panel does not intend to set itself up as an arbiter of the opinions expressed by the scientific community.

8.182 Its role, taking into account the burden of proof, is to determine whether there is sufficient scientific evidence to conclude that there exists a risk for human life or health and that the measures taken by France are necessary in relation to the objectives pursued. The Panel therefore considers that it should base its conclusions with respect to the existence of a public health risk on the scientific evidence put forward by the parties and the comments of the experts consulted within the context of the present case. The opinions expressed by the experts we have consulted will help us to understand and evaluate the evidence submitted and the arguments advanced by the parties. The same approach will be adopted with respect to the necessity of the measure concerned.

8.183 In proceeding with this exercise, the Panel will have to make a pragmatic assessment of the scientific situation and the measures available, as would the decision-makers responsible for the adoption of a health policy. In this connection, it notes that the determination of the existence of other measures consistent or less inconsistent with the GATT largely depends on a scientific assessment of the risk. In any event, this determination cannot be interpreted as restricting the freedom of Members to take certain measures rather than others under Article XX(b), in the absence of a measure that would be consistent or less inconsistent with the GATT 1994.
In accordance with the approach defined by the Panel in *United States – Gasoline*, we must first establish whether the policy in respect of the measure for which the provisions of Article XX(b) were invoked falls within the range of policies designed to protect human life or health. As we have already pointed out, the use of the word “protection” implies the existence of a risk. Accordingly, we must begin by identifying a risk for public health. In the light of the comments of the panel in *United States – Gasoline* and our own remarks in paragraph 8.182, we must also take into account the fact that it is a public health policy that we have to assess.

First of all, we note that the EC argue that in prohibiting the placing on the market and use of asbestos and products containing it, the Decree seeks to halt the spread of the risks due to asbestos, particularly for those exposed occasionally and very often unwittingly to asbestos when working on asbestos-containing products. France considers that it can thereby reduce the number of deaths due to exposure to asbestos fibres among the French population, whether by asbestosis, lung cancer or mesothelioma.140

In principle, a policy that seeks to reduce exposure to a risk should fall within the range of policies designed to protect human life or health, insofar as a risk exists. According to the EC, the international scientific community appears to be generally of the opinion that chrysotile fibres as such are carcinogens. In this connection, we note the EC’s argument that, since 1977, the International Agency for Research on Cancer (IARC) has classified chrysotile among the proven carcinogens.

Canada does not dispute that chrysotile asbestos causes lung cancer. However, Canada argues that the mechanism that could give rise to an increased risk of lung cancer has not yet been fully explained and that the link with chrysotile might only be indirect.141 This risk depends on the intensity and duration of the exposure. On the other hand, according to Canada, there is a great deal of scientific evidence to support the thesis according to which chrysotile does not cause mesotheliomas.142 In particular, the mesotheliomas linked to asbestos could be the result of exposure to low-density products containing amphiboles. It has not been established that, in their uses, chrysotile fibres pose the same risk as amphiboles, whose chemical composition, in particular, is different.

First of all, we note that the carcinogenicity of chrysotile fibres has been acknowledged for some time by international bodies.143 This carcinogenicity was confirmed by the experts consulted by the Panel, with respect to both lung cancers and mesotheliomas.144 Even though the experts appear to acknowledge that chrysotile is less likely to cause mesotheliomas than amphiboles.145 We also note that the experts confirmed that the types of cancer concerned had a mortality rate of close to 100 per cent.146 We therefore consider that we have sufficient evidence that there is in fact a serious carcinogenic risk associated with the inhalation of chrysotile fibres. Moreover, in the light of the comments made by one of the experts147, the doubts expressed by Canada with respect to the direct effects of chrysotile on mesotheliomas and lung cancers are not sufficient to conclude that an official responsible for public health policy would find that there was not enough evidence of the existence of a public health risk.

We note, however, that Canada makes a distinction between chrysotile fibres and chrysotile encapsulated in a cement matrix. In fact, Canada challenges the Decree insofar as it prohibits, *inter alia*, the use of chrysotile-cement products. In this connection, we note that the experts consulted by the Panel agreed that the risks of fibres being dispersed due to the degradation of chrysotile-cement were limited. However, the experts acknowledged that working with non-friable products containing chrysotile might result in the dispersion of large quantities of fibres and that those fibres pose a definite health risk.148 The experts also noted that even though the risk might be lower than for production or processing workers, it concerned a much larger group.149
8.190 In this respect, the Panel notes that the European Communities have stated that
the Decree is intended, in particular, to protect categories of workers or consumers down-
stream of the asbestos mining or processing stage, whatever the frequency and level of
their exposure. Canada considers that below a certain exposure threshold there is no de-
tectable health risk. Accordingly, Canada believes that people only occasionally exposed
are not running a detectable risk.

8.191 The data submitted to the Panel by the EC show that the use of tools not specifi-
cally designed to prevent the release of fibres, which cannot be excluded, especially in
the case of DIY enthusiasts or professionals who work only occasionally in an environ-
ment where asbestos is present, can result in an exposure in excess of the statutory limits
under ISO 7337, which are themselves higher than those of the WHO (0.2 fibre/ml) or
those applied by France before the ban (0.1 fibre/ml). The Panel also notes the position
of the experts consulted on this point. All agree that building workers now count among
those most exposed to chrysotile fibres and hence to the risk of mesothelioma, but they
also mention cases of mesothelioma in patients who had been only incidentally exposed,
without any relation to their occupational activity. The scientists consulted by the Panel
also considered that the existence of a threshold below which exposure does not present
any risks had not been established for any of the diseases attributable to chrysotile, except
perhaps for asbestosis.

8.192 The Panel took note of Canada’s argument according to which there has been no
study specifically concerned with the occupational sectors to which the EC refer. It also
notes that Canada disputes the relevance of the data of the studies of Charleston textile
factory workers (United States) as compared with the Canadian studies of Quebec as-
bestos sector miners and workers, which are said to show the limited impact of chrysotile
on public health. However, the scientific experts consulted stressed the relevance and
quality of the Charleston study. On the other hand, doubts were expressed with regard to
the reliability of certain exposure data in the studies carried out in occupational and non-
occupational environments in Quebec and invoked by Canada. Canada also refers to a
study concerned with car brake maintenance. We note that the scientists consulted drew
attention to the limits of this study and produced statistical data which, on the contrary,
confirmed the impact of chrysotile on mechanics exposed to that material in a car brake
maintenance context.

8.193 The Panel therefore considers that the evidence before it tends to show that han-
dling chrysotile-cement products constitutes a risk to health rather than the opposite. Ac-
cordingly, a decision-maker responsible for taking public health measures might reason-
ably conclude that the presence of chrysotile-cement products posed a risk because of the
risks involved in working with these products.

8.194 Accordingly, the Panel concludes that the EC has made a prima facie case for the
existence of a health risk in connection with the use of chrysotile, in particular as regards
lung cancer and mesothelioma in the occupational sectors downstream of production and
processing and for the public in general in relation to chrysotile-cement products. This
prima facie case has not been rebutted by Canada. Moreover, the Panel considers that the
comments by the experts confirm the health risk associated with exposure to chrysotile in
its various uses. The Panel therefore considers that the EC have shown that the policy of
prohibiting chrysotile asbestos implemented by the Decree falls within the range of poli-
cies designed to protect human life or health. On the other hand, Canada has not suc-
cceeded in rebutting the presumption established on the basis of the evidence submitted by
the EC and confirmed by the experts. The Panel concludes therefore that the French policy
of prohibiting chrysotile asbestos falls within the range of policies designed to protect hu-
man life or health, within the meaning of Article XX(b) of the GATT 1994.

8.195 Accordingly, the Panel will now turn to the question of whether the measure is
"necessary" within the meaning of Article XX(b).
“Necessary”

The ban on chrysotile asbestos in its various forms

8.196 According to the European Communities, the danger of inhaling asbestos at levels above 0.1 fibre/ml concerns not only the asbestos mining and processing sectors but, more especially, secondary (textile, building and automobile industry, for example), para-occupational (servicing, maintenance) and domestic (DIY) workers whom Canada mentions only in part or not at all. Even in production and processing, which in principle are easier to monitor, there are limits to the controlled or safe use of asbestos, which does not halt the spread of the risks. So-called “controlled” or “safe” use is a fortiori completely ineffective in cases of occasional exposure to asbestos. The 1997 INSEERM report indicates that the risk occurs mainly among those who work with materials containing asbestos. The encapsulation of asbestos in a matrix cannot be guaranteed to make asbestos-cement products harmless, inasmuch as any subsequent working of the product will release large numbers of carcinogenic fibres in the form of dust. Controlled use is impossible to implement where hundreds of thousands of people are involved in sectors as unregulated in terms of health as the building industry. The 1984 ISO standard is inadequate in relation to the French health objective. The EC also note that, once asbestos is on the market, there is no reasonable way of controlling its use and, in particular, of controlling the everyday operations that many people are likely to perform. Moreover, the numerous, particularly legal obstacles which confront the victims of current exposure seeking redress in the courts constitute an additional social justification for resorting to a total ban.

8.197 With regard to the test of necessity, Canada takes the same approach as in connection with Article 2.2 TBT, considering that in many respects the test of necessity is similar in the two provisions. According to Canada, the measure must not be an excessive or over-reaching means to achieve a legitimate end. The two factors to be considered are, on the one hand, the risks that the absence of a technical regulation would create and, on the other, the existence of a less trade-restrictive alternative measure that would make it possible to fulfill the stated objective. As far as the second test is concerned, Canada notes that the controlled use of asbestos fibres allows fulfillment of the French objective of protecting human health while authorizing certain safe or controlled uses of chrysotile and products containing it. As controlled use is a less trade-restrictive alternative based on scientific data and having international support, a total ban on asbestos is not necessary. Canada is of the opinion that, today, high-density non-friable products do not pose a detectable risk. The risks which existed in the past and in certain cases still exist today are associated with past uses, very often of amphiboles in friable materials. In Canada’s view, the EC are trying to mislead the Panel by invoking the risks of the asbestos mining and processing industry, even though they have already recognized that controlled use is effective here in eliminating the risk. The EC also show bad faith in citing the risks for building maintenance workers and mechanics. The EC do not explain that these exposures are essentially to friable materials very often containing amphiboles with a high pathogenic potential. Controlled use is effective and can be accepted by professionals. According to the data of the United States Occupational Safety & Health Administration (OSHA), the institution of control measures lowers the average exposures of workers handling asbestos-cement pipes to 0.00253 f/ml and those of workers handling asbestos-cement sheets to 0.00727 f/ml. The average exposure of mechanics handling friction products is 0.00294 f/ml. Moreover, the use of pre-machined parts and fittings has been a big success. It reduces field activities and, at the same time, exposure rates.

8.198 We note that in Thailand—Cigarettes the Panel defined the test of necessity applicable under Article XX(b):

“The import restrictions imposed by Thailand could be considered to be ‘necessary’ in terms of Article XX(b) only if there were no alternative measure consistent with the General Agreement, or less inconsistent with it, which Thailand could reasonably be expected to employ to achieve its health policy objectives.”
This test has been applied in other disputes\textsuperscript{199} in order to apply the test defined in Thailand – Cigarettes, we must (a) establish the scope of the health policy objectives pursued by France and (b) consider the existence of measures consistent, or less inconsistent, with the GATT 1994.

First of all, we note that the risk due to chrysotile is important to the extent that, as confirmed in the previous section, it can generate lung cancers and mesotheliomas which are still difficult to cure or even incurable.\textsuperscript{167} The populations potentially at risk in France are very numerous, since products containing chrysotile, in particular, chrysotile-cement, have many applications in industrial, commercial and residential buildings. The fields of activity concerned include building workers (several hundred thousand) and DIY enthusiasts\textsuperscript{104}. These are areas in which health controls are difficult to apply, as the comments of the experts have shown.\textsuperscript{198}

The experts also confirmed that the intensive use of asbestos (in France, mainly chrysotile) over several decades has resulted in the risks of exposure being displaced from the mining and processing industry towards other sectors further downstream and, indeed, the general public.\textsuperscript{170} In this context, the Panel finds that the European Communities have shown that a risk exists for a very broad sector of the French population.

The Panel notes that the exposure of these groups is generally lower. However, the experts confirm the position of the European Communities according to which it has not been possible to identify any threshold below which exposure to chrysotile would have no effect.\textsuperscript{171} The experts are also agreed that the linear relationship model, which does not identify any minimum exposure threshold, is appropriate for assessing the existence of a risk.\textsuperscript{172} We find therefore that no minimum threshold of level of exposure or duration of exposure has been identified with regard to the risk of pathologies associated with chrysotile, except for asbestosis. Consequently, the possibility remains that low exposure over a fairly long period of time could lead to lung cancer or mesothelioma. Similarly, high-level exposure over a short period could also result in lung cancer or mesothelioma. These two possibilities were confirmed by the experts.\textsuperscript{173} The Panel therefore concludes that even though some trades or the French population in general are only intermittently exposed to low levels of asbestos, a decision-maker responsible for public health policy might reasonably conclude that there was nevertheless a real risk for these categories.

In the light of the above, the Panel concludes that, in addition to the risk presented by low-density friable products, there is an undeniable public health risk in relation to the chrysotile contained in high-density chrysotile-cement products. This risk exists even at low or intermittent exposure levels and can affect a broad section of the population.

We also note that France’s objective is to halt the spread of this risk which, considering the risk identified and its extent, could in principle justify strict measures. However, it is necessary to consider whether there is not, as Canada alleges, a measure that would be consistent, or less inconsistent, with the GATT 1994 and would allow the objective pursued by France to be achieved. Canada refers to the possibility of controlled use which consists in taking precautionary measures to restrict the release of fibres (use of special tools, high-density products and special methods of handling asbestos products) and protect the airways and in adopting methods of decontaminating equipment and work clothing. This controlled or safe use would be based on international standards.

We note that, according to the EC, controlled use does not work in certain occupational sectors such as those connected with building.\textsuperscript{174} Moreover, if the international standards suggested by Canada (in particular, ISO 7337) were applied, the exposure rate would still be higher than the level of risk that France considers acceptable.\textsuperscript{175} The EC also stress that the consistent or less inconsistent measures must be “technically and economically feasible.”\textsuperscript{176}

We note that the EC do not dispute that controlled use could constitute a measure consistent, or less inconsistent, with the GATT 1994. They nevertheless consider that it
would not allow the public health objectives pursued by France to be achieved. Insofar as Canada refers solely to controlled or safe use as an alternative to outright prohibition, we will focus our attention on this possibility. However, before continuing with our analysis of whether measures consistent, or less inconsistent, with the GATT are available in the present case, we consider it pertinent, in the light of the contradictory arguments put forward by the parties, to return to the question of the applicability of the feasibility test suggested by the EC.

8.207 We note that in United States – Section 337, the Panel stated that:

“A contracting party cannot justify a measure inconsistent with another GATT provision as ‘necessary’ in terms of Articles XX(d) if an alternative measure which it could reasonably be expected to employ and which is not inconsistent with other GATT provisions is available to it. By the same token, in cases where a measure consistent with other GATT provisions is not reasonably available, a contracting party is bound to use, among the measures reasonably available to it, that which entails the least degree of inconsistency with other GATT provisions.”

We therefore find that in order to determine whether a measure is necessary it is important to assess whether consistent or less inconsistent measures are reasonably available. The term “reasonably” has not been defined as such by the panels that have referred to it in the context of Article XX. It suggests, however, that the availability of a measure should not be examined theoretically or in absolute terms. Nevertheless, in the light of the reasoning of these panels, we find the word “reasonably” should not be interpreted loosely either. The fact that, administratively, one measure may be easier to implement than another does not mean that the other measure is not reasonably available. We consider that the existence of a reasonably available measure must be assessed in the light of the economic and administrative realities facing the Member concerned but also by taking into account the fact that the State must provide itself with the means of implementing its policies. Thus, the Panel considers that it is legitimate to expect a country such as France with advanced labour legislation and specialized administrative services to deploy administrative resources proportionate to its public health objectives and to be prepared to incur the necessary expenditure.

8.208 After clarifying this point, we will now proceed to examine whether controlled use (a) is sufficiently effective in the light of France’s health policy objectives and (b) whether it constitutes a reasonably available measure.

8.209 In relation to the first of these considerations, we note, first of all, that although controlled use is applied in some countries, such as the United States or Canada, and has also been applied by France, in general in certain sectors its efficacy still remains to be demonstrated. This is confirmed by a number of studies, as well as by the comments of the experts. Thus, even though it seems possible to apply controlled use successfully upstream (mining and manufacturing) or downstream (removal and destruction) of product use, it would seem to be much less easy to apply it in the building sector, which is one of the areas more particularly targeted by the measures contained in the Decree. The Panel therefore concludes that, in view of the difficulties of application of controlled use, an official in charge of public health policy might reasonably consider that controlled use did not provide protection that was adequate in relation to the policy objectives.

8.210 Moreover, Canada refers to the existence of international standards for the protection of workers in contact with chrysotile. First of all, we find that the international standards cover only the precautions to be taken if a worker has to handle asbestos. They contain neither a guarantee of free access for asbestos nor an incentive to use asbestos. On the contrary, the international conventions suggest that, as far as possible, asbestos should be replaced by less hazardous materials. Next, we note that the levels of protection obtained by following international standards, whether it be the ISO standard or the WHO Convention, are lower than those established by France, including those applicable before the introduction of the Decree. Considering the high level of risk identified, France’s objective – which the Panel cannot question – justifies the adoption of exposure ceilings
lower than those for which the international conventions provide. We therefore find that controlled use based on international standards would not seem to make it possible to achieve the level of protection sought by France.

8.211 The Panel is aware that in some sectors controlled or safe use could be envisaged with greater certainty that it would prove effective. However, as confirmed by the experts, the circumstances of use must be controllable. These circumstances are extremely varied and we note that the safety measures that would make possible results at least equivalent to the exposure level (0.1 f/ml) applied by France before the ban (restrictions on the number of workers and working areas and total containment of the product) exceed the requirements of the international standards and considerably limit the number of industrial sectors that could apply them. Even in these cases, according to one of the experts, the level of exposure is still high enough for there to be a significant residual risk of developing asbestos-related diseases. According to another, it is not possible to guarantee that fibre concentrations will never exceed 0.1 f/ml. In addition, we note that for the application of controlled use to satisfy France’s public health objectives, mined or processed products should never be handled by anyone outside the mining and processing industries. If these products were subsequently to be handled by unprotected persons, the fact that they could be mined and processed and then destroyed using controlled use techniques would not be sufficient to meet those objectives. We therefore find that a decision-maker responsible for establishing a health policy might have reasonable doubts about the possibility of ensuring the achievement of France’s health policy objectives by relying on controlled use, even in sectors which might lend themselves more readily to these practices.

8.212 *A fortiori* and for the following reasons, we consider that controlled use is not a reasonably available alternative in all the other sectors in which workers may be exposed to chrysotile.

8.213 The Panel notes that Canada’s arguments under Article III:4 are limited to chrysotile and chrysotile-cement products. In fact, chrysotile-cement is mainly used in the building sector. As the experts have confirmed, because of the mobility of the workers and their sometimes inadequate training, as well as the large number of sites and therefore of people liable to exposure, it is very difficult to impose on the building sector sophisticated occupational safety practices of the type that can be applied in sectors with smaller numbers of workers concentrated in well-defined areas.

8.214 Moreover, while controlled use may seem difficult to apply in the building sector, it is even less feasible in the case of DIY enthusiasts or undeclared workers operating outside any proper framework or system of controls. France’s objective is to halt the spread of the risks associated with chrysotile. At least as far as DIY enthusiasts are concerned, controlled use is not a reasonably available option. In this context, the fact that controlled use might be reasonably available in other sectors is irrelevant. Here, we are concerned with a product that could be installed in people’s homes, not one limited to a restricted use in areas in which only professionals would be required to work. Insofar as, once the products have been installed, it is impossible to guarantee that they will not be handled by someone who will not follow controlled use practices, it seems to us that a decision-maker responsible for adopting a health policy might well conclude that there was still a flaw in the health protection system if only controlled use measures were applied.

8.215 Canada also points out that, even though their use might not be limited to occupational activities, some products will never be handled by unqualified people. The Panel notes, however, that, as it found in paragraphs 8.200-8.202 above, exposure to asbestos fibres is not restricted exclusively to workers or DIY enthusiasts. Others present while the work is going on, in some cases even spouses, could be directly or indirectly affected. As one of the experts has shown, exposure to asbestos may be the result of pure chance.

8.216 Moreover, the Panel notes that products containing chrysotile may remain in place for very long periods of time. Thus, every time they are handled, the handler could
be exposed. The continued marketing of products containing chrysotile asbestos multi-
plies the situations in which workers are routinely exposed and exposes them to concent-
trations of asbestos which have already been related to pathologies in humans. In other
words, even if the exposure rates are very low, the multiplication of sources of exposure
may lead to concentrations already found to have caused disease.

8.217 We therefore conclude that the European Communities have shown that con-
trolled use is neither effective nor reasonably available, at least in the building sector and
for DIY enthusiasts. Accordingly, controlled use does not constitute a reasonable alter-
ative to the banning of chrysotile asbestos that might be chosen by a decision-maker respon-
sible for developing public health measures, bearing in mind the objectives pursued by
France.

Recourse to substitute fibres and products

8.218 Canada argues that France is creating a false sense of security by requiring
chrysotile, whose effects are well known, to be replaced by products whose effects on health
are uncertain.

8.219 The European Communities argue that the Decree does not recommend the in-
discriminate use of substitute products. It leaves it to businesses to replace asbestos by
whichever product they choose. In view of the procedure laid down in Article 2 of the
Decree, the replacement of asbestos fibres by substitute fibres is the result of a reasonable
and justified process.

8.220 The Panel notes, first of all, that the risk posed by chrysotile is recognized inter-
nationally, which in itself may justify the taking of measures to restrict its use. On the
other hand, we find that the substitute fibres examined in the context of this case (PVA,
cellulose and glass) are not classified by the WHO at the same level of risk as chrysotile.
Moreover, the experts consulted by the Panel who commented in detail on the risks associ-
ated with the substitute fibres examined in the context of this case (PVA, cellulose and
glass) and other fibres, (in particular aramid and ceramic) confirmed that these fibres did
not present the same risk to health as chrysotile. In particular, animal studies do not
show the fibres in question to be carcinogenic.

8.221 Canada’s approach seems to be based on the fact that chrysotile can be used
safely. As we have already concluded, this does not appear to be a reasonably available
possibility, at least as far as building workers and DIY enthusiasts are concerned. If we
were to follow Canada’s reasoning, then substitute fibres could not be used until a degree
of certainty equivalent to that which exists with respect to chrysotile had been established.
In the opinion of the Panel, to make the adoption of health measures concerning a definite
risk depend upon establishing with certainty a risk already assessed as being lower than
that created by chrysotile would have the effect of preventing any possibility of legislat-
ing in the field of public health. In fact, it would mean waiting until scientific certainty, which
is often difficult to achieve, had been established over the whole of a particular field before
public health measures could be implemented.

Conclusion

8.222 In the light of France’s public health objectives as presented by the European
Communities, the Panel concludes that the EC has made a prima facie case for the non-
existence of a reasonably available alternative to the banning of chrysotile and chrysotile-
cement products and recourse to substitute products. Canada has not rebutted the pre-
sumption established by the EC. We also consider that the EC’s position is confirmed by
the comments of the experts consulted in the course of this proceeding.

8.223 At this stage, we conclude that the Decree satisfies the conditions of Article XX(b)
(d) Application of the introductory clause (chapeau) of Article XX of the GATT 1994 to the application of the Decree

(i) “Means of arbitrary or unjustifiable discrimination between countries where the same conditions prevail”

8.224 With regard to the introductory clause of Article XX, the EC point out that it applies to the manner in which the measure is applied. In the present case there is nothing to support the contention that France acted “in bad faith” or in an “unreasonable”, “improper” or “abusive” manner in exercising its right under Article XX(b). The ban is not a means of imposing arbitrary or unjustifiable discrimination between countries where the same conditions prevail. It covers products originating in any country, including France, where the same conditions prevail. The European Communities consider that, as the Decree does not meet the definition of discrimination laid down by the Appellate Body, it cannot be applied in such a manner as to constitute a means of arbitrary or unjustifiable discrimination between countries where the same conditions prevail.

8.225 Canada claims that it has already shown, in respect of Article III:4, that the initial Decree was discriminatory. The Decree is also arbitrary and unjustified because it is not motivated by the objective of protecting human life or health but rather by the desire to reassure a panicked population.

8.226 The Panel notes that under the first of the alternatives mentioned in the introductory clause of Article XX it is required to examine whether the application of the Decree constitutes a means of arbitrary or unjustifiable discrimination between countries where the same conditions prevail. The Panel considers that, within the context of this alternative, its first step should be to determine whether the measure is “discriminatory” in its application. If the application of the measure is found to be discriminatory, it still remains to be seen whether it is arbitrary and/or unjustifiable between countries where the same conditions prevail. It is in this context, and not in the stage of the existence of discrimination – which is an objective fact, that we shall determine whether the measures falling within the particular exceptions of Article XX(b) – in this case the initial Decree as applied – have been applied reasonably, with due regard to both the legal duties of the party claiming the exception and the legal rights of the other parties concerned.

8.227 The Panel also notes that, in United States – Gasoline, the Appellate Body stated that the word “discrimination” in the introductory clause of Article XX covers both discrimination between products from different supplier countries and discrimination between domestic and imported products. Finally, in the same case, the Appellate Body ruled that “the provisions of the chapeau [of Article XX ] cannot logically refer to the same standard(s) by which a violation of a substantive rule has been determined to have occurred.” In other words, we cannot conclude that discrimination exists on the basis of the violation of Article III:4 identified above. This means that the less favourable treatment of asbestos as compared with substitute fibres identified by Canada is not relevant for establishing the existence of discrimination under Article XX. The question is therefore confined to the suppliers of asbestos, whether domestic or foreign. However, we understand that another form of discrimination, for example between supplier countries, not invoked by Canada as its principal argument, could be taken into consideration under the introductory clause of Article XX. It is on this dual basis that we shall examine whether the measure is being applied in a “discriminatory” manner.

8.228 The Panel notes that the European Communities consider the Decree to concern products originating in any country, including France, where the same conditions prevail. The text of the Decree as such appears to confirm this. Only the product in question is mentioned, without any reference to its origin. Even Articles 2 and 3, which concern the establishment and administration of exceptions, do not contain any expressly discriminatory provision. It should be borne in mind, however, that the introductory clause of Article XX concerns the application of the measure. It would therefore be possible for Canadian exports of chrysotile or products containing it to receive less favourable treatment than
imports from other countries or French production with respect to the way in which the exceptions are administered by the French authorities. For example, the minister responsible might use the powers of notice under Article 3:III of the Decree to discriminate against an operator qualifying for an exception who imports chrysotile fibres from Canada. However, we note that Canada has not argued that this was or had been the case. Canada merely recalls that it has demonstrated the existence of discrimination under Article III:4. For the reasons set out above, we consider that this demonstration is not relevant in terms of the introductory clause of Article XX.

8.229 We therefore conclude that, although this is a heavier task than that involved in showing that an exception falling within one of the paragraphs of Article XX encompasses the measure at issue, the EC have made a prima facie case for their argument that the Decree does not constitute, in its application, arbitrary or unjustifiable discrimination. We do not consider that Canada has rebutted the presumption established by the prima facie case made by the EC, according to which the Decree does not introduce discrimination.

8.230 In accordance with our approach, since discrimination has not been established in relation to the application of the Decree, there is no need to consider the question of its arbitrariness or unjustifiability.

(ii) “Disguised restriction on international trade”

8.231 According to the EC, the Decree does not constitute a “disguised restriction on international trade”. The EC consider that the fact that the Decree is not applied in a manner which would constitute a means of arbitrary or unjustifiable discrimination is proof enough that it is not being applied in a manner which constitutes a disguised restriction on international trade. In particular, the EC point out that the restriction was published, that many other Members of the WTO also apply restrictions on these products and that the restriction takes international standards as a base. Any other approach would imply that all legislation on asbestos and asbestos-containing products, in so far as it imposes restrictions, amounts to a “disguised restriction on international trade”.

8.232 Canada considers that the Decree is a “disguised restriction on international trade”. The Appellate Body has excluded a narrow reading of the term “disguised restriction”. The fact that the measure was published does not prevent it from being a disguised restriction on international trade. The Decree is contrary to the introductory clause of Article XX in the sense that, under the cover of a public health decision, it favours the French national industry of substitute products for chrysotile and products containing it.

8.233 The Panel notes, first of all, that the actual scope of the words “disguised restriction on international trade” has not been clearly defined. Under the GATT 1947, panels seem mainly to have considered that a disguised restriction on international trade was a restriction that had not been taken in the form of a trade measure or had not been announced beforehand or formed the subject of a publication, or even had not been the subject of an investigation.

8.234 In United States – Gasoline, the Appellate Body considered “that concealed or unannounced restriction … in international trade does not exhaust the meaning of ‘disguised restriction’. This seems to imply that a measure that was not published would not satisfy the requirements of the second proposition of the introductory clause of Article XX. We note that the Decree was published in the Official Journal of the French Republic on 26 December 1996 and entered into force on 1 January 1997. We also note that it applies unequivocally to international trade, since as far as asbestos is concerned both importation and exportation are prohibited. In this sense, the criteria developed in United States – Tuna (1982) and in United States – Automotive Springs have already been satisfied.

8.235 However, the remark made by the Appellate Body in United States – Gasoline also implies that the expression "disguised restriction on international trade" covers other requirements. In the same case, the Appellate Body mentions that “disguised restriction includes disguised discrimination in international trade”. This appears to signify that the word “restriction” should not be given a narrow meaning. In paragraph 8.229 above, we
found that the Decree did not constitute discrimination within the meaning of the first condition of the introductory clause of Article XX. We consider that our conclusion can also be used for determining whether the Decree constitutes a disguised restriction. In this respect, we note that in the above-mentioned case the Appellate Body considered that:

“'disguised restriction', whatever else it covers, may properly be read as embracing restrictions amounting to arbitrary or unjustifiable discrimination in international trade taken under the guise of a measure formally within the terms of an exception listed in Article XX. Put in a somewhat different manner, the kinds of considerations pertinent in deciding whether the application of a particular measure amounts to ‘arbitrary or unjustifiable discrimination’, may also be taken into account in determining the presence of a ‘disguised restriction’ on international trade. The fundamental theme is to be found in the purpose and object of avoiding abuse or illegitimate use of the exceptions to substantive rules available in Article XX.”204

8.236 Referring also to the remark made by the Appellate Body in the same case according to which “the provisions of the chapeau [of Article XX] cannot logically refer to the same standard(s) by which a violation of the substantive rule has been determined to have occurred”, we consider that the key to understanding what is covered by “disguised restriction on international trade” is not so much the word “restriction”, inasmuch as, in essence, any measure falling within Article XX is a restriction on international trade, but the word “disguised”. In accordance with the approach defined in Article 31 of the Vienna Convention, we note that, as ordinarily understood, the verb “to disguise” implies an intention. Thus, “to disguise” (déguiser) means, in particular, “conceal beneath deceptive appearances, counterfeit”, “alter so as to deceive”, “misrepresent”, “dissimulate”.205 Accordingly, a restriction which formally meets the requirements of Article XX(b) will constitute an abuse if such compliance is in fact only a disguise to conceal the pursuit of trade-restrictive objectives. However, as the Appellate Body acknowledged in Japan – Alcoholic Beverages, the aim of a measure may not be easily ascertained.206 Nevertheless, we note that, in the same case, the Appellate Body suggested that the protective application of a measure can most often be discerned from its design, architecture and revealing structure.207

8.237 It is on this basis that we shall analyse Canada’s argument according to which the Decree favours the French domestic industry producing substitutes for chrysotile and chrysotile-containing products.208 We have already found that the Decree was necessary to achieve a public health objective and did not, in its application, constitute arbitrary or unjustifiable discrimination. We recall that in United States – Gasoline, the Appellate Body considered that the kinds of considerations pertinent in deciding whether the application of a particular measure amounts to “arbitrary or unjustifiable discrimination” may also be taken into account in determining the presence of a “disguised restriction” on international trade. Since we have not identified discrimination, we consider it is unnecessary to determine whether we are faced with discrimination that might constitute a disguised restriction on international trade.

8.238 As far as the design, architecture and revealing structure of the Decree are concerned, we find nothing that might lead us to conclude that the Decree has protectionist objectives. Canada implicitly admits as much when it asserts that the Decree is a response of the French authorities to panicked public opinion and other health scares implicating officials and members of the government.209 If this was the case, it seems difficult to reconcile the Decree being adopted in great haste with the idea that it was the result of a premeditated intention to protect French industry.

8.239 Admittedly, there is always the possibility that measures such as those contained in the Decree might have the effect of favouring the domestic substitute product manufacturers. This is a natural consequence of prohibiting a given product and in itself cannot justify the conclusion that the measure has a protectionist aim, as long as it remains within certain limits. In fact, the information made available to the Panel does not suggest that the
import ban has benefited the French substitute fibre industry, to the detriment of third country producers, to such an extent as to lead to the conclusion that the Decree has been so applied as to constitute a disguised restriction on international trade.210

8.240 Consequently, we conclude that the Decree satisfies the conditions of the introductory clause of Article XX.

5. Conclusion

8.241 In the light of the above, the Panel concludes that the provisions of the Decree which violate Article III:4 of the GATT 1994 are justified under Article XX(b).

F. ALLEGATION OF NULLIFICATION OR IMPAIRMENT OF A BENEFIT UNDER ARTICLE XXIII:1(b) OF THE GATT 1994

1. Arguments of the parties211

8.242 Having determined that the Decree was justified in the light of Article XX(b) of the GATT 1994 and consistent in its application with the introductory provisions of that Article, we must now examine Canada’s allegations and arguments based on Article XXIII:1(b) of the GATT 1994.

8.243 The main arguments of the parties with regard to the existence of non-violation nullification or impairment of a benefit (hereinafter “non-violation nullification”) can be summarized as follows.

8.244 Canada considers that, in accordance with panel practice under the GATT 1947, as confirmed in Japan – Measures Affecting Consumer Photographic Film and Paper212, three conditions must be met for a case of non-violation nullification: (i) the negotiation of a tariff concession; (ii) the subsequent adoption of a governmental measure that unfavourably disrupts conditions of competition between the imported product for which the concessions were granted and the like or directly competitive domestic products; and (iii) the fact that the adoption of the measure in question could not reasonably have been foreseen at the time of the tariff concession negotiations.

8.245 Canada considers that these three conditions are present in this case. Asbestos and many asbestos-containing products are subject to tariff concessions by France and the European Communities starting in 1947 and 1960-1961, respectively, as well as during the Uruguay Round negotiations. By establishing a total ban, the Decree disrupted the competitive relationship in the French market between, on the one hand, chrysotile asbestos fibre and products containing it and, on the other, like and competitive French products, thus creating a monopoly for substitute fibres. Finally, at the time the tariff concessions concerning asbestos were negotiated, Canada could not reasonably have foreseen that France was going to abandon its policy of controlled use and compromise the value of its commitments by implementing a total ban on chrysotile and any possible use thereof by means of a measure as excessive as the Decree. Furthermore, this measure was inconsistent with the type of regulatory intervention then in place and still in effect today concerning products equally harmful, if not more harmful, than chrysotile. Canada’s expectations were that this type of measure would not be adopted unless there were exceptional circumstances, but there had been no new scientific developments that had changed anything in terms of managing the risks or the effects associated with chrysotile. Finally, Canada argues that it could not reasonably have expected that chrysotile would be banned in favour of substitute products without those products having been subjected to a rigorous process of examination to prove that their use satisfied the public health objectives invoked by France.

8.246 By way of a preliminary remark, the European Communities point out that in the context of a non-violation allegation the burden of proof is especially onerous as a result of, in particular, Article XXVI:1(a) of the Understanding. They also note that in Japan – Film, the Panel stated that Article XXIII:1(b) of the GATT 1994 should be approached with caution and that non-violation nullification or impairment of benefits should be treated
as an exceptional concept. The European Communities also claim that the rules on non-violation nullification or impairment apply only if the measure in question does not fall under other provisions of the GATT and that, while there may be “legitimate expectations” in connection with a purely commercial measure, there can be no such expectations with respect to a measure taken to protect human health and which can therefore be justified with regard to Article XX:(b) of the GATT 1994.

8.247 The European Communities maintain that Canada fails to demonstrate how the French measure could not reasonably have been anticipated. In substance, according to the European Communities, since 1977, chrysotile has been classified as a category I carcinogenic product by the WHO. Thus, when the tariff negotiations took place, Canada knew that there was a danger that the product under negotiation could at any time be prohibited by Members of the WTO, particularly if non-hazardous or less hazardous substitutes could be used. In 1986, ILO Convention 162 on asbestos stated that national legislation should provide wherever possible for the replacement of asbestos or of certain types of asbestos or products containing asbestos by other materials or products or the use of alternative technology, scientifically evaluated by the competent authority as harmless or less harmful. Since 1990, the European Communities, under Directive 90/394/EEC, has provided for the replacement of asbestos. This Directive recommends the principle of replacing a dangerous substance or process with a non-dangerous or less dangerous substance or process, where one exists. In 1983, WTO Members began to ban the use of asbestos, including chrysotile. Among EC Members, Austria banned chrysotile in 1990, followed by Finland and Italy in 1992, and Germany in 1993. Canada might have expected France to follow suit.

8.248 Furthermore, the European Communities assert that there are no provisions in the GATT or the TBT Agreement requiring consistency in the application of health measures against substances that pose a carcinogenic risk for human health. Accepting Canada’s argument that it could not legitimately have anticipated the ban imposed by the Decree because France did not simultaneously ban other potentially dangerous substances (such as lead and copper) would be equivalent to preventing Members entirely from taking measures to protect human health on their territory.

8.249 Moreover, according to the European Communities, Canada cannot claim a legitimate expectation of “improved” market access with respect to a product which entails risks for human health when, as the facts show, the tendency is for exports to the industrialized countries to fall. Canada must give detailed reasons as to why it could legitimately have expected that France would not adopt measures restricting or eliminating the use of any asbestos product after the Uruguay Round negotiations, given the growing scientific evidence that all types of asbestos and asbestos-containing products are carcinogenic to humans.

8.250 The European Communities then argue that Canada fails to demonstrate how the Decree upsets the competitive relationship between asbestos and fibrous or non-fibrous substitute products. It has not established a “clear correlation” between the two. The issue is not whether equality of competitive conditions exists but whether the relative conditions of competition which existed between domestic and foreign products as a consequence of the relevant tariff concessions have been upset. The EC consider that the products for which the competitive conditions must be examined are those covered by the tariff concession. If a tariff concession is granted for asbestos, the competitive conditions to be examined are those concerning Canadian asbestos and French asbestos. It is irrelevant to compare chrysotile with French substitute products because such products cannot be considered in terms of the same relevant tariff concession.

8.251 Canada rejects the EC’s claim that an examination of the impact of the Decree’s effect on competitive conditions must be limited to Canadian asbestos and French asbestos. This approach is contradicted by two panel reports which have clearly established that Article XXIII:1(b) can be invoked in the case of a measure which upsets the competitive relationship between two non-identical products.
2. Analysis by the Panel

(a) Preliminary issues

(i) Questions before the Panel

8.252 Article XXIII:1(b) of the GATT 1994 (Nullification or impairment) reads as follows:

“1. If any contracting party should consider that any benefit accruing to it directly or indirectly under this Agreement is being nullified or impaired or that the attainment of any objective of the Agreement is being impeded as the result of

[...]

(b) the application by another contracting party of any measure, whether or not it conflicts with the provisions of this Agreement,

[...]

the contracting party may, with a view to the satisfactory adjustment of the matter, make written representations or proposals to the other contracting party or parties which it considers to be concerned. Any contracting party thus approached shall give sympathetic consideration to the representations or proposals made to it.”

8.253 The Panel notes, first of all, that the EC have put forward two arguments which, if well-founded, should lead us to conclude that Article XXIII:1(b) does not apply to the facts in this case. If we find that these arguments are not pertinent, we shall examine the substance of the Canadian complaint.

8.254 The EC’s two arguments which could determine the applicability of Article XXIII:1(b) are as follows:

8.255 Firstly, the EC claim that the rules on non-violation nullification apply only if the measure in question does not fall under other provisions of the GATT 1994. Article XXIII:1(b) is applicable only if the Panel reaches the conclusion that the Decree is consistent with Article III of the GATT 1994. Otherwise there cannot be “non-violation”. In the light of the introductory clause of Article XX of the GATT 1994, the European Communities conclude that if the French measure is considered “necessary” for the protection of human health by the Panel and hence if specific rules have been applied in this respect, the provisions of Article XXIII:1(b) of the GATT are inapplicable.

8.256 Canada replies that the cases Uruguay – Recourse to Article XXIII and United States – Trade Measures Affecting Nicaragua do not support the EC’s interpretation. Moreover, inasmuch as Article 26:1(b) of the Understanding provides for the granting of compensation rather than the withdrawal of a measure, recourse in non-violation affects neither the adoption nor the application of the contested measure.

8.257 The EC’s second argument is that while it is possible to have legitimate expectations in connection with a purely commercial measure, it is not possible to claim legitimate expectations with respect to a measure that is taken to protect human health and can therefore be justified, particularly in the light of Article XX(b) of the GATT 1994. The protection of human health is a fundamental duty and cannot be compromised or restricted by the concept of non-violation.

8.258 For Canada, the distinction made by the EC between measures of a purely commercial nature and measures which have health-protection related aspects has no basis either in the texts of the WTO Agreement or in case-law. A legitimate expectation does not
in any way concern a particular measure adopted by a Member, but rather the opportunities for competition agreed during multilateral trade negotiations on a given product. The Community reasoning is also wrong because it does not concur with the preparatory work on the GATT 1947, which shows that the objective of Article XXIII:1(b) is to prevent abuse of the provisions of the General Agreement.

8.259 The EC consider that Canada has taken a selective look at the preparatory documents. As the Appellate Body pointed out in the United States – Shrimp case, the conditions laid down in the chapeau of Article XX(b) are meant precisely to address situations in which a Member applies in bad faith and in an abusive manner the exceptions laid down in Article XX. There cannot be two sets of provisions which address the same problem twice.

(ii) The EC’s argument according to which the rules on non-violation nullification apply only if the measure in question does not fall under other provisions of the GATT.

8.260 The EC seem to believe that the fact that a measure is “justified” on the basis of Article XX creates a legal situation different, on the one hand, from the situation in which the measure violates a provision of the GATT 1994 and, on the other, from the situation in which the measure does not fall under the provisions of the GATT 1994. In support of their position, the EC cite a passage from the Panel Report in Japan – Film which mentions that Article XXIII:1(b) provides “the means to redress government actions not otherwise regulated by GATT rules ... ”. The Communities also refer to the introductory clause of Article XX which states that “nothing in this Agreement shall be construed to prevent the adoption or enforcement by any contracting party of measures” necessary to protect human life or health.

8.261 The Panel recalls, first of all, that both the preamble to Article 26.1 of the Understanding and Article XXIII:1(b) use the words “measure, whether or not it conflicts with the provisions [of the particular agreement]” (emphasis added). To begin with, it should be noted that the wording of Article XXIII:1(b) shows unequivocally that this provision applies both in situations in which a measure conflicts and in situations in which it does not conflict with the provisions of the GATT 1994. Above, we found that the treatment accorded by the Decree to chrysotile asbestos fibres violated Article III-4 of the GATT 1994 as such, in as much as these products were like the substitute fibres mentioned by the parties and the treatment of products containing chrysotile asbestos and products containing the substitute fibres mentioned by the parties was discriminatory. Accordingly, the Decree conflicts with the provisions of Article III-4, in the sense in which that word is used in Article XXIII:1(b). However, we note that the introductory clause of Article XX states that “nothing in this Agreement shall be construed to prevent the adoption or enforcement by any contracting party of measures” necessary to protect human life or health, which might suggest that a provision consistent with the requirements of Article XX no longer conflicts with Article III-4, because Article III-4 cannot be construed as preventing this kind of measure. However, whether a measure justified on the basis of Article XX of the GATT 1994 is considered still to be in conflict with Article III-4 or is considered no longer to conflict with Article III-4 because justified under Article XX, under the terms of Article XXIII:1(b) the latter continues to be applicable to it.

8.262 We also note, firstly, that the introductory clause to Article XX, to which the EC refer, concerns the adoption or enforcement of measures necessary to protect health. The application of Article XXIII:1(b) does not prevent either the adoption or the enforcement of the Decree concerned. Article 26:1(b) stipulates that even where a measure has been found to nullify or impair benefits under, or impede the attainment of objectives, of the GATT 1994 without violation thereof, there is no obligation to withdraw the measure. Accordingly, there is no contradiction between the invocation of Article XX and the application of Article XXIII:1(b). However, that Article must be applied in such a way as to protect the balance of rights and duties negotiated. Accordingly, we do not consider that the text of Article XXIII:1(b) or that of Article XX or, finally, that of Article 26.1 of the Understanding supports the EC’s interpretation.
8.263 Secondly, we do not consider that the passage from the Japan – Film report cited by the EC supports its interpretation either. Admittedly, the words used by the panel, taken in isolation, might at first glance appear to confirm the EC’s position, insofar as it refers to “government actions not otherwise regulated by GATT rules” (emphasis added). The use of the word “regulated” could signify that the field of application of Article XXIII:1(b) covered only situations in which no provision of the GATT was applicable. First of all, it is our opinion that the fact that a measure does not violate Article III:4 does not necessarily mean that the latter is not applicable to it. Article III:4 applies to any law, regulation or requirement affecting the internal sale, offering for sale, purchase, transportation, distribution or use of imported products and like products of national origin. Consequently, even if the EC’s interpretation were correct, it would not apply in the present case insofar as Article III:4 continues to be applicable to the Decree. Next, it should be noted that the panel in Japan – Film refers, in the footnote at the end of the sentence cited by the EEC219, to the EEC – Oilseeds report which states, in particular, that:

“the Panel noted that these provisions, as conceived by the drafters and applied by the CONTRACTING PARTIES, serve mainly to protect the balance of tariff concessions. [footnote omitted] The idea underlying them is that the improved competitive opportunities that can legitimately be expected from a tariff concession can be frustrated not only by measures proscribed by the General Agreement but also by measures consistent with that Agreement”.220

8.264 We consider that a “measure which is not otherwise regulated by GATT rules”, that is to say to which the GATT does not apply, is, a fortiori, “not in conflict” with the GATT within the meaning of Article XXIII:1(b) or “consistent” within the meaning of the EEC – Oilseeds report. Consequently, we find that the passage in the Japan – Film report cited by the EC, far from supporting their position, confirms the opinion according to which Article XXIII:1(b) applies to a measure whether it is consistent with the GATT because the GATT does not apply to it or is justified by Article XX.221

8.265 For these reasons, we do not allow the EC’s first argument.

(iii) The EC’s argument according to which there cannot be a “legitimate expectation” in the case of a measure that concerns the protection of human health.

8.266 With regard to the EC’s second argument, we note, first of all, that neither the text of Article XXIII:1(b) of the GATT 1994 nor that of Article 26:1 of the Understanding expressly incorporates the separation suggested by the EC between measures of a purely commercial nature and measures designed to protect human health. Although these articles require the existence of a measure – which neither of the Parties disputes - they do not distinguish between different types of measures. We have also found, on the basis of Article XX, that the application of the latter does not a priori exclude the application of Article XXIII:1(b).222 We therefore find that the terms and the context of Article XXIII:1(b) do not support the interpretation proposed by the EC.

8.267 Canada cites the preparatory work on the GATT 1947. On the basis of Article 32 of the Vienna Convention, we consider that it is not necessary to have recourse to the preparatory work unless, in particular, the interpretation based on the criteria of Article 31 leaves the meaning of the terms ambiguous or obscure or leads to a manifestly absurd or unreasonable result. Such is not the case. However, recourse to the preparatory work also makes it possible to confirm the meaning resulting from the application of Article 31.

8.268 In this respect, we note that the EC consider that Canada’s reading of the preparatory work is selective. According to them, the potential problems of abuse and bad faith to which Canada alludes are adequately covered by the chapeau of Article XX. For the EC, there cannot be two sets of provisions which address the same problem twice.

8.269 Although it is not necessary to take a position on the content of the preparatory work, we consider that the EC’s argument tends to confuse two aspects: the first is abuse resulting from the application of a measure falling within one of the paragraphs of Article
XX. If a measure necessary to protect human health is applied in a manner that conflicts with the provisions of the introductory clause of Article XX, the measure will still be in conflict with the provisions of the GATT whose violation Article XX is supposed to justify. This aspect is very different from the situation in which a measure is perfectly justified in itself in relation to the GATT (as in the case of a measure which satisfies all the conditions of Article XX), but which, viewed in a given context, could give rise to a situation of nullification or impairment of a benefit under a tariff concession.

8.270 It remains, however, for us to discuss the EC’s argument to the effect that the fundamental duty to protect human health cannot be compromised or restricted by the concept of non-violation nullification. We must begin by acknowledging that all the cases examined by panels so far have concerned situations in which the measure adopted following the negotiation of a concession was purely commercial in nature, generally a subsidy, a tariff preference or a measure relating to product distribution. Accordingly, we have no precedents to guide us. However, a preliminary remark, similar to that made in paragraph 8.262 above, is called for. A finding based on Article XXIII:1(b) of the GATT 1994 and Article 26.1 of the Understanding never results in an obligation not to apply or to withdraw the measure in question. The Member concerned can only be asked to make “a mutually satisfactory adjustment”. Article 26:1(b) also specifies that compensation may be part of a mutually satisfactory adjustment as final settlement of the dispute. The Member adopting a public health protection measure is totally free to continue to apply the measure concerned as it stands while offering in exchange compensation for the benefits nullified or impaired.

8.271 The Panel also considers, as did the panel in Japan – Film, that non-violation should be approached with caution and treated as an exceptional instrument of dispute settlement. It appears that Members which have negotiated a set of rights and obligations would only exceptionally expect to be challenged for actions not in contravention of those rights or obligations.

8.272 Moreover, the Panel is of the opinion that even if the justification of a measure by Article XX does not, in principle, make it impossible to invoke Article XXIII:1(b) in relation to the application of the measure justified, the situation of a measure falling under Article XX with respect to Article XXIII:1(b) cannot be quite the same as that of a measure consistent with another provision of the GATT 1994. This is because Article XX, which is headed “General Exceptions”, is intended, in particular, to ensure the protection of public health or, as stated by the Appellate Body in United States – Gasoline, to “permit important State interests – including the protection of human health … to find expression”. The Panel considers that in accepting the WTO Agreement Members also accept a priori, through the introduction of these general exceptions, that Members will be able, at some point, to have recourse to these exceptions. Moreover, Members have attached to the use of these exceptions a certain number of conditions contained either in paragraphs (a) to (j) or in the introductory clause of Article XX. These conditions have generally been narrowly interpreted. The result is that

(a) both the intended objective of these exceptions (pursuit of interests recognized a priori as being of greater importance than Members’ commercial interests, since they can outweigh the latter) and

(b) the specific conditions that must be satisfied by Members invoking these exceptions

mean that, while recognizing that Article XXIII:1(b) applies to measures that fall under Article XX, we are justified in treating recourse to Article XIII:1(b) as particularly exceptional in relation to measures justified by Article XX(b).

8.273 All this leads the Panel to consider that, in practice, even if in a particular case a mutually satisfactory adjustment may be made under Article XXIII:1(b), in general, the risk of an effective increase in the cost of measures necessary to protect public health because of the applicability of Article XXIII:1(b) to measures justified under Article XX can only be very marginal. In fact, considering the criteria mentioned in the previous paragraph, very few measures of this kind could give rise to the application of Article XXIII:1(b).
8.274 For these reasons we do not subscribe to the interpretation proposed by the European Communities. Accordingly, we will continue our examination of the measure in the light of Article XIII:1(b) of the GATT 1994.

(b) Examination of the substantial aspects of Canada’s arguments under Article XXIII:1(b) of the GATT 1994

(i) Burden of proof

8.275 The Panel has already discussed the question of the burden of proof in general in Section VIII.E.1(b) above. However, it is important, in the light of the arguments of the parties, to clarify the application of the burden of proof in the context of non-violation nullification or impairment of a benefit.

8.276 Where the application of Article XXIII:1(b) is concerned, Article 26.1(a) of the Understanding and panel practice in the context of the WTO Agreement and the GATT 1947 confirm that this is an exceptional course of action requiring the complaining party to carry the burden of presenting a detailed justification in support of its complaint. Thus, Article 26.1(a) stipulates that:

“Where the provisions of paragraph 1(b) of Article XXIII of the GATT 1994 are applicable to a covered agreement, a panel or the Appellate Body may only make rulings and recommendations where a party to the dispute considers that any benefit accruing to it directly or indirectly under the relevant covered agreement is being nullified or impaired or the attainment of any objective of that Agreement is being impeded as a result of the application by a Member of any measure, whether or not it conflicts with the provisions of that Agreement. Where and to the extent that such party considers and the panel or the Appellate Body determines that a case concerns a measure that does not conflict with the provisions of a covered agreement to which the provisions of paragraph 1(b) of Article XXIII of the GATT 1994 are applicable, the procedures in this Understanding shall apply, subject to the following:

(a) the complaining party shall present a detailed justification in support of any complaint relating to a measure which does not conflict with the relevant covered agreement;”

8.277 In Japan – Film, the Panel specified how the burden of proof should be applied in this context:

“Consistent with the explicit terms of the DSU and established WTO/GATT jurisprudence, and recalling the Appellate Body ruling that ‘precisely how much and precisely what kind of evidence will be required to establish … a presumption [that what is claimed is true] will necessarily vary from … provision to provision’, we thus consider that the United States, with respect to its claim of non-violation nullification or impairment under Article XXIII:1(b), bears the burden of providing a detailed justification for its claim in order to establish a presumption that what is claimed is true. It will be for Japan to rebut any such presumption.”

8.278 We conclude that, with respect to its claims of non-violation, Canada bears the primary burden of presenting a detailed justification for its claims.

8.279 Canada maintains that, when a complainant proves that it enjoys a tariff concession and the respondent subsequently adopts a measure that affects the value of this concession, the complainant benefits from the presumption that it could not reasonably anticipate that this concession would be nullified or otherwise impaired by this measure. In such circumstances, it is up to the respondent to prove that the complainant should have
anticipated the possibility of such a measure being adopted. Canada refers to the Appellate Body report in *India – Patent (United States)* and to the Panel report in *Japan – Film*.

8.280 We do not consider that Canada has correctly interpreted the Panel report in *Japan – Film*. First of all, the presumption to which the Panel refers is that, if it is shown that a measure has been introduced after the conclusion of the tariff negotiations in question, then the complainant should not be considered as having anticipated that measure, which is only one of the tests applied by the Panel. Moreover, if the interpretation of the burden of proof suggested by Canada were followed, the obligation to present a detailed justification for which Article 26.1(a) provides might in certain cases be evaded. Accordingly, we do not follow the interpretation proposed by Canada but the rule laid down in *Japan – Film*.

8.281 Furthermore, in the light of our reasoning in paragraph 8.272 above, we consider that the special situation of measures justified under Article XX, insofar as they concern non-commercial interests whose importance has been recognized *a priori* by Members, requires special treatment. By creating the right to invoke exceptions in certain circumstances, Members have recognized *a priori* the possibility that the benefits they derive from certain concessions may eventually be nullified or impaired at some future time for reasons recognized as being of overriding importance. This situation is different from that in which a Member takes a measure of a commercial or economic nature such as, for example, a subsidy or a decision organizing a sector of its economy, from which it expects a purely economic benefit. In this latter case, the measure remains within the field of international trade. Moreover, the nature and importance of certain measures falling under Article XX can also justify their being taken at any time, which militates in favour of a stricter treatment of actions brought against them on the basis of Article XXIII:1(b).

8.282 Consequently, the Panel concludes that because of the importance conferred on them *a priori* by the GATT 1994, as compared with the rules governing international trade, situations that fall under Article XX justify a stricter burden of proof being applied in this context to the party invoking Article XXIII:1(b), particularly with regard to the existence of legitimate expectations and whether or not the initial Decree could be reasonably anticipated.

(ii) Examination of the conditions

8.283 As recalled by the Panel in *Japan – Film*, the text of Article XXIII:1(b) establishes three elements whose existence a complainant must prove in order to be able legitimately to invoke that provision, namely: (1) the application of a measure by a Member of the WTO; (2) the existence of a benefit accruing under the applicable agreement; and (3) the nullification or impairment of a benefit as a result of the application of the measure.

8.284 The existence of a *measure* is not in dispute. In this particular case, the measure in question is Decree No. 96-1133 of 24 December 1996 which, moreover, forms the subject of a violation complaint on the part of Canada and which we have examined in this respect in previous sections. We also note that, apart from stressing that its objective is to protect human health, the European Communities do not claim that the nature of the measure (an import and marketing ban) is such that it is not covered by the definition of the term “measure” in Article XXIII:1(b). We therefore conclude that a “measure” exists.

8.285 With regard to the existence of a *benefit*, we note that the Panel in *Japan – Film* recalled that, with only one exception, in all the previous cases in which Article XXIII:1(b) was invoked the benefit claimed consisted in the legitimate expectation of improved market access opportunities resulting from the relevant tariff concessions. We first need to know what benefit Canada could legitimately have expected from the Community concessions on chrysotile asbestos. We note, however, that previous panels approached the question differently, insofar as they appear to have assumed the existence of a benefit in the form of improved market access opportunities and then considered whether a party could have had a *legitimate expectation* of a given benefit.
In the present context, there would be a certain logic in making a distinction between the concept of legitimate expectation of a benefit and that of the reasonable foreseeability of a measure. In fact, it could be argued that, in the circumstances, Canada’s expectations with regard to the benefits it could derive from the concession could not be as high as for a product that posed no known risk to health. In such a context, in which it might be argued that, at best, Canada could anticipate a gradual decline in its chrysotile exports, an “upsetting of the competitive relationship” as a result of the measure in question would seem to be more difficult to establish. The situation is not one in which more or less promising prospects of market access disappear, as in the previous cases considered, but of a market in decline, as confirmed by the trend in the amounts produced and the number of countries which, in the last 25 years, have restricted or banned the use of chrysotile and by the increasing reliance on substitute products. In these circumstances, the Decree could not be considered to have upset the competitive relationship between chrysotile and products containing it, on the one hand, and substitute fibres and products containing them on the other.

However, as this element would be difficult to quantify and in the light of our reasoning in paragraphs 8.272 and 8.282 above, we consider it more appropriate to discuss the circumstances which led France to adopt a measure justified under Article XX in the context of the examination of the question of whether Canada could reasonably have anticipated the French measure. These elements seem to relate more particularly to the measure at issue. We will therefore now proceed to examine the criteria developed by previous panels.

According to the criteria developed in previous cases, (a) the Decree must have had the effect of upsetting the competitive relationship between Canadian asbestos and products containing it, on the one hand, and substitute fibres and products containing them, on the other, and (b) the Canadian Government must not have been able reasonably to anticipate the Decree when it was negotiating the various tariff concessions covering the products concerned.

At this stage, the Panel finds it appropriate to consider that in view of the type of measure in question the “upsetting of the competitive relationship” can be assumed. By its very nature, an import ban constitutes a denial of any opportunity for competition, whatever the import volume that existed before the introduction of the ban. We will therefore concentrate on the question of whether the measure could reasonably have been anticipated by the Canadian Government at the time that it was negotiating the various tariff concessions covering the products concerned.

With regard to the tariff concessions to be taken into account in assessing the “predictability” of the measure, the Panel notes that Canada refers expressly to the concessions made by France in 1947, by the European Economic Community in 1962 and by the European Communities at the end of the Uruguay Round. Canada notes, in particular, that the concessions included in the 1962 Protocol have been renewed until now. The European Communities do not dispute the information submitted by Canada, but recall that the Panel in Japan – Film pointed out that the establishment of a case based on expectations from rounds concluded 18 or 30 years ago might be difficult. The EC consider that Canada should provide detailed explanations of why it could legitimately have expected that France would not adopt measures restricting the use of any asbestos product after the Uruguay Round negotiations, given the growing scientific evidence that all types of asbestos were carcinogenic to humans.

With regard to the factors to be taken into account determining whether the measure in question could reasonably have been anticipated, previous panels found that a number of elements were not relevant. We consider it necessary to assess their applicability in relation to the circumstances of the present case.

(a) First of all, we note that the reports in Japan – Film and EEC – Oilseeds concluded that a specific measure could not be considered foresee-
able solely because it was consistent with or a continuation of a past general government policy. However, we note that, in contrast to the two cases mentioned above, France had already developed a specific policy in response to the health problems created by asbestos before the adoption of the Decree. This factor must certainly be taken into account in our analysis.238

(b) The Panel in Japan – Film, also concluded that it would not be appropriate to charge the United States with having reasonably anticipated all GATT-consistent measures. Consequently, we do not consider that Canada reasonably anticipated all GATT-consistent measures, or even possible measures justifiable under Article XX.

(c) Finally, insofar as the Decree postdates the most recent tariff negotiations, we could apply the presumption applied by the Panel in Japan – Film, according to which normally Canada should not be considered to have anticipated a measure introduced after the tariff concession had been negotiated. However, we do not consider such a presumption to be consistent with the standard of proof that we found to be applicable in paragraph 8.272 above in the case of an allegation of non-violation nullification concerning measures falling under Article XX of the GATT 1994.

Moreover, the circumstances of the present case seem to us to be different from the situation envisaged in Japan – Film. In that case, the measures in question concerned the organization of the Japanese domestic market. They were therefore economic measures of a kind that a third country might find surprising and, accordingly, difficult to anticipate. Here, it is a question of measures to protect public health under Article XX(b), that is to say, measures whose adoption is expressly envisaged by the GATT 1994. We therefore consider that the presumption applied in Japan – Film is not applicable to the present case.

8.292 Canada mentions, first of all, the concessions made by France in 1947 and by the EEC in 1962. However, we consider that in view of the time that elapsed between those concessions and the adoption of the Decree (between 50 and 35 years), Canada could not assume that, over such a long period, there would not be advances in medical knowledge with the risk that one day a product would be banned on health grounds. For this reason, too, we also consider that the presumption applied in Japan – Film cannot be applied to the concessions granted in 1947 and 1962. Any other interpretation would extend the scope of the concept of non-violation nullification well beyond that envisaged by the Panel in Japan – Film. On the contrary, it is for Canada to present detailed evidence showing why it could legitimately expect the 1947 and 1962 concessions not to be affected and could not reasonably anticipate that France might adopt measures restricting the use of all asbestos products 50 and 35 years, respectively, after the negotiation of the concessions concerned. In the present case, the burden of proof must be all the heavier inasmuch as the intervening period has been so long. Indeed, it is very difficult to anticipate what a Member will do in 50 years time. It would therefore be easy for a Member to establish that he could not reasonably anticipate the adoption of a measure if the burden of proof were not made heavier.

8.293 In the present case, we do not consider that Canada has provided a detailed explanation of why it could reasonably expect France not to adopt measures restricting the use of any asbestos products 50 and 35 years respectively, after the concessions concerned.

8.294 We therefore now turn to the question of whether, at the conclusion of the Uruguay Round, Canada could reasonably have expected France (a) not to adopt a measure restricting the use of asbestos and (b) not to do so by introducing a total ban.
8.295 As we have found (in paragraph 8.28), the presumption applied by the Panel in Japan – Film cannot be applied to the present case. Unlike Canada, which claims that no recent scientific development could have made the measure foreseeable, we consider that there is evidence to show that regulations restricting the use of asbestos could have been anticipated. First of all, the hazardous nature of chrysotile has long been known. Thus, since 1977, chrysotile has been classified by the WHO as a category I carcinogen. In 1986, ILO Convention 162 required national legislators to make provision, whenever possible, for the replacement of asbestos or of certain types of asbestos or of products containing asbestos by other materials or products or the use of alternative technology, scientifically evaluated by the competent authority as harmless or less harmful. In 1990, the European Communities issued Directive 90/394/EEC providing for the replacement of asbestos. This Directive advocates the principle that a dangerous agent or procedure should be replaced by a harmless or less harmful agent or procedure insofar as one exists.

8.296 Moreover, in the light of the information submitted by the parties and the experts, we consider that the study of the diseases associated with the inhalation of asbestos is a field of science in which any possible conclusion would appear to be based on the observation of pathological cases day by day. In the Panel’s opinion, the numerous constraints under which the scientists must work, such as the need to limit themselves to a given number of individuals, the establishment of the history of exposure and the period over which the health effects of asbestos develop, make the impact of these studies largely relative. It is the accumulation of data that gradually establishes scientific certainty. This impression is confirmed by the experts consulted by the Panel whose opinions are based on a large number of studies, including recent ones. Consequently, even supposing that there has been no major recent discovery, Canada cannot argue that it could not anticipate the adoption of a measure restricting the use of asbestos.

8.297 On the other hand, the accumulation of international and Community decisions concerning the use of asbestos, even if it did not necessarily make it certain that the use of asbestos would be banned by France, could not do other than create a climate which should have led Canada to anticipate a change in the attitude of the importing countries, especially in view of the long-established trend towards ever tighter restrictions on the use of asbestos. We also note that the use of chrysotile asbestos was banned by Members of the WTO well before it was banned by France. Admittedly, in Japan – Film the Panel considered that the adoption in other Members’ markets of measures similar to the measures in question could not make the latter foreseeable. However, here again it was a question of commercial measures. We consider that in the present case the situation is different since it concerns public health and the competent international organizations have already taken a position on the question. The adoption, in an already restrictive context, of public health measures by other States, faced with a social and economic situation similar to that in France, creates an environment in which the adoption of similar measures by France, is no longer unforeseeable.

8.298 Moreover, as noted above, at the end of the Uruguay Round France already had in place a number of measures regulating the use of asbestos. These included, in particular, measures relating to the exposure of workers taken after asbestos was recognized as a carcinogen by the IARC (Decree 77-949 of 17 August 1977) and the adoption of ILO Convention 162, as well as for the purpose of implementing Community directives applicable. The Panel also notes that Decree 88-466 of 28 April 1988 on products containing asbestos had prohibited the use of chrysotile asbestos in the manufacture of certain products.240

8.299 In these circumstances, it is difficult for Canada to claim that it had legitimate expectations of maintaining or even developing its exports of chrysotile and products containing it. If there was any benefit for Canada, the Panel does not consider Canada to have presented a detailed justification to show that that benefit was anything other than precarious.

8.300 We thus conclude that at the conclusion of the Uruguay Round Canada could reasonably have anticipated that France might, in the short term, adopt more restrictive measures on the use of asbestos.
8.301 The question that remains to be decided is whether Canada could reasonably have anticipated the introduction of a ban on chrysotile asbestos by France. We consider that the scientific literature and the international regulations, as well as Members’ public policy\(^{231}\), gave numerous indications such as to create a context in which Canada could not reasonably not have anticipated that sooner or later chrysotile would be banned by France. We also recall that the legitimacy of measures justified under Article XX justifies a stricter burden of proof on the party invoking Article XXIII:1(b). In the present case, Canada has not presented a detailed justification in support of its claim that it could not reasonably have anticipated that France would, in the light of developments in the scientific evidence and the choices made by other Members, also decide to apply a total ban on asbestos on its market in the short term.

8.302 We note Canada’s argument according to which it could not reasonably have anticipated the adoption of the Decree by France insofar as France failed to display the necessary consistency in its approach to chrysotile and other hazardous products, such as lead and copper. We do not consider this a pertinent argument. Although it is true that lead and copper, like asbestos, are hazardous and that France has not yet taken any measure with respect to lead and copper, essentially this means that Canada cannot legitimately expect the lead market not to be the subject of some public health measure, even though France has not yet taken any such measure. It is our opinion that each Member is free to adopt the health policies it deems appropriate and to give each such policy the priority it deems necessary. To accept Canada’s argument would not only conflict with that principle by requiring an “all or nothing” approach impossible to realize in practice but would considerably facilitate the task of a Member invoking Article XXIII:1(b) and required to prove the existence of a legitimate expectation. Thus, it would be sufficient to identify a product which had not yet been regulated even though it was recognized as being hazardous. There would then be no need to provide detailed evidence of the existence of non-violation nullification or impairment.

8.303 In the present case, we do not consider that Canada has presented a detailed justification of why it could legitimately have expected that France would not adopt measures restricting the use of any asbestos product in a context in which measures to limit the use of asbestos were being proposed at the international level and when countries at the same level of social and economic development as France had already banned the use of chrysotile asbestos by the end of the Uruguay Round.

(c) Conclusion

8.304 On the basis of the above, we conclude that Canada has not established the existence of nullification or impairment of a benefit within the meaning of Article XXIII:1(b) of the GATT 1994 as a result of the application of the measure in question.

IX. CONCLUSIONS

9.1 In the light of the above, the Panel concludes as follows:

(a) On the basis of its findings in Section VIII.D above, the Panel concludes that the “prohibition” part of the Decree does not fall within the scope of the TBT Agreement. The part of the Decree relating to “exceptions” does fall within the scope of the TBT Agreement. However, as Canada has not made any claim concerning the compatibility with the TBT Agreement of the part of the Decree relating to exceptions, the Panel refrains from reaching any conclusion with regard to the latter.

(b) On the basis of its findings in Section VIII.E.2 above, the Panel concludes that chrysotile asbestos fibres as such and fibres that can be substituted for them as such are like products within the meaning of Article III:4 of the GATT 1994. Similarly, the Panel concludes that the
asbestos-cement products and the fibro-cement products for which sufficient information has been submitted to the Panel are like products within the meaning of Article III:4 of the GATT 1994.

(c) With respect to the products found to be like, the Panel concludes that the Decree violates Article III:4 of the GATT 1994.

(d) However, on the basis of its findings in Section VII.E.4 above, the Panel concludes that the Decree, insofar as it introduces a treatment of these products that is discriminatory under Article III:4, is justified as such and in its implementation by the provisions of paragraph (b) and the introductory clause of Article XX of the GATT 1994.

(e) Finally, the Panel concludes, on the basis of its findings in Section VIII.F above, that Canada has not established that it suffered non-violation nullification or impairment of a benefit within the meaning of Article XXII:1(b) of the GATT 1994.
NOTES

1 Pursuant to Article 15.3 of the Understanding on Rules and Procedures Governing the Settlement of Disputes (hereinafter the “Understanding”), “The findings of the final panel report shall include a discussion of the arguments made at the interim review stage”. The following section entitled “interim review” therefore forms part of our report’s findings.

2 The comments received on 4 July 2000 are dealt with where necessary in the sections on the comments from Canada and the European Communities below.

3 The Panel notes that, in the letter attached to Canada’s comments, the latter indicates that its request for review “is without prejudice to Canada’s position on all the aspects of the Panel’s Report”. In this connection, the Panel considers that, if it had misunderstood or misrepresented some of the factual aspects of the case in its findings, the parties would need the interim review stage in order to make the necessary corrections or clarifications because, unlike errors of law, errors of fact cannot usually be modified on appeal. The parties should take advantage of this last opportunity to rectify the factual assessments of the Panel otherwise the Panel could unnecessarily be at risk of being accused of not having made an objective evaluation of the facts. It might be claimed that the fact that a party does not inform the Panel of a factual error in its findings may be contrary to the obligation in Article 3.10 of the Understanding, which provides inter alia that “all Members will engage in these procedures [settlement of disputes] in good faith in an effort to resolve the dispute” (emphasis added).

4 For example, the Panel kept the words “in good faith” in para. 8.29 because the good faith requirement applies both to the person interpreting the text and the person having drafted it (see Ian Sinclair: The Vienna Convention on the Law of Treaties (1980, p. 119).

5 Hereinafter the “EC”.

6 Hereinafter the “Decree”.

7 The change in the numbering of paragraphs is due to the introduction of a paragraph on the Panel’s decision concerning a written submission and a request for a hearing made by a non-governmental organization during the interim review procedure.


9 In these findings, the word “asbestos” is used to describe all varieties of asbestos without distinction (see paras. 2.1 and 2.2 above for a description of the product). “The Panel notes that, although the only variety of asbestos referred to by Canada is chrysotile asbestos, Decree No. 96-1133 does not distinguish among the different varieties of asbestos. Only Article 2 on exceptions specifically mentions chrysotile fibres. Consequently, wherever necessary, the Panel will indicate whether it is referring to asbestos in general or chrysotile in particular.


11 The full text of the Decree is attached to this report as Annex I.

12 The preambular paragraphs of the Decree have not been included here. The full text of the Decree is attached as Annex I to this Report.

13 Ibid.

14 Articles 8 and 9 of the Decree have not been reproduced. The full text of the Decree is attached as Annex I to this report.

15 The full text of the claims by the parties can be found in section III.A above.

16 Hereinafter the “TBT Agreement”.

17 Hereinafter the “EC”.

18 These comments can be found in paras. 5.2 to 5.16 above.

19 The letter contains the following note: “See the reports of the Appellate Body in European Communities – Measures Concerning Meat and Meat Products (Hormones) (WT/DS26-DS48/AB/R, para. 147 (‘… in disputes involving scientific or technical issues, neither Article 11.2 of the SPS Agreement, nor Article 13 of the DSU prevents panels from consulting with individual experts. Rather, both the SPS Agreement and the DSU leave to the sound discretion of a panel the determination of whether the establishment of an expert review group is necessary or appropriate.’) and in Argentina – Measures Affecting Imports of Foot-
wear, Textiles, Apparel and Other Items (WT/DS56/AB/R), para. 84 (‘… Article 13 of the DSU enables a panel to seek information and technical advice as it deems appropriate in a particular case, and (…) the DSU leaves ‘to the sound discretion of a panel the determination of whether the establishment of an expert review group is necessary or appropriate’."

20 The letter contains the following note: “See the Report of the Appellate Body in Guatemala – Anti-Dumping Investigation Regarding Portland Cement from Mexico (WT/DS60/AB/R), paras. 65 and 66.”

21 See paras. 6.1-6.3 above.


23 Hereinafter “the WTO Agreement”. Reference will also be made in these findings to the Marrakesh Agreement Establishing the World Trade Organization without its Annexes using the words “Agreement Establishing the WTO”. The words “WTO Agreements” refer to the Agreements annexed to the Agreement Establishing the WTO.


27 The parties’ arguments are set out in detail in Section III above.

28 The wording of the definition is given in paragraph 8.18 above.


30 The Panel also notes the importance attached by the Appellate Body to the principle of effectiveness (ut res magis valeat quam pereat) concerning the interpretation of the provisions of the WTO Agreement in several cases (see, for example, United States – Standards for Reformulated and Conventional Gasoline, adopted on 20 May 1996, WT/DS60/AB/R (hereinafter “United States – Gasoline”), op. cit., pp.18 and 23; Guatemala – Cement, op. cit., para. 75; Argentina – Safeguards, op. cit., para. 88. The Panel also notes in the Reports of the Commission to the General Assembly, Yearbook of the International Law Commission, 1966, Volume II, A/CN.4/SER.A/1966/Add.1, p. 219, that the International Law Commission indicates that: “[…] in so far as the maxim ut res magis valeat quam pereat reflects a true general rule of interpretation, it is embodied in Article [31], paragraph 1, which requires that a treaty shall be interpreted in good faith in accordance with the ordinary meaning to be given to its terms in the context of the treaty and in the light of its object and purpose. When a treaty is open to two interpretations one of which does and the other does not enable the treaty to have appropriate effects, good faith and the object and purposes of the treaty demand that the former interpretation should be adopted.” [Italics in the original.]

31 See the report of the Appellate Body in United States – Measures Affecting Imports of Woven Wool Shirts and Blouses from India, adopted on 23 May 1997, WT/DS33/AB/R (hereinafter “United States – Shirts and Blouses”), p. 14 and Section VIII.E.1(b) below.

32 The Panel is not, however, obliged to base itself solely on the arguments of the parties. See the report of the Appellate Body in EC Measures Concerning Meat and Meat Products (Hormones), adopted on 13 February 1998, WT/DS26/AB/R, WT/DS48/AB/R (hereinafter “European Communities – Hormones”), para. 156: “Panels are inhibited from addressing legal claims falling outside their terms of reference. However, nothing in the DSU limits the faculty of a panel freely to use arguments submitted by any of the parties – or to develop its own legal reasoning – to support its own findings and conclusions on the matter under its consideration. A panel might well be unable to carry out an objective assessment of the matter, as mandated by Article 11 of the DSU, if in its reasoning it had to restrict itself solely to arguments presented by the parties to the dispute.”

33 With regard to the provisions which the Panel must take into account when considering the Decree, the Panel is of course limited by the provisions in its terms of reference (see the Report of the Appellate Body in European Communities – Bananas, op. cit., para. 141).

34 This is precisely why our analysis lays greater emphasis on the characteristics of the product than on the process and production methods.

35 See footnote 22 above on application of the principle of interpretation *ut res magis valeat quam pereat*.

36 See the Report of the Appellate Body in European Communities – Hormones, op. cit., para. 164.

37 Although the title of Annex 1A is “Multilateral Agreements on Trade in Goods” (emphasis added), we note that the Agreement on Agriculture uses the word “products”, as do the Agreement on Textiles and Clothing and other Agreements in Annex 1A. The scope of Annex 1A therefore covers trade in “products”. See also the Report of the Panel in United States – Restrictions on Imports of Tuna, unadopted, DS21/R, 3 September 1991, para. 5.11: “...Article III covers only measures affecting products as such”.

38 The definition in para. 3.5.1 of the ISO/IEC Guide General Vocabulary Concerning Standardization and Related Activities reads as follows:

“3.5.1 technical regulation: regulation that provides technical requirements, either directly or by referring to or incorporating the content of a standard, technical specification or code of practice” (in bold in the original, note omitted)

39 In relation to the foregoing, we also note that the standard notification forms under the TBT Agreement contain a section to be completed by a description of the product affected by the notification, which in our view seems to confirm the second meaning we have given for the word “product”.


“The Commission, by heading the article “General rule of interpretation” in the singular and by underlining the connexion between paragraphs 1 and 2 [of Article 31] and again between paragraph 3 and the two previous paragraphs, intended to indicate that the application of the means of interpretation in the article would be a single combined operation. […] the Commission desired to emphasize that the process of interpretation is a unity and the provisions of the article form a single, closely integrated rule.”


45 “Desiring, therefore to encourage the development of such international standards and conformity assessment systems;”.

46 “Desiring, however to ensure that technical regulations and standards, including packaging, marking and labelling requirements, and procedures for assessment of conformity with technical regulations and standards do not create unnecessary obstacles to international trade;”.

47 Although technically these are additional ways of interpreting Article 32, we note that the *preparatory work* confirms this objective. Document TRE/W/21 of 17 January 1994 notes that the draft Code (Spec(72)) elaborated in the course of the preparatory work for the Tokyo Round refers to the preparation and adoption of “mandatory standards”. The use of the words “mandatory standards” (one of the definitions of the word “standard” in English is “a document specifying nationally or internationally agreed properties for manufactured goods, principles for procedure, etc.”. The New Shorter Oxford English Dictionary (1993), p. 3028) implies that the purpose of the Agreement was to ensure a product’s conformity with the characteristics of other products marketed.

48 “Recognizing that no country should be prevented from taking measures necessary […] for the protection of human […] life or health […] subject to the requirement that they are
not applied in a manner which would constitute a means of arbitrary or unjustifiable discrimination between countries where the same conditions prevail or a disguised restriction on international trade […]” (identical wording to that in Article XX of the GATT 1994, emphasis added).

49 We note in this connection that the preparatory work on the Agreement on Technical Barriers to Trade in the Tokyo Round show that the TBT Agreement that should have emerged from the Tokyo Round was already seen as being a development of the existing rules of the GATT, notably Article XX. See for example the extract from document MTN/3E/W/26, October 1974, quoted in paragraph 7 of document TRE/W/21, 17 January 1994.


52 According to the definition in Annex 1.1 to the TBT Agreement, the criteria for placing a product on the market must concern its characteristics or the related processes or production methods, including the applicable administrative provisions.


54 See the preamble to the Decision on Notification Procedures, adopted by the Trade Negotiations Committee on 15 December 1993.

55 See the second paragraph in Section I of the Decision on Notification Procedures, adopted by the Committee on Trade Negotiations on 15 December 1993, as annexed to the Final Act embodying the results of the Uruguay Round of multilateral trade negotiations, done at Marrakesh on 15 April 1994.

"Members recall their undertakings set out in the Understanding Regarding Notification, Consultation, Dispute Settlement and Surveillance adopted on 28 November 1979 (BISD 26S/210). With regard to their undertaking therein to notify, to the maximum extent possible, their adoption of trade measures affecting the operation of the GATT 1994, such notification itself being without prejudice to views on the consistency of measures with or their relevance to rights and obligations under the Multilateral Trade Agreements and, where applicable, the plurilateral Trade Agreements, Members agree to be guided, as appropriate, by the annexed list of measures. Members therefore agree that the introduction or modification of such measures is subject to the notification requirements of the 1979 Understanding.” (Emphasis added)

Paragraph 3 of the Understanding Regarding Notification, Consultation, Dispute Settlement and Surveillance, adopted on 28 November 1979 (BISD 26S/210) provides that:

“Contracting parties moreover undertake, to the maximum extent possible, to notify the CONTRACTING PARTIES of their adoption of trade measures affecting the operation of the General Agreement, it being understood that such notification would of itself be without prejudice to views on the consistency of measures with or their relevance to rights and obligations under the General Agreement.”

56 See Canada’s arguments, section III.C.1(a) above, in particular paras. 3.246 and 3.260-3.261.

57 These criteria are by no means exhaustive.

58 According to the definition in Annex 1.1 of the TBT Agreement, the criteria for placing a product on the market must concern its characteristics or the related processes or production methods, including the applicable administrative provisions.

59 See the reply by the EC to question 2 in the questions by the Panel at the second substantive meeting, paras. 220-221.

60 See R. Guillien and J. Vincent: Lexique de termes juridiques (1981), p. 3: in civil law “the subsidiary follows the principal in the sense that the principal good transmits its legal characterization to the good attached to it”. (WTO translation). See also Black’s Law Dictionary (1990), p. 14, regarding the criminal law applicable in certain Members.

61 For example, Canada did not claim that the WTO Agreement had been violated by the way in which France administered the exceptions. Moreover, the Panel notes that they are exceptional and temporary exemptions. They will disappear as and when reliable and effective substitute products are developed.
"The burden of proof rests upon the party, whether complaining or defending, who asserts the affirmative of a particular claim or defence. If that party adduces evidence sufficient to raise a presumption that what is claimed is true, the burden then shifts to the other party, who will fail unless it adduces sufficient evidence to rebut the presumption".


Ibid., para. 128-129.

Ibid., para. 129.

The arguments of the parties are set out in detail in Section III above.

The arguments of the parties are set out in detail in Section III above.

It is only if the Panel rejects Canada’s first interpretation that the Decree comes in part under Article III:4 and in part under Article XI:1 that Canada considers that the whole of the Decree should fall under Article XI:1 or, if the Panel also rejects this approach, Article III:4.


"In the same way as", "to the same extent as" are among the alternative meanings for the word “comme” in the French text (Le Nouveau Petit Robert, op. cit., p. 411). [In the English text the word is “and”].

Emphasis added. In the place of “and” and “comme”, the Spanish version uses the conjunction “y” (“et” in French).

The Panel gave the grounds for its decision in para. 5.10 as follows:

“The fact that Section 337 is used as a means for the enforcement of United States Patent Law at the border does not provide an escape from the applicability of Article III:4; the interpretative Note to Article III states that any law, regulation or requirement affecting the internal sale of products that is enforced in the case of the imported product at the time or point of importation is nevertheless subject to the provisions of Article III. Nor could the applicability of Article III:4 be denied on the ground that most of the procedures in the case before the Panel are applied to persons rather than products, since the factor determining whether persons might be susceptible to Section 337 proceedings or federal district court procedures is the source of the challenged products, that is whether they are of United States origin or imported. For these reasons, the Panel found that the procedures under Section 337 come within the concept of ‘laws, regulations and requirements’ affecting the internal sale of imported products, as set out in Article III of the General Agreement.”


We note in this connection that the Report of the Panel in Canada – Import, Distribution and Sale of Certain Alcoholic Drinks by Provincial Marketing Agencies, adopted on 18 February 1992, BISD 39S/27, in para. 5.28 specifically examines the question of whether minimum prices fell under Article XI:1 or Article III:4 and noted the following:

“The Panel first examined whether the minimum prices fell under Article XI:1 or Article III:4. The Panel noted that, according to the Note Ad Article III, a regula-
tion is subject to the provisions of Article III if it ‘applies to an imported product and to the like domestic product’ even if it is ‘enforced in case of the imported product at the time or point of importation’. The Panel found that, as the minimum prices were applied to both imported and domestic beer, they fell, according to this Note, under Article III.”

81 The arguments of the parties are set out in detail in Section III above.
84 Ibid., p. 22.
86 Bearing in mind Canada’s reply to question No. 15 by the Panel, paras. 74-76, we conclude that Canada does not raise the issue of the likeness of non-fibrous products, whether substitute products or products containing them, as compared to chrysotile or products containing it.
87 See the EC’s reply to question 30 by the Panel, Annex II, para. 113 below.
88 See in particular the reports of the Appellate Body in Australia – Salmon, op. cit. and Japan – Agricultural Products, op. cit., paras. 136.
89 We are aware that Canada’s claims (see para. 3.1 above) state that the Decree is incompatible with Article III-4 of the GATT 1994 because it “favours the national industry of products like chrysotile cement and chrysotile-cement products …” Moreover, our terms of reference are not confined to PVA, cellulose and glass fibres (see WT/DS135/3). We note, however, that Canada does not ask for findings on other fibres. We therefore limit our findings to the products for which we consider Canada had requested findings, namely, PVA, cellulose and glass fibres.
90 See Annex II, Canada’s reply to question 15 by the Panel, paras. 74 et seq.,
91 See the Report of the Appellate Body in European Communities – Bananas, op. cit., para. 136, where it is mentioned that the potential export interest by the United States could not be excluded even though at that time the United States did not export bananas from their territory to the EC. See also the Report of the Panel in Indonesia – Certain Measures Affecting the Automobile Industry, adopted on 23 July 1998, WT/DS54, 55, 59, 64/R, para. 14.113.
94 See, for example, the arguments of the parties on the concept of “like products”, Section III.C. 2(b)(i) above.
95 In this connection, we note the observation by one of the experts consulted by the Panel that the characteristics of some artificial substitute fibres can be altered by industry if necessary (Dr. Henderson, para. 5.383 above).
96 Regarding the chemical composition of asbestos, see the table provided by the EC in reply to question 29 by the Panel, para. 106.
99 In this connection, we note that the English text of the Report on Border Tax Adjustments, op. cit., para. 18, uses the words “properties” and “quality”.
100 See the EC’s arguments in para. 3.44 above concerning the Report of the Panel in EEC – Measures on Animal Feed Proteins, adopted on 14 March 1978, BISD 255/49. The Panel does not consider that the factual elements peculiar to this affair and the conclusions of the Panel (see para. 4.2) make it possible to draw the conclusions suggested by the EC in the present case.
101 Adopted on 10 November 1987, BISD 34S/84, hereinafter “Japan – Alcoholic Beverages (1987)”.
102 Ibid., para. 5.6.
103 We note that the EC draw attention to the risk posed by chrysotile fibres not only in connection with the properties, nature and quality criterion but also in relation to the end-
use and the tastes and habits of consumers. In this subsection, however, we examine the
criterion of the risk of a product inasmuch as the EC, in its arguments, referred to this
criterion mainly in relation to the criterion of the properties, nature and quality of asbestos
fibres. The conclusions of our examination, however, will apply to all the circumstances in
which the EC refer to the risk of a product in relation to the determination of likeness
within the meaning of Article III:4 of the GATT 1994.

104 Mesothelioma is a form of pleural cancer. See for example the description by Dr.
Henderson, para. 5.29 above.

105 See the report of the Panel in Canada – Measures on Export of Unprocessed Herring and

106 See the discussion on the concept of effectiveness, footnote 22 above.

107 See in particular Argentina – Safeguards, op. cit; Brazil – Desiccated Coconut, op. cit.

108 See para. 8.112 above.

109 See para. 8.125 above.

110 These cases are cited in para. 3.446 above and in the corresponding footnotes.

111 In the cases cited by the EC – to which Canada also refers – we do not find any legal
reasons relating to Article III:4 to justify excluding the criterion of consumers’ tastes and
habits. In Measures Affecting Alcoholic and Malt Beverages (adopted on 19 June 1992, BISD
395/206), para. 5.73, the panel indeed did not agree that low alcohol beer and high alcohol
beer were like products on the basis of consumers’ tastes and habits, but it did nevertheless
use this criterion. In EEC – Measures on Animal Feed Proteins (adopted on 14 March 1978,
BISD 255/49), para. 4.2, the panel did not exclude the applicability of consumers’ tastes,
noting:

"[...] In this case, such factors as the number of products and tariff items carrying
different duty rates and tariff bindings, the varying protein contents and the
different vegetable, animal and synthetic origins of the protein products before
the Panel – not all of which were subject to the EEC measures”.

In view of the terms used by the panel in this case, it does not seem that the panel in
principle rejected the relevance of consumers’ tastes and habits. In United States – Gasoline,
op. cit., even though the panel did not mention the criterion of consumers’ tastes and hab-
its in its conclusions in para. 6.9, in para. 6.8 it nonetheless recalled the applicability under
Article III:4 of the criteria summarized by the Working Party in Border Tax Adjustments, op.
cit., which include consumers’ tastes and habits. We have not found any reason either to
restrict the meaning of the word “consumers” solely to the end-consumers of chryso-
tile-cement products, i.e. individual consumers.


113 Hereinafter “Harmonized System” or “HS”.

114 Asbestos in its natural state falls in heading 25.24. Polyvinyl alcohol falls in heading
39.05, cellulose in 39.12 and glass fibre in 70.19. We note that in Japan – Alcoholic Beverages
(1987), the panel referred to the “Nomenclature of the Customs Cooperation Council (CCCN)
for the classification of goods in customs tariffs”. We also note that the Appellate Body in
Japan – Alcoholic Beverages op. cit. p. 24 reaffirmed that that tariff classification, if suffi-
ciently detailed, was a useful basis for confirming the likeness of products. We note that in
this case the parties based themselves on the tariff classification in the Harmonized Sys-
tem, and not on tariff bindings, which the Appellate Body urges should be used with cau-
tion. We do not consider however, that the particular circumstances of this case justify
that, when determining the likeness or absence of likeness of asbestos, PVA, cellulose and
glass fibres, overriding significance should be attached to the fact that the products fall in
different tariff headings of the HS.

115 In fact, the HS Explanatory Notes describe the products that fall within heading 68.11
(“Articles of asbestos-cement, of cellulose fibro-cement or the like”) as follows:

"This heading covers hardened articles consisting essentially of an intimate mix-
ture of fibres (for example, asbestos, cellulose or other vegetable fibres, synthetic
polymer, glass or metallic fibres) and cement or other hydraulic binders, the
fibres acting as strengthening agents. [...]"

The heading includes sheets of all sizes and thicknesses, obtained as described
above, and also articles made by cutting these sheets or by pressing, moulding
or bending them before they have set, e.g., roofing, facing or partition sheets and
tiles; sheets for making furniture; window sills; sign-plates, letters and num-
bers; barrier bars; corrugated sheets; reservoirs, troughs, basins, sinks; tubing joints; packing washers and joints; panels imitating carving; ridge tiles, gutters, window frames; flower-pots; ventilation or other tubing, cable conduits; chimney cowls, etc.
All these articles may be coloured in the mass, varnished, printed, enamelled, decorated, drilled, filed, planed, smoothed, polished or otherwise worked; they may also be reinforced with metal, etc."

116 The arguments of the parties are set out in detail in Section III above.
117 We note, incidentally, that Canada has not made any allegation concerning the less favourable treatment of Canadian chrysotile fibres as compared with domestic chrysotile fibres. Accordingly, we shall not make any finding in this respect.
118 See the Report of the Panel in United States – Sections 301-310 of the Trade Act of 1974, op. cit., para. 7.27.
119 The arguments of the parties are set out in detail in Section III above.
120 See the Report of the Appellate Body in United States – Gasoline, op. cit., pp. 14-15 and 17-18. In the present case, it is the part of the Decree relating to the prohibiting of the placement on the French market of chrysotile asbestos and products containing it that has been found contrary to Article III:4 of the GATT 1994 (see paras. 8.155 and 8.158 above). It is this same “measure” that we shall examine in the light of Article XX(b).
123 In this connection, we note that it is not for the Panel to examine on its own initiative the exceptions for which Article XX provides unless they have been invoked (see Report of the Panel in United States – Restrictions on Imports of Tuna, DS21/R (unadopted), 3 September 1991, BISD 39S/155, para. 5.27, citing the Report of the Panel in EEC – Regulation on Imports of Parts and Components, adopted 16 May 1990, 37S/132, para. 5.11). However, it is only necessary to determine that the measure in question satisfies the conditions of one of the paragraphs of Article XX and the introductory clause of the same article for that measure to be justified under Article XX of the GATT 1994.
127 Our approach does not affect the possibility of each Member determining what level of risk it wishes to assure. Guaranteeing a very low risk of exposure when the health risks are high might require strict measures, but when the health risk is low a very low level of exposure might be obtainable with less severe measures. This comment is made in full awareness of the fact that particularly serious risks can sometimes be neutralized by very simple means.
128 In this case, the justification for less severe measures would not be the existence of less inconsistent measures that meet the objectives originally set by the EC, but the fact that the risk was actually lower than the EC had estimated.
129 See Section VIII.E.1(b) above.
130 Op. cit., pp. 15-16: “We acknowledge that several GATT 1947 and WTO panels have required such proof of a party invoking a defence such as those found in Article XX(12) or Article XI:2(c)(i)(12) to a claim of violation of a GATT obligation, such as those found in Articles I:1, II:1, III or XI:1. Articles XX and XI:2(c)(i) are limited exceptions from obligations under certain other provisions of the GATT 1994, not positive rules establishing obligations in themselves. They are in the nature of affirmative defences. It is only reasonable that the burden of establishing such a defence should rest on the party asserting it.”
133 Hereinafter the “SPS Agreement”.

922
See, in particular, Articles 2.2, 3 and 5 of the SPS Agreement.

In European Communities – Hormones, op. cit., para. 115, the Appellate Body itself expressed its reluctance to adopt within the context of the SPS Agreement a standard of review not clearly rooted in the text of the SPS Agreement, for fear of disturbing the balance between the rights and obligations negotiated.

Report of the Appellate Body in Japan – Agricultural Products, op. cit., para. 129. At this point, we recall that the experts were selected in consultation with the parties and that the latter did not challenge the appointment of any of them, although they reserved the right to comment on their statements. With respect to the various stages in the selection and consultation of the experts, see Section V.A and B.


See paras. 8.173-176.

With regard to the group of pathologies that asbestos can cause, see Dr. Henderson, para. 5.28.

See Annex II, Canadian reply to the EC’s question No. 4, para. 180.

See Annex II, Canadian replies to the EC’s questions nos. 1-4, paras. 175-181.

Since 1977 by the IARC (see List of Agents Carcinogenic to Humans, Overall Evaluations of Carcinogenicity to Humans, Monographs of the International Agency for Research on Cancer, Volumes 1-63), see also WHO, IPCS Environmental Health Criteria (203) on Chrysotile, Geneva (1998), cited in para. 5.584 above. On the development of knowledge of the risks associated with asbestos, see Dr. Henderson, para. 5.595.

See, in particular, Dr. Henderson, paras. 5.29 to 5.34; 5.142 to 5.165; Dr. Infante, paras. 5.267, 5.290-5.298; Dr. de Klerk, para. 5.288.

See, for example, the ratios suggested by Dr. Henderson, paras. 5.103, 5.141, 5.415, 5.589 and his remarks, paras. 5.265-5.266; see also Dr. de Klerk, para. 5.264; Dr. Infante, paras. 5.267-5.268 and Annex VI, para. 19 of the transcript of the meeting with experts.

Dr. Henderson, meeting with experts, Annex VI, para. 182.

See the comments made by Dr. Henderson, paras. 5.153-5.157 concerning the link between fibrosis and lung cancer.

See the replies of the experts to the Panel’s question 1.(b), paras. 5.196-5.209.

See Dr. Henderson, paras. 5.176, 5.183; Dr. de Klerk, para. 5.185.

In reply to a question by Canada (see Annex II, para. 167) the EC indicated that the question of the exposure associated with working intermittently with material such as chrysotile-cement had been analysed in the report of the Institut national de la science et de la recherche medicale (INSERM) entitled Rapport sur les effets sur la santé des principaux types d’exposition à l’amiante (Report of the health effects of the main types of exposure to asbestos, INSERM joint report, 1997, pp. 193-214). The EC appended to their first written submission to the Panel exposure values measured during such working (see Note de presentation des orientations du Conseil supérieur de prévention des risques professionnels, 3 July 1995). They also give the example of a roofing worker using a grinder in the open air to repair corrugated roof sheeting made of asbestos-cement and exposed to a peak level of 41f/ml. In this connection, the Panel also notes the reference by Dr. Henderson, in para. 5.199 above, to a study by Kamugai S, Nakachi S, Kurumatani N, et al. Estimation of Asbestos Exposure Among Workers Repairing Asbestos Cement Pipes Used for Conduits, Sangkyo Igaku 1993; 35.

By "DIY enthusiast" (bricoleur) the Panel means someone who does small repair and renovation jobs (Le Nouveau Petit Robert (1994), p. 261) without engaging in these activities in a professional capacity.


See Decree 96-98 of 7 February 1996. The European Communities, in para. 3.134, note, for example, that in the case of a handsaw application of the ISO standard leaves the worker exposed to a level 30 times in excess of the maximum limit of 0.1 f/ml. In this connection, the Panel notes that the experts agree that at least some of the fibres released during operations on products containing chrysotile present the same carcinogenicity as chrysotile fibres not incorporated in cement (see Dr. de Klerk, para. 5.220; Dr. Henderson, paras. 5.221-5.224; Dr. Infante, paras. 5.225-5.226).

See Full Public Report: Chrysotile Asbestos – Priority Existing Chemical No. 9, National Industrial Chemicals Notification and Assessment Scheme (NICNAS), National Occupa-
tion Health and Safety Commission (NOHSC), Australia 1999 (hereinafter “NICNAS 99”), cited by Dr. Henderson, which shows that exposure is spreading. Originally confined to workers in the traditional industries, exposure to asbestos now extends to products and the domestic and external environment (see para. 5.179). Similar observations have been made in the United Kingdom with regard to the patterns of exposure and the resultant diseases (para. 5.180).

See the example of the fireman and the lecturer whose exposure was not connected with their occupations, given by Dr. Henderson, para. 250 of the transcript of the meeting with experts.

Concerning the opinion of the experts with regard to the absence of a threshold, see paras. 5.306 et seq. and, more particularly, Dr. Infante, para. 5.315.

The absence of a study on the impact of chrysotile on building workers was confirmed by the scientists consulted by the Panel (see Dr. Infante, para. 137; Dr. Henderson, para. 140; Dr. Musk, para. 202 of the transcript of the meeting with experts, Annex VI). Concerning Canada’s arguments and the comments of the experts, see the discussion of the relevance of the study of Charleston textile factory workers (hereinafter “Charleston study”), see footnote 150) as compared with the study of Quebec asbestos mine and mill workers (hereinafter “Quebec workers study”), see footnote 151), paras. 135–153 of the transcript of the meeting with experts, Annex VI.


See the comments of Dr. Infante on the Quebec workers study by McDonald et al, para. 19 of the transcript of the meeting with experts, Annex VI, and those of Dr. Henderson, paras. 5.118 and 5.158–5.162 above.


See Dr. Henderson, paras. 59, 89 and 101 of the transcript of the meeting with experts, Annex VI. We note that Canada has not formulated a specific request with regard to friction products, such as car brakes, for example.

INSERM: Rapport sur les effets sur la santé des principaux types d’exposition à l’amiante, INSERM joint report, 1997. This report was submitted by the European Communities as one of the underpinnings of the measure adopted by France.

See Canada’s arguments, paras. 3.279 et seq above.

BISD 375/200, para. 75.


See Dr. Henderson, para. 5.29 and the transcript of the meeting with experts, Annex VI, end of para. 182.

See Dr. Infante, para. 5.183; Dr. de Klerk, para. 5.185.

See Dr. Infante for the United States (para. 161 of the transcript of the meeting with experts); Dr. de Klerk and Dr. Henderson for Australia (paras. 222 and 225, respectively, of the transcript of the meeting with experts).

See Dr. Henderson, paras. 5.174–5.181; Dr. Infante, paras. 5.182-5.183, 5.190.

See Dr. Henderson, para. 5.312; Dr. Infante, paras. 5.313-5.315.

See Dr. de Klerk, para. 5.317; Dr. Henderson, para. 5.318; Dr. Infante, paras. 5.321-5.323; Dr. Musk, para. 5.324.

See, for example, Dr. Infante, para. 5.304.

See, in particular, Annex II, reply by the EC to Canada’s question No. 6, para. 168

See Annex II, reply by the EC to Canada’s question No. 5, para. 167.

The European Communities refer to Article 5.6 of the Agreement on the Application of Sanitary and Phytosanitary Measures. They consider that the principle incorporated in
that provision should also be applied in the context of Article XX.

177 In this respect, we note that the experts likewise did not mention any alternatives other than controlled use. Moreover, in the light of the approach defined by the Appellate Body in *Australia – Salmon*, op. cit. (see Section VIII E.1.(b) above), we consider that we do not need to ascertain whether other measures were possible.


179 See *United States – Gasoline*, op. cit., paras. 6.26 to 6.28.

180 See *Peto et al: Continuing Increase in Mesothelioma Mortality in Britain, The Lancet*, volume 345, 535-539 (1995). See also Dr. Henderson, citing EHC 203, paras. 5.365-5.368; Dr. Infante, paras. 5.351-5.352 and 5.369.

181 See the comments of the experts on controlled use and its feasibility, paras. 5.335-5.373.

182 See, for example, ISO 7337.


184 See *United States – Gasoline*, op. cit., para. 6.22. The Panel also notes that XX(b) does not impose the same constraints as Article 3.3 of the SPS Agreement on Members wishing to apply measures that involve a level of protection higher than that which would be obtained with measures based on the relevant international standards.

185 Dr. Henderson, paras. 5.336-5.341.

186 Ibid., paras. 5.337-5.339.

187 Dr. Infante, para. 5.343 referring to an opinion given by the United States Occupational Safety and Health Administration (OSHA). See also Dr. Infante, paras. 5.358-5.362.

188 Dr. Henderson, paras. 5.355-5.357.

189 In any event, in the light of the opinion of one of the experts (Dr. Henderson, para 5.658 and footnote), the sectors in which effective controlled use is feasible appear to be extremely few and concern applications for which, at this stage, there are no chrysotile substitutes, an area in which the Decree provides for exceptions to the prohibition on the use of chrysotile.

190 We note that the experts’ comments relate to countries with a level of economic development equivalent to that of France and with similar administrative resources.

191 See the experts’ comments on this subject: Dr. de Klerk, para. 5.335; Dr. Henderson, para. 5.341; Dr. Infante, para 5.343.

192 See the studies cited by Dr. Infante, para. 5.304.

193 See the examples of the lecturer and the fireman mentioned by Dr. Henderson during the meeting with experts, para. 250, Annex VI.

194 See Dr. Infante, para. 5.315.

195 See IARC classification, op. cit.

196 See Dr. de Klerk, paras. 5.375-5.377; Dr. Henderson, para. 5.385; Dr. Infante, para. 5.388.

197 See Dr. Musk, para. 5.390.


199 The Panel considers that what is prohibited by the introductory clause to Article XX is a particular form of discrimination (that which is arbitrary or unjustifiable between countries where the same conditions prevail) and not all forms of discrimination. If the measure is not discriminatory in general in its application, then a fortiori it cannot constitute arbitrary or unjustifiable discrimination between countries where the same conditions prevail. See also the Report of the Panel in *United States – Imports of Certain Automotive Spring Assemblies*, adopted on 26 May 1983, BISD 305/107 (hereinafter “United States – Automotive Springs”), para. 55.


201 Ibid.


> “[The Panel] furthermore felt that the United States action should not be considered to be a disguised restriction on international trade, noting that the United States prohibition of imports of tuna and tuna products from Canada had been taken as a trade measure and publicly announced as such. The Panel therefore considered it appropriate to examine further the United States import prohibition of tuna and tuna produce from Canada in the light of the list of specific
types of measures contained in Article XX and notably in Article XX(g).”

203 The Report of the Panel in United States – Automotive Springs, op. cit., states in para. 56:

“The Panel then considered whether or not the exclusion order was “applied in a manner which would constitute … a disguised restriction on international trade”. The Panel noted that the preamble of Article XX made it clear that it was the application of the measure and not the measure itself that needed to be examined. Notice of the exclusion order was published in the Federal Register and the order was enforced by the United States customs at the border. The Panel also noted that the ITC proceedings in this particular case were directed against the importation of automotive spring assemblies produced in violation of a valid United States patent and that, before an exclusion order could be issued under Section 337, both the validity of the patent and its infringement by a foreign manufacturer had to be clearly established. Furthermore, the exclusion order would not prohibit the importation of automotive spring assemblies produced by any producer outside the United States who had a licence from Kuhlman Corporation to produce these goods. Consequently, the Panel found that the exclusion order had not been applied in a manner which constituted a disguised restriction on international trade.”

204 United States – Gasoline, op. cit., p. 25.
207 Ibid. Although this approach was developed in relation to Article III:4 of the GATT 1994, we see no reason why it should not be applicable in other circumstances where it is necessary to determine whether a measure is being applied for protective purposes.
208 In this respect, we consider the EC to have satisfied the burden of proof as regards the introductory clause of Article XX. We shall therefore treat Canada’s argument as an attempt to rebut the presumption established in favour of the EC.
209 See Canada’s arguments, para. 3.27. Canada refers to a study entitled L’amiante dans l’environnement, ses conséquences et son avenir, Office parlementaire d’évaluation des choix scientifiques et technologiques, National Assembly No. 329/Senate No. 41, p. 57.
210 See, in particular, the reply of the European Communities to the Panel’s question No. 30, paras. 113-115, Annex II.
211 The arguments of the parties are set out in detail in Section III above.
212 Adopted on 22 April 1998, WT/SD44/R, hereinafter “Japan – Film”.
213 Ibid., para. 10.36.
215 Adopted on 16 November 1962, BISD 115/95.
216 L/6053 (1986), unadopted.
217 This remark is made without prejudice to Canada’s allegations based on Article XXIII:1(a) (see Section VIII.A.2(a) above).

“… the recognition of the legitimacy of an expectation relating to the use of production subsidies therefore in no way prevents a contracting party from using production subsidies consistently with the General Agreement, it merely delineates the scope of the protection of a negotiated balance of concessions.”

We also note that the Appellate Body in United States – Gasoline, op. cit., p. 25, stated, with regard to the application of Article XX, that “the measures falling within the particular exceptions must be applied reasonably, with due regard both to the legal duties of the Party claiming the exception and the legal rights of the other Parties concerned”.
219 See Japan – Film, para. 10.50, footnote 1214.
221 In this connection, we note that in the context of the examination of a United States measure granted a waiver under Article XXV of the GATT 1947, the Panel in United States
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– Restrictions on the Importation of Sugar and Sugar-containing Products applied under the 1955 Waiver and under the Headnote to the Schedule of Tariff Concessions, adopted on 7 November 1990, BISD 375/228, noted, in para.5.21, that:

"... Article XXIII:1(b) applies whether or not the measure at issue conflicts with the General Agreement and that therefore, the question of whether a measure inconsistent with Article XI:1 remains inconsistent with the General Agreement even if covered by a waiver cannot, by itself, determine whether it nullifies or impairs benefits accruing under the General Agreement within the meaning of that provision".

222 In this respect, we note that panel practice cannot be regarded as falling within Article 31.3(a) or (b) of the Vienna Convention (see Article 3.2 of the Understanding, Article IX: 2 of the Agreement Establishing the WTO and the Report of the Appellate Body in Japan – Alcoholic Beverages, op. cit., pp.13-5). Moreover, the GATT 1947 and WTO panels have never discussed the situation with respect to a measure designed to protect public health. All previous cases have concerned situations in which a measure of a purely commercial nature was adopted pursuant to the negotiation of a concession.


226 Op. cit., para. 10.36. Referring to the Report of the Panel in EEC – Oilseeds, footnote 1205, the Panel in Japan – Film noted the following:

"In EEC – Oilseeds, the United States stated that it concurred in the proposition that non-violation nullification or impairment should remain an exceptional concept. Although this concept had been in the text of Article XXIII of the General Agreement from the outset, a cautious approach should continue to be taken in applying the concept. EEC – Oilseeds, BISD 375/S/86, para. 114. The EEC in that case stated that recourse to the “non-violation” concept under Article XXIII:1(b) should remain exceptional, since otherwise the trading world would be plunged into a state of precariousness and uncertainty. Ibid., para. 113.”


228 In Thailand – Cigarettes, op. cit., para. 73, the panel considered that Article XX “clearly allowed contracting parties to give priority to human health over trade liberalization”.

229 See, in particular, United States – Article 337, op. cit., para. 5.9; Canada – Foreign Investment Review Act, adopted on 7 February 1984, BISD 305/S/140, para. 5.20, and United States – Prohibition of Imports of Tuna, op. cit., para. 5.22.

230 Japan – Film, op. cit., para. 10.32.

231 Previous Panels have not defined the precise scope of the concept of detailed justification.


235 See para. 3.22 in which the European Communities point out that world production of
asbestos reached a peak of 5.2 million tonnes, since when production has steadily declined, falling to 1.92 million tonnes in 1997. With regard to the countries that have restricted or banned the use of asbestos, see the description given by the European Communities in paras. 3.31 et seq. above.

The Panel does not consider that the EC’s argument based on a comparison between products corresponding to the same tariff line are relevant. The concept of competition is not limited to products that fall in the same tariff heading and was established in the earliest cases based on Article XXIII:1(b). See, in particular, the Report of the Working Party on Australian Subsidy on Ammonium Sulphate, adopted on 3 April 1950, II/188, para. 12, the Report of the Panel on Complaints in the case Germany – Sardines, op. cit., para. 16, and the Report of the Panel in Japan – Film, op. cit., para. 10.73.

The concessions concerned do not seem to have been affected by the enlargement of the European Communities to include Austria, Finland and Sweden in 1996.

In our opinion, there is a difference between, on the one hand, an import ban following upon a series of national measures gradually reinforcing, since 1977, the measures taken to protect public health against the effects of asbestos and, on the other, the relationship which the EC tried to establish in EEC – Oilseeds between the existence in 1962 of oil-seeds subsidies in certain member States of the European Communities and the development of a subsidy programme insulating oil-seed producers from competition from imports (see para. 149 of the panel report).

Even if it were applicable, we consider that the EC rebutted this presumption by their references to the systems established at international and Community level concerning the use of asbestos.

See Annex II, reply of the European Communities to the Panel’s question No. 4 at the Second Meeting with the Parties, paras. 254 to 261.

EUROPEAN COMMUNITIES – MEASURES AFFECTING ASBESTOS AND ASBESTOS-CONTAINING PRODUCTS

Report of the Panel

Addendum
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This addendum contains the annexes to the Report of the Panel to be found in document WT/DS135/R.
ANNEX I

DECREE NO. 96-1133 OF 24 DECEMBER 1996

Concerning the ban on asbestos, implemented pursuant to the Labour Code and the Consumer Code

(Official Journal of 26 December 1996)

The Prime Minister,

Acting on the report of the Minister for Justice, the Minister for Equipment, Housing, Transport and Tourism, the Minister for Labour and Social Affairs, the Minister for the Economy and Finance, the Minister for Industry, Postal Services and Telecommunications and the Minister for Agriculture, Fisheries and Food,


Having regard to the Labour Code, in particular, Articles L.231-1, L.231-6, L.231-7 and L.263-2 thereof;

Having regard to the Consumer Code, in particular Article L.221-3 thereof;

Having regard to the Penal Code, in particular, Article R. 610-1 thereof;

Having regard to the Customs Code, in particular Article 38 thereof;

Having regard to the Traffic Law;

Having regard to Decree No. 88-466 of 28 April 1988 as amended, concerning asbestos-containing products;

Having regard to Decree No. 96-98 of 7 February 1996 concerning the protection of workers against the risks associated with asbestos dust inhalation;

Having regard to the referral of the matter to the Commission of the European Communities by the French Government on 29 October 1996, under the emergency procedure established under Article 9.7 of Directive (EEC) 83/189 as amended, providing for a notification procedure in the sphere of technical standards and regulations;

Having regard to the opinion of the National Commission for Occupational Health and Safety in Agriculture of 26 September 1996;

Having regard to the opinion of the Consumer Safety Commission of 2 October 1996;

Having regard to the opinion of the Senior Council for the Prevention of Occupational Hazards of 16 October 1996;

Upon consultation with the professional employer and employee bodies concerned;

Having heard the Council of State (Social Affairs Section),

Decrees:
Article 1

I. For the purpose of protecting workers, and pursuant to Article L. 231-7 of the Labour Code, the manufacture, processing, sale, import, placing on the domestic market and transfer under any title whatsoever of all varieties of asbestos fibres shall be prohibited, regardless of whether these substances have been incorporated into materials, products or devices.

II. For the purpose of protecting consumers, and pursuant to Article L. 221.3 of the Consumer Code, the manufacture, import, domestic marketing, exportation, possession for sale, offer, sale and transfer under any title whatsoever of all varieties of asbestos fibres or any product containing asbestos fibres shall be prohibited.

III. The bans instituted under Articles I and II shall not prevent fulfilment of the obligations arising from legislation on the elimination of wastes.

Article 2

I. On an exceptional and temporary basis, the bans instituted under Article 1 shall not apply to certain existing materials, products or devices containing chrysotile fibre when, to perform an equivalent function, no substitute for that fibre is available which:

- On the one hand, in the present state of scientific knowledge, poses a lesser occupational health risk than chrysotile fibre to workers handling those materials, products or devices;
- on the other, provides all technical guarantees of safety corresponding to the ultimate purpose of the use thereof.

II. The scope of application of paragraph I of this Article shall cover only the materials, products or devices falling within the categories shown in an exhaustive list decreed by the Ministers for Labour, Consumption, the Environment, Industry, Agriculture and Transport. To ascertain the justification for maintaining these exceptions, the list shall be re-examined on an annual basis, after which the Senior Council for the Prevention of Occupational Hazards and the National Commission for Occupational Health and Safety in Agriculture shall be consulted.

Article 3

I. The manufacture, processing, importation and domestic marketing of any of the materials, products or devices falling into one of the categories mentioned on the list envisaged under Article 2 shall be subject to a statement, signed, as appropriate, by the head of the business establishment, the importer or the party responsible for domestic marketing, which should be addressed to the Minister for Labour. This statement shall be filed in January of each year or, as appropriate, three months before the start of a new activity or the alteration of an existing production activity, by means of a form decreed by the Ministers for Labour, Consumption, Industry and Agriculture.

The statement shall be accompanied by all the supporting documents in the possession of the declaring party making it possible, considering the state of scientific and technological progress, to determine that as of the date of signature of the statement, the activity covered by the statement meets the conditions set forth in Article 2.1.

II. Activities that have not been the subject of a full statement submitted within the set time-frame may not benefit from the exception granted under Article 2.
III. The Minister for Labour may at all times convey to the author of the statement such information as may seem to him to establish that the material, product or device in question, although falling into one of the categories on the list mentioned in Article 2, does not meet the conditions laid down in paragraph I of that same Article. After requesting comments from the declaring party, he may serve notice to said party to cease manufacture, processing, importation or domestic marketing and to observe the ban instituted under Article 1. He may make such notification public.

Article 4

The manufacture and processing of the materials, products and devices falling into the categories on the list mentioned in Article 2 of this Decree must conform with the rules laid down under Chapters I and II and Chapter III, Section 1 of the aforementioned Decree dated 7 February 1996.

Labelling and marking shall conform with the requirements of Article L. 231-6 of the Labour Code and the rules established by the aforementioned Decree dated 28 April 1988.

Article 5

Without prejudice to the application of the penalties envisaged under Article L. 263-2 of the Labour Code in the event of violation of the provisions of Article 1.I of this Decree, the act of manufacturing, importing, introducing into the domestic market, exporting, offering, selling, transferring under any title or possessing for sale all varieties of asbestos fibres or any product containing asbestos fibres, in contravention of the provisions of Article 1.II shall be punishable by the fine prescribed for 5th class offences.

Article 6

I. Articles 1, 2, 3 and Article 6.I of the above-mentioned Decree No. 88-466 of 28 April 1988 are hereby repealed.

II. In the first subparagraph of Article 4 of the same Decree, the words: “bans envisaged in Article 2 above” shall be replaced by the word: “bans”.

III. In Article 6.II of the same Decree, the words: “other than those envisaged under Article 2” shall be replaced by the words: “which are not subject to bans”.

Article 7

Until 31 December 2001 and on a transitional basis, the ban on possession for sale, offering for sale and transfer under any title shall not apply to the used vehicles nor to the agricultural or forestry machinery put into circulation before the effective date of this Decree, and covered by Article R.138 of the Traffic Law.

Article 8

This Decree shall become effective on 1 January 1997.

Article 9

The Minister for Justice, the Minister for Equipment, Housing, Transport and Tourism, the Minister for Labour and Social Affairs, the Minister for the Economy and Finance, the Minister for the Environment, the Minister for Industry, Postal Services and Telecommunications, the Minister for Agriculture, Fisheries and Food, the Deputy Minister for the Budget, the Government Spokesperson, and the Deputy Minister for Finance
and Foreign Trade shall be responsible, each in his own sphere of competence, for implementing the present Decree, which shall be published in the Official Journal of the French Republic.


By the Prime Minister

ALAIN JUPPE

The Minister for Labour and Social Affairs,
JACQUES BARROT

The Minister for Equipment, Housing, Transport and Tourism
BERNARD PONS

The Minister for the Environment,
CORINNE LEPAGE

The Minister for Agriculture, Fisheries and Food,
PHILIPPE VASSEUR

The Deputy Minister for Finance and Foreign Trade,
YVES GALLAND

The Minister for Justice,
JACQUES TOUBON

The Minister for the Economy, and Finance,
JEAN ARTHUIS

The Minister for Industry, Postal Services and Telecommunications
FRANCK BOROTRA

The Deputy Minister for the Budget, Government Spokesperson,
ALAIN LAMASSOURE
ANNEX II

QUESTIONS – REPLIES

At the First and Second Substantive Meetings
(1-2 June 1999 and 20-21 January 2000)

I. QUESTIONS TO THE PARTIES

A. QUESTIONS AT THE FIRST SUBSTANTIVE MEETING (1-2 June 1999)

1. Questions by the Panel to Canada

Question 1: Canada states that “the risks to health associated with modern chrysotile products are undetectable”. Does the concept of “undetectable” risk mean the same for Canada as no risk?

The term “undetectable” should not be interpreted as a subjective judgement with respect to risk management, but rather as a scientific term related to quantification of the risk. In the specific context of its submission, Canada could just as easily have used the expression “below detection limits” (BDL), which is commonly used by scientists. This expression means that as determined using the latest methods and techniques and the most rigorous statistical analysis, the risk (effect) related to exposure conditions (type of fibre, dose, duration) is so slight, if it exists at all, as to be “below detection limits”. Scientists generally do not use the expression “zero effect” or “no risk” or any other similar expression to describe a level of risk. Rather, they use “below detection limits” (BDL). The term “undetectable” used by Canada and the scientific community at large should be interpreted in this specific sense. Canada considers that it is inappropriate to use expressions such as “zero risk” or “no risk”. Canada thus adheres strictly to the scientific definition of the expression used, i.e., not “no risk”, but “undetectable risk”, as indeed do the European Communities. In fact, the European Communities themselves corroborate the validity of this concept when they state, with respect to ambient concentrations of asbestos in buildings, that “it is clear that the risk [thereof] is undetectable”.

Question 2: In its oral submission (1 June) Canada indicated, with respect to encapsulated products, that chrysotile fibres are only released under certain conditions. What are those conditions?

Products in which chrysotile is an encapsulated component release virtually no fibres naturally, or, if they do so, in such minimal concentrations that they are below detection limits. In other words, only aggressive operations on products containing encapsulated chrysotile can lead to the release of a detectable quantity of fibres. Such operations may occur during installation, renovation or removal of such products.

Question 3: Do operations by professional workers or private individuals on asbestos-cement products (such as sawing, sanding or demolition) and erosion of the matrix over time have the effect of releasing chrysotile fibres at levels that present a “detectable risk”?

When asbestos products are installed in the workplace, be they asbestos-cement products (pipes, panels or tiles), friction products or other products containing encapsulated chrysotile, small quantities of fibres may be released. To mitigate this potential risk, numerous codes of practice such as Standard 7337 of the International Organization for Standardization (ISO) or the American Water Works Association Code have been developed in order to minimize the release of dust. These codes of practice cover the tools and procedures for use with these products in detail. Recommended installation methods can eliminate the need to cut or drill basic chrysotile products at construction sites, since those
products are distributed in various pre-cut and pre-drilled formats to buyers’ specifications. Where products do have to be drilled or cut, the use of appropriate tools can minimize the release of dust and keep it well within the level considered safe by WHO. This has been confirmed by laboratory tests and testing at construction sites of release of fibres during the installation of asbestos products.¹

4. These workers are subject to exposure peaks, the magnitude of which depends on the protective measures in place. However, aside from the intensity of exposure, the duration of exposure is also critical, for the risk is essentially determined by cumulative exposure. In the United States, according to the studies by CONSAD for OSHA (Occupational Safety and Hygiene Agency) between 1985 and 1990, the average annual exposure for such workers was 10 to 100 times higher than for occupants of buildings with asbestos insulation, which is corroborated by the WHO report EHC-203 (exposure of 0.002 to 0.02 f/ml). It can be inferred that the cumulative lifetime exposure for such workers is 2 to 30 times higher than for occupants of such buildings. These workers therefore appear to have a lifetime risk of some 20 to 300 per million, which is an “undetectable” risk, i.e. it cannot be shown or measured empirically.²

5. Unlike professional workers, private individuals probably work only very sporadically with chrysotile cement products. The exposure incurred by these do-it-yourself enthusiasts will only be a fraction of that of professional workers. Consequently, if professional workers working on a daily basis with chrysotile-cement products are not subject to any detectable risk, logically private individuals will be even less so. Generally speaking, private individuals will not perform any major operations such as sawing, sanding or demolition of materials. More likely, do-it-yourselfers will occasionally perform minor operations such as drilling a hole for a cable, for example. Finally, whenever they work with cement products, whether or not they contain chrysotile, do-it-yourselfers should use simple protective measures, if only because of the presence of other carcinogenic substances such as crystalline silica.

6. In the context of the general human environment, small quantities of fibres can also be released through natural wear and tear of the product during its life cycle, either by mechanical erosion of the product as a result of water or wind action or by the physicochemical effects of changes in temperature. There has been extensive study of this matter, and it has been recognized that the release of fibres is at levels which do not measurably add to the chrysotile naturally present in the environment. The European Communities recognize that at such levels “it is clear that the risk is undetectable”.

Question 4(a): What exactly does Canada mean by “modern chrysotile products” or “modern asbestos products”?

7. By “modern chrysotile products”, Canada means the range of non-friable products where: (i) only chrysotile asbestos is used, but no amphibole asbestos (crocidolite and amosite); and (ii) in which the fibres are firmly bonded physically and chemically into the matrix (cement, asphalt, resins, plastic, etc.) of the compound (chrysotile-cement, friction material, asphalt road surfacing, etc.) and cannot easily be released in biologically significant concentrations. In this respect, the following observation seems germane:

“Once an asbestos-containing product has been manufactured, whether or not it constitutes a source of asbestos in the environment will depend to a great extent on whether or not the asbestos is firmly “locked-in” the product with a binder, saturant, coating or bonding agent such that normal handling, application and use do not release it. Asbestos-cement products are a good example of “locked-in” products which probably do not constitute a significant source of asbestos to the environment under normal conditions of use.”³

Question 4(b): Since when have “modern” asbestos processing techniques been in use?

8. The date of the shift to exclusive use of chrysotile asbestos in non-friable (“locked-in”) products varies from country to country, as do the dates on which different countries
banned asbestos flockings or the use of amphibole asbestos. ILO Convention 162 concerning Safety in the Use of Asbestos has now been adopted by over 20 countries. It should also be noted that many countries have banned or severely restricted and regulated the use of amphiboles or barred the spraying technique as required by Convention 162, without actually acceding to it.

Question 5: Canada observes that “the removal of most modern asbestos products should present little difficulty when a building is demolished”. Can Canada expand on this assertion, and in particular develop the concept of “little difficulty”? In “everyday life”, how can an individual who demolishes or works on a building containing asbestos-cement detect the presence of this material?

9. Demolition of a building is an operation which normally requires a permit and which is usually performed by professional workers. The issue of permits allows the authorities to ensure that the demolition is carried out by persons who are familiar with the appropriate working methods necessary to control all the risks associated with demolition activities including those involving the dust of all kinds which may be released during the works.

10. In the case of demolition of buildings, the first concern of the person responsible for the work should be to identify the presence of friable materials such as amphibole flocking, whether that person is a professional worker or a private individual. Indeed, when the demolished building contains flockings, their release into the open air will be the main source of contamination of the site by asbestos, much more so than the presence of asbestos-cement materials. The latter may be broken or crushed to a greater or lesser degree, but the concentration of respirable fibres thus released will be very much lower than concentrations of fibres originating from flockings.

11. This observation demonstrates the importance of producing a register of buildings containing flockings and introducing regulations requiring consultation of the register and/or inspection of buildings to be demolished to determine whether or not flockings are present before a demolition permit is issued.

12. When it is determined that a building contains flockings, specific control measures must be followed if demolition is considered necessary.

13. Otherwise, demolition of structures containing products in which chrysotile is encapsulated (generally flat or corrugated sheeting and roof tiles) requires only elementary precautions since chrysotile-cement debris remains largely inert. In Quebec, for example, demolition is regulated by the Code of Safety for the Construction Industry issued by the Occupational Health and Safety Commission (Commission de la Santé et de la Sécurité du Travail). The debris resulting from demolition is disposed of in public landfill sites for solid wastes in the same way as other construction waste and coated with covering materials as a preventative measure, superfluous according to some, in order to ensure that no chrysotile dust is left suspended in the air. Under these conditions, chrysotile-cement debris causes little or no increase in natural concentrations of asbestos in the environment, levels which the European Communities, it should be recalled, consider “undetectable.” Moreover, as chrysotile asbestos is not soluble and in any case constitutes no risk when ingested, it has no effect on the local or nearby water-table.

14. Private individuals who work on a construction containing chrysotile asbestos in everyday life run little risk since such work is very sporadic and generally of short duration, while chrysotile-related risks are primarily associated with prolonged occupational exposure to high concentrations of fibres.

15. It should also be noted that individuals should exercise equal care when working with any cement material, whether or not it contains chrysotile, because they are exposed to risks of the same magnitude due to the presence of other dust, such as crystalline silica or substitute fibres which may be released during the work. Crystalline silica is
classified as a Group I carcinogen by the IARC and no study has yet been carried out on the health risks associated with the inhalation of dust from substitute fibres which may be released into the air during work of this type.

16. Wearing a “surgical” type mask would be a wise precaution whenever work by an individual on any form of material might result in the formation of respirable dust.

**Question 6:** International institutions such as WHO or ILO are encouraging a gradual switch to substitute products (see for example ILO Convention 162 concerning Safety in the Use of Asbestos; IPCS Environmental Health Criteria (203) on Chrysotile, WHO 1998). Does Canada subscribe to this approach?

17. Canada has itself ratified Convention 162 concerning Safety in the Use of Asbestos. The Convention should, however, be considered in its entirety, and it should be recalled that in Article 10, ILO encourages a gradual switch to substitute products “where necessary to protect the health of workers and technically practicable ... by other materials or products or the use of alternative technology, scientifically evaluated by the competent authority as harmless or less harmful.”

18. To date, no comparative scientific study has conclusively shown that, under similar conditions of production, manufacture or use, substitute products are harmless or less harmful than chrysotile asbestos. Indeed, some recent studies show that chrysotile displays lower biopersistence than the main fibrous substitutes such as refractory ceramic fibres, glass fibres, aramid fibres and cellulose fibres. There are also numerous scientific studies in existence which show that, at the low levels of exposure currently observed in the chrysotile products industry (generally less than 1 fibre/ml.), there is no measurable increase in the risk to human health.

19. The same argument applies to the IPCS Environmental Health Criteria (203) on Chrysotile, WHO 1998, which states: “where safer substitute materials for chrysotile are available, they should be considered for use.”

**Question 7:** What criteria should be used to determine the relative risk associated with substitute products and chrysotile asbestos?

20. Canada has referred to the consensus that the relative risk of fibrous materials varies according to three factors (“3 D”): dimension, durability and dose.

21. The dimension (length and diameter) affects “respirability”. This is the factor which determines whether a fibre can actually penetrate the confines of the respiratory system: the alveoli.

22. However, the dimension factor is a necessary but not sufficient condition. An inhaled fibre must stay in the system long enough i.e. it must have sufficiently long (biopersistence) to exert its pathogenic effect. This is the durability factor. In this respect, Canada has submitted (and the latest data, which Canada can provide, confirms this) that in inhalation experiments with animals, chrysotile is very quickly eliminated from the lungs (within 24 to 48 hours), while amphiboles persist practically indefinitely, and then trigger the range of inflammatory reactions which precede and herald the known pathologies. The small quantity of data available on the biopersistence of certain substitute fibres (for example aramid fibres are more biopersistent than chrysotile) suggest that the durability factor should be seriously considered in evaluating the relative risk associated with fibres of substitutes for chrysotile asbestos. In this connection, the following quotation from a Scandinavian study bears repeating:

“(…) adverse effects are associated rather with the fibres that are retained (amphiboles), than with the ones being cleared (largely chrysotile).”

23. Another study, published in 1995, indicates that: “biopersistence of inhaled fibrous materials is a critical factor in determining carcinogenic potency.”
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24. A recent report by Bernstein (1997) for the Joint Research Centre, Environmental Institute, European Chemical Bureau in Ispra (Italy) under the title Correlation Between Short Term Biopersistence and Chronic Toxicity Studies confirmed the relevance of the durability factor in evaluating the risk associated with substitute fibres and chrysotile asbestos.7

25. We had good reason to draw attention to the importance of the third factor, the dose. It introduces the phenomenon of the threshold of exposure above which harmful effects begin to appear, and below which (with obvious differences according to the type of fibrous material) the risk, if any, becomes undetectable.

26. Although the “3 Ds” are the recognized critical factors in the risk associated with respirable fibres, other factors; Such as the capacity to induce the production of reactive molecular forms may influence the degree of risk. Indeed, as several fibre toxicity mechanisms remain uncertain, it must be ensured that the experimental and epidemiological studies to evaluate the risk posed by the fibres to humans are equally valid and comparable for chrysotile and its substitutes. It must also be ensured that the main comparability and validity criteria and principles for the design of such studies are established by recognized international bodies. We will not list these principles here, but we wish to emphasize that comparisons of the risk associated with different fibres need to satisfy the following conditions in order to be valid, i.e. unbiased: (i) analysis of the structural characteristics must be based on the “3 Ds”; (ii) the dose or exposure must be similar and must not overload the pulmonary macrophages; (iii) the observation periods must be similar and long enough to observe the effect of differences in biopersistence; (iv) the number of subjects and observations/studies must be sufficiently large to be able to detect even a slight risk; (v) the characteristics of the fibres studied must represent in the same way the real uses of chrysotile fibres and their real substitutes; and (vi) animal studies must study the effects of fibre inhalation in the same concentrations, under the same conditions of exposure and using the same experimental method.

27. Only on the basis of such criteria can the relative risk of chrysotile fibres and their substitutes be determined, and only a comparative risk analysis such as the one defined by the National Research Council in the United States can assess the quality and comparability of the data concerned and estimate the relative risk of the substances. For example, recent and incomplete toxicological data on substitute fibres cannot be compared with epidemiological studies based on 100 years of human exposure. It is necessary to compare data of the same type and of the same quality in order to have confidence that a proposed substitute is safer than chrysotile.

Question 8: What occupations are at greatest risk of exposure to asbestos at the present time? Can protection measures be instituted - and enforced - for all these occupations? Is controlled use of chrysotile asbestos and products containing it possible for occupations exposed to occasional but potentially high exposure, and exposure of a para-occupational and domestic nature?

28. In Canada’s view, the occupations which run the greatest risk of exposure to chrysotile asbestos, are, in descending order: (i) chrysotile miners and employees of chrysotile-ore processing plants (mills); (ii) workers employed in the manufacture of chrysotile fabrics; (iii) workers employed in the manufacture of friction linings (brakes, clutches, etc.); (iv) workers employed in the manufacture of chrysotile cement products; (v) workers employed in asbestos removal; and (vi) workers employed in construction, renovation, maintenance and insulation.

29. While all workers in the above categories are more likely to be exposed to different types of asbestos fibres, it is most important to note that those in categories five and six may be exposed to amphiboles, while the others are not. In the case of workers in areas where chrysotile is used, effective protection measures have been put in place since the 1970s by the competent authorities, with the collaboration of manufacturers and trade unions, (ventilation, filtration, wet processes, mechanization, etc.). The methods used for this purpose rely on relatively simple technologies. They include the following: better ventila-
tion of working areas; more effective filtration of dust-bearing air; crushing and processing under negative pressure to prevent the escape of dust; exhaust hoods in work stations directly exposed to fibre; wet manufacturing processes; mechanization, etc..

30. While workers in category 6 are exposed to friable asbestos products (of all kinds) on quite a regular basis in the course of their work, Canada is nevertheless of the view that the same information and training measures for employers and workers, combined with regular inspections and checks by occupational health and safety agencies, together with the compilation of registers of buildings containing asbestos, should give results similar to those obtained following the introduction of similar measures for the other five categories of workers. Prevention and protection measures (masks, dampening, ventilation, etc.) can considerably reduce occupational exposure to asbestos. It should be reiterated that these working conditions stem from uncontrolled use of asbestos (mainly flockings) which has not occurred since the early 1980s, and a ban on chrysotile would not in any way alter the situation.

31. Controlled use of chrysotile asbestos and its modern products is possible for all occupations, even those where the workers may be subject to occasional exposure. It is a matter of establishing and enforcing an appropriate set of rules for their use, in the same way as is done for many other dangerous substances used in the workplace.

32. There are no conclusive studies of the risks associated with occasional but potentially high exposure, even the French study by Iwatsubo et al. quoted by the European Communities, concurs "(...) subjects with sporadic exposure were not at greater risk of mesothelioma than were controls."8

Question 9: Does Canada agree with the French experts’ estimate (Evolution of the annual incidence of mesothelioma in Canada and Quebec, Section III.B.4), of the annual incidence of mesothelioma in Canada and Quebec? If not, what are the Canadian figures? Does Canada have statistics for deaths from lung cancer caused by chrysotile asbestos? If yes, what do they show?

33. The estimates of the incidence of mesothelioma in Canada and Quebec in the first written statement of the European Communities match the Canadian data, precisely because the International Agency for Research on Cancer data are taken from Canadian cancer records. The French experts’ analysis however, requires further explanation in order to be correctly understood.

34. Unlike France whose cancer records cover only 9.5 per cent of the population (INSERM page 173), Canada is one of the few countries in the world whose cancer records cover 100 per cent of the population, with the consequence that its data are more reliable than those of France or most countries. For this reason, and because Canada has a large number of studies on cohorts of asbestos workers, compiling a special registry of mesothelioma cases was less pressing than in other countries. It should be noted that France also does not have a central registry of mesothelioma.

35. Three caveats are necessary in order to interpret correctly the Canadian data summarized in the table produced by the European Communities: (i) Quebec cancer reports and likewise the Canadian totals that include Quebec are only reliable since 1984; (ii) as in most countries, mesothelioma was considerably underdiagnosed and under-reported until the mid-1980s, which falsely inflates the increase in the incidence of mesothelioma during the 1970s and 1980s9; (iii) data and trends prior to 1984 should therefore be treated with caution.

36. Recent analyses of Canadian data on mesothelioma in Canada10, British Columbia11 and Quebec 13-14 agree that rates of incidence of mesothelioma have been stable in women of all age groups since 1984. In Quebec, the rates are 70 per cent higher than elsewhere in Canada, probably because of more frequent and more intense exposure in the workplace. In fact, Quebec produced about half of the world’s commercial chrysotile until
the 1950s. Quebec also used large quantities of amphiboles in various sectors, particularly in certain sectors in which many women were employed, especially during the Second World War.

37. According to a study by Schanzer, Semenciw and Ugnat (Health Canada, 1997), the incidence of mesothelioma in men in Canada increased by 22 per cent from 1984 to 1993, half the increase estimated by the French experts, and also half that in France. Over the same ten-year period, the incidence of mesothelioma rose by 45 per cent in Quebec (the same as in France), 34 per cent in Ontario, 0 per cent in British Columbia and less than 10 per cent for the rest of Canada. These rates reflect wide disparities in Canada, contrary to the European Communities’ assertion. The incidence in men also levelled off in 1984 in British Columbia (according to Coleman and Philips), and seems to have levelled off in Quebec after 1990.

38. Finally, the analysis of Canadian rates between 1973 and 1992 (Schanzer and colleagues, 1997) estimates that the risk is four times higher for men born before 1940 than for those born between 1951 and 1955. These analyses therefore suggest that the incidence of mesothelioma has levelled off in Canada is declining in British Columbia, and has levelled off in Quebec.

39. The following table, adapted from the one submitted by the European Communities, shows that despite the much higher asbestos production in Canada and Quebec than in France, the incidence of mesothelioma and the increase in that incidence were lower in Canada than in France. Thus, although Quebec produced almost half the chrysotile used in the world, the incidence of mesothelioma in Quebec was, surprisingly, no higher than in France.

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<th>Period</th>
<th>Canada M</th>
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<td>1978-1982</td>
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<td>1983-1987</td>
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<td>11</td>
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<td>1988-1992</td>
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40. With respect to the proportion of lung cancer cases attributable to occupational exposure to asbestos, no country in the world possesses such statistics. However, case-control studies of a number of cancers and types of exposure in the general population in Canada (some metropolitan regions or provinces) suggest an upper limit to the proportion of lung cancer cases attributable to asbestos, but they suffer from flaws which make it impossible to determine the specific proportion that is due to asbestos, independently of other major risk factors.

41. A case-control study of cancers in Montreal suggests that the specific association between lung cancer and asbestos would explain at most 7.6 per cent of lung cancers in men between 1979 and 1985; but the real proportion is lower because part of the 7.6 per cent is attributable to concomitant exposure: PAHs (polycyclic aromatic hydrocarbons), solvents, alkanes, welding fumes, tobacco (residual effects after statistical control), etc.

42. Case-control studies in the United States, the United Kingdom and Sweden provide other estimates of the percentage of lung cancers attributable to asbestos. As the review of the studies cited by INSERM (pages 10 and 179 of the Report) indicates, the enormous heterogeneity of these studies (e.g. the attributable percentage ranges from 0.6 per cent to 16.6 per cent in the United Kingdom alone) stems from variations in the prevalence of asbestos exposure (types of industry and proportion of workers exposed), which prevents any extrapolation of their findings to Canada (or to France).

Question 10: Canada refers to the WTO report Environmental Criteria 53: Asbestos and Other Natural Mineral Fibres, published in 1986, but does not mention a subsequent report by the same organization, IPCS Environmental Health Criteria (203) on Chrysotile,
in 1998. In Canada’s view, to what extent does the 1998 report confirm the findings of the 1986 report, particularly with regard to the dangers of chrysotile and risk management methods?

43. The reply to this question should be prefaced by two preliminary remarks. Firstly, it should be recalled that the forewords to the above-mentioned reports read as follows: “[T]his report contains the collective views of an international group of experts and does not necessarily represent the decisions or the policy of the United Nations Environment Programme, the International Labour Organization, or the World Health Organization.” Secondly, Canada wishes to emphasize that the interpretation of the two reports, each over 150 pages long, in the limited context of the reply to this question will necessarily be incomplete.

44. That being so, this reply will address the question of the dangers of chrysotile before turning to risk management methods. Report 203 surveys the data on exposure to chrysotile and suggests among other things that: (i) in the mines and mills in Quebec, “the average fibre concentrations (…) are now generally well below 1 f/ml”; (ii) in the production of asbestos cement in Japan, concentrations range between 0.05 and 0.45 f/ml in 1992; (iii) in the production of textiles in Japan between 1984 and 1986, the concentrations found were between 0.1 and 0.2 f/ml; (iv) in other countries and various other sectors: vehicle maintenance: “practically all measured levels after 1987 were less than 0.2 f/ml”; buildings maintenance: “[I]n buildings with control plans, personal exposure of buildings maintenance personnel in the USA … between 0.002 and 0.02 f/ml.” The trend towards lower concentrations is thus confirmed, and demonstrates the feasibility and benefits of applying controlled use policies.

45. Report 203 also recognizes the difference in the dangers associated with chrysotile and amphiboles:

“The mechanisms of the relatively more rapid clearance of chrysotile fibres compared to those of amphiboles (…)”

“The more rapid removal of chrysotile fibres from the human lung is further supported by findings from animal studies (…)”

46. Report 203 recognizes that the presence of chrysotile fibres in water is harmless: “… it was concluded that there was little convincing evidence of an association between asbestos in public water supplies and cancer induction. More recent identified studies do not contribute additionally to our understanding of health risk associated with exposure to chrysotile in drinking water.”

47. None of the IPCS reports (EHC No. 53 and 203) suggests any measures whatsoever with respect to risk management methods:

The EHC monographs are intended to assist national and international authorities in making risk assessments and subsequent risk management decisions. They represent a thorough evaluation of risks and are not, in any sense, recommendations for regulation or standard setting.”

48. Thus among the conclusions in the 1998 report, we read: “[W]here safer substitute materials for chrysotile are available, they should be considered for use.” It is clear from this latter extract that the scientists who wrote the EHC 203 report refer to the consideration of safer and available substitute material. On the issue of proven safer substitute fibres, Canada has already indicated in its written and oral submissions that none of the fibrous materials that have been proposed as substitutes have undergone rigorous tests to show that they are harmless. The limited data at present available suggests that premature replacement could be a serious risk management error. This last extract refers to available substitutes. Experience shows that the mere fact that such materials are available does not mean that they are technically adequate. Finally, the recommendation does not say that
asbestos should be replaced, rather than replacement “should be considered”. The 1998 WHO report thus confirms the 1986 report. Both emphasize a reduction in fibre concentrations, recognize the difference between chrysotile and amphibole and suggest replacing chrysotile by safer substitutes. However, as Canada submits, the fibre substitutes have never been subjected to rigorous tests that demonstrate them to be harmless.

**Question 11:** Does Canada accept the logic that an increase in a country’s imports of chrysotile asbestos is accompanied by a proportional increase in asbestos-related pathologies?

49. Canada does not accept the “logic” that an increase in a country’s imports of chrysotile asbestos is accompanied by a proportional increase in asbestos-related pathologies. Indeed, this “logic” only holds up if one ignores the marked differences in risk between different types of asbestos, between friable and non-friable products, between sanitary and unsanitary conditions and between the production, use and presence of asbestos products.

50. It is recognized that the health risks associated with asbestos are primarily a function of the “three Ds”: dose, dimension, durability (biopersistence). The volume of asbestos imports bears little or no relation to these risk factors. Rather, specific information should be provided for each country on the following: the proportion of workers exposed, the conditions of exposure (types of industry and products, workplace hygiene regulations) product uses and distribution in various population groups (brakes, flockings, construction materials, cement, public buildings, etc.), the types of asbestos used and their applications. These conditions vary greatly from country to country and over time.

51. No analysis can distinguish the impact of asbestos imports by type of fibre for the following reasons: (i) the available data do not normally allow us to distinguish retrospectively between the mineralogical types of imported asbestos; (ii) even if such information were available, it would not be possible to distinguish the effects based on mineralogical type of asbestos because the volumes of imports of chrysotile and amphiboles are closely correlated (i.e. historically, the more chrysotile that was imported, the more amphiboles were imported too); and (iii) there are paradoxes, such as countries which produced chrysotile while also importing amphiboles to satisfy their industrial needs of the time.

52. Furthermore, importing a small quantity of amphiboles for uncontrolled use, in flockings, for example, would probably lead to a measurable increase in pathologies. On the other hand, an increase in the chrysotile imports for controlled use in the manufacture of non-friable products in a context of controlled utilization would not result in any detectable additional risk.

53. As for the type of fibre, three recent studies examined the relationship between asbestos imports and mesothelioma in three countries: the United Kingdom24, the United States25 and France.26 None of these studies addressed the link between chrysotile imports and mesothelioma. Moreover, in the most recent of these studies, conducted by French researchers, the authors explain that their estimate of the incidence of mesothelioma in France is much lower than that predicted using Peto’s model for Great Britain because of the type of asbestos and not the imports:

> “a possible explanation can be found in the different fibres used in this country; France used a much smaller proportion of amphiboles than Great Britain.”27

54. As for the level of exposure and the difference between workers and the general population, the French experts explain that there was no increase in mesothelioma among women, despite an increase in production and imports over time, because of the low levels of environmental exposure:

> “This does not support the hypothesis that current environmental exposure to asbestos could be associated with detectable risk of death.”28
55. It should be noted, incidentally, that, like Canada, the French experts use the concept of “detectable risk”. This “logic” also does not take account of the variety of ways that asbestos is used or the way its use has evolved. For example flockings in the 1960s and 1970s, must have significantly increased the risk of mesothelioma for a given volume of consumption or imports. Neither does this “logic” hold good for countries/regions such as South Africa, Brazil, Quebec and Western Australia, which are asbestos producers but import little, since the proportion of their populations exposed in the workplace and the levels of exposure were higher than in most importing countries and often led to higher risks of pathologies than in countries/regions that import more asbestos.

56. Technically speaking, what the Panel and the European Communities call “logic” consists of an error of inference which is classic in epidemiology, the social sciences and biostatistics: the ecological fallacy. This fallacy consists of confusing observations of large groups (aggregates) showing non-specific aggregate correlations with real individual effects. In particular, it consists of ignoring the fact that the conditions of exposure and “co-factors” (other risk factors) for individuals in the various large aggregates vary enormously. These big observation units are so broad that they are in fact not comparable or similar when it comes to the factors which actually determine the risk.

57. Thus, in the case of chrysotile, the study and graph by Takahashi et al cited by the European Communities do not distinguish between the effects of chrysotile and those of asbestos; separate graphs are needed for the consumption of chrysotile and amphiboles. In fact, similar graphs and correlations to those of Takahashi would be obtained for the consumption of artificial fibres or consumption of cement. What should one deduce from that?

58. Crude ecological correlations lend themselves to arbitrary and contradictory interpretations. There are only ten countries in the graph, which is too few: the sample, selection criteria or their characteristics skew the analysis. If another twenty countries were added, including Quebec, South Africa and Western Australia, the picture would be more representative but much less coherent, the correlation would be weaker and the line would be flatter. Indeed, to fit a straight line between these ten points is arbitrary. The best correlation with Takahashi’s ten points is not a straight line but rather an “S-shaped” curve (e.g. logistic or cumulative normal curve), with a practical risk threshold (no risk for consumption below 1 kg./capita/year) and a risk ceiling (no additional risk for consumption over 2.5 kg./capita/year). It is no more absurd to argue for this interpretation than for a linear relationship. We are not saying that these data show a risk threshold, but wish to underline the arbitrary nature of the European Communities’ interpretation of these ecological data.

59. For lung cancer, the significance of smoking and other risk factors, means that there is even less reason to deduce a proportional risk relationship with a country’s asbestos imports or consumption levels.

60. In short, there is no clear-cut link between asbestos imports (particularly chrysotile) and the incidence of asbestos-related pathologies.

Question 12: Does Canada agree with the European Communities that cellulose, polyvinyl alcohol and aramid fibres have been shown to be safe or of low toxicity?

61. No. Canada does not agree that cellulose, PVA and aramid fibres have low toxicity.

62. The European Communities claim that the diameter of these fibres is generally greater than that of chrysotile fibres. This can probably be explained by the data on “nominal diameter” supplied by the substitute fibre industry. For example, in the case of aramid fibres, the “nominal diameter” is from 12 to 15 microns. However, there are data which show that in commercial preparations of fibres labelled “nominal diameter: 12-15 microns”, there is a significant proportion of fibres with a diameter within the range of respirable...
diameters. The presence of such fibres has even been observed in the pulmonary alveoli of laboratory animals which have been exposed to commercial preparations.

Furthermore, as the IPCS-EHC Report No. 151 states on page 76:

“All fibres that are respirable and biopersistent must undergo testing for toxicity and carcinogenicity. Exposures to these fibres should be controlled to the same degree as that required for asbestos until data supporting a lesser degree of control become available. The data available suggest that para-aramid fibres fall within this category. Furthermore, other respirable organic fibres should be considered to fall within this category until data indicating a lesser degree of hazard become available.”

In addition, the claim that the fibres are less “biopersistent” is arguable. As the authors of the INSERM report recognize, there are few verifiable data on this subject. As stated in INSERM’s summary report - *Effects on Health of Substitute Fibres for Asbestos*:

“The results of many studies suggest that the ‘fibrous’ structure of asbestos is a significant pathogenic factor in the same way as some of its chemical characteristics. Consequently, any new fibre proposed as a substitute for asbestos or for any other purpose, must be suspected, a priori, of being pathogenic because of its structure, which does not preclude the analysis of other possible consequences of its physical and chemical characteristics.”

This view is echoed by the Scientific Committee on Toxicity, Ecotoxicity and the Environment of the European Commission (DG XXIV) which stated: “for the substitute materials, with the exception of vitreous fibres, there is no significant epidemiology base to judge the human health risks.”

Canada is prepared to prove that, in fact, aramid fibres are more durable and biopersistent than chrysotile fibres.

Canada shares the opinion of the experts that fibres replacing chrysotile should not be assumed to be harmless until all relevant data have been collected. The authors of the INSERM Report are equally circumspect on this point.

**Question 13:** In paragraph 238 of its first written submission, Canada states that Article 2.4 of the Agreement on Technical Barriers to Trade lays down the principle that a Member “shall use relevant international standards, or the relevant parts of them, as a basis for their technical regulations”. In paragraph 249 of its submission, Canada further states that the Decree does not comply with international standards (…). Does Canada consider the concept of regulations “based on” or “in accordance with” an international standard to be equivalent to the concept of using international standards or relevant parts of them “as a basis” for their technical regulations, within the meaning of Article 2.4 of the Agreement on Technical Barriers to Trade?

**No.** Under Article 2.4 of the TBT Agreement, France is required to use relevant international standards or parts of them as the basis or foundation of its technical regulations. In paragraph 249 of its first written submission, Canada merely observes that the Decree does not comply with international standards. In paragraph 249 of its first oral submission presented at the first substantive meeting, Canada maintained that “Article 2.4 should not be interpreted as allowing a Member to circumvent a complete series of international standards that provide the foundations – sometimes quite specific – for a measure enabling Members to fulfil the alleged objective”. In paragraph 250 of the same oral submission, Canada adds the following: “the French measure deviates significantly from the precepts of international standards”. In other words, the French measure deviates to such an extent from the fundamental principles of international standards that the latter are no longer recognizable.
Question 14: Is there any difference between the concept of using international standards “as a basis” for a technical regulation within the meaning of Article 2.4 of the Agreement on Technical Barriers to Trade and the concept of a technical regulation “in accordance with relevant international standards” within the meaning of Article 2.5 of the same Agreement?

69. Yes. As the Appellate Body reported in its Report on European Communities Measures Concerning Meat and Meat Products (Hormones), there is a difference between the meaning of the terms “based on” and “conform to”. In paragraph 163 of the Report, the Appellate Body stated that: “[…] the ordinary meaning of ‘based on’ is quite different from the plain or natural import of ‘conform to.’ A thing is commonly said to be ‘based on’ another thing when the former ‘stands’ or is ‘founded’ or ‘built’ upon or ‘is supported by’ the latter. In contrast, much more is required before one thing may be regarded as ‘conform[ing]’ to ‘another’: the former must ‘comply with’ ‘yield or show compliance’ with the latter. The reference of ‘conform to’ is to ‘correspondence in form or manner’ to ‘compliance with’ or ‘acquiescence’, to ‘follow[ing] in form or nature.’ A measure that ‘conforms to’ and incorporates a [given standard] is, of course, ‘based on’ that standard. A measure, however, based on the standard might not conform to that standard, as where only some, not all, of the elements of the standard are incorporated into the measure.” [Footnote omitted]

70. Article 2.4 of the TBT Agreement lays down the obligation to use relevant international standards or relevant parts of them as the basis of technical regulations. According to the ordinary meaning of words, this means that technical regulations must be founded on international standards or relevant parts of them. In other words, technical regulations must have as their fundamental principle or point of departure what is set out in international standards. This does not mean that the technical regulations adopted by a Member must be identical to international standards but where the latter are relevant, the technical regulations should be prepared on the basis of those international standards: the underlying international standard must be recognizable on a reading of the measure. Canada has established that the international standards concerning asbestos are relevant. The European Communities have not offered any convincing evidence to the contrary. In this instance, the French measure strays so far from the international standards that they are no longer recognizable as the foundation. The European Communities have not offered any reason to justify this departure and consequently the disputed measure is in contravention of Article 2.4 of the TBT Agreement.

71. Given that the Decree deviates to such an extent from the international standards that they are not evident as its basis, the European Communities cannot claim that “[…] it must be concluded (sic) that the international texts quoted, or sometimes not quoted, by Canada serve ‘as a basis’ for the French decree.”

72. Furthermore, the obligation to use international standards as the basis for preparing technical regulations should not be interpreted as permission simply to ignore important aspects of those standards, as France has done. France is, in effect, forcing the replacement of asbestos by substitutes, ignoring the fact that it is only in the event of necessity that such replacement is recommended and only where the substitutes are harmless and safe. France has quite simply ignored the test of “necessity” required by the international standards and reports in order to operate a policy of banning asbestos. This is particularly disconcerting when it is considered that these are the same standards and reports that are cited by the European Communities to justify the French position in paragraphs 531 et seq. of their submission and that those standards and reports are very specific in this respect.

73. For its part, Article 2.5 of the TBT Agreement establishes a rebuttable presumption when the measure is in accordance with international standards. The French measure is clearly not in accordance with international standards. In this instance, therefore, the European Communities cannot rely on the presumption in Article 2.5 of the TBT Agreement.
Question 15: Canada states that it “will use the example of chrysotile fibre and chrysotile cement in order to show that chrysotile fibre and substitute fibres are similar and products containing chrysotile and those containing substitute fibres are like products”. In the light of this paragraph and the overall arguments presented by Canada in its analysis of the similarity of products under Article III:4 of GATT 1994, is our understanding correct that Canada does not allege any likeness between chrysotile fibres and products containing chrysotile and any possible non-fibrous substitute products?

74. Yes. Both under Article III:4 of GATT 1994 and Article 2.1 of the TBT Agreement, Canada is not invoking the argument of the likeness of non-fibrous substitute products (PVC, ductile iron). Nor does Canada extend the likeness argument to substitute fibres other than glass, cellulose and PVA fibre or fibro-cement products containing these types of fibre.

75. Canada does not need to invoke all products like to chrysotile or chrysotile cement products to demonstrate a violation of Article III:4 of GATT 1994 and Article 2.1 of the TBT Agreement. For there to be a violation, it simply has to show that for a given product or series of given products there are one or more like products which enjoy more favourable treatment.

76. We wish to point out that in rejecting the analysis of likeness advanced by Canada, the European Communities go to great pains to affirm that PVC and ductile iron are not products “like” chrysotile fibre and chrysotile cement products. Interesting as it may be, however, this analysis is in no way relevant to the issue of whether glass fibre, cellulose fibre, PVA fibre and fibrous-cement products incorporating these types of fibres are “like” chrysotile fibre and chrysotile cement products.

Question 16: In its argument concerning Article 2.2 of the Agreement on Technical Barriers to Trade, Canada maintains that the ban on asbestos and products containing asbestos has no “rational link” with the objective pursued by France. Could Canada provide additional clarification on the distinction between this concept of “rational link” and the test of necessity which it defines as the second test of a measure’s conformity under Article 2.2?

77. Canada maintains that a technical regulation creates an unnecessary obstacle to international trade if the objective of the regulation is not legitimate or if the regulation is more restrictive than necessary to fulfil its objective, taking account of the risks non-fulfilment would create.

78. Once an objective is recognized as legitimate, it is then a matter of determining whether the technical regulation is a rational and necessary measure. The nature of the “rationality” and “necessity” of a technical regulation may vary according to the circumstances.

79. In our view, rationality and necessity entail the following elements. Firstly, the technical regulation must be carefully conceived to fulfil the objective concerned. It must be neither arbitrary, nor founded on irrational considerations. That is the rational link. Secondly, even supposing that there is a rational link, the technical regulation must be such that it is not more trade-restrictive than necessary to fulfil a legitimate objective, taking account of the risks non-fulfilment would create. These are the risks entailed by the absence of a technical regulation and minimum prejudice to trade.

80. With regard to the rational link, the point is whether the technical regulation is rationally linked to the fulfilment of the objective. In other words, is the technical regulation carefully designed to fulfil the objective in question. We feel that this stage of the analysis is essential in order to avoid justification of technical regulations which have tenuous links with the intended objective or even none at all. If this were not a necessary stage, a great many technical regulations could be justified without the slightest link with the objective.
81. With regard to the risks of having no technical regulations and minimal effect on trade, it is first necessary to try and establish if the absence of a technical regulation would create risks that the objective would not be fulfilled. To evaluate these risks, the relevant factors to be considered include scientific and technical data available and the intended end-uses of the products. One must then ask whether the prejudicial effects of the technical regulation are appropriate given the objective, in short, whether there is a less trade-restrictive alternative solution which would allow the intended objective to be fulfilled equally well.

82. It is only once the rational link between a technical regulation and the legitimate objective has been established that there is reason to examine the risks that might be created by the absence of that technical regulation and the existence of an alternative measure.

2. Questions by the Panel to the European Communities

Question 17: The EC states that “between 1945 and 1988, some 97 per cent of the asbestos consumed by France was chrysotile asbestos. From 1988 onwards, all the asbestos consumed in France was chrysotile asbestos”. What has been the pattern of consumption of hard and friable forms of chrysotile asbestos in France since 1945?

83. The question contains an ambiguity that needs to be removed. Thus, it is necessary not to confuse: (i) the variety of the fibres: there are different natural varieties of asbestos, in particular, the chrysotile variety which should be distinguished from the amphiboles. These fibres are incorporated in materials: (ii) the “fraility” of materials containing asbestos: these materials may be hard or friable, in varying degrees, depending on the application. In the French regulations, the fraility of a material containing asbestos is defined as follows: “A friable material is any material liable to emit fibres under the influence of impact, vibration or air currents.” Flocking (material composed of a mat of blown fibres) and lagging consisting of asbestos wadding or felt are typical friable materials. Asbestos-filled plastics and asbestos-cement products are non-friable materials. Before amphiboles were banned in the countries of the European Union, as in many other countries, there may have been friable products based on chrysotile and non-friable products based on amphiboles, and vice versa. It would be quite mistaken to associate amphiboles exclusively with friable products such as flocking and lagging and chrysotile exclusively with non-friable products such as asbestos cement. Since amphiboles were banned, all products containing asbestos - both friable and non-friable - have been based on chrysotile.

84. With regard to trends in asbestos fibre consumption and the division of the market between friable and non-friable products, the data supplied by the asbestos-processing industry indicate that over the period 1970-1975 (when both amphiboles and chrysotile were in use): (i) asbestos intended for very friable products (flocking, wadding), which do not form part of the statistics of the asbestos-processing industry, accounted for 10 to 20 per cent of imports; (ii) the remaining imports (80 to 90 per cent of the total), intended for processing, can be broken down as follows:

Non-friable products:

- 73 per cent for asbestos cement (compared with over 75 per cent in 1950);
- 8.4 per cent for floor coverings (production increased seven-fold between 1950 and 1975);
- 3 per cent for brakes (production increased seven-fold between 1950 and 1975);
- 1.9 per cent for mouldings (stable between 1950 and 1975);
- 1.3 per cent for gaskets;
- 2.4 per cent for miscellaneous applications (adhesives, mastics, mortars, etc.).
Friable products:
- 7 per cent for asbestos board and paper;
- 3 per cent for textile products (braiding, tape, coverings, etc.).

The use of very friable products (asbestos flocking and wadding) ended in France in 1978. No more asbestos fibres were incorporated in floor coverings after 1984. In the 90s (before the ban), asbestos-cement products, brakes and mouldings accounted for more than 90 per cent of the asbestos imported. Friable products such as textiles and board consumed less than 10 per cent.

85. The use of very friable products (asbestos flocking and wadding) ended in France in 1978. No more asbestos fibres were incorporated in floor coverings after 1984. In the 90s (before the ban), asbestos-cement products, brakes and mouldings accounted for more than 90 per cent of the asbestos imported. Friable products such as textiles and board consumed less than 10 per cent.

**Question 18:** The European Communities state that Canada “consumes little asbestos and thus exports the bulk of its production”. The European Communities also note, in paragraph 53: “... the fact that the increase in the frequency of mesothelioma-type cancers can be seen throughout Canada shows that the risk of death from chrysotile is not confined to the asbestos mining industry ... but that it affects all sectors of the economy”. To which sectors of the economy do the European Communities refer? Are statistics and studies available?

86. Per capita consumption in Canada is among the highest in the industrial world. Canada exports most of its chrysotile asbestos production (400,000 tonnes out of 450,000 in 1990). Thus, the proportion reserved for home use is very small. Nevertheless, Canada’s consumption is very considerable compared with other industrialized countries. Thus, in 1990, per capita consumption in Canada was distinctly higher than in France: (i) Canada: 2.05 kg per capita; (ii) Brazil: 1.26 kg per capita; (iii) France: 1.11 kg per capita; (iv) United States: 0.13 kg per capita. This high level of asbestos consumption in Canada explains the high and steadily increasing incidence of mesothelioma in that country.

**Question 19:** With respect to substitute products, INSERM has stated, in particular, that “suitable research work should be carried out and developed as a matter of urgency, before substitute fibres are generally introduced” (Effects on health of the main types of asbestos exposure, INSERM, 1997, p. 434) and that “taking into account the present uncertainties concerning the effects on humans of exposure to fibres used as a substitute for asbestos, it is important to ensure that the levels of exposure among users of products containing fibre substitutes for asbestos are as low as possible” (Effects on health of fibres used as a substitute for asbestos, INSERM, 1998, p. 34). In the light of these observations, can the European Communities explain the statement in paragraphs 140 et seq. of their written submission according to which “there are no data giving rise to concern as to the carcinogenicity of fibres used as a substitute for asbestos in cement fibres”?

87. There are studies available which show that in Quebec and the other areas investigated a wide range of economic sectors is affected by mesothelioma. Thus, a study by Siemiatycki in Montreal shows that the workers mainly at risk of mesothelioma are those exposed while working with materials containing asbestos. The results of the study show that construction sector workers run a risk of developing mesothelioma almost 12 times higher than the average. Another study by the Quebec Occupational Health and Safety Commission (CSST), reveals that the risk of mesothelioma has increased steadily in Canada since 1967. The study notes that the incidence of this disease is increasing especially rapidly in the maintenance sector.

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88. There are no data giving rise to concern as to the fibres used as a substitute for asbestos in cement fibres. First of all, it should be noted that asbestos cement is often replaced by products from another branch of technology (PVC plastics, ductile iron, various metals, etc.). This is the case, in particular, with pipes and roofing. In France, industry had decided to stop producing asbestos-cement pipes before the ban, because of the competition from PVC and ductile iron. When asbestos is replaced in fibro-cement, it is replaced by PVA, para-aramid or cellulose. It is never replaced in fibro-cement by man-made mineral fibres. PVA, cellulose and para-aramid fibres have been used for a very long time.
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without any sort of health warning having been given by occupational health specialists. PVA fibres have been used since 1930, para-aramid fibres for 30 years or so. As for cellulose, it has been in use for several centuries.\(^{25}\)

89. In 1996, these substances gave no cause for concern, and this has since been confirmed by studies conducted by CSTEE DG XXIV of the Commission of the European Communities and COC\(^{36}\) in the United Kingdom.\(^{27}\) Moreover, by the time the decision to ban asbestos was taken (July 1996), key facts had become available as a result of the G2SAT report on the comparative harmfulness of asbestos and man-made mineral fibres.\(^{38}\) This report established a hierarchy of risks as between chrysotile asbestos, ceramic fibres and mineral wools (glass wool, rock wool, slag wool) which made it impossible to avoid a global ban with exceptions.\(^{29}\) Thus, in 1996, France had to choose between asbestos, a known and proven human carcinogen, and substances used for decades without any problem ever having been reported.

90. INSERM’s concerns relate to certain man-made mineral fibres rarely used as substitutes for asbestos and France has taken INSERM’s recommendations into account. The expert opinion requested from INSERM related mainly to the fibres most under suspicion, i.e. man-made mineral fibres (ceramic fibres and mineral wools), whose harmfulness had been stressed by the opponents of a ban on asbestos. It confirmed the results of the G2SAT Report, as well as the classification adopted at the European level for ceramic fibres and mineral wools. France took into account the recommendations to proceed with caution and proposed an action plan for consideration by the social partners as soon as the results were published: (i) reminder of the regulations applicable, given the European classification; (ii) controls on fibre labelling; (iii) package of measures to monitor exposure levels, in particular among secondary users; (iv) establishment of groups of workers exposed to man-made mineral fibres for epidemiological follow up purposes.

Question 20: Why did asbestos cement remain outside the scope of Directive 91/659/EEC (paragraph 82 of the first submission of the European Communities)?

91. The purpose of the 1991 Directive 91/659/EEC was to ban, on the basis of the epidemiological data available at the time\(^{40}\), all varieties of asbestos other than chrysotile, which was then considered less dangerous than amphiboles. This directive also restricted the use of chrysotile to those products which were incapable of releasing asbestos fibres spontaneously into the air without special intervention and for which the possibilities of replacement had not yet been fully established. Since the publication of that directive, our understanding of the risk and the international database have expanded considerably\(^{41}\) and research on the replacement of the asbestos in asbestos cement products has led to solutions that pose no threat to the health of the user and are technically and economically viable.

Question 21: Is it possible to estimate the number of deaths, since 1945, due to chrysotile asbestos occurring in the categories which the submission of the European Communities described as “secondary” occupational, para-occupational and domestic? What type of exposure is responsible for the 25 per cent of mesothelioma cases in the building sector mentioned in paragraph 413 of the submission of the European Communities (construction, occasional handling, de-flocking, demolition, etc.)?

92. The number of deaths by mesothelioma among “secondary” users can be estimated at between 10 and 15 per million inhabitants per year. For mesothelioma, the spontaneous death rate, i.e. other than by exposure to asbestos, is about one to two cases per million inhabitants per year. It is true that, up to about the 1970s, the great majority of these deaths occurred among primary users (asbestos mining and processing).\(^{42}\) During that period the rate reached about five cases per million inhabitants per year in most of the industrial countries for which reliable health data are available (see Table 4, page 166 of the INSERM Report. All the studies show that those affected were almost exclusively so-called “secondary” users.\(^{43}\) The study by J. Peto et al.\(^{44}\) analyses mesothelioma mortality in England and Wales during the years 1979 to 1990. It shows that about 95 per cent of all the
deaths that occurred during that period concerned workers belonging to the “secondary” user group. In the industrial countries, this proportion of 95 per cent of all asbestos-related deaths has applied to this category of workers since about the 1970s. At present, among men, the mesothelioma death rate stands at about 15 to 20 deaths per million inhabitants per year: thus it may be estimated that in these countries 10 to 15 deaths per million inhabitants per year occur among male workers in the “secondary” user category. Even if a small fraction of these deaths can be attributed to exposure to amphiboles, it is none the less true that the number of deaths due to chrysotile is quite considerable.

93. The overwhelming majority of mesotheliomas occurring among construction workers are the result of occasional exposure to high peaks of asbestos. This can be illustrated by reference to the table headed “Distribution of Deaths from Mesothelioma by Occupation (Section III.B.4)” which gives a breakdown of the occupations most affected: the construction sector trades most affected are the carpentry, plumbing, electrical and related trades. Typically, workers in these trades have to handle asbestos-containing materials only intermittently. This is because of the numbers involved. To the workers in these construction trades it is necessary to add the workers in many other occupations: welders, dockers, laboratory technicians, fitters, upholsterers, power station workers, etc., who are only occasionally exposed to high peaks of asbestos and, taken together, account for the majority of deaths by mesothelioma.

Question 22: The article annexed by the European Communities (A. Gilg, et al., Estimation of the Past and Future Burden of Mortality from Mesothelioma in France, Occupational Environmental Medicine, 1998; 55: 760-765) estimates that between 1996 and 2020, about 20,000 men will die of mesothelioma. Is it possible to determine the different circumstances of exposure to asbestos that will have induced these 20,000 cases of mesothelioma?

94. Occupational exposure to asbestos in France concerns the approximately 20-25 per cent of men who have been exposed at least once in the course of their working life. This reflects the enormous variety of exposures. In France about 85 per cent of all the men exposed work in industrial production (mainly metallurgy, machines and appliances), building and public works or services, sectors characterized by occasional exposure to asbestos. It is these types of exposure that will cause the 20,000 deaths expected to occur among men in France between now and the year 2020.

Question 23: Have cases of death clearly attributable to occasional, but high exposure (“of the exposure peak” type) been documented?

95. All the studies based on the individual examination of mesothelioma cases, documented by a detailed study of the patient’s work history, have shown that since the 1970s the great majority of cases have occurred among workers who have never had any type of exposure other than occasional exposure with pollution peaks. The studies are mainly of two kinds:

- Studies relating to cases of acknowledgement of occupational illnesses, which always involve an in-depth examination of the circumstances of exposure, in particular, because of the financial consequences of acknowledgement (payments of benefits and pensions). Thus the above-mentioned study by the Quebec Occupational Health and Safety Commission (CSST) shows that during recent times the cases of mesothelioma acknowledged as being an occupational illness generally correspond to short-term exposure and most frequently involve servicing and maintenance workers.

- Epidemiological studies of the “case-control” type, in which each case of mesothelioma considered forms the subject of an in-depth examination of the victim’s entire working life by industrial health experts. Because of their high scientific quality, studies of this kind are particu-
larly suitable for assessing the circumstances of exposure. The recently published study by Iwatsubo et al.,50 concerning cases that occurred between 1987 and 1993 in France, is of this type. It, too, shows that almost all the mesotheliomas reported in France were observed in patients who had been subjected to only occasional exposure with pollution peaks. The previously mentioned study by Siemiatycki51 in Montreal, which is of the same case-control type, yielded similar results.

Question 24: The Comparative Table of the Characteristics of the Fibres Studied by the CSTEE (Section III.B.6) submitted by the European Communities, seems to indicate that the diameter of PVA, para-aramid and cellulose fibres could be greater than their length. Could the European Communities clarify these figures.

96. The fibres which must be taken into account in making a metrological assessment of a working environment have been defined by WHO52 in accordance with the following dimensional parameters: (i) more than 5 microns in length; (ii) less than 3 microns in diameter; (iii) ratio of length to diameter greater than 3 microns. The table in question is intended to show that the polyvinyl alcohol, cellulose and para-aramid fibres generally used in France for replacing chrysotile asbestos all have lengths distinctly greater than the 5 microns indicated by WHO; in any event, all the fibres in question have a length that is greater than the diameter. The diameters of these fibres are greater than 10 microns. This physically prevents them from penetrating into the pulmonary alveoli, which are accessible only to fibres less than 3 microns in diameter. Hence this characteristic of the substitute fibres eliminates the risk of their penetrating deep into the lungs. It is important to note that, generally speaking, unlike chrysotile fibres which have a diameter of between 0.1 and 1 micron and which separate lengthways into even finer crystalline fibrils (0.020 microns), the synthetic fibres used as substitutes, whether of organic or mineral origin, retain the diameter resulting from the manufacturing process throughout the life cycle of the fibre, even when released into the air from a material undergoing machining.

Question 25: The European Communities indicate that the substitute products using fibres shown to be harmless or less harmful include cellulose, PVA or aramid fibres. Could the European Communities give further details of the harmlessness or low toxicity of these products?

97. Cellulose, polyvinyl alcohol and aramid fibres have been used for a very long time, since well before asbestos was banned in France. There has been nothing to give cause for concern, (in particular, reports of cases of cancer among the workers exposed) which might have led to extensive research, whereas asbestos has been the subject of numerous scientific studies because of the large number of cases of disease found in workers over the last 70 years. The CSTEE committee of DG XXIV of the Commission of the European Communities and COC in the United Kingdom have made in-depth assessments of the comparative risks of asbestos and the fibres used to replace it, particularly in asbestos cement.

98. On the basis of an analysis of the studies and reports supplied by the Health and Safety Executive, COC53 concluded: (i) with regard to polyvinyl alcohol fibres, “these fibres will have no potential for the induction of lung cancer or mesothelioma” (…). The information on PVA suggests a low carcinogenic hazard”; (ii) with regard to para-aramid fibres, “although there is some evidence of adverse biological effects with para-aramids, there is no convincing evidence to suggest a carcinogenic hazard (…). The evidence suggests a lower carcinogenic risk than chrysotile”; with regard to cellulose fibres, “a recent investigation with cellulose fibres had documented evidence of a long biopersistence in the rat lung. However, the COC agreed that this study was not relevant to consideration of the question […] and that it was unlikely [to] identify a carcinogenic response attributable to cellulose fibres.” The COC concluded from its study that “the evidence presented to the committee on fibre dimensions, studies in animals including that of biopersistence in the lung, indicate that the carcinogenic risk posed by PVA fibres, para-aramid fibres or cellu-
lose fibres is likely to be less than that posed by chrysotile”. These conclusions are shared by CSTEE of DG XXIV 54 which found that “there is sufficient evidence that all forms of asbestos, including chrysotile, are carcinogenic in humans. There is no evidence of the occurrence of cancer induced by fibres in humans with respect to any of the three substitute products [investigated]”. CSTEE also notes that “pulmonary fibrosis is a well-known consequence of exposure to chrysotile [whereas], so far, no case has been reported among workers exposed to any of the three substitute products”.

Question 26: The European Communities maintain that the Agreement on Technical Barriers to Trade does not cover general prohibitions on the use of the product. In the present case, Decree No. 96-1133 provides for the possibility of exceptions to the ban. Do the European Communities consider that the Agreement on Technical Barriers to Trade is also inapplicable to these exceptions and to the provisions relating to them?

99. The French Decree provides that in certain circumstances a limited number of products may contain asbestos (see Art. 2(1) of the Decree). The European Communities consider that, like the general ban, the exceptions to the general ban do not constitute “technical regulations” in the meaning of the TBT Agreement. The reason is that the definition of a “technical regulation” requires that the document concerned must “lay down product characteristics or their related processes and production methods.” Article 2(1) of the French Decree does not lay down any particular characteristics. In particular, it does not stipulate any performance or design characteristics that such products must meet. All it does is to permit, provisionally and under justified conditions, the use of asbestos (in a general sense, not as a specific characteristic), in the absence of a substance that ensures equivalent performance. The exceptions in the French Decree deal with or are based on the specific characteristics of asbestos, not of the products that are allowed provisionally to contain it.

100. The provisions of the French Decree, which lay down the procedures for establishing which products may be allowed under an exception, also do not qualify as a technical regulation. Contrary to what Brazil argues, the inclusion of the term “applicable administrative provisions” in the definition of a “technical regulation” does not give independent meaning and existence to the term “applicable administrative provisions”, but simply clarifies that the definition includes such legislative provisions.

Question 27: The European Communities state that “many substitute products are not fibrous in texture”. Could the European Communities indicate, within the context of their analysis of “likeness” within the meaning of Article III:4, which parts of their argument relate to substitute products containing fibres and which to non-fibrous substitutes?

101. See paragraphs 102 and 103 below.

Question 28: The European Communities indicate that “Canada mentions only fibrous substitute products (cellulose, para-aramid, PVA), making no reference at all to non-fibrous products.” In the opinion of the European Communities, is it appropriate to take into account non-fibrous substitute products for the purpose of examining, in the present case, the likeness of the products under Article III:4 of GATT 1994 or Article 2.1 of the Agreement on Technical Barriers to Trade?

102. The European Communities have explained in their submissions (and in their reply to written question No. 29 of the Panel) that it is essentially the morphology of the chrysotile fibres (in terms of length and diameter/thickness) that determines their carcinogenic effects. As the mode of action of the fibres is through inhalation, the fibrous nature of the chrysotile asbestos is essential. In legal terms, the fibrous nature relates directly to the product’s properties, nature and characteristics and, probably, end-use. That probably explains why Canada has claimed “likeness” only with regard to substitute fibrous products. It is, therefore, probably not necessary for the Panel to determine whether non-fibrous substitute products are “like” products to chrysotile or chrysotile containing products.
103. The arguments in the European Communities’ first written submission relate primarily to the fibrous nature of chrysotile and chrysotile-containing products, on the one hand, and the fibrous substitute products, on the other. The European Communities have demonstrated that the fibrous substitute products are not like products to chrysotile or chrysotile-containing products on the basis of the four criteria normally used by Panels. It follows that, *a fortiori*, non-fibrous substitute products are also not “like” products, an additional reason being the even more marked difference in the physical characteristics or nature of the products concerned.

3. Questions by the Panel to both parties

Question 29: Does the term “chrysotile” cover a product of uniform quality (in particular, in terms of the length and quality of the fibres) or should it be assumed that there are different qualities of chrysotile and, therefore, different levels of risk?

(i) Reply by Canada

104. In strictly technical and commercial terms, there are various groups of chrysotile fibres which are classified according to their length. The method developed by the Association of Asbestos Mines of Quebec has been adopted by the majority of producer countries for the classification of chrysotile and lists the groups of fibres from the longest fibres (Group 1) to the shortest (Group 7). The length of fibre is normally measured in millimetres (10⁻³ metres). This classification is used to determine the strength of the fibre used; the longest groups of fibres are used for the manufacture of chrysotile cement products, while the shortest fibre groups are used for the manufacture of friction products, mastics and roof coatings.

105. This technical and commercial classification, however, has nothing to do with the level of risk to health, which should be evaluated on the basis of the criteria of durability (biopersistence), dose and dimension. The pathogenicity of the fibres is examined on the basis of their dimension: for a natural or synthetic fibre to be potentially harmful, it must have a length greater than five microns, a diameter less than three microns and a length to diameter ratio over 3:1 (a micron is equal to 10⁻⁶ metres).

(ii) Reply by the European Communities

106. Chrysotile, a natural mineral of the asbestos group, is a magnesium silicate of crystalline structure with the empirical formula “Mg₃(Si₂O₅)(OH)₄”. In chrysotile, as in all silicates, chemical elements other than magnesium may be present in low concentrations and variable proportions. Thus the chemical analysis of different samples of chrysotile may yield results which vary over the following ranges:

<table>
<thead>
<tr>
<th>Component</th>
<th>Range</th>
</tr>
</thead>
<tbody>
<tr>
<td>Silica (SiO₂)</td>
<td>38 to 42%</td>
</tr>
<tr>
<td>Alumina (Al₂O₃)</td>
<td>0 to 2%</td>
</tr>
<tr>
<td>Ferric oxide (Fe₂O₃)</td>
<td>0 to 3%</td>
</tr>
<tr>
<td>Ferrous oxide (FeO)</td>
<td>0 to 3%</td>
</tr>
<tr>
<td>Magnesium oxide (MgO)</td>
<td>88 to 42%</td>
</tr>
<tr>
<td>Calcium oxide (CaO)</td>
<td>0 to 3%</td>
</tr>
<tr>
<td>Sodium oxide (Na₂O)</td>
<td>0 to 1%</td>
</tr>
<tr>
<td>Nitrous oxide (N₂O)</td>
<td>11.5 to 13%</td>
</tr>
</tbody>
</table>

107. Finally, at the same geological site, the purity of a vein of chrysotile in the process of being mined may vary from one point to another. It may also differ between sites. A deposit of chrysotile asbestos (like any natural one) is never perfectly pure and may contain various types of impurities which will still be present, in whole or in part, in the asbestos-containing end-products. This applies, in particular, to quartz (silica) and other non-fibrous or fibrous silicates, such as amphibole asbestos.
108. Incidentally, chemical composition alone is not sufficient to identify a mineral; the physical structure is equally important. Thus, minerals such as lizardite and antigorite have exactly the same chemical composition as chrysotile but are not fibrous.

109. Under the heading “chrysotile” it is possible to distinguish between the forms that contain impurities (that is, a small proportion of tremolite, an asbestos of the amphibole type) and those that contain no impurities or only very small amounts. Despite attempts to show that only the traces of tremolite present in Canadian chrysotile are responsible for the numerous mesotheliomas among workers exposed to it, no difference in toxicity between these two forms of chrysotile has ever been proved. This theory of chrysotile being “inoffensive” with respect to mesothelioma has, moreover, been rejected by the scientific community as noted in the written submission of the European Communities. As far as we know, there is only one epidemiological study concerning Zimbabwe’s chrysotile which is reputed not to contain amphiboles: it shows a net excess of mesothelioma among the workers in chrysotile mines and mills. Thus, from the standpoint of the composition of chrysotile, there is no reason to believe that the toxicity of the pure forms with respect to mesothelioma would be any different from that of the forms that contain impurities.

110. It appears to be clearly established that it is essentially the morphology of the fibres, rather than their geological origin, that determines their carcinogenicity with respect to the lungs. These conclusions stem from observation of the very high risks to which groups of textile asbestos workers handling fine fibre chrysotile are exposed. (It should be noted that the asbestos used by some of these groups comes from Canada.) For their part, chrysotile “mine and mill” workers exposed to thicker fibres present clear but much lower risks.

111. As Landrigan notes, the risks of cancer of the lung can vary by a factor of 50 between a group of chrysotile miners and a group of industrial workers handling chrysotile asbestos from the same mines. Reference may also be made to the INSERM Report which compares the risks of cancer of the lung according to the industry using the chrysotile. The very considerable differences in risks observed are mainly attributable to differences in the morphology of the fibres, which are subjected to different forms of physical processing depending on the end-use: generally short and thick when mined, they may undergo changes in morphology and be made finer to adapt them better to particular uses.

**Question 30:** What are the output and import and export volumes of substitutes for chrysotile asbestos products, particularly asbestos cement, in France and Canada respectively?

**(i) Reply by Canada**

112. Without wishing to offer a full review of all the substitute products individually, the following is an indicative summary of the value of the products used as substitutes for chrysotile asbestos products (1997 data):

<table>
<thead>
<tr>
<th>PRODUCT</th>
<th>PRODUCTION</th>
<th>IMPORTS</th>
<th>EXPORTS</th>
</tr>
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(ii) Reply by the European Communities

113. In France, more than 90 per cent of imported chrysotile asbestos fibre was used for making asbestos-cement products. Since asbestos was banned, the market in asbestos cement products and materials has shifted towards: (i) materials and technologies which have existed for a very long time (for example, PVC and ductile iron for making pipes, aluminium for roofing); (ii) fibre substitutes for asbestos: mainly cellulose fibres and, to a lesser extent, polyvinyl alcohol and polypropylene fibres.

114. The first submission of the European Communities provides some statistical data on cellulose and polyvinyl alcohol fibres. Cellulose fibres are mainly intended for the textile and paper (pulp) industries. Each year, the fibro-cement industry uses less than one per cent of the total volume of cellulose fibres consumed in the various industrial sectors. In 1998, France produced 2,660,000 tonnes of cellulose fibres and imported 2,030,000 tonnes (including 660,000 tonnes from countries of the European Union and 740,000 tonnes from North America, of which 380,000 tonnes from Canada). It should be noted that imports of cellulose fibre from Canada increased by 14,000 tonnes between 1996 and 1998.

115. Polyvinyl alcohol fibres, which are mainly used in the textile and packaging industries, are made by only two factories in the world, one in China and the other in Japan. In 1998, France recorded a trade deficit of 700 tonnes or 115 million francs. In 1998, deliveries from French manufacturing units amounted to 4,200 tonnes and imports to 11,400 tonnes. As far as other products are concerned (the remaining ten per cent of chrysotile asbestos fibres consumed), the most popular substitute has been man-made mineral fibres of the glass or rock fibre type (the great majority of applications), with very limited use being made of para-aramid and ceramic fibres (these two products are very expensive as compared with asbestos fibres). With regard to para-aramid fibres, in 1998, France recorded a trade deficit of 400 tonnes or 26 million francs. Man-made mineral wools and fibres are mainly produced in France and principally intended for insulating buildings. It should be noted that less than one per cent of the output of these man-made mineral wools and fibres is now being used in products previously manufactured with asbestos.

Question 31: Are there statistical data available which indicate what are the positive effects of controlled use, the use of encapsulated products and the banning of chrysotile asbestos, respectively, as regards reducing the number of mesotheliomas and cancers of the lung?

(i) Reply by Canada

116. The sources of asbestos-related pathologies have been identified as utilization of amphibole varieties in friable products such as flocking or cases of exposure to high quantities of chrysotile. Controlled use, a practice that was gradually introduced in the 1970s, has eliminated these dangerous sources. However, since there is a latency period of some decades between exposure to asbestos and the appearance of disease, pathologies related to uncontrolled exposure occurring up to the 1970s will continue to appear for a few decades yet, even if the sources of that exposure have now disappeared. Furthermore, as long as the competent authorities do not ensure appropriate and systematic management of the flockings installed during the years of uncontrolled use, the associated problems will not be completely eliminated.

117. There are few statistics on the prohibition of flockings and the exclusive use of chrysotile at low levels of exposure. Data collected and compiled by the Asbestos International Association (AIA) in 1995 cover 28 countries with some 25,000 workers employed.62 In these countries, because of the introduction of controlled-use procedures and measures, the A.I.A. surveys show that 97.3 per cent of workers are exposed to less than 1.0 f/ml. These exposure levels satisfy the recommendations on worker protection made by the Group of Experts which met under the auspices of the WHO in Oxford, England, in April 1989. The findings of a similar survey conducted in 1997 will be available in the summer of 1999.
118. It should also be made clear that there is little data for assessing the positive effects of controlled and exclusive use of chrysotile asbestos, and that the reference period does not begin until the late 1970s when the current workplace controls were implemented (plants in a few countries for the manufacture of asbestos cement and friction materials).

119. Other data have also been published on groups of workers in chrysotile asbestos mines in Quebec, which feature both a sufficiently high number of workers and a sufficiently long observation period to measure the effects of asbestos exposure. The study covers cohorts formed in 1966 and monitored since then, including more than 11,000 workers born between 1891 and 1920. The most recent update, published in 1997, reports data until 1992. The findings for workers in this group who were exposed to chrysotile concentrations of up to 22 f/ml for 40 years led the authors to conclude “from the point of view of mortality that exposure in this industry to less than 300 mpcf. years [i.e. equal to about 22 f/ml over 40 years] has been essentially innocuous.”

(ii) Reply by the European Communities

120. To the best of our knowledge, the only published study that enables the effectiveness of controlled use to be measured is that issued in 1996 by the Health and Safety Executive in the United Kingdom concerning the risks of cancer incurred by asbestos workers after 1969, when the “safe” use of asbestos was adopted by the United Kingdom. This study shows that, despite strictly “controlled” use (since the study relates exclusively to workers making asbestos-based products), there is a significant net excess of cancers among those who worked only under the “controlled” use regime. It follows from this finding that “controlled” use does not make it possible to prevent deaths from cancer, even in specific industrial manufacturing sectors with a workforce which is relatively small and a priori easy to train and supervise.

121. For at least 40 years, the manufacture of asbestos cement has involved “encapsulating” asbestos in cement. This encapsulation does not guarantee the harmlessness of asbestos cement during use: in practice, when asbestos cement is used, in an occupational, para-occupational or domestic environment, it is generally sanded, crushed or sawn and releases carcinogenic fibres in the form of dust. The occurrence of mesotheliomas among workers exposed to asbestos fibres which, in the manufacturing stage, were encapsulated in cement clearly shows that this process offers absolutely no protection against the carcinogenic effects of chrysotile fibres released when products and materials containing “encapsulated” asbestos are worked. This is perfectly understandable considering the very high levels of exposure measured under these conditions. It is possible to encounter concentrations several tens, even several hundreds of times higher than the national statutory limits and the internationally recommended levels.

122. The number of asbestos-related illnesses in a country is very strongly correlated with the amount of asbestos imported into that country. The most effective way of reducing, in the future, the number of asbestos-related illnesses is therefore to reduce asbestos imports. A ban with exceptions is the most effective means of achieving that result. France has demonstrated this, since imports fell very rapidly after the implementation of the ban (an estimated 1,200 tonnes in 1997, 200 tonnes in 1998, and a projected 55 tonnes in 1999 as against 35,000 tonnes in 1995). Thus, France could not afford to wait another 30 years in order to verify whether tighter controls on so-called “safe” use would make it possible to achieve the same results.

Question 32: With regard to substitute products is there a difference in potential or proven risk between fibrous and non-fibrous products?

(i) Reply by Canada

123. The replacement of fibre products with products containing no fibre might imply that the manufacture and use of the latter are risk free. This is not the case, at least for certain non-fibrous products being proposed as substitutes for chrysotile asbestos prod-
ucts. Only a comparative, case-by-case evaluation can yield relevant information. For example, let us take the alternative of PVC pipes. It should be realized that manufacture of the vinyl chloride monomer (a proven carcinogen which is later polymerized in PVC) involves the use of chlorine, an element which is later incorporated in an organic molecule. It has been well established that the synthesis of organo-chlorines is a significant source of dioxins, substances whose medium to long-term effects are very harmful, and regarding which the WHO has recommended a threshold of exposure not exceeding 10pg/kg./day (approximately 220ng/year).

124. The fact that the production of ductile iron also poses health risks recognized by the IARC as Group I is something which should not be concealed, since it entails greater energy consumption leading to the emission of carcinogens such as polynuclear aromatic hydrocarbons (PAH), etc.. These two examples clearly show that there are virtually no products or technologies that offer zero risk: one must learn to control and manage the risks within the framework of a policy of controlled and disciplined use. In any case, only a case-by-case analysis can answer the question asked.

(ii) Reply by the European Communities

125. The carcinogenic effect of asbestos is linked with its inhalation. In fact, not only does asbestos have a length to diameter ratio that enables it to penetrate into the pulmonary alveoli but, in addition, asbestos fibres are all capable of separating lengthways into even finer crystalline fibres, including inside the bronchial tree, which helps them to penetrate into the deepest recesses of the lungs. Many of the products used to replace asbestos cement rely on alternative technologies which do not use fibres. Accordingly, these products, which cannot be inhaled, do not carry any threat of carcinogenicity. Non-fibrous products such as ductile iron and PVC have never been suspected of being carcinogenic and no case of cancer among the workers exposed has been recorded, although the products have been used for a very long time for a wide variety of purposes. The dust released when they are machined (against which protection should in any case be provided) is considered to be without specific toxicity for the human organism (ductile iron or plastic dust).

126. In assessing the carcinogenicity of fibrous products, WHO takes into consideration, in particular, the ratio of length to diameter which makes it possible or impossible for the fibres to penetrate into the pulmonary alveoli. Fibres are used to replace asbestos in asbestos cement. With the exception of man-made mineral fibres, none of the fibres used to replace chrysotile has a length to diameter ratio that enables them to penetrate into the lungs. In fact, polyvinyl alcohol and para-aramid fibres are 2 to 8 mm. (i.e. 2,000 to 8,000 microns) in length and 10 to 16 microns in diameter. Only cellulose fibres, which are 12 to 40 microns in diameter, can give rise to finer particles which are said to irritate the respiratory pathways, but which are never carcinogenic. None of the fibres sometimes used to replace chrysotile asbestos (glass or rock man-made mineral fibres, organic fibres such as aramid fibres, plant fibres such as cellulose) are capable of separating lengthways: fibres extracted from the material retain their original diameter, which is almost always greater than the basic diameter of asbestos fibres.

127. Thus, scientifically, a consensus exists in favour of the following classification based on risk: (i) asbestos, of both the amphibole and chrysotile varieties, is a proven human carcinogen; (ii) some fibrous substitute products, such as ceramic fibres, are suspected of being carcinogenic: their carcinogenicity has been revealed by animal studies, but there is no evidence of their being carcinogenic in humans; (iii) the other fibrous products used instead of asbestos (PVA, cellulose, para-aramids, glass and rock fibres) are not human carcinogens. There have been no studies designed to determine whether or not they are carcinogenic in animals; (iv) the non-fibrous products have never been suspected of being carcinogenic. It should be noted that the use of the substitute products which are suspected of being carcinogenic (ceramic fibres) is strictly limited to applications for which there is at present no less dangerous substitute that would ensure equivalent performance. In any case, their use is regulated.
Question 33: Is the concept of necessity in Article XX:(b) of GATT the same as that in Article 2.2 of the Agreement on Technical Barriers to Trade?

(i) Reply by Canada

128. With regard to ‘necessity’, the text of Article 2.2 of the TBT Agreement is not identical to the text of Article XX:(b) of GATT 1994. Article XX:(b) uses the phrase, “necessary to protect [...]”. Article 2.2 of the TBT Agreement evokes the concept of necessity in a more specific context: “not more trade-restrictive than necessary to fulfil a legitimate objective, taking account of the risks non-fulfilment would create”. Article 2.2 in particular prescribes that account be taken of available scientific and technical information and the intended end uses of the products.

129. Despite the differences in textual formulation, the case law that has developed around analysis of Article XX:(b) can be helpful in determining the substance of the concept of necessity in Article 2.2. Conversely, the avenues for evaluating the ‘necessity’ of the measure contained in 2.2 (i.e. use of scientific information and consideration of end uses) may be relevant to analysis of necessity under Article XX:(b).

(ii) Reply by the European Communities

130. All the Panel reports that have examined the concept of necessity in the context of Article XX:(b) of GATT have concluded that it was not the necessity of the policy goal pursued by the measures at stake that was to be examined, but whether or not it was necessary to subject the imported products to the legal regime of the contested measure. In all cases a measure inconsistent with another provision of GATT was found to be not “necessary” if an alternative measure which the defending Member could reasonably be expected to employ and which was not inconsistent (or less inconsistent) with other GATT provisions was available to it. The same should apply as regards the concept of necessity in the context of Article 2.2 of the TBT Agreement. This results clearly from the ordinary meaning of the terms of Article 2.2 in their context, which state that: “For this purpose, technical regulations shall not be more trade-restrictive than necessary to fulfil a legitimate objective, taking account of the risks non-fulfilment would create”. The sixth preambular paragraph to the TBT Agreement clarifies that Members have the autonomous right to determine the level of human health protection they consider appropriate in their territory. The necessity test, therefore, in Article 2.2 may only be applied in order to examine whether the technical regulations have been prepared, adopted or applied “with a view to or with the effect of creating unnecessary obstacles to international trade”. An obstacle to trade will be found to be unnecessary, if another measure consistent (or less inconsistent) with GATT is reasonably available to the defending Member, to enable it to achieve the level of health protection which it has determined.

131. The fact that, in substance, the necessity test should be applied in the same way in both provisions does not, however, have the same procedural implications as regards, for instance, the burden of proof. Unlike Article XX:(b) of GATT 1994, where initially the burden is on the Member claiming its application to prove it, in the context of Article 2.2 of the TBT Agreement, the burden is initially on the plaintiff Member to establish that there has been a violation. Indeed, the plaintiff Member must first demonstrate the availability of an alternative consistent or less inconsistent measure which can achieve the level of health protection determined by the defending Member:73 The European Communities have already explained that so-called “controlled use” is not “an alternative measure” because, inter alia, it does not provide the same level of protection as the French Decree. The European Communities would, moreover, recall that, as is clear from the factual part of their two written submissions, it is “in the real world” not feasible to make sure that controlled use is applied in all circumstances where people work with asbestos in such a way as to achieve the level of sanitary protection determined by France.

Question 34: Is the notion of “like product” within the meaning of Article III:4 of GATT 1994 identical with that of “like product” within the meaning of Article III:2, first sentence of GATT 1994?
The case law of GATT and the WTO indicates that the concept of “like product” in Article III:2 is to be construed narrowly. However, this narrow interpretation does not apply to Article III:4 where the concept of likeness must be construed more broadly, given the purpose and context of Article III:4. Consequently, Article III:4 encompasses a more extended “range of like products” than does Article III:2, first sentence.

The report of the Appellate Body in Japan – Taxes on Alcoholic Beverages, in its discussion of Article III:2 commented precisely on the relative character of likeness under various articles and under various agreements.

“No one approach to exercising judgement will be appropriate for all cases. The criteria in border tax adjustments should be examined, but there can be no one precise and absolute definition of what is “like”. The concept of “likeness” is a relative one that evokes the image of an accordion. The accordion of “likeness” stretches and squeezes in different places as different provisions of the WTO Agreement are applied. The width of the accordion in any one of those places must be determined by the particular provision in which the term “like” is encountered as well as by the context and circumstances that prevail in any given case to which that provision may apply.”

This approach should be helpful in identifying on a case-by-case basis the range of “like products” that fall within the narrow limits of Article III:2, first sentence in GATT 1994. Yet this approach will be most helpful if decision-makers keep ever in mind how narrow the range of “like products” in Article III:2, first sentence, is meant to be, as opposed to the range of “like” products contemplated in some other provisions of GATT 1994 and other multilateral trade agreements of the WTO Agreement.

Using the analogy of the accordion of likeness, the Panel in Japan – Taxes on Alcoholic Beverages pointed out that the same interpretation of the term “like product” in Article III:2 and Article III:4 would give a different scope to two paragraphs in the same Article. Two paragraphs of the same Article designed to protect the conditions of competition for products imported into the territory of a WTO Member should not have a different scope. Since the terms “directly competitive” or “directly substitutable” add to the term “like product” in defining the scope of Article III:2, interpreting “like product” in III:4 in the same way as in III:2 would give a narrower overall scope to Article III:4 than to Article III:2. In Japan – Taxes on Alcoholic Beverages, the Panel wrote:

“6.20 The Panel noted that the term “like products” appears in various GATT provisions. The Panel further noted that it did not necessarily follow that the term had to be interpreted in a uniform way. In this respect, the Panel noted the discrepancy between Article III:2, on the one hand, and Article III:4 on the other: while the former referred to Article III:1 and to like, as well as to directly competitive or substitutable products (see also Article XIX of GATT), the latter referred only to like products. If the coverage” of Article III:2 is identical to that of Article III:4, a different interpretation of the term “like product” would be called for in the two paragraphs.”

What emerges from this Panel report is that the overall scope of paragraphs 2 and 4 of Article III should be identical. The Panel continues in paragraph 6.20:

“Otherwise, if the term “like product” were to be interpreted in an identical way in both instances, the scope of the two paragraphs would be different. This is
precisely why, in the Panel’s view, its conclusions reached in this dispute are relevant only for the interpretation of the term “like products” as it appears in Article III:2.”

137. It is therefore clear that the analysis of the term “like product” was valid only for III:2, because the Panel did not want to give a different overall scope to the two paragraphs. The concept of “likeness” must therefore have a broader scope under Article III:4 than under III:2. To maintain the same scope in the two paragraphs, the accordion of “likeness” must be tightly squeezed in Article III:2 and stretched in Article III:4. In Japan – Taxes on Alcoholic Beverages, the Appellate Body supports the Panel’s analysis and rejects the appellants’ interpretation that the scope of Article III:2 (first and second sentences) and of Article III:4 is not identical:

“We note the argument on appeal but the Panel suggested in paragraph 6.20 of the Panel report that the product coverage of Article III:2 is not identical to the coverage of III:4. That is not what the Panel said.”

138. This interpretation of Canada’s is consistent with doctrine on the subject. For example, Edmund McGovern observed:

“[…] the Appellate Body in the 1996 Japanese Alcohol case doubted whether [similar products] has the same meaning even among the various paragraphs of Article III. In particular, the narrow interpretation appropriate in the first sentence of paragraph 2 (concerning internal taxation) does not necessarily apply in the context of paragraph 4 (concerning internal regulations), and it is possible that “like products” in paragraph 4 might even have the same scope as “directly competitive and substitutable products” in paragraph 2. […] Consequently, the point has to be discussed separately with regard to each provision.”

139. Canada therefore maintains that the test of likeness in Article III:4 is different from and broader than that developed in Article III:2.

(ii) Reply by the European Communities

140. The European Communities consider that for the purpose of the present dispute the reply to this question is of theoretical relevance only. Canada does not claim a violation of Article III:2 of GATT and it is clear from the submissions of the European Communities that only paragraph 4 of Article III of GATT might be of relevance in this case. The European Communities have demonstrated, however, that the French Decree does not discriminate either de jure or de facto between national and imported products. It has been consistently held by Panels and the Appellate Body that the concept of “like product” does not necessarily have the same meaning in all GATT provisions in which it occurs and that its meaning “must be determined by the particular provision in which the term “like” is encountered as well as by the context and the circumstances that prevail in any given case to which that provision may apply.” The European Communities have explained in their submissions that asbestos and asbestos-containing products have very different physical characteristics from products that do not contain asbestos and, therefore, cannot be considered “like products.” Article III:4 does not cover “directly competitive or substitutable” products and, according to the Appellate Body, such an omission must have some meaning.

Question 35: What is the relationship between GATT 1994 and the Agreement on Technical Barriers to Trade? Does the nature of this relationship have any effect on the order in which a panel should examine a measure alleged to constitute a violation of both GATT 1994 and the Agreement on Technical Barriers to Trade? Does this relationship have any particular bearing on the present dispute?
(i) Reply by Canada

141. The nature of the relation between the different agreements in Annex 1A to the Agreement Establishing the World Trade Organization can be determined on the basis of the wording of the General Interpretative Note to Annex 1A (Annex 1A). This Interpretative Note was introduced in order to reflect the relation between GATT and the 12 other Agreements constituting Annex 1A. This Note is directly modelled on the customary rule of interpretation of international public law whereby, in the event of conflict, specific provisions prevail over more general rules. The Note recognizes that the 12 Agreements are, relative to GATT, provisions concerned with the detailed preparation and application of the more general provisions of GATT. These Agreements constitute the most recent and most specific expression of the WTO Members as to what the interpretation and application of these disciplines should be.

142. In the case of the Agreement on Technical Barriers to Trade, this is confirmed by the wording of its Preamble, which states, among other things: “Desiring to further the objectives of GATT 1994”,80 In this sense, then, the TBT Agreement is the most recent and the most specific expression of the WTO Members as to the necessary interpretation and application of the general disciplines of GATT (e.g. the obligations of Article I, III, X and XI in view of the exceptions provided for in Article XX) in the context of technical regulations and standards. The more specific character of the TBT Agreement does not have the effect of setting aside the more general applicable disciplines of GATT, and hence the two Agreements apply simultaneously and must be examined separately. This rule was clearly recognized by the Appellate Body in the Bananas III case where it stated that:

“Although Article X:3(a) of GATT 1994 and Article 1.3 of the Licensing Agreement both apply, the Panel, in our view, should have applied the Licensing Agreement first, since this Agreement deals specifically, and in detail, with the administration of import licensing procedures”.

143. The same idea was revisited by the Panel in the Indonesian Automobiles case, when it was dealing with the issue of simultaneous application of the Agreement on Trade-Related Investment Measures and Article III of GATT (see the report, paragraph 14.62). With regard to the order in which the Agreements should be considered by the Panel, since Article 2 of the TBT Agreement deals in a more specific and detailed fashion with the preparation, adoption and application of technical regulations and standards, giving due regard to what WTO Members consider to be an acceptable balance between the various interests protected by the general provisions of GATT 1999, it seems to us that the TBT Agreement must be examined first. This position is in keeping with the approach adopted by the Appellate Body in the Bananas III case and by the Panel in the Indonesian Automobiles case.82 Therefore the TBT Agreement should be examined first.

(ii) Reply by the European Communities

144. GATT and the TBT Agreement are two legally distinct Agreements. The General Interpretative Note to Annex 1A to the WTO Agreement clarifies that, in the event of conflict between the two, the provisions of the TBT shall prevail to the extent of the conflict. The object and purpose of the TBT Agreement, like its predecessor Agreement, is “to further the objectives of GATT 1994” (second preambular paragraph) in the areas of international standards and conformity assessment systems so as to ensure that technical regulations and standards do not create unnecessary obstacles to international trade (third to fifth preambular paragraphs). The European Communities consider that the legal relationship of the two agreements as explained above does not dictate any particular order in which the Panel should examine the claims and arguments of the parties in this dispute. The two options of examining first GATT and then the TBT Agreement or vice versa are both theoretically available.

145. The European Communities note, however, that several important concepts (such as the concept of like products, the principle of non-discrimination or the concept of neces-
ity) are found in both Agreements, but there is very little case law and practice of the Members under the TBT Agreement on which the Panel may draw. It may therefore be more prudent, from the interpretative point of view, to proceed first with an analysis under GATT, especially when the two parties disagree on the applicability of one of the two agreements, in this instance the TBT Agreement. This approach is, in any case, not unusual, as is shown by the US Gasoline Panel and Appellate Body reports. The choice of the order by which the claims of the parties under these two Agreements will be examined does not appear to have any particular or significant implication for this dispute, other than that the Panel should ensure consistent interpretation of the provisions of the two Agreements. As regards the separate issue of allocation of the burden of proof, see the European Communities’ reply to Canada’s written question No. 8.

Question 36: Do the exceptions provided for in Article XX of GATT 1994 apply to violations of provisions of the Agreement on Technical Barriers to Trade? Could this question have any bearing on the present dispute?

(i) Reply by Canada

146. No, the exceptions provided for in Article XX of GATT are not intended to apply to violations of the provisions of the TBT Agreement. Article XX is not applicable outside GATT 1994 unless there is a specific stipulation to the contrary, as in the TRIMs Agreement. The text of the TBT Agreement makes no reference to Article XX of GATT.

(ii) Reply by the European Communities

147. Yes, the European Communities consider that the basis of the exceptions in Article XX are applicable under the TBT Agreement. This means that the substantive grounds on which an exception can be based under Article XX(b) of GATT 1994 are also available under the TBT Agreement. This derives from a systematic interpretation of the TBT Agreement and GATT, in particular the sixth preambular paragraph of the TBT Agreement, and the history of the preparatory work. The European Communities consider that the availability under the TBT Agreement of the reasons behind the exceptions in GATT Article XX(b) does not appear to have any particular or significant implication for this dispute, other than that the Panel should ensure consistent interpretation of the provisions of the two Agreements. As regards the separate issue of allocation of the burden of proof, see the European Communities’ reply to Canada’s written question No. 8.

Question 37: What are the factors that determine the “relevance” of an international standard within the meaning of Article 2.4 of the Agreement on Technical Barriers to Trade?

(i) Reply by Canada

148. The “relevance” of an international standard is a question of fact which must be determined on a case-by-case basis.

149. The ordinary meaning of the word “pertinent” is given in Larousse as “approprié, qui se rapporte exactement à ce dont il est question.” The English version of the TBT Agreement uses the word “relevant”, which is defined by the Concise Oxford Dictionary as: “bearing on or pertaining to the matter at hand.”

150. In this case, the international standards cited by Canada are relevant since they relate to the same product, namely asbestos, and to the same regulatory purpose, namely protection of the health of workers and individuals. The international standards all deal with the use of chrysotile in a controlled and safe manner.

151. The Panel must also be aware that the relevance of the international standards invoked by Canada has never been challenged by the European Communities. Rather, the European Communities confine themselves to attempting to demonstrate that the interna-
tional standards cited by Canada are not “international standards” within the meaning of the TBT Agreement.

(ii)  Reply by the European Communities

152. The phrase in question reads: “[w]here technical regulations are required and relevant international standards exist or their completion is imminent (…)”. The ordinary meaning of “relevant” is bearing on or pertaining to the matter at hand. The context of this provision, however, makes it clear that the substantive requirement to judge the relevance of an international standard is that the latter should meet the requirements of the definition of “technical regulation” in Annex 1 to the TBT Agreement (see Article 1.2 of the TBT Agreement).

153. In the present case, none of the international standards invoked by Canada define the characteristics which asbestos or products containing asbestos must have. They deal, for instance, with the way in which asbestos and asbestos-containing products must be handled in the workplace and the relationship between employers and employees. They are, therefore, not “relevant” in the meaning of Article 2.4 of the TBT Agreement. In any case, even if they were relevant (which they are not) they would be completely ineffective and inappropriate to fulfil the legitimate objective of France, which is to protect human health in its territory.

Question 38: In the context of the Agreement on Technical Barriers to Trade, is a Member free to determine, for the purpose of drafting and adopting a technical regulation, the level of protection it considers appropriate?

(i)  Reply by Canada

154. Yes, but in compliance with the obligations of the TBT Agreement. For example, a Member’s freedom to adopt a regulation for the protection of human health, at the levels it considers appropriate, is mentioned in paragraph 6 of the preamble to the TBT Agreement. However, this freedom is circumscribed. First, it is subject to the requirement that the technical regulation does not constitute either a means of arbitrary or unjustifiable discrimination between countries where the same conditions prevail or a disguised restriction on international trade. Second, it is subject to the technical regulation being otherwise in accordance with the provisions of the TBT Agreement.

(ii)  Reply by the European Communities

155. Yes. Article 2.2 of the TBT Agreement provides that protection of human health and safety is a legitimate objective. The preamble to that agreement confirms that “no country should be prevented from taking measures necessary to … protect human … life or health, at the levels it considers appropriate (…)”.

Question 39: Article 2.4 of the Agreement on Technical Barriers to Trade envisages the situation in which the relevant international standards or the relevant parts of them would be “an ineffective or inappropriate means for the fulfilment of the legitimate objectives pursued, for instance because of fundamental climatic or geographical factors or fundamental technological problems”. What other types of situation would be liable to make the standards or some of their parts “ineffective or inappropriate” within the meaning of this article?

(i)  Reply by Canada

156. Situations that might render international standards or certain parts of them ineffective or inappropriate, other than those situations explicitly mentioned in Article 2.4, should be examined in the light of the ordinary meaning of the terms in this article, taken in their context. The text of Article 2.4 states that they must be “fundamental” factors or problems.
157. For Article 2.4 not to be stripped of its meaning, it requires that, in order to set aside an otherwise relevant international standard, the Member must provide real and tangible evidence of a “fundamental” consideration and not a mere allegation that certain standards are not appropriate.

(ii) Reply by the European Communities

158. The most obvious examples are when the latest scientific evidence suggests that the scientific basis of an international standard is inaccurate or obsolete, or when the level of protection that could be achieved by the international standard is lower than that determined by a Member in its territory. In the present case, the relevant scientific evidence that became available, in particular in the late 1980s and early 1990s, indicates that: (i) there is no doubt at all that chrysotile asbestos is a proven carcinogen; (ii) there is no safe exposure limit (threshold) for chrysotile asbestos and products containing asbestos; (iii) so-called “controlled use” is not applicable in all circumstances and for all types of persons that may come in contact with asbestos or asbestos-containing products and, in addition, does not eliminate all the risks; and (iv) there are substitute products that are safe, or safer than chrysotile asbestos.

159. Under these circumstances it is clear that an international standard that would permit the use of asbestos or asbestos-containing products or would set an exposure limit or would recommend “controlled use” would be ineffective or inappropriate to achieve the level of health protection determined by France.

4. Questions by Canada to the European Communities

Question 1: The first written submission of the European Communities states that “mesothelioma is a pleural cancer for which the only known cause is the inhalation of asbestos”. Are we to take this assertion to mean that the European Communities are unaware of the scientific data identifying x-rays and erionite, inter alia, as causes of mesothelioma?

160. Erionite is indeed an indisputable cause of mesothelioma (as was made clear in the INSERM Report: see pages 125-126): however, exposure to the fibres of erionite, a mineral of natural origin, has to our knowledge been documented only in the Turkish region of Cappadocia. For that reason, in Section 4(a) of the first submission by the European Communities, a more complete form of words was used: “apart from exposure to asbestos, no other causal factor present in the industrialized countries has been established or even seriously suspected”. As for the other factors mentioned by Canada, such as x-rays, none has been positively confirmed up to now; even though presumptions exist in the case of some; none of them is classified in Group 1 of the International Agency for Research on Cancer as a proven carcinogen with regard to mesothelioma.

Question 2:

(a) In paragraph 238 of their first written submission, the European Communities conclude that the controlled use policy is inapplicable. Was France applying a policy of controlled use of asbestos at the time when the workers referred to in the study by Y. Iwatsubo were exposed? If not, how can this policy be found to be inapplicable?

161. The study by Iwatsubo et al., concerns workers who developed mesothelioma in recent years (1987-1993) and who have therefore been exposed to asbestos during a period dating back at least 20 or 30 years, when so-called controlled use was not applied. However, the study in question (like others cited in the European Communities’ submission, which were carried out in different countries), confirms that the vast majority of such cases of mesothelioma occurred in a wide variety of trades, particularly in the construction sector, where workers are in most cases unaware of their sporadic exposure due to having to handle a wide range of materials that have frequently been in place over a very long period of time without anyone knowing of their asbestos content.
162. The study by Iwatsubo et al., like similar studies, has the advantage of showing clearly that “controlled use” procedures would have to be applied on so wide a scale (several hundred thousand workers a day in France carry out isolated operations on materials containing asbestos) that they are in practice largely unrealistic although every effort must be made to encourage such procedures given the enormous quantities of asbestos that have been imported for decades and which remain in place. Moreover, the prevailing unawareness that the materials concerned contain asbestos makes it even more difficult to introduce such work procedures systematically: it would mean in practice putting a supervisor behind every worker in the construction trade and numerous other sectors of activity which use heavy equipment covered by the ISO 7337 Standard, and prohibiting any operation until checks have been carried out on the presence of asbestos. Specifically, that would mean having to send a sample of the material on which a sometimes very brief operation is to be carried out (such as drilling or sawing) to an approved laboratory and awaiting the laboratory findings. It is clear that the procedures described above cannot be applied on such a broad scale, uniformly and continuously, i.e. over a period of decades given the continued presence of asbestos.

Question 2:

(b) Could the European Communities specify the varieties of asbestos to which the persons covered by the study by Y. Iwatsubo et al. were exposed?

163. The study by Iwatsubo et al. does not distinguish between the varieties of asbestos to which workers suffering from mesothelioma were exposed. However, France has used chrysotile almost exclusively: the share of amphiboles in asbestos imports into France has not exceeded 3 per cent since 1945, and France has never produced amphiboles domestically. Amphiboles were used mainly for specific purposes: it is therefore highly likely that the great majority of workers with mesothelioma covered by the study by Iwatsubo et coll. were never exposed to any variety of asbestos other than chrysotile.

Question 3: The European Communities acknowledge in Section 4(a) of their first written submission that there is a difference between the toxicity of chrysotile and that of amphiboles. Do the European Communities therefore acknowledge that assessment of the risk from exposure to chrysotile exclusively must be based solely on data concerning chrysotile exposure, and not on data concerning exposure to amphiboles or mixtures of asbestos containing amphiboles?

This question calls for two preliminary comments:

- The European Communities pointed out in their first written submission that, while it is true that amphiboles appear to be a more important cause of mesothelioma than chrysotile, that is not the case for lung cancer. However, lung cancer has caused a higher number of asbestos-related deaths than mesothelioma (many authors consider that for every death from mesothelioma there is at least one or even possibly two deaths from cancer of the lung due to asbestos). It is therefore impossible to accept the wording of the question by Canada which, once again, seems to be seeking to ignore this fact.

- Canada does not explain what is meant by “exposure to chrysotile exclusively”: the chrysotile produced by Canada is contaminated by tremolite (a variety of amphibole asbestos with a very strong carcinogenic potential in respect of mesothelioma). This argument has also been widely used by the “defenders” of chrysotile to dispute the fact that the latter could cause mesothelioma (the reasoning being that the traces of tremolite contained in Canadian chrysotile were solely responsible for the many cases of mesothelioma observed among workers exposed to Canadian chrysotile asbestos). This theory of the harmlessness of chrysotile in relation to mesothelioma has been rejected by
the scientific community, as is pointed out in the submission of the European Communities. It is hard to see what Canada is recommending when it suggests that assessments of the risk associated with “exposure to chrysotile exclusively” should not be based on exposure to mixtures of asbestos containing amphiboles.

164. Canada suggests in its question that the assessments of the risk from exposure to asbestos carried out up to now (by INSERM and by all the other official bodies which have performed such assessments) are wide of the mark as they are all based on increases in cancer risk observed in different studies where workers were exposed to different types of asbestos. The models used are in fact based on the average dose-risk ratios observed in the main studies available, the validity of which has been deemed adequate. Those ratios differ widely in “extreme” studies, and this may reflect the statistical uncertainty associated with each study and/or genuine differences in risk due, for example, to the conditions of exposure or the nature or morphology of the fibres.

165. The decision to use a single average value in order to set up a dose-dependent risk model is the most realistic option when it is wished to assess the risk to the general population of a country, which is exposed under highly variable conditions, particularly in terms of the nature and morphology of the asbestos fibres encountered. While a “detailed” assessment may be justified in specific and well-known exposure situations, a “universal” risk assessment is broadly speaking a more plausible option in most situations. If it was nevertheless wished to assess the risk associated with “exposure to chrysotile exclusively”, difficult problems would be faced. The dose-risk ratios observed in the main studies available on exposure to chrysotile are extremely variable: for example, in the case of lung cancer, the dose-risk ratios are more than twenty times higher in the textile asbestos industry than in the asbestos mining and milling industry. Generally speaking, higher lung cancer risks are found in studies concerning exposure to chrysotile than in those concerning amphiboles.84 Which of the dose-risk ratios does Canada consider should be chosen for risk assessment purposes?

Question 4: Do the data from the study by Peto et al. (1998) cited by the European Communities concern exposure to chrysotile only or exposure to amphiboles or mixtures containing amphiboles?

166. The data from the study by Peto et al. concern none of the specific varieties of asbestos: they are based exclusively on statistical models for mesothelioma mortality data actually observed in the European countries concerned. The study therefore deals with the fatal effects of asbestos in all its forms. However, in order to evaluate the role of each of the different varieties of asbestos in this health catastrophe, it should once again be recalled that chrysotile accounts for the overwhelming majority of asbestos imports, particularly in France where the share of chrysotile has never been lower than 97 per cent of all asbestos. For that reason, chrysotile must be considered to be responsible for most cases of mesothelioma, as is demonstrated in detail in the article by Smith et al.85, the very title of which is unambiguous: “Chrysotile asbestos, the main cause of pleural mesothelioma”.

Question 5: Did the INSERM researchers study the question of exposure from sporadic work on materials in which chrysotile fibres are firmly embedded in a binding agent, so that no dust can be formed, such as chrysotile cement?

167. This type of exposure was analysed in the INSERM report86, together with the range of occupational exposures to sporadically high levels of pollution, which are responsible for the vast majority of cancers due to asbestos. Moreover, an Annex87 to the first written submission of the European Communities sets out exposure values measured in connection with such work, which show that levels tens or even hundreds of times higher than the prescribed limit values can be encountered. For example, a roofing worker using a grinder out of doors to repair corrugated roof sheeting made of asbestos-cement is subjected to a maximum exposure level of 41 f/ml, 410 times the limit value. It is worth noting that the authorized limit values are substantially exceeded when the ISO 7337 standard is applied.
Question 6: Since the INSERM Report acknowledges a latency period of some decades for asbestos-related illnesses, how can the European Communities assert that controlled use does not work, when fewer than 25 years have elapsed since the effective implementation of controlled use?

168. The assertion that “controlled use does not work” is based essentially on the finding that extremely high levels of asbestos fibre are released into the atmosphere during operations on asbestos-containing materials carried out in accordance with the ISO 7337 Standard, as well as the practical impossibility of ensuring that the Standard is observed at all times in operations on such materials. As it is clearly established that such levels of exposure provoke cancers, it is obvious that fatal illnesses will occur in the future as a result of such exposures. What Canada is suggesting in this question is a waiting period of 25 years (and a considerable number of deaths) to confirm this patent fact which is based on indisputable data. Furthermore, it should be noted that the HSE (Health and Safety Executive) study carried out in the United Kingdom on a population of workers in the asbestos-processing industry, which was subjected to strict rules of “safe use” from 1969, shows that the application of those rules does not enable a significant excess of cancer to be avoided, even in a sector which appears to be easy to demarcate and control.

Question 7: Did INSERM itself estimate the risk level for lung cancer or did it use the 1986 EPA estimates?

169. It should first of all be pointed out that the “risk levels” referred to indicate the increase in the relative risk of lung cancer mortality for each additional unit of exposure to asbestos. These levels were estimated by neither INSERM nor the EPA: they were actually observed in the epidemiological studies carried out among workers in the asbestos industry. Canada no doubt wishes to draw attention to the “average” level (derived from all those observed in different epidemiological studies) which is used for risk assessments and which represents the average risk. This value serves to calculate “theoretical” numbers of deaths that would be caused under various scenarios of exposure to asbestos in a population; the higher the level, the greater the calculated number of deaths. The aim therefore is to use epidemiological models to measure the theoretical impact of such levels of exposure in the population, for scoping purposes. Such models based on uncertain scenarios are commonly used in many areas relating to health or the economy, for instance in order to inform decision makers about the possible consequences of their decisions. This is what was done by the INSERM researchers.

170. It is important for the Panel to be aware that, prior to the INSERM study in 1996, the very basis of which is challenged in the Canadian submission, six other official expert groups (including a Canadian group) had done similar work:

- Consumer Product Safety Commission (USA, 1983)
- National Research Council (USA, 1984)
- Ontario Royal Commission (Canada, 1984)
- Health and Safety Commission (United Kingdom, 1985)
- Environmental Protection Agency (EPA, USA, 1986)
- Health Effects Institute (USA, 1991)

171. INSERM carried out a thorough review and careful analysis of the risk levels used by these different groups of experts, who had already made assessments of the risk of lung cancer caused by exposure to asbestos. Selecting an average level is a complex matter given the sharp variations observed in the different epidemiological studies (once again, it should be noted that the highest values are usually observed in studies concerning workers exposed to chrysotile asbestos). It is observed that four of them selected the same value as INSERM (+1 per cent); another selected a lower value, albeit combined with a range of variation with a very high upper limit; another selected a higher value (+ 2 per cent). It can be seen, therefore, that the choice made by INSERM is consistent with that of the expert groups of other countries, and no attempt was made to exaggerate the risks from asbestos.
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Question 8: Do the European Communities acknowledge that they bear the burden of proof under Article XX(b) of GATT 1994 and Article 2.2 of the TBT Agreement?

172. The burden to establish a prima facie violation of a provision of the GATT Agreement is on Canada. When the prima facie case has been made, the burden of proof passes to the defending Member, which must in turn counter or refute the alleged inconsistency, for instance by claiming the application of Article XX(b) of GATT. The burden of proof and the concomitant burden of persuasion continue, however, to shift back and forth (“like a pendulum”) throughout the entire Panel proceedings. The case law on the burden of proof that has been developed under Article XX(b) is not, however, applicable in the context of the TBT Agreement (in particular Article 2.2 thereof), as Canada’s question seems to suggest. Article 2.2 cannot be described as an exception to another provision of the TBT Agreement. The case law of the Appellate Body in the Hormones case (paragraphs 99 to 109) is more relevant in this context, taking into account the structure and context of Article 2.2 of the TBT Agreement.

Question 9: Do the European Communities maintain that the chemical composition of fibres is a relevant criterion in determining the likeness of products under Article III:4 of GATT?

173. Chemical composition is certainly relevant to the extent it affects or influences the nature, properties and qualities of the product in question. According to GATT practice and case law, a product’s properties, nature and quality are extremely important in determining “likeness” under Article III:4 of GATT. Since the chemical composition of the fibres contained in a product is almost certain to influence its characteristics (including its potential health effects), the European Communities consider that criterion indeed relevant. In addition, it is not unreasonable to assume that the chemical composition may also affect or influence the nature or quality of the product and, consequently, consumers’ tastes. It is hardly open to doubt that any informed consumer would very probably refuse a product proved to be carcinogenic. Please see also the European Communities’ reply to question 27 of the Panel.

Question 10: Do the European Communities recognize the relevance of international standards for the establishment of technical regulations on the use of asbestos fibres?

174. Please refer to the European Communities’ reply to question 37 of the Panel.

5. Questions to Canada by the European Communities

Question 1: On what facts does Canada base its claim that cases of mesothelioma in France are due solely to amphibole asbestos, not to chrysotile asbestos?

175. Canada has never stated that cases of mesothelioma in France are due solely to amphibole asbestos and not to chrysotile asbestos; science has not yet proved beyond a shadow of a doubt that chrysotile cannot cause mesothelioma. There is, however, a great deal of scientific evidence that this is, in fact, the case. On the other hand, the link between exposure to amphibole fibres, particularly crocidolite, and development of mesothelioma has been clearly established. Since the disease was identified in the late 19th Century, there have been many reported cases which could not be linked to exposure to asbestos. According to Dr. Premysl V. Pelnar in a study published in the Scandinavian Journal of Work, Environment & Health in 1988, it has been proved that many agents other than asbestos cause mesothelioma in laboratory animals, including biological agents (e.g. the SV 40 virus), chemicals (e.g. polysilicone plastics and diatomaceous earth), physical chemical products (glass fibre, erionite-zeolite, tremolite and attapulgite), and physical agents such as x-rays. It has also been proved that exposure to some of these agents (erionite-zeolite, tremolite and x-rays) causes mesothelioma in humans. See also the answer to question 3 by the European Communities to Canada.
Question 2: Can Canada give its figures for amphibole asbestos imports and consumption before the ban?

176. First of all, the question contains an error of fact concerning the ban on use of amphiboles in Canada that needs to be rectified. Under the Hazardous Products Act/Hazardous Products (Crocidolite Asbestos) Regulations, “The importation of crocidolite asbestos fibres shall be (... restricted and regulated”), but not banned. Under the regulations, the importation of crocidolite asbestos fibres for the manufacture of diaphragms, chloralkali and certain acid and temperature resistant products (e.g. waterproof seals) and of certain products containing the fibres (e.g. asbestos cement pipes) is permitted subject to the following conditions: (i) a notice must be sent to the Federal Department of Industry, which informs the provincial organization responsible for enforcing the Occupational Health and Safety Act; (ii) the notice must include the import date, the port of entry, the quantity being imported, the address of the premises where the product is to be used, etc.; (iii) the product must be labelled appropriately.

177. However, Canadian statistics on asbestos imports do not differentiate among the various types of asbestos fibres, so imports of chrysotile, amosite and crocidolite are combined in one figure in the available data. Data on Canadian consumption of asbestos fibre have been available since 1982, the year that annual surveys began. Low levels of amosite fibre consumption were reported for the period 1982-85, but no crocidolite consumption was reported.

Question 3: Does Canada have any scientific evidence to suggest that materials other than asbestos may cause mesothelioma in industrialized countries?

178. Canada has access to the same scientific data that is available to the international community at large, including France on “other potential causes of mesothelioma in industrialized countries”. Canada feels compelled to inform the European Communities of the considerable body of evidence contradicting their statement that “asbestos in all forms (amphibole and chrysotile) is the only known factor that can cause mesothelioma or pleural cancer”. A number of studies suggest other potential risk factors that may have been underestimated in epidemiological studies in industrialized countries. We take this opportunity to correct the European Communities’ simplistic definition of mesothelioma: “mesothelioma is a cancer of the pleura …”. In fact, malignant diffuse mesothelioma is a cancer of the mesothelial cells of the pleura, the pericardium and the peritoneum. Furthermore, peritoneal mesothelioma is much more typical of exposure to amphiboles than pleural mesothelioma.

179. A number of artificial fibres cause mesothelioma when they are injected into the pleura and peritoneum of laboratory animals. It should also be noted that the International Agency for Research on Cancer (IARC) has classified refractory ceramic fibres as probable carcinogens, partly because of instances of mesothelioma induced by inhalation and injection in animal studies. The SV 40 virus readily induces mesothelioma when injected into animals; studies suggest that the virus contaminated anti-polio (poliomyelitis vaccines) from 1955 to about 1963 and may induce mesothelioma with or without the help of asbestos fibres. Some studies of humans report the presence of the simian SV 40 virus in the biological tissue of mesothelioma victims. Ionizing radiation used in cancer therapy, as well as, perhaps, occupational exposure to radiation have induced mesothelioma. In addition, a large proportion of mesothelioma cases (as high as 25 per cent) are not attributable to exposure to asbestos; they may have been caused by hidden (unidentified but real) exposure to asbestos, but the figures suggests that there are other significant causes that cannot be countered by targeting exposure to asbestos alone. For example, erionite has been shown to be even more toxic than crocidolite in causing mesothelioma: it has killed large numbers of villagers in Turkey. Erionite is a mineral fibre but does not belong to the asbestos family, which suggests that fibres with similar physical characteristics could pose a health threat to other population groups. Finally, we wish to remind the European Communities that the issue at hand is the ban on chrysotile products and that there are mes-
othelioma risk factors other than chrysolite, particularly all the amphibole fibres and flockings placed around furnaces, boilers and high-temperature pipes.

Question 4: Does Canada dispute that chrysotile asbestos is a proven carcinogen in humans - in general? In the case of lung cancer? In the case of mesothelioma?

180. Canada does not dispute that chrysotile causes lung cancer. However the way in which exposure to chrysotile asbestos may increase the risk of lung cancer has not yet been fully explained, and might only be indirect. First of all, Canada maintains that the risk is dependent on the intensity and duration of exposure and that there is a level of risk below which the risk, if any, is undetectable. According to Churg on this subject:

"As a practical matter, the data indicate that chrysotile will not produce mesotheliomas in those exposed to any current or recently regulated numbers of chrysotile, and certainly not in those exposed to chrysotile encountered at environmental levels".96

181. The risk may become detectable in cases of long-term exposure to high levels, but it is then by no means certain whether the chrysotile acts as a direct carcinogen or by causing the formation of pulmonary fibrosis, which would be a precursor of neoplasia. In other words, exposure must be sufficient in intensity and duration to induce pulmonary fibrosis, which predisposes the pulmonary parenchyma to a higher risk of cancer. Regarding asbestos-related mesothelioma, a number of studies have demonstrated cogently that this type of cancer is almost exclusively linked to exposure to amphiboles. Cases of mesothelioma in chrysotile asbestos miners in Quebec are relatively rare - in a cohort of 11,000 workers who were very carefully monitored (in the McDonald study), there were at most fifty or so cases over several decades. Exhaustive research into their employment history revealed that most of the cases were related to short-term exposure to commercial amphiboles. For example, during the Second World War, some of the miners with mesothelioma had worked in plants manufacturing products for the allied forces and amphiboles imported into Canada had been used to make a variety of products, including gas masks, to assist in the war effort.97

Question 5: How does Canada explain the fact that the risk of mesothelioma is seven times higher than average for women living near chrysotile mines in Quebec? (Note that this population group was the subject of a special study because, in its case, the exposure is environmental, not directly occupational).

182. The women in the towns of Thetford Mines and Asbestos have been continuously exposed from birth to asbestos concentrations of 1 f/ml. or more. Given that the risk of mesothelioma increases by approximately the power of 3 in relation to the time that has elapsed since exposure, these women have had more time to develop a detectable risk of mesothelioma than many workers in other parts of the asbestos industry. Even so, their risk level seems to be at least 20 times lower than predicted by the Environmental Protection Agency (USA) and INSERM models.

183. More specifically, it should be borne in mind that some 75 per cent of the women in this study98 lived with an asbestos worker, of whom a small minority worked with amphiboles, particularly crocidolite. Furthermore, some 5 per cent had themselves already worked in the asbestos industry as “sheddeuses” or “gobeuses” or had worked in workshops for repairing jute sacks which had contained crocidolite or had worked at home for the asbestos industry repairing jute sacks used to transport amphiboles. Finally, the results to which the EC refers concern deaths from pleural cancer. Since then, researchers have compiled and checked the diagnosis in pleural mesothelioma cases, thus providing a better measurement of the risk arising from asbestos. The authors of the study have collected but not yet analysed the exposure histories of cases of pleural mesothelioma in women (more than ten so far, and not merely the seven mentioned in the 1997 publication).
Preliminary analyses indicate that a certain number of mesothelioma cases were exposed to crocidolite or amosite more than thirty years before the onset of the disease (Camus and Siemiatycki, personal communication, 1999). The other mesotheliomas may be attributable to a long period of induction, following a massive build-up since childhood, of continuous environmental exposure or exposure in the home (living with an asbestos worker) to commercial chrysotile exceeding 1 f/ml. The women's cumulative exposure was equivalent to between 100 and 300 fibre years/ml for asbestos workers.

Question 6: Does Canada disagree that most cases of mesothelioma occur in industrialized countries in occupations involving intermittent exposure to asbestos, specifically those where the worker has to work with asbestos containing materials?

Nowadays, most cases of mesothelioma in the industrialized countries occur in occupations other than mining and manufacture of asbestos products because workers in those occupations are exposed to amphiboles in poorly controlled work areas. The workers who remain at risk, working in the construction sector, are specifically those who are most exposed to flocking and friable products made of asbestos mixes containing amphiboles or composed entirely of amphiboles. This situation is completely different from the case of workers in chrysotile mines and in manufacturing plants where only chrysotile is used.

Question 7: Because Canada did not provide the health statistics requested at the consultative meetings in July 1998, France used gross data on the worldwide situation published by the WHO to calculate the incidence of mesothelioma in Canada and Quebec from 1978-1992. Could Canada provided its own recent health statistics (since 1992) on mesothelioma deaths in Canada and Quebec?

See above, reply to question 9 by the Panel to Canada.

Question 8: Can Canada explain how the safe use of chrysotile can have resulted in the following observations from a study conducted in the Montreal area:

- A risk of mesothelioma more than 14 times higher than the average for persons “significantly” exposed to chrysotile asbestos?
- A risk of lung cancer 2.3 times higher than the average for people “significantly” exposed to chrysotile asbestos?
- A risk of mesothelioma almost 12 times higher than the average for construction workers?

The purpose of this major case-control study, which was conducted on 4,500 cancer cases diagnosed in hospitals in the Montreal area between September 1975 and June 1985, was to identify potential associations for follow-up from among the tens of thousands of possible associations between 23 groups of cancer, 98 occupational groups and 293 substances. As the authors state, the study and the analysis were designed to establish hypotheses, not to verify specific hypotheses. The downside of this greater sensitivity is a tendency to overestimate associations and to obtain more “false positives.” The authors therefore recommend that each reported association should be interpreted with prudence and caution (page 304).

The main limitation of the reported analyses is that they are “univariate”: they consider only one occupational exposure at a time and do not evaluate the effects of other concomitant occupational exposures (page 119). This problem is particularly significant in the case of chrysotile. The workers in the study who were exposed to chrysotile were also exposed to amphibole fibres, crystalline silica, cement dust, PAH’s and alkanes, wood dust, solvents and pyrolysis fumes (pages 50-51). Seven of the ten substances in the study were associated with mesothelioma and all ten were associated with lung cancer. Thus the reported associations between mesothelioma, lung cancer and chrysotile do not isolate the level of risk associated with chrysotile itself from the compound effect of several other risk factors.
factors. The authors thus caution against oversimplified interpretations (page 301): “[A] much more important problem is the fact that the association between cancer and one occupational substance may be confounded by other occupational substances.” The problem may be even more serious in respect of mesothelioma, because the study included only a very small number of cases (12 cases).

189. Regarding lung cancer, the study also recognizes the possibility of a residual effect of smoking and a paradox that the researchers cannot explain: the absence of an association between lung cancer and exposure to amphiboles. This anomaly illustrates the difficulty in population case-control studies of distinguishing between past exposure to chrysotile and amphibole. The same problem was underlined in the article by Iwatsubo et al., cited by the European Communities99 and in the preface100 by Siemiatycki and Boffetta (IARC) to the Iwatsubo study. Finally, because the study was restricted to cancers occurring between 1979 and 1985, and the induction/latency periods for lung cancers and mesothelioma are several decades, the exposures that may have caused the cancers occurred before controlled use measures were applied. This data can, therefore, in no way provide a basis for assessing the affect of controlled use of chrysotile which was introduced in the 70’s.

Question 9: Why does Canada think that the controlled use of amphiboles is not technically feasible (since it has banned this type of asbestos)?

190. The distinction is based on the proved lower toxicity of chrysotile compared with amphiboles and the application of principles inherent in international standards on the controlled use of asbestos.

Question 10: In its submission, Canada acknowledges that “certain uses for which exposure cannot be controlled to an acceptable degree should be banned”. Could Canada specify to which uses it is referring?

191. The guiding principle in controlled use is the elimination of any use involving a risk that cannot be adequately controlled. Thus the Canadian Hazardous Products Act prohibits the use of the following:

(a) Textile fibre products that are to be worn on the person and that contain asbestos fibres, other than products that are designed for the purpose of affording protection from fire and heat hazards and constructed in such a way that ensures that the asbestos fibres will not, on reasonably foreseeable use, become separated from the products;

(b) asbestos in spray form except products composed of a mixture of asbestos fibres and bituminous and resinous-based binding materials where the fibres are encapsulated with the binding during spraying and the resulting materials are not friable after drying;

(c) products composed of or containing asbestos for use by a child in learning and playing and made in such a way that asbestos may become separated from the products;

(d) products composed of or containing asbestos for use in modelling or sculpture;

(e) dry-wall joint cements or compounds or sprackling or patching compounds, composed of/or containing asbestos, that are made in such a way that airborne asbestos may become separated from the products during the preparation of the products, other than preparation at the manufacturing level, or during the application of the products or at any time thereafter up to and including the repair and removal of the products;
(f) products composed of or containing asbestos for use to simulate ashes or embers.

Question 11: The enormous amount of work in France on airborne particles in the workplace has shown no evidence of chemical changes in chrysotile asbestos fibres released during high-speed machining of materials containing asbestos. Can Canada provide evidence of changes in the chemical composition of chrysotile asbestos during high-speed machining?

192. Canada is surprised that this question should be asked, because the answer was given at the first substantive meeting of the Panel, with relevant supporting references. In any case, given that the question concerns fibres released during dry, high-speed machining of high-density materials, the operations concerned clearly do not comply with controlled-use safety procedures.

Question 12: Can Canada state exactly when the “modern” use of asbestos began? How does Canada characterize the so-called “modern” characteristics of chrysotile cement products?

193. The advent of the use of “modern” asbestos products was a gradual process that began in the 1970s with the phasing out of uses in which asbestos fibres could readily be separated from the finished product, as in the case of friable insulating materials and flockings, toys and unprocessed textiles. The manufacture of products containing amphibole-type fibres was also phased out because they were recognized as more harmful. In short, a distinction must be made between two periods: the first is characterized by the use of amphibole fibres and uses in which asbestos fibres could be easily separated from the finished product; the second is characterized by the prohibition or restricted use of amphibole asbestos and by the advent of non-friable products, i.e. products in which fibres are firmly bound to a matrix and are highly unlikely to be released in biologically significant concentrations. The distinction is a fundamental one. It should nonetheless be noted that non-friable products in which chrysotile is encapsulated in a matrix, as in the case of chrysotile cement, were manufactured and used during both periods.

194. The industrial diseases attributable to asbestos that are now occurring in Europe and North America are associated with so-called “old-style” products and amphiboles. This fundamental distinction has been recognized by the International Labour Organization in Convention 162 concerning Safety in the Use of Asbestos, by the World Health Organization and in the regulations of most countries. Even though they have been manufactured for nearly 100 years, chrysotile cement products meet the “modern use” criterion because the fibres are encapsulated in a cement matrix and are not released in significant quantities as long as simple controlled-use procedures are followed. More recently, new procedures have been introduced to eliminate the use of crocidolite fibres in the manufacture of pipes, and countries such as Japan and France have developed new chrysotile cement products manufactured by extrusion or covered with a lustre coating. These advances have in no small measure helped to make products not only safe but also more attractive.

Question 13:

(a) Has Canada established a monitoring enforcement mechanism for the application of “controlled” use measures, including maintenance work?

195. In Canada, controlled use has entailed the prohibition of friable products containing asbestos and the adoption of measures to protect the health and safety of workers exposed to asbestos. In general, the protective measures are a provincial responsibility, and in Quebec they come under the Regulation on the Quality of the Work Environment and the Safety Code for the Construction Industry, which are administered by the Occupational Health and Safety Commission (CSST). Both sets of regulations cover working con-
ditions in chrysotile mines, asbestos product manufacturing plants, and firms specializing in maintenance and removal of asbestos flockings in Quebec.

196. Regarding the health and safety of workers specializing in maintenance and repairs (plumbing, electricity, air-conditioning, etc.) in buildings containing asbestos flockings, the CSST is in the process of implementing a programme for the prevention of occupational diseases associated with exposure to asbestos. The programme is aimed at management and employees in this sector of activity and its purpose is to inform them about the required prevention measures and give them the appropriate training.

197. In addition, the Regulation on Air Quality and the Regulation on Solid Waste, issued by the Department of the Environment of Quebec, state respectively the environmental measures to be taken regarding airborne asbestos and the standards governing burial of waste containing asbestos. Officers of the CSST and the Department of the Environment monitor and enforce the application of all these measures as part of their regular duties, in the same way as they enforce provisions for the control of other substances considered to pose a public health risk and a threat to the health of workers. The measures make it possible to control asbestos throughout its useful life, from the time it is mined to the time it is buried.

(b) What monitoring mechanism did Canada set up to check whether the applicable ISO standards were being followed in Canada and in countries with companies that signed the “Agreement”?

198. In their respective areas of jurisdiction, the regulations of the Canadian Government and the provincial governments provide that the use of chrysotile, at all stages from extraction to use of the finished product, is to be controlled in such a way as to minimize exposure to fibres. Furthermore, they cover the maintenance of buildings containing various types of asbestos and provide for a variety of measures based on the type of work being done and the type of asbestos fibre involved. In most provinces and in buildings under the jurisdiction of the Canadian Government, property owners are required to maintain friable materials in good condition and to inform employees in advance in the event that they may be exposed to such products during maintenance work, repairs or renovations.

199. In the Agreement on the responsible use policy, there is no question of any intent to check how each business uses its products containing chrysotile. The aim of the Agreement is to ensure that users manufacturing finished products, where the health risk is the highest, comply with national and international occupational health standards. The relevant passage of the Canadian submission, concerning standards for installing asbestos cement products, was intended to show that the safe use of asbestos cement products is possible and that this fact is recognized by international organizations such as the International Organization for Standardization. So Canada is surprised at European claims that safe use is impossible, particularly in a country like France.

Question 14: Could Canada clarify whether asbestos producers have already had occasion to stop exports to countries after determining that businesses using their products were not complying with the “Agreement”?

200. Stopping chrysotile sales by producers from signatory countries is the measure of last resort in cases where a consumer is not complying with national regulations and makes no effort to improve working conditions in the business concerned. Under the Agreement, producers must first provide businesses with technical support in order to reduce concentrations of airborne respirable fibres. Measures include the implementation of working procedures, the installation of dust collectors, and the purchase of air sampling and analysis equipment. To enhance the effectiveness of their actions, Canadian chrysotile producers have also used the services of international organizations such as the Asbestos Institute and the Asbestos International Association for plant inspections, training and the collection of data on airborne fibre rates on a continuing basis. In this context, more than
100 technical visits have been made over the last five years, along with training courses in Canada and in a number of consumer countries.

201. Numerous contacts have been established with the governments of chrysotile importing countries to raise their awareness of the work being done by the producers, often with the assistance of trade unions and the Government. In this way, signatories to the Agreement can be assured that non-compliant businesses cannot obtain supplies from other sources. As a result, some countries have decided to issue import licences so as to ensure that every user complies with national regulations. However, in the few cases where users do not follow controlled-use procedures, Canadian producers refuse to sell or stock shipments of the fibre. This has been the case, in particular, of certain users in Mexico, Argentina, Korea, China and Egypt.

Question 15: How can Canada claim, on the one hand, that amphibole-asbestos and chrysotile-asbestos are very different products and, on the other, claim at the end of its submission, that chrysotile-asbestos and substitute products are like products?

202. The distinction made by Canada between amphibole fibres and chrysotile fibres is intended to differentiate between the fibres according to their pathogenicity, on a medical and scientific basis. The objective is not to determine whether they are “like” for the purposes of any provision in the TBT Agreement or GATT but to determine whether one is more dangerous than the other. In this context, the capacity for inducing mesotheliomas and the carcinogenicity of the different types of fibres are clearly relevant to determining the respective pathogenicity levels of amphiboles and chrysotiles.

Question 16: If Canada were to admit that there was a difference in toxicity between two products, would it accept the idea that they were not like products?

204. No. First, it should be noted that the toxicity of a product is not a recognized criterion for analysis of likeness. In Canada’s opinion, the fact that two products have similar toxicities is not an element of their likeness under Article III:4 of GATT or Article 2.1 of the TBT Agreement. Benzene, for example, is not “like” crystalline silica just because they are both carcinogens.

Question 17: On what scientific basis did Canada decide not to follow the recommendations made by the ILO in 1986 and those made by WHO in 1998 for asbestos, including chrysotile asbestos, to be replaced with less hazardous products?

206. To answer this question, we would first like to quote Article 10 of ILO Convention 162, to which the European Communities refer in discussing substitute products: “where ... technically feasible, national laws or regulations shall provide for ... the replacement of asbestos or of certain types of asbestos or products containing asbestos by other materials or products or the use of alternative technology, scientifically evaluated by the competent authority as harmless or less harmful; ...”. The concept of substitution
approved by WHO is identical to the one conveyed here: it must be proven that the substitute products are harmless or less harmful. The fact is that the scientific community still questions the harmlessness of most of the substitutes. Thus, INSERM stated that “any new fibre proposed as a substitute for asbestos or for any other use must be suspected, a priori, of being pathogenic because of its structure but that this does not preclude [the requirement] for analysis of the possible consequences of its physico-chemical characteristics.”

207. Given the substantial latency period (between 15 and 45 years for asbestos) between exposure to a pathogen and the development of disease, it is impossible to make a definitive assessment of the carcinogenic potential of substitute fibres which came on to the market only recently. However, a number of recent studies cast doubt on the harmlessness of these fibres. As long ago as 1993, in a report entitled *Selected Synthetic Organic Fibres*, the World Health Organization’s International Programme for Chemical Safety (IPCS) identified the para-aramid fibre as being respirable and biopersistent, two criteria associated with toxic potential in a substance. The same applies to most replacement fibres. Given the lack of knowledge on the subject, it would be premature to press on blindly with substitution.

**Question 18:** Why has Canada not instituted similar proceedings against other countries that have imposed similar bans on asbestos, such as Iceland, Norway, Denmark, Switzerland and New Zealand?

208. Canada could have taken action and is entitled to take action against similar measures previously introduced by other countries, and in particular by seven other members of the European Communities. However, it has decided, for the time being, to confine itself to the “model case” of France, basically for the following reasons:

- France is the first European country to have banned asbestos since the creation of the WTO and the introduction of the new dispute settlement rules;
- France has switched from a policy of controlled-use of asbestos (precisely the one advocated by Canada) to a diametrically opposite one – a total ban – even though there is no new scientific data; and
- the procedures under the Agreement do not include any mechanism for combining similar or identical legal action.

**B. QUESTIONS AT THE SECOND SUBSTANTIVE MEETING (20-21 JANUARY 2000)**

1. **Questions by the Panel to both parties**

**Question 1:** Is the concept of “like product” in the meaning of Article III:4 of GATT identical to that contained in Article 2.1 of the TBT Agreement? In this context, what precisely, in your opinion, is the relevance of the criterion of the effects of the product on human health in the context of the two provisions?

(i) **Reply by Canada**

209. Article 2.1 of the TBT Agreement reaffirms the obligation concerning national treatment (Article III:4 of GATT) and the obligation of most-favoured-nation treatment (Article I:1 of GATT). In the case of the obligation concerning national treatment, we consider that Article 2.1 of the TBT Agreement and Article III:4 of GATT contain the same type of prescription: not to treat imported products less favourably than like national products. The only difference between the two provisions is that Article 2.1 applies only to technical regulations while Article III:4 of GATT applies to “all laws, regulations and requirements […]”. From that it follows that the concept of like products in the meaning of Article 2.1 of the TBT Agreement is identical to that contained in Article III:4 of GATT and that the criteria which are used to identify the range of like products under Article III:4 of GATT are the
same as those which are used to identify the range of like products in Article 2.1 of the TBT Agreement. These criteria do not include the effects of the product on human health and we consider that they are not relevant in this case.

210. According to the Appellate Body in the case of Japan – Taxes on Alcoholic Beverages, the report of the working group on Border Tax Adjustments set out the principle for interpretation of the formula “like products” in general in the various provisions of GATT 1947. It must be interpreted on a case by case basis using criteria such as the product’s end-use in a given market, consumers’ tastes and habits, which change from country to country, and the products properties, nature and quality. This approach has been followed in almost all Panel reports which have been adopted since the one concerning border tax adjustments. We have followed this principle in developing our arguments that chrysotile fibre and cellulose, PVA, glass fibres as well as chrysotile-cement and fibro-cement are like products pursuant to Article III:4 of GATT and Article 2.1 of the TBT Agreement. In the same case, Japan Taxes on Alcoholic Beverages, the Appellate Body implied that other criteria can also be utilized to identify the range of like products within the context of a particular provision of the multilateral trade agreements of the WTO Agreement.

211. The example of another criterion frequently used in previous Panel reports to determine whether products are like is the tariff classification. We have followed this example in the present case and invoked the uniform classification in the tariff nomenclatures of the harmonized system as a criterion to confirm that chrysotile fibre and cellulose, glass and PVA fibres, as well as chrysotile-cement and fibro-cement, are like products. The toxicity of a product, however, has never been taken as a criterion for determining whether products are like. Moreover, in this case, the effects of the substitute fibrous products on human health are too little known and uncertain to constitute a criterion which could assist the Panel in identifying the range of like products. Consequently, we are of the opinion that the Panel should only examine the criteria defined by previous practice of GATT on which we have relied in our arguments. We refer the Panel to our second written submission in which we stated that the effect of a product on human health is not a criterion that should be used to determine whether products are like. Two products may be similarly toxic and not be like in the meaning of Article III:4 of GATT or Article 2.1 of the TBT Agreement. Conversely, assuming that it is clearly established that two products do not share the same toxicity, they may still be like for the purposes of Article III:4 of GATT or Article 2.1 of the TBT Agreement.

212. We reiterate our replies to questions 15 and 16 by the European Communities in which we explain that the scientific question of the pathogenicity of the fibres has no place in the context of an argument to show what is like pursuant to Article III:4 of GATT 1994 or Article 2.1 of the TBT Agreement. Instead, the broadest and most general criteria contained in the case law, such as characteristics, nature and quality of the product, tariff classification and product’s end use should be applied. Chrysotile fibre is indisputably different from amphibole fibres when it comes to toxicity, but we consider that chrysotile fibre and amphibole fibres are like products. Just as chrysotile fibre and amphibole fibres are like products although amphibole fibres are much more toxic, so PVA, glass and cellulose fibres and chrysotile fibres are like products by virtue of their characteristics, nature and quality, even if we do not know, given the present state of scientific research, the actual toxicity of each of these substitute fibres. In our view, the effects of chrysotile fibre on human health, in the same way as those of substitute fibres, which in most cases are not known, are not relevant to the issue. The Panel should not take them into account in the context of its examination of the various characteristics of these products to establish whether they are like pursuant to Article III:4 of GATT or Article 2.1 of the TBT Agreement.

(ii) Replies by the European Communities

(a) No, the concept of “like products” in Article III:4 of GATT is not identical to the concept of “like products” contained in Article 2.1 of the TBT Agreement for the following reasons.
213. The concept of “like products” has to be interpreted in accordance with the customary rules of interpretation of public international law, that is in accordance with the ordinary meaning to be given to this concept in its context and in the light of its object and purpose. It should also be recalled that the Appellate Body has held that the concept of “likeness” is a relative one that evokes the image of an accordion. The accordion of “likeness” stretches and squeezes in different places as different provisions of the WTO Agreement are applied. The width of the accordion in any one of those places must be determined by the particular provision in which the term “like” is encountered as well as by the context and the circumstances that prevail in any given case to which that provision may apply. No one approach to making a judgement will be appropriate for all cases.

214. First of all, the context in which this term appears is different in the two Articles. The context of GATT Article III.4 is clearly different from that of Article 2.1 of the TBT Agreement. The TBT is a specific Agreement that elaborates further on the objectives of GATT 1994. Whereas the obligation of national treatment and the concept of “likeness” in Article III.4 of GATT 1994 have a fairly wide application “in respect of all laws, regulations and requirements affecting internal sale, offering for sale, purchase, transportation, distribution or use”, the obligation of national treatment and the concept of “likeness” in Article 2.1 are confined by the specific object and purpose of the TBT Agreement, that is it applies only “in respect of technical regulations”. Thus, Article 1.2 of the TBT Agreement provides that for the purposes of that Agreement the meaning of the terms given in Annex 1 thereto applies. Also the title of Article 2 reads “Preparation, Adoption and Application of Technical Regulations by Central Government Bodies”. These phrases may not be read so expansively as to subvert the purpose and object of Article 2.1 and of the TBT Agreement in general. It follows that because the TBT Agreement deals only with technical regulations, standards and conformity assessment procedures related to products or processes and production methods, the concept of “likeness” is by definition narrower than that of Article III.4 of GATT as regards the legal context within which and the object and purpose for which the determination of “likeness” is to be made. In other words, the legal context as well as the object and purpose determine inevitably the coverage of the term “like”: not all like products are covered by Article 2.1 of the TBT Agreement, but only those to which the technical regulation was intended to apply.

215. Secondly, the European Communities dispute Canada’s claim that the French Decree in question lays down a technical regulation in the sense of the TBT Agreement. This is clearly not the case of the general, horizontal prohibition of any kind of asbestos as well as of the limited and transitional exceptions laid down in Article 2 thereof, as shown in our written and oral submissions. However, if we suppose, for the sake of argument, that the French Decree did lay down a technical regulation, such a regulation would only have been applicable to asbestos as such and to asbestos-containing products. Indeed, the French Decree in question has laid down no technical regulation whatsoever for the so-called “substitute” products as claimed by Canada. Canada’s argument that the so-called “substitute” products are, for the purpose of Article 2.1 of the TBT Agreement, “like” asbestos and asbestos-containing products runs counter to the very object and purpose of the TBT Agreement and leads to unacceptable results from the regulatory point of view. This is because such an interpretation is likely to restrict unreasonably the regulatory freedom of the WTO Members and would introduce uncertainty and unpredictability into international trade for no valid reason. As a general rule, Members introduce technical regulations, standards and conformity assessment procedures for the purpose of achieving a legitimate objective (e.g. safety). The object and purpose of the TBT Agreement is to guard against “unnecessary obstacles to international trade” (fifth preambular paragraph) in the products covered by the technical regulation, not in the potentially vast category of alternative/substitute products, because normally no government is in a position to know in advance the category of products that are likely to be affected by the adoption of the technical regulation. The concept of “like” products cannot be given so broad a reach as effectively to underline the autonomous right of Members to determine their level of health protection. In other words, to interpret so broadly the concept of “like” products is bound to present Members’ regulatory authorities with the dilemma of abandoning the pursuit of a legitimate objective (e.g. safety) for the sake of avoiding a possible adverse finding under
Article 2.1 of the TBT Agreement, even if there exist only a few alternative/substitute products of national origin. Such a paradoxical result can surely never have been intended by the WTO Members. It is contradictory and unreasonable to judge “likeness” on the basis of the possible effects of the technical regulation on the imported (prohibited) product, as compared to the alternative/substitute products of national origin, because a technical regulation by definition, almost always, entails different effects (negative or positive) on those products (of whatever origin) which do not conform with the technical regulation in question.

216. Thirdly, since the context, object and purpose of Article 2.1 of the TBT Agreement is different from that of Article III:4 of GATT, the four criteria usually used by panels to judge “likeness” on a case-by-case basis in the context of Article III of GATT (i.e. the product’s end uses, consumers’ tastes and habits, the product’s properties, nature and quality, and tariff classification) may be applied in the context of Article 2.1 of the TBT Agreement but only in the light of the purpose of the technical regulation under consideration, i.e. in order to identify only whether the imported asbestos and asbestos-containing products receive, in respect of the technical regulation in question, treatment less favourable than that accorded to various asbestos (different types of amphibole asbestos, chrysotile, etc.) and asbestos-containing products (brakes, various asbestos-containing cement products, etc.) of national origin.

(b) As the European Communities have explained when it has been scientifically established that a product is dangerous to human health, this finding attributes a special internal and/or external characteristic to the product in question that should be taken into account in defining “likeness” in both Articles. Indeed, the scientifically established dangerous nature of a product (like that of any kind of asbestos) necessarily affects the normal criteria used by panels to define “likeness”, because:

- The product’s very properties, nature and quality are necessarily affected, in the sense that they are not the same as those of a safe or safer identical or alternative or substitute product. For example, an apple that contains a poisonous pesticide can never be “like” a safe apple of the same or different variety or a safe orange, when one would otherwise consider apples and oranges “like” products for the purpose of Article III:4 of GATT.

- The product’s end uses are also different, because a product that has been scientifically shown to be dangerous to human health (e.g. an apple that contains a poisonous pesticide or any type of asbestos) usually does not have the same end uses as those of a safe or safer identical or similar or substitute product.

- The consumers’ or end-users’ tastes and habits are also bound to differ in the case of a product that has been scientifically shown to be dangerous to human health (e.g. an apple that contains a poisonous pesticide or any type of asbestos) and a safe or safer identical or similar or substitute product.

217. In reality, a product that has been scientifically shown to be dangerous to human health attributes to that product a specific characteristic that makes it unlike any other product of the same family or category (e.g. apple or asbestos) or similar or alternative or substitute products (e.g. orange or PVC fibrous product). That kind of dangerous product should not be put into international trade and no importing country should have to rely on Article 2.2 of the TBT Agreement, for example, or the exception provisions of Article XX to justify any restriction on imports. It follows that the dangerous character of a product to human health is of particular relevance in the context of Article III:4 of GATT in deciding “likeness”. In the context of Article 2.1 of the TBT Agreement, the dangerous character of a product to human health is also highly relevant because, in the light of the object and purpose of a technical regulation, the protection of human health is a legitimate objective.
that may be taken into account in the course of its preparation. This is particularly true, if the concept of “like” products in the context of Article 2.1 of the TBT Agreement is to be given the very wide meaning advocated by Canada.

Question 2: Is it possible for a measure to fall partly within the scope of the TBT Agreement while other elements of the same measure would come under GATT?

(i) Reply by Canada

218. As the Appellate Body indicated in EC - Regime for the Importation, Sale and Distribution of Bananas\(^{119}\) and in Canada - Certain Measures Concerning Periodicals\(^{120}\), it is possible that a measure, because it has several aspects, could be examined under more than one WTO agreement. Conceptually, then, it is possible that certain aspects of a particular measure are covered by the TBT Agreement, while other aspects of the same measure, which are not covered by the TBT Agreement, are covered by GATT. However, taking account of the facts of this case, Canada is of the opinion that the French measure is a technical regulation and that all aspects of it come under the TBT Agreement. It seems to us, that in terms of methodology, the whole French measure should initially be examined in relation to the TBT Agreement; subsequently, it could be examined in relation to GATT.

219. In the above cases, the question of whether the General Agreement on Trade in Services (GATS) and GATT 1994 were mutually exclusive was raised because GATS does not cover the same subject as GATT. The Appellate Body decided at that time that the agreements were not mutually exclusive. The situation in this case is different. Indeed, the respective areas of application of the TBT Agreement and GATT 1994 overlap. In our reply to question 35 by the Panel, we said that “the TBT Agreement is the most recent and most specific expression of the WTO Members as to what the interpretation and application of the disciplines under GATT 1994 […] should be in the context of technical regulations and standards”\(^{121}\). The two agreements cover the same subjects, the TBT Agreement being a development of GATT disciplines. It is therefore difficult to see how in practice certain specific aspects of the French measure would come under the TBT Agreement while other aspects would be governed by GATT. The TBT Agreement is more specific than GATT 1994 in the area of technical regulations. As the French measure is a “technical regulation” in the meaning of the TBT Agreement, it would seem to us appropriate first to examine the measure in the light of the disciplines in that agreement, with particular reference to the GATT disciplines. This methodological approach flows from the most recent case law on relations between the WTO Agreements, which requires that claims should first be examined under a more specific agreement rather than GATT 1994.\(^{122}\) Canada maintains that the Panel should first examine all the disputed elements and aspects of the French measure in the light of the TBT Agreement.

(ii) Reply by the European Communities

220. In general, the question of deciding which WTO Agreement applies to a given measure is a problem of determining the character of the measure. In the absence of a specific provision in the relevant WTO Agreements\(^{123}\), one has to examine essentially the aim and content of the measure in question. A measure that has a single and uniform aim and content should fall under one agreement. Equally, a measure that pursues several distinct or heterogeneous aims can potentially fall under several, different agreements. But not every element in a measure attributes a different, multiple aim to the measure under examination. Secondary, accessory or ancillary provisions do not necessarily alter the essential aim of the measure for the purpose of determining its legal character.\(^{124}\) Therefore, the essential or principal purpose of the measure, as this is objectively expressed in its design, architecture and structure, determines its character and the applicable agreement. In defining the essential or principal objective of a measure, one has to look at the centre of gravity or predominant component of the aim pursued by the measure in question, as this emerges or becomes apparent from the measure itself.\(^{125}\)
221. Therefore, the reply to the question would depend on how significant and independent the “other elements” of the measure are, in other words, whether they are capable of attributing a distinct, different and additional aim to the one principally pursued by the measure in question. Only in such a situation may some elements of a measure fall under the provisions of GATT and other elements of the same measure fall under the provisions of the TBT Agreement.

222. Applying the above principles to the French Decree in question, the Panel should conclude that it pursues only one, single, uniform aim, that is to ban the use of any kind of asbestos for the purpose of protecting human health. As the European Communities have explained at length in their written and oral submissions, this sole object of the Decree flows from the ordinary meaning of its terms in their context, the design and structure of the measure and the history of the preparatory work. The essential object and purpose of the Decree is to lay down a general, horizontal ban on the use of asbestos and asbestos-containing products. It does not relate to the preparation, adoption, and application of a technical regulation of any kind, in the sense of the TBT Agreement. Neither does it lay down a process and production method for asbestos and asbestos-containing products, simply because it bans their use.

223. One may wonder whether the limited and temporary exceptions laid down in Article 2 et seq. are a sufficiently important or independent element as to attribute another, separate and different aim to the Decree. As the European Communities have already explained, this is clearly not the case. Article 2 et seq. of the Decree state expressly that “exceptionally and on a temporary basis” certain products may continue to use chrysotile asbestos in order to ensure “an equivalent function” and so long as there exists “no substitute” for chrysotile which can ensure a lower level of risk and guarantee the same level of safety to users. This is an exception to the general ban, and exceptions by definition are to be interpreted narrowly. Article 2 et seq. explicitly state that the exception is temporary and will be phased out when substitute products become technically available. As the European Communities have already shown, the practice followed since the adoption of the Decree in 1996 confirms the ephemeral nature of these exceptions. Consequently, these exceptions are not an essential, but rather an ancillary, accessory or subsidiary element of the Decree. They lay down no technical regulations or standards in the sense of the TBT Agreement and, therefore, cannot render the entire Decree subject to the TBT Agreement. And even if we were to assume that the exceptions do lay down technical regulations (which they do not), at best only those exceptions would fall under the scope of the TBT Agreement. But Canada does not claim that those exceptions constitute a violation of the TBT Agreement. Therefore, the question of whether some elements of the Decree (in this case the temporary exceptions) can fall within the TBT Agreement is, from a strict legal point of view, totally irrelevant to the outcome of this case.

Question 3:

(a) In assessing the conformity of a measure with Article XX of GATT, what would be the relevance of any practical difficulties encountered by a country in implementing a measure, in assessing whether the measure was reasonably capable of achieving the objectives of its health policy?

(b) In assessing the conformity of a measure with Article 2.2 of the TBT Agreement, what would be the relevance of any difficulties in the practical implementation of a measure in evaluating whether it was less trade-restrictive than the measure in place?

(i) Reply by Canada

Reply 3(a)

224. Canada is of the opinion that any claims of practical difficulties inherent in the implementation of a measure are not a factor to which any weight should be given in assessing the reasonable availability of a measure as a less trade-restrictive alternative. In
considering a measure in the light of the criteria of necessity in Article XX of GATT, the practical difficulties of implementing an alternative to the measure under dispute – if such exist – do not detract from the alternative or exclude it as a less trade-restrictive alternative. In the context of the test of necessity, assessing the legality of a measure in the light of so-called practical difficulties encountered in implementing it would be to reward laxity in implementing a measure and compensate administrations for their inefficiencies. It would then become easy for Members to evade the GATT disciplines, because they would simply have to allege practical difficulties in implementation – i.e. the inefficiency of their own administration – to exclude a less trade-restrictive alternative measure, and thus escape the GATT disciplines. In the light of the application of the test of necessity in Article XX of GATT, a Member that generally devotes significant resources to overcoming practical difficulties in implementing a regulation and which can pride itself on the efficient administration of its regulatory framework would be penalized vis-à-vis a Member who decided to deploy few resources to implement regulations or which proved to be lax in that regard. To assess, in the context of analysing Article XX of GATT, whether an alternative measure is reasonably available in the light of the existence or otherwise of practical difficulties in implementing it would be a dangerous precedent which would weaken the test of necessity, produce Kafkaesque results and lead to pernicious effects on a systemic scale. In this respect, the decisions of the Panel and the Appellate Body in the case of United States – Standards for Reformulated and Conventional Gasoline support Canada’s position. In that case, Venezuela and Brazil complained about the discriminatory nature of the American standards on gasoline, which were aimed at environmental protection. The United States unsuccessfully used as a defence the general exceptions in Article XX of GATT. It is clear from both the decision of the Panel and of the Appellate Body that under the regime of the test of necessity in Article XX of GATT, practical difficulties in the implementation of an alternative measure do not exclude that measure as a less trade-restrictive measure.

225. In the case of United States – Standards for Reformulated and Conventional Gasoline, the United States pleaded, as in this case do the European Communities, that the alternative measure invoked – the establishment of individual baselines for foreign refiners – was not reasonably available and could not be upheld because it presented practical difficulties in implementation. In support of their argument under Article XX of GATT, the United States submitted to the Panel that the application of an individual baseline to foreign refiners was not feasible for three reasons: (i) the impossibility of determining the refinery of origin of each imported cargo; (ii) the temptation for exporters and importers to “manipulate” the system; (iii) the difficulty for the United States of assuring compliance by foreign refineries with the Gasoline rule, since, to be effective, that, it was supposed, would require penal and civil sanctions. In its communication to the Appellate Body, the United States reiterated its positions:

“The impracticability of verification and enforcement of foreign refinery baselines in this instance shows that the ‘discrimination’ is based on serious, not arbitrary or unjustifiable concerns stemming from different conditions between enforcement of its laws in the United States and abroad. (Footnote omitted).”

226. The Panel rejected the American argument based on the practical difficulties of implementing individual baselines for foreign refiners. The Appellate Body, for its part, although its decision mainly related to Article XX(g) of GATT and the introductory paragraph of Article XX of GATT, cited and approved the reasoning of the Panel concerning the incidence of practical difficulties in implementing individual baselines for foreign refiners on the application of the test of necessity in Article XX. Thus, the Appellate Body states:

“The United States stated that verification and enforcement of the gasoline rules requirements for imported gasoline are ‘much easier when the statutory baseline is used’ and that there would be ‘a dramatic difference’ in the burden of administering requirements for imported gasoline if individual baselines were allowed.
While the anticipated difficulties concerning verification and subsequent enforcement are doubtless real to some degree, the Panel viewed them as insufficient to justify the denial to foreign refiners of individual baselines permitted to domestic refiners. (Footnotes omitted).227

It is interesting to note that the Appellate Body, like the Panel, recognizes that the practical difficulties of implementing the alternative measure are indeed real. That does not prevent them from considering it as a measure reasonably available to achieve the environmental objectives pursued by the United States. The Appellate Body thus confirms the reasoning and conclusions of the Panel whereby the existence of practical difficulties in implementing an alternative measure do not disqualify it as a less trade-restrictive alternative, under the test of necessity.133 For the Appellate Body and the Panel, an alternative measure fulfils the test of necessity even if the measure preferred by the defending party is "much easier" to implement than the alternative measure, and that there is "dramatic difference" in the administrative burden between the measure under consideration and the alternative. For the Appellate Body and the Panel, the alternative measure will be excluded only if it is clearly impossible to implement it.134 The decision of the Appellate Body and the Panel in United States – Standards for Reformulated and Conventional Gasoline clarify the principles outlined by the Panel in United States – Section 337 of the Tariff Act of 1930.135 The decisions of the Appellate Body and the Panel in United States – Standards for Reformulated and Conventional Gasoline can be seen as an extension of the Panel's decision in United States – Section 337 of the Tariff Act of 1930. It should be recalled that in that case, the Panel had rejected the basic American claims that the disputed provision met the test of necessity under Article XX(d) of GATT 1947 "because of difficulties with service of process on and enforcement of judgement against foreign manufacturers."136

Concerning the dispute between Canada and the European Communities regarding measures concerning asbestos and products containing asbestos, Canada’s position in relation to the problem of practical difficulties in implementation hinges on two points. Firstly, we maintain that to achieve its objective of protecting human health, the French Government had available to it an alternative less trade-restrictive measure. Instead of acting in haste and under pressure from its public opinion, it could have introduced a regulatory framework under which prohibitions and authorizations of products containing asbestos would have been established in a rational manner on the basis of two guiding principles: (i) assessment of the risks on a product-by-product and use-by-use basis; (ii) analysis of the feasibility and effectiveness of controlled use for each product. Canada maintains that such a regulatory framework would not be difficult to implement in practice and that it would not be impossible to introduce. In that light, Canada maintains that there is a less trade-restrictive and reasonably available alternative measure to achieve the objectives pursued by France. Secondly, Canada maintains that in any case and as a general rule, any practical difficulties in implementing a measure are not a factor to be considered in assessing whether the measure is reasonably available and applying the test of necessity under Article XX of GATT, unless it is clearly established that these practical difficulties make implementation absolutely impossible. Canada’s position relies partly on the cases of United States – Standards for Reformulated and Conventional Gasoline and United States – Section 337 of the Tariff Act of 1930 and, secondly, on the conviction that taking account of the practical difficulties of implementation in applying the test of necessity would weaken the GATT disciplines by indirectly rewarding laxity and inefficiency.

Reply 3(b)

The European Communities and Canada agree that: (i) Article 2.2 of the TBT Agreement contains a test of necessity; (ii) the analysis involved in this test of necessity is similar to that relating to the test of necessity in Article XX(b) of GATT. Consequently, our reply to question 3(a) on Article XX of GATT applies equally to Article 2.2 of the TBT Agreement. In particular, the principles formulated by the Panel and the Appellate Body in United States – Standards for Reformulated and Conventional Gasoline cited in Canada's reply to question 3(a) are wholly applicable in the context of the test of necessity in Article 2.2 of the TBT Agreement.
(ii) Reply by the European Communities

230. All the Panel and Appellate Body reports that have examined so far the test of "necessity" under Article XX:(b) of GATT have come to the conclusion that a restrictive measure taken by a Member is not necessary if an alternative measure, which that Member could reasonably be expected to employ and which is not inconsistent with other GATT provisions, is available to it.137 A careful look at the rationale of the Panels demonstrates that the alternative measure must be: (i) effectively available to the Member in question; (ii) reasonably expected to be employed; (iii) not inconsistent or less inconsistent with other GATT provisions; (iv) capable of ensuring the Member’s desired level of health protection. Thus, in the Section 337 report the Panel held that "neither Article III:4 nor Article XX:(d) puts obligations on contracting parties specifying the level of protection that they should accord to patents or the effectiveness of procedures to enforce such protection."138 Consequently, the Panel held that:

- A different scheme for imports alleged to infringe process patents established by Section 337 was not necessary because the alternative of granting jurisdiction to civil courts over imports of products manufactured abroad was available and actually applied both in third countries and by the United States (paragraph 5.28);

- Presidential review in order to secure compliance with United States patent legislation and the difficulties with service of process on and enforcement of judgments against foreign manufacturers was not objectively necessary mainly because no equivalent requirements were applicable involving products of domestic origin (paragraphs 5.29 and 5.30);

- A system for the enforcement of in rem orders was necessary because it would “generally be more difficult” and “seldom feasible” to secure enforcement of the rulings of a court in the country of production. The alternative measure (i.e. an action in personam) was judged to be not “an adequate substitute ... in all cases” because, inter alia, importers might be “very numerous and not easily” brought into a single judicial proceeding (paragraph 5.31). For these reasons the panel concluded that there could be “an objective need” in terms of Article XX:(d) to apply limited in rem exclusion orders to imported products only. On this point, therefore, the lack of an alternative feasible and equally efficient measure played an important role in deciding "necessity";

- exclusion orders which were automatically enforced by the United States customs service were also found to be necessary because such enforcement at the border was considered to be "a means necessary to render such orders effective" (paragraph 5.33).

231. In the Thai Cigarette case the Panel found that the reasons advanced by Thailand to justify the import restrictions at issue were to protect the public from harmful ingredients in imported cigarettes and to reduce the consumption of cigarettes. So the measures were intended to ensure the quality and reduce the quantity of cigarettes sold in Thailand. But the Panel found that the Thai concerns about the quality of cigarettes consumed could be met “with strict, non-discriminatory labelling and ingredient disclosure regulations ... coupled with a ban on unhealthy substances” (paragraph 77). As regards the Thai concerns about the quantity of cigarettes consumed, the Panel noted the view expressed by WHO that demand for cigarettes, in particular by the young, was influenced by cigarette advertisements and that bans on advertisements could therefore curb demand, and that Thailand could also restrict the supply of cigarettes by maintaining governmental monopolies on the importation and domestic sale of products. Consequently, the Panel concluded that Thailand’s practice of permitting the sale of domestic cigarettes while not permitting the importation of foreign cigarettes was not necessary within the meaning of Article XX:(b) of GATT (paragraphs 78-81).
232. In the Reformulated Gasoline case, the Panel examined carefully all the arguments of the United States in order to determine whether it was practically feasible to assign to foreign producers an individual baseline so as to enable imported gasoline to benefit from the same favourable sales conditions under the Gasoline Rule as domestic gasoline. The Panel held that preventing imported gasoline from benefiting from as favourable conditions of sale as domestic gasoline was not necessary to achieve the stated goals of the Gasoline Rule, that is to reduce air pollution resulting from the consumption of gasoline. The reasons on which the Panel based its findings were that the United States had not satisfied its burden of proving that it was "not feasible" to establish individual baselines for foreign producers, or that there were any reasons that "precluded the effective use" of individual baselines, or that there were "any particular difficulty" sufficient to warrant the method of establishing baselines used by the United States (paragraphs 6.23-6.26). It also found the that United States did not meet its burden of showing whether the "gaming" concern "would actually occur", and that slightly stricter overall requirements on non-degradation of gasoline could not be implemented by the United States "at any time" (paragraph 6.27). The Panel also found that "the imposition of penalties" on importers was "an effective enforcement mechanism" used by the United States in other settings, because the United States had not demonstrated that the data available from foreign refiners was "inherently less susceptible" to established techniques of checking, verification, assessment and enforcement than data for other trade in goods subject to United States regulation (paragraph 6.28).

233. In consequence, in order to decide whether an alternative measure is reasonably available, panels have looked at the specific facts of each individual case. In all the three Panel Reports mentioned above, the Panels identified specific alternative measures that were objective, effective and constantly available to the Member applying the inconsistent measure in question. They also found that theoretical or potential measures, i.e. measures that were not real, feasible and as effective in practice as the measure applied, were not enough. The Panels and the Appellate Body appear, almost invariably, to have judged a measure not necessary, simply on the grounds that the same or an equivalent measure was not applied to products of domestic origin. They also found that the effective alternative measure should be clearly capable of achieving the pursued legitimate objective (i.e. achieve the desired level of health protection). It follows that objective difficulties in the effective application of the possible alternative measure that risk compromising the desired level of protection render the measure in question not reasonably available and one not reasonably expected to be employed by the Member.

234. To reply specifically to the Panel’s question, objective difficulties in the practical application of a measure play a crucial and determining role in deciding whether that measure is a reasonably available alternative measure to the one actually applied by a Member. As the European Communities have already explained in their written and oral submissions, the possible difficulties in the practical application of a measure may be of a wide and diverse nature, e.g. practical, technical, legal, economic, scientific or a combination of two or more of these reasons. The European Communities have already identified in its submissions a large number of such objective difficulties. The scientific experts have also confirmed those difficulties, in writing and orally. The outcome of their evaluation was that all these difficulties render “controlled use” not feasible or practicable. In deciding whether an alternative measure is reasonably available, Panels have to examine whether it is objectively available, feasible, effective and proportional to the pursued legitimate objective of protecting human health. In so deciding, one should always keep in mind that Article XX:(b) of GATT clearly allows contracting parties to give priority to human health over trade liberalization. Therefore, the result in this particular case is that so called “controlled use” does not achieve the level of health protection desired by France.

235. The same applies in the context of Article 2.2 of the TBT Agreement. Possible objective difficulties in the practical application of a measure play an equally crucial and determining role in deciding whether that measure is less restrictive on trade than the one actually applied. The text of Article 2.2 is clearer on this point because it provides that when applying the necessity test (“creating unnecessary obstacles to international trade”)
and “more trade-restrictive than necessary”), panels should take into account “the risks that non-fulfilment of the legitimate objective would create”. This clarification imparts meaning to the concept of necessity and indicates the nature of the measure under examination, because the mere existence of a possible alternative measure will not render the measure actually applied more trade-restrictive if the former cannot achieve the desired level of health protection.

236. Under Article 2.2 of the TBT Agreement, as under Article XX:(b) of GATT, the possible objective difficulties in the practical application of a measure may be of a wide and diverse nature, e.g. practical, technical, legal, economic, scientific or a combination of two or more of these reasons. Article 2.2, in fine, provides some examples, inter alia, of the elements that Members (and Panels) may take into account in deciding the less trade-restrictive nature of the measure actually applied and whether an alternative measure, which can fulfil the pursued legitimate objective, is reasonably available to the Member in question.

237. Without going as far as Article 5 of the SPS Agreement, Article 2.2 of the TBT Agreement nevertheless goes textually further than Article XX:(b) of GATT, because it explicitly links the necessity test with an assessment of the risks to human health and the design and structure of the measure under examination. Moreover, by analogy, Article 5.6 of the SPS Agreement and the notes thereto clarify that “a measure is not more trade-restrictive than required unless there is another measure, reasonably available taking into account technical and economic feasibility, that achieves the appropriate level of sanitary or phytosanitary protection and is significantly less restrictive on trade”. Although the principles and provisions of the SPS Agreement are not applicable in the context of GATT and the TBT Agreement, the underlying philosophy of these provisions may nevertheless assist us in interpreting Article XX:(b) of GATT and Article 2.2 of the TBT Agreement as regards the need, for the purpose of judging the necessity of a measure, to take into account the risks to human health arising from difficulties in the practical application of a measure.

238. Applying the above principles to the facts of the present case, it is clear that a number of practical, technical, legal, economic and scientific reasons do exist such that there is no alternative measure reasonably available to France to the total ban on all kinds of asbestos. The European Communities have discharged their burden of proof, because the reports of numerous competent international institutions (e.g. the WHO Environmental Health Criteria Report Number 203 and the IARC Monographs) clearly confirm the findings of the INSERM Report, and the scientific experts chosen by the Panel fully endorsed INSERM’s findings and the arguments of the European Communities. In the case of a substance that has been classified officially as a proven human carcinogen and for which it is impossible to establish scientifically an exposure threshold and when there is a vast category of persons that may be exposed to it because of the diverse nature of their activities, difficulties in the practical application of so-called “controlled use” make it clearly not a reasonable alternative measure for France, in the sense explained above, to the total ban on any kind of asbestos. Canada’s belated attempt to limit the scope of this dispute to high density chrysotile-containing cement products flies in the face of the available scientific evidence.\textsuperscript{141} For example, the WHO Report Number 203 states:

“Some asbestos-containing products pose particular concern and chrysotile use in those circumstances is not recommended. These uses include friable products with high exposure potential. Construction materials are of particular concern for several reasons. The construction industry work force is large and measures to control asbestos are difficult to institute. In-place building materials may also pose risk control to those doing alternations, maintenance and demolition. Minerals in place have the potential to deteriorate and create exposures.”\textsuperscript{142}

239. These WHO findings have been fully endorsed by all four scientific experts chosen by the Panel. The available evidence and the experts consulted by the Panel concur with the European Communities in saying that so-called “controlled use” is not really fea-
sible or practicable. Moreover, it is important to emphasize that the available evidence and the experts also confirm that even if “controlled use” were feasible in practice, in those cases where it might be applicable it is still inherently not effective because it cannot eliminate all risks. Canada has never argued that there was “no risk” at low levels of exposure, but merely that it becomes “undetectable”. The available evidence and the experts consulted by the Panel (including one of the experts of Canada, Dr. McDonald) confirmed during the meeting with the experts on 17 January that the linear model is appropriate and that there is no scientifically established threshold of exposure below which there is no risk. The experts also agree with the European Communities that to apply “controlled use” would entail very serious health risks as well as technical, administrative and economic costs to France. They also agree that the application of administrative controls and fines do not act as a real deterrent, because too many and serious violations still occur very frequently in countries which do not prohibit asbestos completely but simply try to control its use. Therefore, the risks associated with asbestos in place and the measures taken to deal with them are not relevant in deciding whether the “controlled use” proposed by Canada is a reasonable alternative measure to the total prohibition of asbestos. In the absence of any discrimination in the application of the total ban, the legitimate health objective pursued by France by far outweighs any economic benefits to Canada. Once again, the European Communities would like to emphasize that under Article XX(b) of GATT a WTO Member has the right to determine its level of protection as it considers appropriate. The European Communities would also like to emphasize that under Article 2.2 of the TBT Agreement a WTO Member is entitled to take measures “taking account of the risks non-fulfilment would create”, and that the ultimate risk in the case of asbestos is death. The European Communities have established in their submissions and oral statements that lung cancers, mesotheliomas and other asbestos-related diseases occur despite the appropriate use of adequate protection. Hence, the European Communities' objective to eliminate the risks induced by asbestos cannot be met by so-called “controlled use”, even if it were feasible in practice.

Question 4: Can Canada and the European Communities expand on their arguments concerning the implementation of measures relating to the use of chrysotile asbestos and its various applications (i) in Canada; (ii) in France prior to the prohibition imposed by the Decree. In particular, for France and Canada respectively, are there any data on the effectiveness of inspection and surveillance measures designed to ensure compliance with the regulations concerned, in particular in the various activities relating to the building industry? If yes, what do they show?

(i) Reply by Canada

240. Controlled use in Canada involves strict control of average concentrations of asbestos fibres in the workplace, the prohibition of friable products containing asbestos and the adoption of measures to ensure the health and safety of workers exposed to friable asbestos in place and chrysotile in high density products. Thus, controlled use consists of regulation, workplace inspections (especially construction sites), prevention through information and training and registration and monitoring the health of workers exposed to asbestos. In Canada, these measures normally come under the responsibility of the provinces and as the only Canadian asbestos mines are in Quebec and that is where the bulk of the chrysotile industry is located, our analysis focused on the situation in Quebec.

Existing legislation and regulations

241. The Quebec system of health and safety at work is the result of a broad consensus; it is the social contract which binds over two million workers and their employers in relation to occupational health and safety. In order to establish the rights and obligations of each, and to put in place the means to exercise them, Quebec passed the Occupational Health and Safety Act (LSST)\(^{43}\), which covers prevention and the Occupational Accidents and Diseases Act (LATMP)\(^{44}\), covering workers compensation and rehabilitation. This legislation makes employers and workers responsible for health and safety in their workplace. The Occupational Health and Safety Commission (CSST) is responsible for enforcing it. There are several implementing regulations under this legislation concerning the whole raft of
pollutants present in the workplace. In the case of the LSST, we would draw attention in particular to the Regulation on the Quality of the Workplace (Regulations)\(^{145}\) and the Code of Safety for Construction Works (the Code).\(^{146}\) In 1989, the Quebec Government amended all the legislation and regulations covering the workplace in order to take account of the provisions of Convention 162 of the International Labour Organization (ILO) concerning Safety in the Use of Asbestos.\(^{147}\) The Regulations and the Code govern working conditions in chrysotile mines, in plants manufacturing asbestos products, in enterprises specializing in maintenance of buildings and removal of asbestos flockings in Quebec and in the construction sector. Among other things, Section 3 of the Regulations governs air quality by prescribing the applicable standards, protective equipment and control measures for some 600 air pollutants in the workplace, including chrysotile asbestos. Annex A of the Regulations sets out the acceptable levels of exposure to dust. The weighted average exposure level for chrysotile asbestos is 1 f/ml. Article 5 of the Regulations provides that any establishment whose operations may lead to the emission of dust in the workplace must be operated so as to ensure that the concentration of dust, in the workers’ respiratory area, does not exceed the levels laid down in the regulations for any indicated period of time. Article 5 of the Regulations also provides that the use of crocidolite, amosite or a product containing either of these substances, is prohibited unless there is no reasonable and practicably feasible alternative.\(^{148}\) Subsection 3.23 of the Code applies to any construction site where work is carried out that may lead to the emission of asbestos dust.

242. The Code prohibits the spraying of a surface with a mixture of friable materials containing asbestos and the installation of friable insulation materials containing asbestos. In order to ensure the application of adequate protection measures, construction sites are divided into three categories: (1) low risk; (2) moderate risk; and (3) high risk. Sites where low risk work is carried out include installation, handling or removal of manufactured articles containing asbestos, provided that they are and remain in a non-friable condition, such as an asbestos-cement product. They also include sawing, cutting, machining and drilling of an asbestos-cement product with manual tools or electric tools equipped with a dust extraction system with a high performance filter. The Code provides that on a site where low risk works are carried out, such as sawing, cutting, machining, drilling of an asbestos-cement product with manual tools or electric tools equipped with a dust extraction system with high-performance filter, the employer must ensure that any worker present in the working area wears a breathing mask. The wearing of breathing masks is, however, not required for work such as installation, handling or removal of an asbestos-cement product. Sites where high risk work is carried out, include, for example, handling or removal of friable materials containing asbestos, by spraying a binding agent, and utilization of electric tools, not equipped with a dust extraction system with a high performance filter, for grinding, cutting, drilling or sanding an asbestos-cement product. On sites where high-risk work is carried out on friable materials in place, the employer must comply with much stricter requirements. These include wearing breathing apparatus of the semi or full mask type, sampling of the concentration of respirable asbestos fibres in the air in the working area at least once a shift, provision of protecting clothing to workers, providing workers with lockers for work clothing and personal clothing, provision of a shower room, isolation of the working area and the locker room for work clothes from the remainder of the building by a sealed partition equipped with an extraction ventilation system. The Code also provides that before work is commenced that might result in the release of asbestos dust, the employer must train and inform workers on the risks, methods of prevention and safe working methods. The training and information programme must include the employers general obligations, the effects of asbestos on health, the applicable standards and the sampling to be carried out, the workers rights and obligations, personal and collective protection measures and equipment, tasks to be carried out and the equipment or tools to be used, safe working methods and procedures, and methods of prevention and control.

Role of the Occupational Health and Safety Commission (CSST)

243. The Occupational Health and Safety Commission (Commission de la Santé et de la Sécurité au Travail) fulfils its administrative functions in a variety of ways. It is concerned,
inter alia, to prevent occupational injuries while at the same time playing the role of public guarantor for employers and workers alike. In addition, the CSST provides workers and employers with the services to which they are entitled. In the case of prevention it is involved in promoting occupational health and safety, assistance to workers and employers in their efforts to improve their working environment and eliminate hazards, and workplace inspections. The role of the CSST is mainly focused on prevention, specifically through the comprehensive analysis of the causes of occupational accidents and diseases. Consequently, when it investigates a workplace, all the chemical and physical pollutants, including chrysotile asbestos, and all working constraints are taken into account. The CSST requires the employer to implement a prevention programme. The prevention programme also includes a health programme, which involves monitoring workers' health for the prevention and early detection of any medical condition caused or aggravated by work.

Implementation of controlled use

244. In workplaces where, inter alia, chrysotile asbestos is found, a variety of actions for implementing the legislation and regulations under ILO Convention 162 has been carried out in recent years. More specifically, we present data for which we have performance indicators relating to inspection, health services, training and information for workers in a variety of environments, in particular, the construction industry.

245. Inspection: When an inspector investigates a construction site or an industrial establishment, he opens an investigation file and visits the work site on one or more occasions. During his visit, he may find various breaches of the regulations concerning chemical and physical pollutants and other provisions on occupational health and safety. He may take a number of actions, including closing the site which means halting the work until the appropriate remedial measures are put in place. The record of activities of the inspection service of the CSST in the construction sector show that in Quebec in 1999, 14,928 inspections were carried out in all areas of economic activity, including 5,171 inspections of construction sites. These inspections gave rise to 234 occupational health and safety proceedings relating to asbestos. The main reasons for proceedings were the following: lack of lockers or showers, failure to isolate the working area and the locker room from the remainder of the building by means of a sealed partition equipped with a dust extraction and ventilation system (37 cases), failure to dampen friable materials containing asbestos during removal works and failure to use a dust extraction system equipped with a high-performance filter to remove debris containing asbestos (28 cases), failure to provide disposable protective clothing or reusable protective clothing (27 cases), and failure to take daily samples of the concentration of respirable asbestos fibres in the air in the working area (5 cases). Eighteen sites were closed for non-compliance with standards. We should emphasize that in 1999, all asbestos-related occupational health and safety proceedings on building sites concerned exclusively friable asbestos products and not asbestos-cement products.

246. Training and information: The various training sessions developed and delivered by partner bodies in the occupational health and safety network are targeted at several types of client and, for that reason, the content and length varies. In 1997, the CSST took stock of its continuous training programmes for its inspectors and that led to the development of a specific training module on asbestos. As a result, in 1998 and 1999, the CSST delivered two types of training on chrysotile asbestos, a three-day course aimed at inspectors in the construction sector and specialists in the health network, and a two-day course for inspectors in industrial establishments. So far, the CSST has delivered eight two-day courses in which 77 inspectors participated. The three-day training sessions provided training for 80 construction sector inspectors and some 30 people from the health network. At the beginning of June 1999, in collaboration with its partners, the CSST launched the programme for the prevention of occupational diseases related to asbestos exposure. Aimed this time at workers and employers, the programme focuses on prevention measures to be taken not only on renovation and demolition sites, but also in repair and maintenance work. The objectives are to inform about appropriate working methods and suitable protective equipment for all work, which may give rise to the release of asbestos dust and ensure that they are used.
247. For its part, the joint sectoral association for occupational health and safety in the construction sector (ASP-Construction) has, since 1992, been offering a four-hour training course for construction workers. The course on asbestos safety meets the requirements on Article 3.23.7 of the Code. The participants in this course are mainly general labour, insulation appliers, pipe fitters, electricians, fire protection engineers, demolition workers etc.. The following table shows the annual distribution of data for this course provided by the ASP-Construction Consultants:

<table>
<thead>
<tr>
<th>Year</th>
<th>Number of Courses</th>
<th>Number of Participants</th>
</tr>
</thead>
<tbody>
<tr>
<td>1999</td>
<td>67</td>
<td>946</td>
</tr>
<tr>
<td>1998</td>
<td>34</td>
<td>509</td>
</tr>
<tr>
<td>1997</td>
<td>60</td>
<td>532</td>
</tr>
<tr>
<td>1996</td>
<td>31</td>
<td>350</td>
</tr>
<tr>
<td>1995</td>
<td>36</td>
<td>407</td>
</tr>
<tr>
<td>1994</td>
<td>12</td>
<td>136</td>
</tr>
<tr>
<td>1993</td>
<td>39</td>
<td>698</td>
</tr>
<tr>
<td>1992</td>
<td>16</td>
<td>245</td>
</tr>
<tr>
<td>TOTAL</td>
<td>295</td>
<td>3,828</td>
</tr>
</tbody>
</table>

248. Concerning information activities, the following table shows the annual breakdown of ASP-Construction data for the distribution of the Prevention Guide for Asbestos and notices to be placed at the entrance to every site when moderate or high risk works are carried out:

<table>
<thead>
<tr>
<th>Year</th>
<th>Number of Guides</th>
<th>Number of Notices</th>
</tr>
</thead>
<tbody>
<tr>
<td>1999</td>
<td>2,044</td>
<td>931</td>
</tr>
<tr>
<td>1998</td>
<td>936</td>
<td>410</td>
</tr>
<tr>
<td>1997</td>
<td>1,342</td>
<td>811</td>
</tr>
<tr>
<td>1996</td>
<td>857</td>
<td>415</td>
</tr>
<tr>
<td>1995</td>
<td>855</td>
<td>415</td>
</tr>
<tr>
<td>1994</td>
<td>614</td>
<td>560</td>
</tr>
<tr>
<td>1993</td>
<td>1,002</td>
<td>349</td>
</tr>
<tr>
<td>1992</td>
<td>1,272</td>
<td>723</td>
</tr>
<tr>
<td>TOTAL</td>
<td>8,922</td>
<td>4,614</td>
</tr>
</tbody>
</table>

249. A few firms specializing in occupational health and safety also provide eight-hour training sessions on chrysotile asbestos. The data for 1997, 1998 and 1999 are as follows:

<table>
<thead>
<tr>
<th>Number of Sessions</th>
<th>Number of Participants</th>
<th>Type of Enterprise</th>
</tr>
</thead>
<tbody>
<tr>
<td>350</td>
<td>4,600</td>
<td>GOVERNMENTAL INSTITUTIONS, BOARDS OF EDUCATION, UNIVERSITIES, HOSPITALS, ALUMINIUM WORKS, PULP AND PAPER FACTORIES, PRIMARY SECTOR FACTORIES, SECONDARY SECTOR FACTORIES</td>
</tr>
</tbody>
</table>

250. Health services: In the Montreal region, in September 1998, there were 23 establishments using chrysotile in their processes or operating plants which could cause occasional exposure of workers to chrysotile.
The preventative measures in place in these establishments include regular medical supervision, environmental monitoring, respiratory protection and information on health risks. All of these actions are encompassed in each establishment’s prevention programme.

Controlled use in France and the United States

252. We note that the French system for registering asbestos in place does not require the inclusion of chrysotile cement products. It appears that France, in its approach to management of the risks related to the use of asbestos, is not concerned with chrysotile cement products which do not present any detectable risk to health. The INSERM data are quite clear about the efficiency of controlled use in the French manufacturing industry. Of 2,480 people working in the asbestos processing sector in France in 1994, only two (0.1 per cent of workers) were exposed to concentrations of over 0.6 f/ml. Consequently, it is difficult to talk of the impossibility of controlled use in the processing industry when compliance with exposure standards throughout the territory of France required only one intervention concerning two workers. The American experience also shows the effectiveness and practicability of implementing a controlled use policy. As pointed out by Canada at the second substantive meeting, the OSHA in the United States only recorded 16 cases of exceeding the maximum exposure levels of 0.1 f/ml throughout America in the year 1998-1999. It should be noted, furthermore, that those exposures were related to friable materials, whose use is proscribed under the principles of controlled use as set out in international standards and by Canada. Yet again, hard to speak of ineffectiveness and the impossibility of controlled use.

Conclusion

253. All the above data shows that here have been many and varied preventative activities relating to chrysotile asbestos used in various working environments, including the construction sector. They effectively take account of the various provisions contained in the legislation and regulations. It is difficult in the case of asbestos to use medical performance indicators because of the latent period of the various pathologies related to exposure to chrysotile. But all these preventative measures make it possible to monitor exposure and to react in cases where limits are exceeded as the result of an incident, of whatever nature, in the working environment.

(ii) Reply by the European Communities

254. In France, the implementation of measures relating to the use of chrysotile asbestos and its various applications was completed in several stages, starting from the date...
when the carcinogenic nature of asbestos was recognized by the IARC (International Agency for Research on Cancer), in 1977. The first phase, covering the 1970s, was centred on the production sector, in the course of which France: (i) laid down specific rules for the use of asbestos in the manufacturing sector and transformation of asbestos-based products; (ii) restricted its use with a view to protecting the population. A second phase, covering the period of the 1980s and the first half of the 1990s centred on restrictions on use, during which the European directives came in: (i) strengthening workers’ protection in the manufacturing and transformation sectors; (ii) setting down rules for protection in the case of removal of asbestos; (iii) progressively eliminating the most dangerous varieties of asbestos; (iv) prohibiting the most dangerous uses; (v) restricting the use of asbestos in general, including the use of chrysotile. A third phase, from 1995 onwards, is centred on the repair and maintenance sectors, in the course of which France is concerned to: (i) take into account the risks faced by repair and maintenance workers; (ii) strengthen workers’ protection on sites where asbestos is removed and in certain residual asbestos processing centres; (iii) strengthen protection of the public.

First phase (1970s)

255. Following the recognition by the IARC, in 1977, that asbestos was carcinogenic, the French authorities issued specific regulations in order to ensure protection of workers against asbestos-related risks (Decree No. 77-949 of 17 August 1977 on Specific Hygiene Measures Applicable in Establishments where the Workforce is Exposed to the Action of Asbestos Dust, hereinafter “the 1977 Decree”). The provisions of that Decree supplemented the general measures on protection of workers and sanitary conditions in occupational premises and general rules of hygiene applicable to all places of work. During the same period, asbestos flockings were prohibited in buildings (Decree No. 78-394 of 20 March 1978 on Prohibition of Flockings in all Buildings). The overall provisions of the 1977 Decree imposed constraints on employers both in terms of collective protection of workers (for example: encasing of industrial systems and devices, humidification, dust extraction and ventilation of premises, compulsory monthly or quarterly removal of dust, maintenance of common protection equipment), and individual protection of workers (in particular, free provision of breathing apparatus). These provisions applied to all occupational activities involved in handling asbestos or products containing asbestos and compliance was monitored by labour inspectors along with all other measures relating to protection of health in the workplace.

Second phase: the 1980s and the period 1990-1995

256. During the second phase, French regulations developed in line with the provisions of the new European directives on asbestos. This development was also in step with progress in scientific knowledge, and international recommendations (in particular, ILO Convention No. 162 concerning Safety in the Use of Asbestos). In particular, the 1977 Decree was amended twice in order to translate the relevant European directives into French domestic law:

- The first modification of the 1977 Decree, by Decree 87-232 of 27 March 1987 was to incorporate the provisions of European Directive 83/477/EEC of 19 September 1983, which set the following limits for occupational exposure: 0.5 fibre/cm² measured over eight hours for crocidolite, and 1 fibre/cm² for all other varieties of asbestos, including chrysotile.

- Second amendment of the 1977 Decree, by Decree 92-634 of 6 July 1992, to incorporate the provisions of European Directive 91/382/EEC of 25 June 1991, which lowered the limits of occupational exposure to 0.6 fibre/cm² for chrysotile, and 0.3 fibre/cm² for all other varieties of asbestos. The European directive also introduced provisions for protection of workers and the environment specific to sites of removal of asbestos.
Parallel to the translation of the above-mentioned directives into French domestic law, the French authorities adopted measures to restrict sales which progressively prohibited the use of crocidolite, then all varieties of amphiboles. The use of chrysotile was also restricted: the use of chrysotile asbestos was prohibited in the manufacture of certain products such as toys, smokers articles, paint and varnishes, liquid filters, mortars, plaster, mastics, sizes, low-density insulating or soundproofing materials, bituminous roofing felt, and textiles susceptible of releasing fibres (Decree 88-466 of 28 April 1988 on Products Containing Asbestos). In any case, since the beginning of the 1990s, the risks for users of products containing asbestos were clearly emerging from the cumulative data. A database, called “EVALUTIL” was set up to evaluate exposure to asbestos among users of products containing asbestos. This database showed up the very high level of exposure of certain construction workers in the course of operations (“exposure peaks” during operations such as cutting asbestos cladding containing 5 per cent chrysotile, cutting up fireproof doors coated in asbestos, for example). Following the publication of certain studies which showed the increase in the number of mesotheliomas and the significant risk to the population of workers in the maintenance and repair sectors, the French authorities convened an expert group in 1994. The deliberations of these experts highlighted, firstly, a number of scientific doubts and, secondly, a number of loopholes in the French regulations then in force. The expert group’s conclusions led the authorities to draw up its first “asbestos plan” which was presented in outline on 6 July 1995 to the Council for the Prevention of Occupational Risks. The plan reflected the determination of the French authorities to strengthen the existing preventative measures and to equip itself with the means to obtain more detailed knowledge of the risks related to asbestos.

Third phase: the asbestos plan of December 1995

The asbestos plan of December 1995 contained immediate measures for the protection of workers and the public, as well as initiating a joint independent scientific report on the effects on health of various varieties of asbestos. This plan, whose measures are still in force today, is intended to reinforce the protection of workers in the asbestos industry, and also, above all, to take into account the risk inherent in asbestos in place for the public and workers in the repair and maintenance sectors. The asbestos plan of December 1995 contains the following provisions, in particular:

- The requirement for owners of buildings to identify flockings and laggings containing asbestos before 31 December 1999, and to have the condition of the flockings and lagging surveyed by a competent body in order to undertake the necessary works in the event of deterioration;

- reduction of the occupational exposure limits in activities involving work in contact with asbestos, to the lowest level technically possible, i.e. 0.1 fibre/cm³ (Decree 96-98 of 7 February 1996). The difference between the limit for “pure” chrysotile (0.3 fibre/cm³) and amphiboles (0.1 fibre/cm³) which only concerned the production sector, was abolished by Decree 96-1132 of 24 August 1996, with effect from 24 December 1996;

- improved procedures for asbestos removal: the power conferred on labour inspectors to close sites where asbestos is being removed where the protective measures do not seem to them to be adequate (amendment of Article L.231-12 of the Labour Code by Law 92-1446 of 31 December 1992, Article 35 and Law 96-452 of 28 May 1996, Article 39); prohibition on the use of casual workers to carry out the work of removing asbestos, introduction of a procedure of accreditation of firms (Decree No. 96-98 of 7 February 1996);

- elaboration of specific rules for prevention suited to repair and maintenance situations: the need to enquire of the owner whether asbestos
is present, mandatory individual protection whenever the presence of asbestos is suspected (Decree 96-98 of 7 February 1996).

259. These provisions completed the general arrangements for information and training of workers set out in the French Labour Code, by adapting them to the risks related to the inhalation of asbestos dust and the type and method of use of individual and collective means of protection. The regulations thus put in place satisfy the objective of reducing to the lowest level possible the risk presented by asbestos in place. In technical terms, they contain the tightest possible protection measures.

260. The introduction of French regulations on asbestos was accompanied by information campaigns directed at trade federations and, more particularly, the construction sector. All the French regions and all those involved in the prevention of occupational risks prepared numerous methodological guides, information leaflets, audiovisual programmes and training. To increase knowledge about the risks related to asbestos in the repair and maintenance sectors, the “EVALUTIL” database, which can evaluate asbestos exposure of users of asbestos-containing products, has been made accessible via the Internet. The Labour Inspectorate has been heavily involved and specially trained in the context of the Ministry of Labour’s priority actions. Sites where asbestos is being removed or sealed are strictly monitored: all removal plans are studied, all sites where “friable” asbestos and the vast majority of other asbestos removal sites are monitored by the Labour Inspectorate. For example, of 2,344 declared sites, 70 per cent were controlled in situ. Apart from their persuasive effect, the extent of which is difficult to measure, these inspections led to work being stopped on 114 sites because the protection measures were judged by the Labour Inspectorate to be inadequate, and have led to more than 3,000 notifications of failure to comply with French regulations.

261. The labour inspectors observe, in regional reports of priority actions, that the regulations are relatively well respected and that the asbestos risk is investigated and taken into account in major works such as large-scale renovation, demolition and works carried out by large companies. On the other hand, in regional reports of priority actions, the labour inspectors also note, objective and serious difficulties in enforcing the regulations in many small operations, particularly by private individuals. Indeed, the labour regulations are mandatory for employees and self-employed workers but not for individual do-it-yourself enthusiasts. In such situations, which by their nature are extremely diverse, little notice is taken of the need to investigate the presence of asbestos and take the necessary protection measures.

The French decision to impose a ban

262. The INSERM report, which was submitted at the end of June 1996, to the Minister of Labour and Social Affairs, showed the need to supplement the existing protection arrangements, as an essential step to reduce the risk of asbestos in place to the lowest possible level, by a ban which would stop the risk spreading. The reasons for this are as follows: (i) chrysotile is carcinogenic and it has not been shown that there is a threshold below which it is harmless; (ii) the vast majority of mesotheliomas appear among “secondary users”, particularly, in the construction sector, which means a very large number of people exposed in a very wide variety of situations in which the so-called “safe” use is inapplicable. Protection measures in force prior to the decision to impose the ban, even if technically the tightest possible, proved inadequate to reduce the risks in all situations to the lowest possible level. That being the case, if the French authorities had not imposed a ban, they would have knowingly allowed the volume of asbestos in place to increase, thereby increasing the risk to workers’ health, particularly that of “secondary users” (workers in the repair and maintenance sector and do-it-yourself enthusiasts).

2. Question by the Panel to Canada

Question 5: With respect to Articles III:4 of GATT and 2.1 of the TBT Agreement, Canada states that it is not invoking the argument of likeness of non-fibrous substitute prod-
ucts and neither does it extend the argument of likeness to substitute fibres other than glass fibre, cellulose fibre, PVA fibre and fibro-cement products containing such fibres (reply to question 15 of the Panel). Could Canada clarify if it considers that the Panel should limit its findings and conclusions to the effect of the regulations in relation to these three substitute fibres?

(i) Reply by Canada

263. Canada reiterates its reply to Question 15 of the Panel, as set out above in paragraph 74 and 75.

264. In the case of Indonesia – Certain Measures Affecting the Automobile Industry, it was clearly established that it was not necessary, for the purposes of Article III:2 of GATT, to determine whether the Honda Civic was like the Indonesian Timor since the demonstration of the likeness between the Toyota Corolla and the Timor was enough to establish the violation of Article III:2 of GATT.165 We note that the recent INSERM study on substitute fibres, Effects on Health of Fibres Replacing Asbestos, dwells specifically on the fibres mentioned by Canada as being like in the meaning of Article III:4 of GATT and 2.1 of the TBT Agreement, namely, glass fibres, cellulose fibre and polyvinylalcohols (PVA). INSERM states in its study:

“This study covered the main fibres used to replace asbestos: artificial mineral fibres (glass, rock and slag wool, continuous thread glass fibre, glass microfibres, refractory ceramic fibres), organic fibres (para-aramid, cellulose). The case of polyvinylalcohols (PVA) has not been considered because of the dearth of scientific literature on them”.

265. For the purposes of Canada’s argument under Article III:4 of GATT and 2.1 of the TBT Agreement, Canada is of the opinion that the findings and conclusions of the Panel should consider the effect of the regulations in relation to these three substitute fibres and fibro-cement products incorporating them.

3. Questions by the Panel to the European Communities

Question 6: Could you describe the type of measures applicable, in France, to substitute fibres, in particular glass, cellulose, PVA, para-aramid and refractory ceramic fibres?

(i) Reply by the European Communities

266. Substitute fibres are considered to be chemical substances. As such, their manufacture, use and sale are governed by a mass of complex European regulations which have been incorporated into French domestic law. In particular, both consumers and workers must be informed of the intrinsic danger of such fibres by means of labelling and “risk warnings” based on the European classification of their risk to human health.167 This requirement to provide information also exists at international level. Discussions are currently taking place in an endeavour to harmonize classification systems at international level. At the moment, the European Union has a system of classification of chemicals and chemical preparations which is applied uniformly throughout the member States, and consequently in France too. The preventative measures applicable to chemicals and chemical preparations vary according to the level of risk as determined by their classification. The applicable regulations depends on the level and type of hazard. The following regulations, in rising level of risk are applied cumulatively, depending upon the particular characteristics of the chemical or chemical preparation: (i) for chemical agents not classified as dangerous, reference must be made to the traditional regulations on ventilation and air quality in the workplace168; (ii) for chemical agents classified as dangerous the regulation on chemical risks has to be applied169; (iii) for chemical agents classified as proven human or animal carcinogens, the regulations on carcinogenic risks applies.170 Furthermore, breaches of these provisions, like all breaches of the provisions of the Labour Code and implementing legis-
lation are criminal offences pursuant to Article L. 263-2 which lays down financial penalties (25 000FF per employee) and, in appropriate cases, a term of imprisonment.

267. Of the fibres mentioned in question 6, three fibres are not classified as dangerous: PVA, para-aramid and cellulose. Nonetheless, the general regulations on ventilation and air quality in the workplace apply and mean that: (i) systems for ventilation and dust capture at source must be installed; (ii) dust levels must be below the thresholds set out in the Labour Code.

268. Glass fibres are classified as irritant and carcinogenic “Group 3” (i.e. suspect in animals under the IARC classification); in this case the regulations on prevention of chemical risks apply, which means that the employer must satisfy the following requirements: (i) risk assessment (levels of collective and individual exposure, methods envisaged to reduce them); (ii) installation of collective protective equipment; (iii) supply and maintenance of individual protective gear; (iv) training and information for workers; (v) signposting of premises where the substances are used; (vi) an explanatory notice on the risks and means of protection specific to each work place. For this purpose the employer has safety information sheets provided by the manufacturer, which give details of the composition of the product, the hazards and the prevention measures to be taken.

269. As ceramic fibres are classified carcinogenic “Group 2” (proven risk to animals under the IARC classification), their use is subject to the regulations on prevention of carcinogenic risks, which incidentally is more restrictive than that concerning chemical risks mentioned above, because in addition it requires: (i) substitution by a less dangerous product when that is technologically possible; (ii) use of a closed system when that is practicable; (iii) limitation of the quantities used and the number of workers exposed; (iv) reduction of exposure to the lowest level technically possible; (v) special medical monitoring of workers.

270. Following the results of the collective INSERM report on asbestos substitute fibres, the French authorities launched a plan of action on artificial mineral fibres (in particular glass wool and ceramic fibres) which provides for: (i) monitoring of the compliance of labelling of the various fibres, in particular, glass wools; (ii) monitoring of exposure levels; (iii) improving knowledge in the area of toxicology and epidemiology.

Question 7: Canada claims that France should have used two guiding principles to determine which chrysotile asbestos products should be used: (i) risk assessment product by product and use by use, and (ii) demonstration of the feasibility and effectiveness of “controlled use” for each product. Could the European Communities comment on these arguments?

(i) Reply by the European Communities

271. Both Canada’s arguments are incorrect for the following reasons.

272. First, neither GATT nor the TBT Agreement lay down any rule whatsoever on how to perform a risk assessment. Even the SPS Agreement, which is not applicable in this case and which contains specific provisions on risk assessment, does not require the performance of a risk assessment in the way suggested by Canada.

273. Secondly, there are in fact no internationally agreed and binding rules on how to conduct a risk assessment for dangerous substances like asbestos. In addition, neither national nor international practice (e.g. by WHO, IARC, FAO/Codex Alimentarius, etc.) support the views of Canada on the two “guiding principles”. It is common practice to assess the risks posed by chemical substances or other potentially dangerous products by evaluating in general the physico-chemical properties, toxicological and other relevant data on the pharmacologically active parent compound and its metabolites, and the possible ecotoxicological effects where this is likely to be relevant to the assessment of the risks posed by the substance in question. When the risk assessment indicates that an Acceptable
Daily Intake (ADI) and a Maximum Residue Limit (MRL) can be set for the substance or product in question, they can subsequently be used without further examination on a product by product and use by use basis, contrary to what is incorrectly suggested by Canada. Risk assessment, however, is a very complex and interactive process and no one particular technique or methodology is always appropriate for all cases. Epidemiological data and occupational exposure data may also be used to support toxicological data from in vitro or in vivo studies.

274. Thirdly, the WHO and IARC Reports that evaluated asbestos, including chrysotile asbestos, do explain the methodology they used. A close look at them shows that, in fact, they evaluated the risks posed by chrysotile asbestos to a very large extent in a way similar to the one suggested by Canada but have arrived at conclusions opposite to those of Canada.

275. Fourthly, the risk assessment carried out by WHO is very similar, if not identical, to the one performed by INSERM and confirms on all essential points the results of the INSERM Report.

276. Fifthly, as regards substances that have been classified as proven human carcinogens, such as chrysotile asbestos, and for which there is no scientifically established threshold of exposure, the two guiding principles suggested by Canada are in fact totally irrelevant, because any exposure to chrysotile or any kind of chrysotile-containing product may pose a risk to human health (because there is no safe exposure threshold). Therefore, the nature and number of the final products and their use does not matter as such. What is important is the fact that they contain the very carcinogenic substance that has been evaluated, i.e. chrysotile asbestos.

277. Sixthly, it is not correct to suggest that France and the other competent international organisations that have evaluated chrysotile asbestos failed to demonstrate the lack of feasibility and effectiveness of the so-called “controlled use”. As the European Communities have explained in detail in their written and oral statements, cases of lung cancer and mesothelioma have been observed or reported even in cases where strict measures have been taken to control exposure to asbestos. It is interesting to note that during the meeting of the Panel with the scientific experts on 17 January 2000, none of the experts was aware of any specific case where the so-called “controlled use” advocated by Canada was applied. And even Canada, in the same meeting, failed to provide a specific example of a country where such “controlled use” has ever been effectively applied.

278. To sum up, the existing rules permit France to apply its own, customary and normal rules on risk assessment to asbestos. This is what France did in this case. The methodology applied by France is similar, if not identical, to the one usually applied internationally and actually used by WHO and the IARC in the case of asbestos. It is up to Canada, not France, to perform the different, novel type of risk assessment it now advocates in order to demonstrate the accuracy of its claims. That Canada has failed to do.

II. QUESTIONS TO THIRD PARTIES

C. FIRST SUBSTANTIVE MEETING - SESSION WITH THE THIRD PARTIES (2 JUNE 1999)

1. Questions from the Panel to Brazil

Question 1: Could you please develop or expand upon the argument you make concerning developing countries in paragraph 4.23 of your written submission, particularly in light of the fact that this is not a point that has been raised by either party to the dispute?

279. In the Marrakech Declaration of 15 April 1994, adopted at the time of signature of the Uruguay Round Final Act, Ministers welcomed the notably active role of developing countries in the negotiations, declaring that “[t]his has marked a historic step towards a more balanced and integrated global trade partnership.” (Paragraph 4.) Ministers also
recalled that the results of the negotiations embodied provisions conferring differential and more favourable treatment for developing economies. (Paragraph 5.) These Ministerial statements reflect the achievement of the goal set out in Part I, Section B, Paragraph (iv) of the 20 September 1986 Punta del Este Declaration launching the Uruguay Round negotiations. As set out in Brazil’s third party submission,174 this commitment to special and differential treatment is embodied in Article 12 of the TBT Agreement. The issue was not raised by Canada because it is not a developing country. It was not raised by the EC because, as responding party, it has no interest in acknowledging its higher level of obligation to Brazil (and Zimbabwe). But, this does not detract from the EC’s obligation to respect the special and differential treatment obligations established at Articles 12.2 and 12.3. A Member’s obligations do not depend on the procedural aspects of dispute settlement; rather, they are set out in the texts of the WTO agreements, in this case the TBT Agreement, in particular. In imposing the ban, France did not “take into account the special development, financial and trade needs” of Brazil or Zimbabwe, as required by Article 12.2. Nor did France ensure that the ban “[did] not create unnecessary obstacles to exports” of chrysotile from Brazil, as required by Article 12.3. This is why Brazil has asked the Panel to scrutinize the ban closely, especially as it applies to the chrysotile exported by Brazil. The ban’s unwarranted effects on Brazil and Zimbabwe provide a supplemental reason why the ban is inconsistent with France’s obligations under the TBT Agreement.

Question 2: To what extent can modern controlled use in both current and recent past practice ensure adequate safety standards throughout the life-cycle of products containing chrysotile asbestos?

280. Current controlled-use policies are available to guarantee safety throughout the life-cycle of chrysotile products. First, as a preliminary matter, Brazil notes that the most current research indicates that chrysotile alone does not present significant health risks. (Please refer below to Brazil’s response to the EC Questions 22 and 23.) Second, an examination of the life cycle of chrysotile products indicates that procedures can be adopted to guarantee safe use of chrysotile. The following discussion breaks the life-cycle into three stages: (i) manufacture; (ii) use and maintenance; and (iii) disposal. The product used as an example is chrysotile cement because it accounts for such a large portion of the market.

(a) Manufacture

281. The safety of workers manufacturing chrysotile cement and chrysotile cement products can be guaranteed through proper controls. The following controls have been demonstrated to reduce chrysotile concentrations to 0.1 f/ml: (i) wet-process treatment (asbestos is mixed with water to make a slurry from which fibres and dust cannot escape); (ii) closed-process treatment (after workers load airtight bags of asbestos into machines, the process is air-tight through the point where the final product (in which the fibres are encapsulated) is produced); (iii) filtered ventilation systems (to remove from the air those few fibres that escape the wet, enclosed process); (iv) special treatment of workers’ clothing (because those few fibres that do remain often cling to clothing, Brazilian cement plants provide special work clothes and launder them appropriately on-site); and (v) personal hygiene measures (supply shower facilities and double dressing rooms for workers to further address fibre deposition).

(b) Use and Maintenance

282. By “use and maintenance,” Brazil refers to the installation and maintenance of chrysotile cement products. The “waste” from use is discussed below under the heading “Disposal/Recycling.” For chrysotile cement products, the procedures set out in ISO-7337 guarantee a high level of safety in installation and use. ISO-7337 establishes guidelines for breaking/cutting/drilling cement products so that they can be installed with minimal release of fibres. In general, ISO-7337 requires breaking or cutting with special saws, using wet process and/or vacuum dust extractors. Brazil also would recommend proper training. Perhaps licensing is appropriate for certain procedures (e.g., removing flocking). At all other points during use, the fibres are encapsulated. However, in certain climates, regular
inspection (e.g., every 5 years) of exposed products (roofing tiles, in particular) may be appropriate. Any exposure from degradation of the product can be halted by using the spray sealant currently recommended for application to flocking.

(c) Disposal/Recycling

Disposal/recycling of asbestos cement products and “waste” should be handled with measures similar to their manufacture end-use - proper wet controls and cutting or breaking methods. Please recall that the EC focus on remediation of flocking is not at issue. Rather, the issue here is the disposal/recycling of encapsulated used chrysotile cement products and waste. In cement products, the chrysotile remains encapsulated. Indeed, when buried, for example, in a landfill, the fibres are as inert (or more inert) as they are in naturally appearing ores containing asbestos.

(d) Personal Protection Equipment

At each stage, proper controls reduce exposure and health risks to de minimis levels. With the use of personal protection equipment (PPE), exposure can be reduced to zero. PPE absolutely guarantees no exposure and thus no risk to health. PPE could constitute “deep sea diver suits.” However, for any modern application involving chrysotile products, a simple air filter, coupled with proper working procedures, will eliminate exposure. In addition, Brazil refers the Panel to Brazil’s answers to EC Questions 10 (flaws of the linear risk model), 16 (safety of modern uses) and 22 and 23 (chrysotile that is not mixed with or accompanied by any amount, even a trace amount, of amphibole or substitute fibres presents no risk; modern uses of chrysotile present no risk).

2. Questions from Canada to Brazil

Question 1: What is Brazil’s position on the relative health effects of chrysotile, amphiboles and man-made fibres?

In considering the question of relative health effects, one must keep in mind four basic truths. First, as accepted and employed by the EC in its rulemaking (see Brazil’s response to EC Question 7), the toxicity of fibres is defined primarily by reference to the size, shape and durability and duration in the lung of the fibre in question. Thus, experts posit that any fibre with characteristics similar to chrysotile (due to its being engineered to substitute for chrysotile) must be a suspected carcinogen. Chrysotile has been proven safer than amphibole asbestos. As INSERM concedes and the studies cited by Brazil in response to EC Question 5 establish, this fact is not subject to question. Equally incontrovertible is the proposition that some man-made fibres have been proven more dangerous than chrysotile or even amphibole. See the discussion and the studies cited at Brazil’s response to EC Question 7. Recent research demonstrates that chrysotile presents no health risk. Please refer to the discussion and studies presented at Brazil’s responses to EC Questions 22 and 23. Scientists now suspect as a result of recent studies that health effects once associated with chrysotile are due to the fact that the past studies examined subjects exposed to chrysotile and amphibole and that the amphibole was responsible for the health effects.

Question 2: What is Brazil’s position as to whether and how chrysotile use can be controlled to guarantee safety?

For years, Brazil has controlled the mining, production and use of chrysotile and chrysotile products to guarantee safety. As explained by Brazil in response to Panel Question 2, controls exist for every part of the life cycle, from the time chrysotile is mined until chrysotile cement products are disposed of. Even as the EC presents the grossly exaggerated picture of workers equipped as “veritable deep sea divers,” the EC provides the seed of an admission which, germinated, compromises its position. The EC concedes that use can be safe and, in doing so, admits that a ban is not the least trade restrictive remedy required to achieve its desired level of protection. But, in any case, the EC’s picture is quite
distorted. For a country that chooses to disturb flocked asbestos, guaranteeing worker safety by equipping workers as veritable deep sea divers may be rational; however, for modern controlled uses of chrysotile, the only relevant personal protective equipment (PPE) is a simple air filter, which reduces already safe exposure levels to zero.

Question 3: How does Brazil interpret the expression “current controlled use”?

287. The expression “current controlled use” is the key to this proceeding. Brazil is not defending past uses, e.g., flocking, the use of amphibole, or the use of naturally exposed asbestos in solution as a white-wash for buildings. Brazil is defending against France’s ban a limited range of beneficial, safe uses: the use of chrysotile in asbestos cement products and friction products. Brazil has demonstrated that these uses, which compose the majority of the market, are safe. Exposure can be controlled so that PPE is not needed to guarantee worker safety. In these limited applications, safety is guaranteed by controls, but PPE can be used to reduce exposure to zero.

3. Questions from the European Communities to Brazil

Question 1: Brazil alleges in a number of instances (e.g. in paragraph 4.28 of its written submission) that in the past France imported and used mainly amphibole asbestos. On what factual and/or statistical data are these allegations based? Could Brazil provide them to the Panel?

288. Brazil’s understanding that France once imported much larger amounts of amphibole than it now does is based on the history of world use of asbestos, not on French import statistics. For some time, countries did not distinguish between imports of chrysotile and the far more dangerous amphibole asbestos. During this period, the two types were used virtually interchangeably and countries, even Brazil, imported substantial amounts of amphibole. As the EC likely is aware, French import statistics do not distinguish chrysotile and amphibole.

Question 2: Could Brazil define what it means by “uncontaminated” chrysotile?

289. By “uncontaminated”, Brazil means chrysotile that is not mixed with or accompanied by any amount, even a trace amount, of dangerous amphibole or substitute fibres. An attached scientific assay demonstrates that the chrysotile that Brazil mines and exports is uncontaminated. Moreover, as discussed in detail in the answer to EC Question 5, INSERM also recognizes that chrysotile is less hazardous than amphibole. Nonetheless, INSERM did not address the health effects of exposure to chrysotile alone. The distinction is quite important. As McDonald (1998) concludes “… it now seems fairly widely accepted that the carcinogenicity of the amphiboles, crocidolite in particular, is appreciably greater than of chrysotile …” Thus, amphibole is viewed widely as the source of toxicity that at one time in the past was associated with the use of chrysotile.

Question 3: Could Brazil provide concrete written evidence supporting its allegation (e.g. paragraphs 4.3 and 4.9 of its written submission) that the French Decree at issue aims to deal exclusively with a public outcry, public pressure and to appease domestic sentiment and is not based on any scientific evidence, given the volume of internationally available scientific evidence documenting the risks of asbestos to human health?

290. Obviously, the French Government did not accompany the Decree with an official statement admitting the Decree was passed to appease the French public. No government would do such a thing. However, the facts preceding and surrounding the Decree and the INSERM Report, as well as the timing of the decisions taken, demonstrate that the Decree was not aimed primarily at protecting public health (for if it was, why are substitutes that are Class II carcinogens not banned?), but was designed to quiet a disturbed populace. In this regard, Brazil refers to Canada’s arguments (contained in Section III.B.3 of this Report) which set out a chronology showing how a series of unfortunate decisions, coupled with the public’s strong response, led the French Government to ban chrysotile,
without scientific basis. The text of the INSERM Report also supports this conclusion. At page 140, INSERM notes that exposure to flocked asbestos in buildings is the “core of the current concerns,” and emphasizes that the subject is “extremely controversial.” INSERM states that French courts had wrestled with the issue and ignored science in tying sickness to indoor exposure. Id. at pages 141-2. Ironically, INSERM concludes that the data “do not confirm” that maintenance and service personnel in “asbestos-sprayed buildings” (flocked buildings) experienced an increased risk of respiratory abnormalities. Id. at pages 143-4.

Question 4: Could Brazil provide concrete evidence supporting the allegation (paragraph 4.3 of its written submission) that the INSERM report was merely a scientific “cover” for a previously taken political decision?

291. Brazil refers to the answer to EC Question 3. In addition, Brazil notes that the fact that the INSERM Report was merely scientific cover for the political decision is confirmed not merely by reference to the events of the time, but also by a review of the Report itself. Even a cursory review shows that INSERM ignored evidence that did not conform to the opinion it apparently was instructed to reach.

Question 5: Could Brazil provide written evidence supporting the allegation (paragraph 4.3 of its written submission) that “especially the spraying of brittle amphibole” caused the asbestos-related health problems in France?

292. Flocked asbestos led to the health problems that most concerned the French officials contemplating a ban. That is incontrovertible as a review of the INSERM Report and the answer to EC Question 3, above, demonstrates. Likewise incontrovertible is the fact that flocking occurred during the time prior to any ban on amphibole asbestos. Indeed, INSERM states (page 18) that chrysotile fibres commonly were mixed “with up to 40 per cent amphibole fibers,” due to the different properties of the two types of fibres. The Report prepared by INSERM at the request and under the direction of the French Government recognizes that chrysotile is much safer than amphibole and thus that amphibole is of primary concern for negative health effects. INSERM concedes this point throughout the Report. For example, at page 102, INSERM states, “subjects exposed mainly to chrysotile have a lower risk of mesothelioma than subjects exposed to amphiboles or a mix of fibers.” Later, discussing Hughes et al. (1987) and Weill et al. (1977), INSERM reports that the prevalence of radiographic asbestosis was found to be higher in the plant that handles more amphibole asbestos. ” Id. at page 323. INSERM further concedes that "crocidolite has a greater fibrogenic effect than chrysotile." Id. at page 326. INSERM concludes that the studies demonstrate that the dose-effect relationship is much stronger with amphibole fibres than with chrysotile, as regards both asbestosis and the other pathogenic effects of asbestos. Id. at page 327. Anecdotal evidence supports the conclusion. INSERM explains the higher incidence of sickness in Australia and New Zealand by reference to the "wide-spread use of the crocidolite these two countries produce." Id. at page 158; [sic] see also id. at page 171 (concluding that, in Australia, "the incidence of mesothelioma is particularly high due to widespread crocidolite use"). This view is widely shared. In 1996, Health and Safety Executive of Great Britain concluded that:

“[v]ery few cases of mesothelioma can be reliably attributed to chrysotile despite the many thousands of workers who have had massive and prolonged exposures … In contrast, mesotheliomas have been observed among some workers who experienced only brief exposures to amphiboles.”

293. A U.S. Government official has expressed a similar view. According to Malcolm Ross, mineralogist for the U.S. Geological Survey, scientific studies show chrysotile is not as great a health risk as amphibole: “[t]here’s no non-occupational risk with chrysotile. … In the workplace, chrysotile should not show any noticeable increase of disease if it is controlled.” These studies and conclusions demonstrate that the primary concern regarding risk to human health is exposure to amphibole, most of which occurred during past uses.
Question 6: Could Brazil provide a copy of the full American Society of Mechanical Engineers (ASME) study it has cited (paragraph 4.5 of its written submission), as well as a reference to where it has been published?

294. Brazil does not cite the ASME study anywhere in its submission. Rather, Brazil cites the Fifth Circuit’s recounting and analysis of the testimony during the court proceeding of one of the study’s authors.

Question 7: In paragraph 4.6 of its written submission, Brazil claims that “available scientific data” would show that, compared to chrysotile, man-made fibers produced in France present greater risks when use is not controlled? Could Brazil provide copies of the relevant scientific articles and/or other data?

295. The available scientific literature demonstrates that chrysotile is safer than many if not all substitutes. Indeed, recent research suggests that chrysotile poses no health risk. In this respect, Brazil attached the following papers and studies: (i) Cossette (1998), concluding that iron ductile and PVC pipe present health risks far greater than that presented by chrysotile cement pipe; (ii) Hesterberg (1992), reporting that 35 per cent of hamsters treated with a variety of ceramic fibres developed mesotheliomas; and (iii) Peraud et al. (1994), concluding that, as compared to all types of asbestos, the man-made mineral fibres silicon carbide, JM 104/475, B1M, and B3K, have a “moderately higher toxic effect” (pages 570-2). In addition, the scientists and doctors attending The Proceedings of the Workshop on Health Risks Associated with Chrysotile Asbestos (1994) concluded that:

“With the exception of the textile industry, the slopes of the exposure response curves for lung cancer in the various chrysotile industry sectors were shallow, with no detectable risk of an extremely low level of risk of lung cancer associated with exposure to chrysotile asbestos at and below lifetime cumulative exposures of 30 fibres/ml-years. No chrysotile related increased risk was detected at considerably higher exposures in the mining sector.”

296. Moreover, in his report to the EC (EC document number ECB/TM/15(97)), Dr. Bernstein analysed statistically all available biopersistence studies, chronic intraperitoneal [injection] studies (IP) and chronic inhalation studies on fibres. This and subsequent research demonstrates that chrysotile is safer than refractory ceramic fibres (RCF), glass fibres, p-aramid fibres and cellulose fibres. Dr. Bernstein’s analyses for the EC were reviewed and accepted by the Ad-Hoc Group of Experts commissioned by the EC and consisting of scientific experts from EC member states and industries. The results of the analysis clearly showed that one can use inhalation biopersistence clearance half-times to predict IP tumour results, the number of long fibres remaining at 24 months in the Chronic Inhalation Studies and a score of pulmonary fibrosis in the Chronic Inhalation Studies (tumours were not used as an endpoint as only a single fibre type (RCF) was tumorigenic in these studies). Based upon this scientific analysis, the EC Fiber Directive included in Note Q the provision that:

“The classification as a carcinogen need not apply if it can be shown that the substance fulfils one of the following conditions: (i) a short-term biopersistence test by inhalation has shown that the fibres longer than 20 im have a weighted half-life less than 10 days; or (ii) a short-term biopersistence test by intratracheal instillation has shown that the fibres longer than 20 im have a weighted half life less than 40 days, or (iii) an appropriate intra-peritoneal test has shown no evidence of excess carcinogenicity; or (iv) an absence of relevant pathogenicity or neoplastic changes in a suitable long term inhalation test.”

297. The results from Dr. Bernstein’s chrysotile biopersistence study show that chrysotile fibres are rapidly removed from the lung. Fibres longer than 20 im are cleared with a half-time of 1.3 days, most likely by unravelling into shorter fibrils. This is consistent with the known chemistry of chrysotile. Shorter fibres also are cleared rapidly from the lung, with fibres 5-20 im clearing even faster ($T_{1/2} = 2.4$ days) than those < 5 im in length.
These short fibres were never found clumped together but appeared as separate, fine fibrils, occasionally unwound at one end. Short free fibres appeared in the corners of alveolar septa, and fibres or their fragments were found within alveolar macrophages. The same was true of fibres in lymphatics, as they appeared free or within phagocytic lymphocytes. These results support the evidence presented by McDonald and McDonald that the carcinogenicity of chrysotile largely depends on whether it is mixed with amphibole. Thus, chrysotile clears faster than most current glass fibres for which in the EC system, the clearance half-time of long fibres must be less than 10 days in order to be exonerated. In addition, chrysotile clears substantially faster (T½ = 1.3 days) than long p-aramid fibres which following a 90-day inhalation exposure cleared with a half-time of 95 days. Finally, chrysotile clears considerably faster than cellulose fibres for which the clearance half-time of WHO fibres was between 1046 days and infinity. One should note that the INSERM Report ignores the above analysis by Dr. Bernstein, as well as other studies comparing chrysotile to p-aramid and cellulose, even though they pre-date the INSERM Report.

Question 8: In footnote 11 of paragraph 4.10 of its written submission, Brazil states that “because only the INSERM report preceded the Ban, the Ban must be supported by the report alone.” Could Brazil explain why the INSERM report alone should support the ban given the volume of internationally available scientific evidence clearly documenting the risks of asbestos to human health? What provisions of the GATT 94 and/or the TBT Agreement (assuming this is applicable) support this proposition?

298. First, the Government of France ordered the INSERM Report precisely for the purpose of conducting a risk assessment and supporting the ban. In promulgating the ban, it relied on the INSERM Report as the risk assessment. For the ban to be legitimate, it must be supported by the risk assessment upon which it is based. That is a matter of logic, not of WTO rules. More importantly, however, INSERM and the French Government failed to conduct an unbiased risk assessment. The EC now appears before this Panel, suggesting that the actual risk assessment was not published, but occurred behind closed doors in a process in which French Government officials reviewed the INSERM Report and all of the relevant studies the EC concedes with this question that INSERM ignored. Then, and only then, suggests the EC, did the French Government issue its decision to ban chrysotile. There is no evidence that any such complex, thorough process occurred in the 24 hours between delivery of the INSERM Report and the issuance of the ban. Moreover, to countenance the EC’s attempt to insulate the French risk assessment from examination would be to make a mockery of the TBT Agreement. It would allow any Member to conduct a secret analysis and issue its foregone conclusion that a ban was necessary, without any oversight or discipline, whatsoever.

Question 9: In paragraph 4.11 of its written submission, Brazil states: “Finally, INSERM concedes that, although the health data it applied to chrysotile is from past, massive and prolonged exposure to amphibole (…).” Could Brazil provide the exact location (page number) where INSERM has stated this?

299. Throughout the Report INSERM relies on data from human exposure and animal exposure (mostly rats and hamsters). All of the studies of human exposure focus on exposure that occurred 20, 30, 40, 50 and 60 or more (for the earliest studies) years ago. See, e.g., INSERM Report at Tables 1 and 2 to Chapter 13, pages 321 through 323; Table 2 to Chapter 8, page 164; and Table 4 to Chapter 8, page 166, summarizing data on past exposure of workers in various asbestos product plants and other occupations, and comparing the incidence of disease in men and women. This point is not subject to debate. The primary fact differentiating current exposure from these older periods of exposure is that, during the older periods of exposure, amphibole was commonly used, and had not yet been banned. See Brazil’s answer to EC Question 5; INSERM Report at page 18. Nowhere does the INSERM Report distinguish between exposure to chrysotile only and to chrysotile mixed with carcinogenic amphibole. Another difference is that, in contrast to current practices, the work processes were not controlled and workers had little personal protective gear, much less safety accommodations such as showers and special clothing. A review of the dates of the studies summarized in the tables referred to above, and of the ages of the
subjects and the length of exposure indicates that many of the exposures studied were quite dated. See also the discussion of latency periods and disease onset at pages 156-59 of the INSERM Report.

Moreover, even in plants using “chrysotile,” studies suggest that a significant amount of the asbestos used was amphibole. For example, INSERM reports that the asbestos used in the Casale Monferrato plant was “chrysotile, but approximately 10 per cent of the total amount of asbestos was crocidolite.” INSERM Report at page 137. Further to the point, INSERM concludes regarding exposure from industrial sources, that “in all positive studies reported, when asbestos fibre type was specified, the fibres involved were from the amphibole group, or fibres containing some amphiboles (amosite, tremolite or crocidolite),” id. at 140. This applies even to the animal studies INSERM relied on. According to INSERM:

“Animal experiments also make it possible to evaluate the dose-effect relationship. Most of the studies have used massive quantities of fibers at concentrations far higher than seen clinically in humans. Recent work by Quinlan et al (1994) analysed the effects of lower concentrations, ranging from 0.1 to 10 mg of crocidolite per cubic meter of air.”

Question 10: In paragraph 4.12 of its written submission, Brazil criticises the use of the “linear risk model” used by France. Could Brazil provide scientific evidence explaining why this model, which is a very widely used one, is unfounded or inappropriate?

Brazil already has introduced scientific evidence regarding the limitations of the linear risk model as applied to chrysotile. Brazil assumes that a statement by INSERM would qualify, at least for France and the EC as “scientific.” As INSERM itself conceded, but apparently the EC wishes to ignore, the linear risk model cannot produce “scientifically certain knowledge” (INSERM Report at 239). Moreover, much of the research upon which INSERM relies for other propositions demonstrates that a safe threshold exists. For example: (i) at Table 2 to Chapter 6, page 88, INSERM reports that Wistar rats exposed to chrysotile in the amount of 6 fibres mg/m³ developed no lung tumours, in contrast to rats exposed to higher amounts or to other fibres; (ii) at pages 88-89, INSERM recounts that hamsters exposed to various amounts of chrysotile in three separate studies (Lee et al. (1981), Smith et al. (1987) and Hesterberg et al. (1991)) did not develop lung tumours; and at page 104, INSERM recounts a study of intermittent exposure at high concentrations and continuous exposure at lower concentrations (Davis et al. (1980)), and concludes that “the authors found no significant effect on the tumour rate in the different treatment groups.” In spite of these and other studies, INSERM supported the linear risk model. But when it came time to state why, INSERM (page 408), equivocated:

“No argument based on the analysis of existing epidemiological data, direct or indirect, supports the belief that no-threshold linear extrapolation based on data corresponding to higher levels of asbestos exposure (adopted in this report to quantify the risks associated with low-level exposure) is not the most plausible uncertain model.”

The conclusion, confused as it is, is utterly implausible given the evidence cited above from the Report itself, much less that discussed below. Moreover, earlier in the report, INSERM concedes that, even if one adopts a linear risk model, a safe threshold nonetheless exists. According to INSERM (page 104), if a:

“... straight-line relationship is maintained for smaller doses, it means there is no minimum below which all risk disappears. However, the tumour latency period increases in animals as the dose falls. This means that a threshold value should exist in practice, corresponding to a dose for which the latency period is greater than the “natural” life expectancy of the animal in question.”

Thus, INSERM concedes that, even assuming that all exposure is dangerous, at some lower exposure level, the latency period would be so long that no tumour would
occur before death from natural causes. The most recent literature on the subject rejects outright the linear risk model. For example, Mossman and Churg now conclude that “asbestosis does not appear until a threshold exposure level has been reached …” and “[e]pidemiological studies indicate very clearly that the development of asbestosis requires heavy exposure to asbestos and provide strong evidence that there is a threshold fibre dose below which asbestosis is not seen …”. In 1994, Churg et al. concluded that:

“Our results show clearly that, despite known historic exposure to amosite and chrysotile, amosite is by far the predominant residual fiber, and there are correlations between amosite measures and disease. Chrysotile was present inconsistently and in relatively small amounts, and no correlations were found between chrysotile measures and disease.”

304. According to Professor Patrick Brochard of the Pellegrin Hospital in Bordeaux, France, “chrysotile is carcinogenic, but not below a certain dosage level, unlike amphiboles”. Health and Safety Executive concludes in the Review of Fibre Toxicology that, for chrysotile, “the balance of toxicological evidence does not support the non-threshold model for asbestos-induced lung cancer. A practical threshold is likely”. Even the WHO suggests that a threshold exists for chrysotile. In discussing the incidence of mesothelioma in workers exposed to chrysotile, the WHO concluded in 1998 that “[n]one occurred in workers exposed for less than two years.” And this statement was in regard to workers in the mining and milling sectors, likely handling chrysotile and amphibole fibres.

Question 11: Has the article of Brazil’s Law No. 9055, cited in paragraph 4.17 of its written submission, according to which miners and wholesalers are prohibited from supplying chrysotile or substitute fibers to any company that does not comply with any provision of the law, already been put in practice? Could Brazil provide written evidence thereof, including copies of all relevant court or administrative decisions, statistics on the number of cases in which the article has been applied, etc.?

305. There has been no occasion to enforce this or any other article. As demonstrated in a document provided by Brazil to the Panel, supplies are immediately withdrawn as soon as non-compliance is discovered. The supply resumes only after compliance measures demanded by ABRA have been implemented. More importantly, the question is irrelevant to this proceeding. Brazil describes its own practice merely to show the Panel that exposure can be controlled. France’s obligation is to adopt the least restrictive means of achieving its own level of safety. That might require controls or restrictions in addition to those imposed by Brazil.

Question 12: Has the article of Brazil’s Law No. 9055, cited in paragraph 4.17 of its written submission, according to which there should be prompt Department of Justice enforcement against infractions of that law, already been put in practice? Could Brazil please provide written evidence thereof, including copies of all relevant court or administrative decisions, statistics on the number of cases in which the article has been applied, etc.?

306. See Brazil’s response to EC Question 11.

Question 13: Could Brazil please provide all available information on the results of its “research into and confirmation of the health effects of chrysotile and its substitutes” foreseen in its Law No. 2350 (paragraph 4.18 of its written submission)? Please provide copies also of those results.

307. Two types of research are under way: epidemiological research and research on bio-persistence. The epidemiological research is far from completion, but tracks the health of thousands of asbestos workers in Brazil. Results or a summary thereof will be provided as soon as they are available. The preliminary findings on bio-persistence are presented in Dr. Bernstein’s study, provided by Brazil to the Panel. Moreover, as discussed in re-
sponse to EC Question 7, research performed in the EC and elsewhere demonstrates that substitute fibres present significant health risks.

Question 14: The so-called Tripartite Agreements (paragraph 4.19 of its written submission), “make ABRA”, the Brazilian Asbestos Association “responsible for providing the companies with technical assistance regarding controls and preventive measures.” Has ABRA ever provided such assistance to French client companies of Brazilian chrysotile producers? If so, please provide evidence thereof.

308. Yes, ABRA provides assistance to those companies that join as members and request assistance. As the French Government well knows, among ABRA’s members is Brasilit S/A, a French company that is one of the largest cement fibre producers in Brazil. (Other ABRA members are German and United States companies.) Moreover, ABRA is available to assist any French or EC company needing assistance after the ban is lifted.

Question 15: Brazil’s written submission does not clearly explain what the Brazilian legal requirements and practices are regarding the waste generated by end-users of products containing asbestos, for instance when parts of asbestos-containing-products have to be cut off to fit certain uses or when buildings, installations, and other structures containing asbestos are demolished? Could Brazil please provide detailed information, including references to legislation, on this question?

309. Brazilian environmental legislation addresses “residues,” including chrysotile residues. The Tripartite Agreements require zero residue or complete recycling of residues in the industrial processes for producing chrysotile cement products. Also, imports of amphibole were banned years ago and the asbestos Brazil produces is chrysotile. Because Brazilian asbestos use is limited to chrysotile and chrysotile products, no health effects have been associated with disposal of cut-off pieces of asbestos cement pipe or roofing. However, cement companies have adopted procedures where they accept and recycle “waste.” End users are directed to return the waste to the point of purchase, which then returns it to the producer. The producer recycles the waste in its industrial process. Certainly, France has the ability to regulate chrysotile “waste,” just as it regulates wastes from many other production processes.

Question 16: In paragraph 4.33 of its written submission, Brazil claims that what it calls “modern-day” products containing chrysotile do not contain loose, friable chrysotile fibres. Could Brazil provide the data it possesses on exposure levels occurring when these products are actually used in practice (for instance when cutting or sawing these products in workplaces or homes, or when demolishing the buildings and other structures in which they are present)?

310. The Brazilian Government has not conducted or sponsored any studies on workplace exposure levels. However, as the EC well knows, many studies document that exposure levels depend upon the controls used. Brazil directs the EC to the INSERM Report at page 70. There, INSERM sets out exposure levels for uncontrolled activities. The levels range from 10 f/ml for a person silly enough to change a friction element on a machine for making corrugated fibreboard and then clean it with a blow gun, to 0.15 f/ml for a person removing a false ceiling. Do-it-yourself activities such as cutting an asbestos seal, and drilling holes in sprayed asbestos fall in between these levels and for the most part are under 1 f/ml. But these value are for uncontrolled use. Controlled use would yield much lower exposure rates. Moreover, controlled use, plus, as INSERM terms it “protection measures” would yield no or de minimis exposure. Indeed, at page 70, INSERM refers briefly to a study (CORN) in which the concentrations of fibres for “work in buildings” (“dismantling false ceilings, cable passageways, electrical work and encapsulation”), ranged between 0 and 0.228 f/ml.” On the related topic of the safety of controlled use, INSERM makes several interesting points. These data are relevant, INSERM states at page 71, because:

“In the area of para-occupational exposure, related in particular to do-it-yourself activities, there are no data in the literature allowing documentation of the
subject. In each basic operation - welding, cutting asbestos board, cutting or drilling asbestos cement and so on - it seems justifiable to consider the emission peaks to be identical to those found during industrial operations of the same type. The possible differences in exposure levels, in terms of inhaled doses, are eventually to be found in exposure times, as the do-it-yourselfer does not perform such operations as often as occupational workers."

Thus, experience with occupational exposure to asbestos is applicable to do-it-yourselfers, provided one recalls that exposure times are far less for do-it-yourselfers than for asbestos workers. The following discussion recounts INSERM’s conclusion regarding occupational exposure. First, INSERM notes that mesothelioma stems from past occupational exposure. According to INSERM (page 182), "stringent worker protection measures" can eradicate mesothelioma: “[b]ecause of the occupational source of asbestos exposure, mesothelioma incidence is not rising in a few countries that implemented stringent worker protection measures at an early date.” Regarding asbestosis, INSERM (page 327) notes that "[t]he current levels of exposure in industries that use asbestos directly should lead to an end of confirmed cases of asbestosis (Doll et al., 1985)." INSERM (page 327) also cites the Peto study, which the EC heavily relied on, for the proposition that "[e]xposure prevention measures have removed this disease [asbestosis] from the list of the causes of death at a British Textiles firm." Thus, contrary to the conclusion reached by the French Government and the position adopted by the EC in this proceeding, chrysotile can be safely used, and even INSERM concedes this fact.

Question 17: In footnote 42 of paragraph 4.27 of its written submission, Brazil seems to imply that the TBT Agreement would require a “rational link” between the measure and the risk assessment, in apparently the same sense as the case law cited therein from the SPS Agreement requires. Could Brazil identify the provision of the TBT-Agreement that justifies this allegation?

A rational link requirement is implicit in the text of the TBT Agreement. Article 2.2 states that a technical regulation “shall not be more trade restrictive than necessary to fulfil a legitimate objective.” The risk assessment is the basis of the level of safety chosen, which, in turn, is the basis of the measure taken. Absent any rational link between the measure taken and the risk assessment, this chain breaks. Thus, there must be a rational link to ensure that the measure taken is not more trade restrictive than necessary to fulfill the legitimate objective, the general statement of the level of safety, which is based on the risk assessment. Moreover, the very nature of the WTO agreements suggests that a rational link is necessary. One does not find in international agreements a “rationality clause.” Rather, the parties to the agreement understand that the provisions are to be interpreted rationally. Does the EC here suggest that a measure with no rational link to a risk assessment should be found WTO consistent? Brazil notes that it presented the panel reports addressing the SPS Agreement to ensure that the Panel was informed that a similar issue already had been addressed and resolved under the SPS Agreement.

Question 18: Has the study by Dr. David M. Bernstein, cited by Brazil, been published in a peer-reviewed journal?

The study by Dr. D. Bernstein was planned in two phases. The first phase (now complete) involved the evaluation of the biopersistence and morphological disposition of chrysotile fibres following inhalation exposure. Dr. Bernstein presented portions of the first phase 10 November 1998 at the Giornata Scientifica sulle Fibre di Vitro in Rome. The second phase is currently under way and involves a similar study of biopersistence and morphological disposition of both short and long tremolite fibres following inhalation exposure. Dr. Bernstein has confirmed that these studies will be published in a peer-reviewed scientific journal when the results from Phase Two are available. In the interim, Dr. Bernstein has published or presented the following reports: (i) a publication providing the scientific basis leading the European Communities to incorporate fibre biopersistence as a key parameter in assessing fibre toxicity194; a presentation of the chrysotile biopersistence and morphological disposition results at a scientific colloquium presented at the
Universidade Federal de São Paulo (Brazil) 19 March 1999; and an abstract by Dr. Bernstein entitled *The Inhalation Biopersistence and Morphologic Lung Disposition of Pure Chrysotile Asbestos in Rats* has been accepted for presentation at the “7th International Symposium on Particle Toxicology” which will be held in Maastricht 12-15 October 1999.

314. More fundamentally, however, Brazil disputes the relevance of the EC’ question. What matters is the soundness and relevance of the research, not whether it has been published yet. These facts INSERM concedes at page 135 of its Report. There, INSERM discusses a study by Camus et al., describing it as “an as yet unpublished study whose preliminary results have been communicated to us.” INSERM notes that, although unpublished, the study is particularly valuable because it was prepared, in part, by “particularly competent experts.” *Id.* Dr. David M. Bernstein is a particularly competent expert. As the EC well knows, the EC has hired him (as has the Government of Germany) to study and develop recommendations concerning the proper control of fibres. Indeed, he is still in the employ of the EC.

**Question 19:** In paragraph 4.28 of its written submission, Brazil refers to the “modern-day internationally recognised controlled-use level of 1f/ml,” based on a reference to a 1991 document. Is Brazil aware that in 1998 the WHO concluded that no threshold has been identified below which asbestos, including chrysotile, can be considered to be safe (see Annex II-1 of first written EC submission)?

315. Yes. Many different organizations have reached different conclusions about safe exposure levels. The United States and Canada, for example, have adopted exposure levels that are significantly higher than those of Brazil. A careful review of the WHO study indicates that the WHO conclusion was based on concern regarding chrysotile mixed with amphibole:

“The more rapid removal of chrysotile fibres from the human lung is further supported by findings from animal studies showing that chrysotile is more rapidly cleared from the lung than are amphiboles including crocidolite and amosite.”

316. These conclusions led the WHO to recommend research on the effects of exposure limited to chrysotile without any exposure to amphibole. This is precisely the type of research Dr. David M. Bernstein is conducting.

**Question 20:** Could Brazil elaborate on the relationship and possible limits of Article 12, in particular 12.3, of the TBT Agreement with the right of Members to take measures to protect human health in their territory?

317. No relationship exists, but this is not an issue in the present case. Brazil totally agrees that protecting public health is a legitimate objective. However, here France has taken a measure that is more trade restrictive than necessary to fulfil the purported objective of protecting public health. Moreover, France did not even consider the fact that Brazil mines and exports only chrysotile, not amphibole or a chrysotile/amphibole mixture. This is inconsistent with the TBT Agreement. *(Please refer also Brazil’s response to Panel Question 1.)*

**Question 21:** Could Brazil please provide data, including scientific evidence, of the number of cases of mesothelioma observed and their evolution during the last 20-30 years in its territory?

318. No case of lung cancer or mesothelioma from exposure only to chrysotile has been reported (this currently is being confirmed in the epidemiological study). Only three cases total of mesothelioma have been reported in Brazil. All of these were in individuals with substantial exposure to amphibole.

**Question 22:** In paragraph 4.14 of its written submission, Brazil states that “Recent research focusing on uncontaminated chrysotile demonstrates why it presents no health risk whatsoever”. Could Brazil provide copy of this scientific evidence?
Question 23: In its oral presentation, Brazil stated that INSERM has not taken account of studies which show that there is no risk associated with what it calls "modern" use of chrysotile. Could Brazil provide copies of the studies concerned, including references to the relevant paragraphs?

319. Brazil has addressed these questions with one answer because they raise similar issues.

320. Brazil has provided a copy of Dr. Bernstein’s study. Please also refer to three other studies presented by Brazil, which, in sum, "... support the hypothesis that adverse effects are associated rather with the fibres retained (amphiboles), than with the ones being cleared (largely chrysotile)." These studies show that, and explain why, modern uses of chrysotile alone present no health risk. In addition, please recall that in 25 years of operation, the Capivari Chrysotile Cement Plant in Brazil has not experienced one case of mesothelioma (please refer to Brazil’s presentation in Section IV of this Report). This health history is similar to that at other work sites using chrysotile (and even some in which chrysotile was predominant in a mix of chrysotile and amphibole). In the Executive Summary of The Workshop on Health Risks Associated with Chrysotile Asbestos, Dr. Graham W. Gibbs emphasized the importance of the "absence of lung cancer and mesothelioma risks in workers exposed to reportedly high concentrations of chrysotile in a UK asbestos cement plant in which silica was not used, as well as two similar plants in Zimbabwe ...". A clean bill of health also has been assigned to chrysotile as used by shipyard workers and insulators in the Pacific Northwest of the United States. According to Churg and Vedral (1994):

“Our results show clearly that, despite known historic exposure to amosite and chrysotile, amosite is by far the predominant residual fiber and there are correlations between amosite measures and disease. Chrysotile was present inconsistently and in relatively small amounts, and no correlations were found between chrysotile measures and disease.”

4. Questions from the European Communities to the United States

Question 1: In paragraph 4.47 of its submission, the US states that chrysotile asbestos is no less toxic than other forms of asbestos. Could the US please develop this statement further, including, if possible, references to and/or copies of the scientific grounds on which it is based?

321. The United States refers the Panel to its third party presentation in this case (see Section IV of this Report). In addition, the United States points out that a review of lung burden studies in human subjects and mechanistic studies does not provide convincing evidence for the “amphibole hypothesis” that chrysotile may be less potent than amphiboles in the induction of mesothelioma. Animal studies support the conclusion that all types of asbestos should be considered equally potent with respect to the production of either lung cancer or mesothelioma. For instance, in an inhalation study in which groups of rats were exposed to the five UICC asbestos fibre types, amosite, anthophyllite, crocidolite, Canadian chrysotile, and Rhodesian chrysotile, comparable incidences of lung tumours and mesotheliomas were induced by all types of asbestos fibres. While epidemiological studies clearly demonstrate a link between asbestos exposure and increased risk of lung cancer and mesothelioma, most of the studies involve workers that were exposed to mixed fibre types. Nevertheless, there are a number of epidemiological studies of workers that were exposed predominantly to chrysotile and one study in which the exposure was solely to amosite. There is no statistically significant difference in the risk of lung cancer seen in the group of workers exposed only to amosite compared to those exposed predominantly to chrysotile in textile production or to mixed fibres in manufacturing. In addition to evidence showing the causal relationship of exposure to chrysotile asbestos and cancer, ample evidence shows that exposure to chrysotile asbestos poses a significant risk of non-malignant respiratory disease. One study, which reported mortality and assessed dose-response relationship for asbestosis in a cohort of asbestos textile workers exposed...
only to chrysotile, found that 17 (5.5 per cent) of 308 deaths were due to asbestosis or pulmonary fibrosis. A second study reported data showing a linear relationship between cumulative fibre dose and morbidity. A third study reported a linear dose-response relationship between asbestosis and levels of asbestos dust. These data also support the hypothesis of no threshold, or low threshold for asbestos, since there is increased risk at cumulative exposures as low as 37 fiber-years/cc.209

322. Because of the carcinogenicity and asbestosis-producing effects of all asbestos types, the U.S. Environmental Protection Agency (EPA) and the Occupational Safety and Health Administration (OSHA) of the U.S. Department of Labor do not distinguish among asbestos fibre types in their regulations.210 In testimony in 1990, Richard Lemen, Assistant Director of the U.S. National Institute for Occupational Safety and Health, summed up the case against distinguishing among asbestos fibre types in public policy. We quote from his testimony:

"Recent reports have appeared in the scientific literature to suggest that different forms of asbestos are not equally pathogenic. … However, there is a great deal of uncertainty associated with these findings and equally important contradictory evidence. Results from research involving animal bioassays present a strong case that there is no safe form of asbestos. … Not only has chrysotile been found to be as potent as crocidolite and other amphiboles in inducing mesotheliomas when injected intrapleurally. … it has been found equally potent in inducing pulmonary neoplasms through inhalation exposures (Wagner et al., 1974). Chrysotile also appears to be more potently fibrogenic and carcinogenic than amphiboles, in relation to the quantity of dust deposited and retained in the lungs of rats (Wagner et al, 1974).

… In fact chrysotile fibers are much more chemically and biologically reactive than amphibole fibers. … These fibers are less readily detected in the tissue after the damage is done. …

At this time, there is no compelling evidence to justify different public health policy for different asbestos fiber types...".

323. Lemen also notes that the Royal Commission study submitted by Canada to the Panel recommended that textile manufacturing using chrysotile be banned, and concluded that “all fibre types can cause all asbestos-related diseases.”211

Question 2: In paragraph 4.47 of its submission the United States writes: “[s]pecification of work practices and other controls does not avoid all the risks associated with hazardous materials such as chrysotile asbestos.” Could the US please expand on this statement, and if possible provide (references to) the facts and data that support this viewpoint?

324. In response to the statements throughout the Canadian submission that “controlled use” will bring the risk associated with chrysotile asbestos to “undetectable” levels, in its third party’s submission to the Panel the United States has discussed the fact that “controlled use” will not completely eliminate the risk caused by asbestos. In 1994 OSHA reduced its permissible exposure limit for occupational exposure to asbestos to 0.1 fibres per cubic centimeter (f/cc) as a time-weighted average measured over 8 hours. Based on its risk assessment OSHA found that the excess cancer risk at that level would be a lifetime cancer risk of 3.4 cases per 1,000 workers and a 20 year exposure cancer risk would be 2.3 per 1000 workers. OSHA found that this risk is still significant and the United States Court of Appeals for the D.C. Circuit affirmed this finding. OSHA set this limit because it believed that it was the lowest exposure level using feasible engineering and work practice controls. It is not a risk-free limit. In other words, there is a still significant health risk at or below OSHA’s permissible limit. For this reason, and because some employers do not adequately implement the work practices, many asbestos workers rely on respirators.
Asbestos removal work often is done by poorly paid and poorly trained workers. OSHA has found that respirator effectiveness depends on how well the masks fit, how often they are replaced, cleaned and repaired, and how well employees are trained. In spite of rules requiring good respirator practices, studies show that many employers do not comply and that respirator-wearing employees still get high exposures. OSHA recently calculated, based on studies of respirator effectiveness, that up to 40 per cent of employees who wear respirators get essentially no protection when their employers do not adequately fit, clean and replace the respirators they wear. Also, where employees are protected solely by masks, the uncontrolled asbestos dust released during their work is airborne and contaminates the work-site. A major work practice in asbestos handling is wetting the tool used to cut or disturb the asbestos material or wetting the material itself. When the material dries, the dust becomes airborne.

Most asbestos-containing products are installed, disturbed or removed on construction work-sites. Construction jobs typically involve transient employees who may work for many different employers; short term work; and lack of training opportunities compared with factory employment. These factors make enforcing work practices and good respirator practices difficult. Another problem is that renovation work occurs in buildings that may or may not contain asbestos. This uncertainty leads to under-protecting the workers. Also, construction work sites have many immediate safety hazards. Employees and employees may concentrate on avoiding these immediate hazards and pay less attention to the risks of chronic hazards caused by asbestos. The difference between exposure levels when employers use available controls and when they do not is striking. During OSHA’s rulemaking, the U.S. Argonne National Laboratory (ANL) reported to OSHA concentrations during cutting operations of asbestos-cement sheet that ranged from 8 to 14 f/cc without local exhaust as compared to 0.04 to 0.15 f/cc when local exhaust was used. Similarly, sampling data collected by the U.S. National Institute of Occupational Safety and Health (NIOSH) during the installation of asbestos-cement sheet ranged up to 0.32 f/cc when local exhaust ventilation was not being used. Similarly, when cutting asbestos-cement pipe with an abrasive disc saw and cutting and machining pipe without a shrouded tool or without using wet methods, exposures exceeded OSHA’s exposure limit of 0.1 f/cc. Many asbestos product manufacturing and installation jobs expose workers to over the OSHA permissible exposure limit of 0.1 f/cc when dry operations take place. For example, OSHA determined in 1986 that "it is generally not feasible for the dry operations of carding and spinning (in textile operations) to comply with” the then permissible limit of 0.2 f/cc. OSHA also found that in the asbestos reinforced plastic manufacturing process, the “problem exposure areas” appear to be in dry finishing operations. These operations are similar to dry mechanical operations in other asbestos products manufacturing industries and include grinding and sanding, which OSHA has determined may not be feasible to achieve exposure levels below 0.2 f/cc without the use of respirators. Based on OSHA’s risk assessments, lifetime cancer risks at these levels are 6.7 per 1000 exposed workers.

Moreover, as discussed in both the U.S. and EC submissions to the Panel, the steady enlargement of the stock of asbestos and asbestos-containing materials in society, which results from failure to ban further use of asbestos, may engender substantial risks where demolition or maintenance takes place and the composition of materials is not obvious. As indicated in the U.S. third party’s submission, EPA has asbestos regulations in place which govern work practices and other controls to be used during the demolition of buildings, and maintenance activities in schools. These regulations require those engaging in such activities to identify asbestos-containing materials, or to assume that certain materials are asbestos-containing, prior to commencement of the work. If the composition of the material is unknown, in the absence of such regulatory requirements concerning work practices and controls, workers could be unprotected. The United States believes that unprotected workers who undertake building demolition or maintenance activities involving asbestos-containing materials would be subject to the exposure levels set forth by OSHA in Table 6 of the preamble to its final rule governing occupational exposure to asbestos. Table 6, entitled “Estimated Occupational Exposure to Asbestos and Reduction in Cancer Risk in General Industry and Shipyards as a Result of the Final Revision to
the Standard;

presents average baseline exposure levels in the absence of respiratory protection and other primary controls and work practices. These figures are given in the third column of the table, entitled “Potential mean fiber exposure with minimal controls (f/cc).” Such exposure levels would be expected to occur for unprotected workers. Based on this table, the exposure level for an unprotected worker involved in a demolition project would be 9.9 f/cc, or almost one hundred times the permissible exposure limit of 0.1 f/cc established by OSHA.

The 1991 Health Effects Institute-Asbestos Research report estimates that the lifetime risk to workers historically exposed to levels of that magnitude is approximately 200,000 per million, or 1 in 5.

EPA has issued public guidance concerning asbestos-containing products and their management. In addition, EPA has issued lists of suspect asbestos-containing materials. But it cannot be assumed that many “Sunday carpenters” know about such guidance, especially since they may not even be aware that they are working with asbestos-containing materials. The data in Table 6 of the preamble to the OSHA rule could be used to estimate exposures to these unprotected persons as they periodically engage in repair activities such as those involving ceiling tiles, plumbing, roofing, and drywall, based on such factors as the frequency and length of these intermittent exposures.

Question 3: In paragraph 4.51 of its submission the US states: “The use of a linear model is appropriate for a quantitative estimation of the risks associated with low levels of exposure to asbestos ….” Could the US further elaborate on this statement, explaining why it considers the use of a linear model – which implies that there is no threshold below which there is no carcinogenic risk – appropriate?

As indicated in the U.S. third party’s submission (footnote 4, paragraph 8, and Exhibit 15), studies of workers exposed to asbestos in the workplace (occupational exposures) link such exposures to high incidences of lung cancer and mesothelioma. However, since workplace exposures to asbestos are higher than general environmental exposures, these occupational exposures lead to an incidence of disease that is considerably higher than for the general population. The International Agency for Research on Cancer (IARC) and the World Health Organization’s International Programme on Chemical Safety (IPCS) recognize that there is no conclusive evidence to demonstrate that there is an exposure level for asbestos below which there is no risk, i.e., there is no “threshold” for asbestos. Even the Report of the Royal Commission on Matters of Health and Safety Arising from the Use of Asbestos in Ontario stated: “Most epidemiological studies of asbestos workers that have demonstrated an excess lung cancer risk associated with the inhalation of asbestos have produced results consistent not only with a linear relationship between cumulative dose and mortality, but also consistent with the absence of a threshold.” EPA uses a linear model to estimate the risks associated with low levels of exposure to asbestos because of the observed linearity of the response in occupational studies and because of the incomplete understanding of how asbestos causes diseases in humans. In order to estimate and project excess risks at the low level of exposures to which the general public is subject for asbestos, EPA utilizes a mathematical model that shows excess risks as simply proportionate to exposures at low levels (low-dose linearity). This procedure uses a curve to describe the excess incidence of disease observed at higher exposures (in the occupational setting) and takes a straight line to project from this excess to the lower-exposure environment of the general public. EPA believes that this approach is a reasonably protective way to address the issue of effects associated with low levels of exposure to asbestos.

Furthermore, as the United States points out in its submission, the limited data that exist for low levels of exposure to asbestos relating to the incidence of mesothelioma also indicate a linear relationship. In addition, the following should be noted, as indicated in the U.S. submission. Canada’s statement that “no epidemiological study to date has detected a higher health risk [than the linear model] resulting from low levels of exposure” does not take into account that, by their very nature, epidemiological studies are capable of detecting only relatively large incidences of cancer. Null results from epidemiological studies conducted at low exposure levels do not prove the absence of carcinogenic effects at such levels because such results can arise from such factors as inadequate study design or small population size. In 1986, OSHA also published a quantitative risk assess-
ment of asbestos, as a basis for amendments made in 1986 and 1994 to the OSHA asbestos standards. The risk assessment made the following key findings concerning the dose-response relationship in the case of asbestos. With regard to lung cancer, the data from several well-conducted epidemiologic studies show that a linear model best describes the dose-response relationship. With regard to mesothelioma, a linear model is reasonable, fits the data well, and is the best estimator of risk.227 As noted in the U.S. submission, IPCS found a linear dose-response relationship for mesothelioma in its 1998 risk assessment on chrysotile.

330. OSHA conducted a public hearing in 1984 with participation by major scientists and researchers in the field, including Dr. Kenny Crump, who testified that the linear model “has been widely used … for asbestos and lung cancer.” Dr. William Nicholson and Dr. Hans Weill also supported the use of the linear model to predict lung cancer risk. Dr. Weill testified: “ … as far as the shape of the curve for the important malignant consequences of asbestos exposure, I think we are all in agreement . . . that the evidence does not permit us, nor does concern for public health or prudence permit use . . . (of) risk analysis on any basis other than linearity of exposure and response and in a no threshold model.”

Question 4: Does the US consider that any of the currently used substitutes in the US for chrysotile asbestos are as dangerous or more dangerous than chrysotile?

331. The big picture of asbestos is that asbestos-exposed workers are right now being diagnosed with disabling asbestosis and mesothelioma that has clearly been caused by asbestos exposure. Asbestos-exposed workers are also being diagnosed with lung cancer. Because lung cancer is also caused by exposure to other toxic agents, it is sometimes difficult to prove that it was the asbestos exposure that caused the lung cancer in these individuals. However, looking at the epidemiological studies of workers, there is no doubt that the asbestos exposure causes lung cancer, and that the workers’ exposure either caused the lung cancer or played a large part in causing lung cancer (as discussed above, no significant difference exists between the toxicity of chrysotile asbestos and other asbestos). The question above asks whether there is an equivalent or stronger causal relationship to human disease or death in the case of substitute fibres or products containing them.

332. To begin with, as stated in the U.S. submission, in some instances asbestos use in a product (e.g., an asbestos-lined hot pad) is not essential, and the “substitute” consists of simply subtracting asbestos from the product, or using other materials such as iron or ceramic for the same purpose (e.g. an iron or ceramic trivet) or otherwise reformulating the product so that neither asbestos nor substitute fibres are used. In this case the health effects of substitutes are nil or trivial. As indicated in the U.S. submission, there is no evidence that any of the currently used substitutes in the U.S. for chrysotile are as dangerous or more dangerous than chrysotile. The only fibre that has been shown to be more hazardous than chrysotile is another naturally occurring fibre, erionite.228 However, erionite is not currently used in commerce. Based on the current understanding of how fibres can cause adverse health effects, available data indicate that the differences in physical structure and properties between substitute man-made mineral fibres (MMMFs) and chrysotile, such as the larger diameter of MMMFs and the fact that they break transversely rather than longitudinally to shorter fibres, may render some MMMF substitutes less dangerous than chrysotile. No MMMF has the same magnitude of carcinogenic potential as asbestos. A man-made mineral fibre, refractory ceramic fibre, has been shown to have considerable potency in some laboratory animal studies, but the legacy of suffering and death found with asbestos has not occurred. The high cost of refractory ceramic fibres has limited their use to high-temperature industrial operations. As a general matter, exposures to fibreglass have been lower than asbestos fibres. This may be due to the inherent property of fibrous glass to have fewer airborne fibres of a size that is deposited in the lungs.

5. Questions from the European Communities to Zimbabwe

Question 1: In its oral statement, Zimbabwe appears to argue that the so-called controlled use, including all the protective measures relating thereto, is capable of “minimiz-
ing” risk to human health. Could Zimbabwe expand on what it means by “minimize”, i.e. which level of risk – and thus chrysotile-caused diseases – would still remain and how much of that risk would be acceptable to Zimbabwe?

333. Zimbabwe first of all wishes to state that, being a natural product, chrysotile asbestos is present, for example, in the air we breathe and that some exposure is thus inevitable. There is therefore no question of eliminating exposure altogether. This said, Zimbabwe recalls that the 1998 Task Group has clearly stated that the risks to humans are conditional on exposure. The goal for responsible governments must therefore be to reduce exposure. It is the contention of Zimbabwe that as a result of the application of risk control measures the risk of exposure to asbestos dust can be minimized - so much so that the use of chrysotile asbestos can be considered safe. There is therefore no justification for a complete ban on the use of chrysotile asbestos. Zimbabwe submits that this conclusion is valid for both “primary” and “secondary” users of chrysotile asbestos. Regarding the former, Zimbabwe notes that the 1998 Task Group has confirmed that:

“Data from industries where control technologies have been applied have demonstrated the feasibility of controlling exposure to levels generally below 0.5 fibres/ml. Personal protective equipment can further reduce individual exposure where engineering controls and work practices prove insufficient.”

334. In Zimbabwe the fibre levels at workplaces where chrysotile asbestos is mined, milled or processed are indeed below 0.5 fibres/ml/eight-hour period. The aim is to bring those levels down even further to 0.3 fibres/ml. It is thus entirely possible for a developed country like France to achieve its target level of 0.1 fibres/ml. As concerns “secondary” users of chrysotile asbestos, Zimbabwe recalls that it has stated in paragraph 4.82 of its written submission that the:

“Combined use of high-density products made from asbestos-cement, which inherently are low-risk products, coupled with adequate risk control measures minimize the risk of exposure to asbestos dust”.

335. Zimbabwe thus believes that the combined use of inherently safe products and adequate risk control measures would enable France to achieve the maximum exposure level it deems acceptable. Zimbabwe finds itself in agreement on this point with Canada and the supporting evidence adduced by it.229

Question 2: In paragraph 4.98 of its written submission Zimbabwe suggests putting “asbestos warning messages” next to buildings. Can Zimbabwe explain why there would still be a need for this type of warning if, as it alleges, controlled use results in encapsulating indefinitely chrysotile in cement?

336. Zimbabwe wishes to remind the EC that the relevant part of paragraph 4.98 of Zimbabwe’s written submission states as follows:

“While Zimbabwe recognizes that it may not be readily apparent to an inexperienced person whether or not he/she is handling a product containing asbestos fibres, it is by no means justification for instituting a far-reaching ban on products which might contain asbestos fibres. […] Where the materials have already been installed or incorporated, say, in a building, Zimbabwe does not see why there could not be, for instance, an asbestos warning message next to the evacuation instructions on a notice board of that building.”

337. Zimbabwe has also stated in paragraph 4.81 of its written submission that:

“… products made from asbestos-cement are products of high density and thus chrysotile asbestos fibres are firmly blended into the final product [footnote omitted]. This reduces to a minimum the likelihood of fibres being released into the air and thereby posing a health hazard to human beings.”
338. Contrary to what the EC tries to suggest, there is no contradiction between the two above-mentioned statements made by Zimbabwe in its written submission. The EC has contended that no measure was available to France other than a complete ban of chrysotile asbestos and of products containing chrysotile asbestos, if France were to achieve its public health objective. The EC relies, inter alia, on the argument that it may not be readily apparent to an inexperienced person whether he/she is handling or dealing with a product containing asbestos fibres. In its submission, Zimbabwe addressed this argument put forward by the EC and Zimbabwe pointed out to the Panel that France could have very easily required asbestos warning messages to be posted in buildings, for example, so as to alert “secondary users” of the presence in buildings of chrysotile asbestos, if and where appropriate. The EC’s argument does therefore not provide a justification for a ban on chrysotile asbestos. It is true, as the EC rightly points out, that asbestos warnings in buildings are in principle unnecessary in view of the fact that chrysotile asbestos fibres today are encapsulated permanently in cement products. The EC nevertheless asserts that there could still be a risk of exposure to asbestos dust whenever the cement-products are installed, maintained or repaired. Zimbabwe has not contested this. Zimbabwe has, however, pointed out that there is a range of measures which would be available to France to effectively control such risks of exposure. All of these measures stop short of imposing outright bans. Thus, Zimbabwe’s submissions are by no means internally inconsistent.

Question 3: Could Zimbabwe explain whether training courses and the certification referred to in paragraph 4.98 of its written submission are the only elements of the so-called “controlled use” which can minimize risk, or does the postulated “controlled use” require also other types of measures?

339. In a sweeping and grossly exaggerated statement the EC has claimed that “Once on the market, there is no further reasonable means of controlling the use of asbestos and, in particular, controlling commonplace operations (cutting, sawing, …) that many persons may have to carry out.” Zimbabwe refutes this argument in its third party submission and cites specific and less trade-restrictive measures which could have been used by the French Government. Even assuming arguendo that the EC’s concern about control were justified with regard to do-it-yourself users of asbestos-containing products, Zimbabwe has submitted that, if indeed the French Government was so concerned about those users of asbestos-containing products, it could have easily banned the sale of such products in all do-it-yourself outlets. As an additional measure, the French Government could also have restricted the handling of asbestos-made products to certified experts, thus eliminating exposure of inexperienced private users to asbestos. Through certification the Government could ensure that training meets certain minimum standards and thus enables the individuals concerned to engage in safe work practices.

340. Zimbabwe also addressed the question of how the French Government could have sufficiently protected occupational “secondary users” of chrysotile asbestos other than through an outright ban on asbestos. As explained in Zimbabwe third party’s submission, the French Government “could have, for example, required certification, which would only be bestowed upon an individual once he/she had successfully followed information and training courses on the use and handling of asbestos-containing products. The French Government could also have laid down the precise work practices and technical appliances that must be used in all contacts with asbestos-containing products. To ensure compliance, the regulations could authorize the imposition of heavy fines or a custodial sentence in the event of a wilful disregard of the government’s regulations. Needless to say, it is also open to a Member to run information campaigns. Thus, it emerges clearly from this statement that certification, which would be conditional upon completion of, inter alia, training courses, is not the only element of “controlled use”. In any event, this statement must be read together with the paragraphs where Zimbabwe has set out the types of measures that together make up “controlled use”. It might be added here that in Zimbabwe’s view, periodical medical surveillance of workers can and should be an integral part of “controlled use”, notably in the case of “primary” and occupational “secondary” users of chrysotile asbestos.
Question 4: Could Zimbabwe comment on the findings of the study by Cullen et al., *Chrysotile Asbestos and Health in Zimbabwe, 1991 Am. J. Int. Med. 19, 171-182*, which establishes a net excess of mesothelioma cases of workers in mines and mills of chrysotile in Zimbabwe?

341. Zimbabwe disputes the assertion by the EC that the study by Cullen et al. establishes a net excess of mesothelioma cases of workers in chrysotile mines and mills in Zimbabwe. At page 178 of the study referred to it is stated that “[m]ajor x-ray abnormalities were rare in the population, with only four having evidence of TB and three having nodules or masses that potentially represented cancer.” This could hardly amount to conclusive evidence in support of the assertion by the EC. In fact, pleural disease was also surprisingly uncommon in the population which was examined. Eighty-five of the subjects had parenchymal changes of whom 65 were in grade 0/1. The fact that the study in question does show a relationship between parenchymal change and cumulative doses merely confirms the need for risk control measures. In no way does it establish a case for a complete ban on the use of chrysotile asbestos. It is worth noting here that there is another study by Cullen et al. entitled *Chrysotile Asbestos and Health in Zimbabwe - Analysis of Miners and Millers Compensated for Asbestos-Related Disease Since 1980* (1991 Am. J. Int. Med. 19, 161 – 169). This study was not, however, specifically on workers of chrysotile mines and mills in Zimbabwe. This was a case study on cases certified as having pneumoconiosis by the Pneumoconiosis Board in Zimbabwe.

342. The fact that this study showed cases of asbestos-related disease does not mean that the origin of those cases can be traced back to chrysotile mines and mills in Zimbabwe. The two cases referred to in the study (numbers 19 and 20), where the histological examination confirmed mesothelioma, were cases of individuals who had also worked in South African crocidolite asbestos mines. They were thus not exposed exclusively to chrysotile asbestos dust. Another case referred to in the study (number 12) had also worked in a crocidolite mine in South Africa for five years. This finding is confirmed by the records of the National Cancer Registry, which have been carefully maintained since 1940 in Bulawayo. The records show no more than 15 cases of mesothelioma. In a detailed study undertaken by Dr. Baloyi et al. of those 15 cases, 14 were found to involve individuals who have had mixed exposures, i.e. exposure to both crocidolite and chrysotile asbestos dust. In the remaining case, no exposures could be established. It is very important to note in this connection that in any event all the findings referred to were the result of past exposures. The control measures which have been put in place at the chrysotile mines, mills and manufacturing units in Zimbabwe since then mean that comparable exposure levels will never occur again.

6. Questions from Brazil to the European Communities

Question 1: In regard to the INSEMER Report: (a) does it contain any original research (research performed by INSEMER during the time the report was prepared)? (b) does it review any study regarding current controlled uses (post 1990 uses of chrysotile products)? (c) does it purport to review all available scientific studies? (d) does it review the then current research performed on the subject by and for the EC, in particular, that regarding bio-persistence and its relationship to disease? (e) if the answer to any of these questions is “no”, the report cannot support the ban, can it?

343. The INSEMER Report bases its conclusions on the analysis of 1200 international scientific studies, and reviews the status of international scientific knowledge concerning asbestos-related risks at the time of its adoption. It should be pointed out that all the studies on so-called controlled use and on bio-persistence available in 1996 were examined and taken into account in the INSEMER Report. The French ban is thus fully justified scientifically. In addition, we do not understand Brazil’s reasoning, which amounts to saying that one negative reply, for example to the question in subparagraph (a), would deprive the French Decree of a scientific basis in the light of the large volume of scientific data from the international sources existing on the subject.
Question 2: Why has France not banned all Class I carcinogens?

344. Brazil appears to be referring to an obligation of consistency which does not exist either in the GATT 1994 or in the TBT Agreement. It should be noted that, to our knowledge, no country has imposed a general ban on all Class I products (proven human carcinogens), but several have already banned asbestos. There is no international text imposing a general ban on all Class I products. France imposed a general prohibition on asbestos as a result of a risk assessment. Risk assessment must be specific to each product, and the ensuing national risk management decisions are different according to each product concerned. Of the Class I carcinogens, none has been used in such a widespread way, in products, spreading the carcinogenic risk when they are used, and sold to the general public. Most of these products are subject to very strict marketing restrictions and are usually used in an isolated environment as synthesis intermediates. Thus, they are no longer present in the finished products and hence do not spread the carcinogenic risk when the end-product is used. Besides, none of these products has caused as many deaths or occupational illnesses.

Question 3: How will the ban protect handymen from existing asbestos?

345. The French measure by prohibiting the use of any type of asbestos, including chrysotile asbestos, provides effective protection by stopping future human exposure to risk from this substance. The fact that quantities of asbestos, because of past use, remain and thus may continue to provide a source of risk, does not diminish at all the right of France to take the measure in question to stop further exposure to the risk from new applications. The implicit logic in Brazil’s question is that because of possible risks from existing asbestos France should continue adding further risks. This type of reasoning the EC contests. In any case, France has taken indeed very serious and strict measures to protect handymen from existing asbestos. They have been communicated to the Panel.

Question 4: Does not France regulate the use of pesticides through use restrictions, labelling and disposal requirements and the like? Are not many of these pesticides extremely toxic? If these types of dangerous products can be regulated to control risk and avoid improper use and disposal during the entire product life cycle, how can France maintain that chrysotile in contrast cannot be regulated?

346. The rationale of regulating the use of pesticides is completely different from that of the ban on the use of asbestos. Use of pesticides is allowed, after proper evaluation, in order to protect human, animal or plant life or health from pests, diseases, illnesses, etc. So although toxic, they serve a specific purpose and their use is authorized under well-specified quantities and conditions. The use of asbestos is not comparable to the controlled use of pesticides, as its use is proposed by Canada exclusively for commercial and economic reasons. To clarify even better the point, the parallel should be drawn with human medicinal products which are allowed to be used under strictly controlled conditions. Taking Brazil’s logic to its extreme, it would also suggest prohibiting human medicinal products because asbestos is also prohibited.

Question 5: Have substitute fibres been subject to the same close scientific scrutiny as asbestos? Do not the relevant studies conclude that many of the fibres cause cancer in rats and thus are suspected carcinogens? Does not IARC specify several of them as Class II carcinogens, including glass fibres, rock wool and slag wool?

347. The products used as substitutes for asbestos have been the subject of scientific studies like asbestos. Among all the products that may be used as asbestos substitutes, no fibre has been recognized as a proven human carcinogen (Class I - International Cancer Research Centre), unlike asbestos, which is a proven carcinogen. Only ceramic fibres are classed as Class II carcinogens, i.e. proven for animals, but they are used only in very limited and carefully controlled cases.
Question 6: If France were to remove from the first submission all reference studies that do not distinguish chrysotile and amphibole, is it true that no studies would remain? If some studies would remain, how many would remain?

Question 7: If France were to remove from the first submission all references to studies that do not directly address the risk of current controlled use of chrysotile, isn’t it true that no studies would remain? If some studies would remain, how many would remain?

348. Those questions of Brazil are purely rhetoric. Brazil appears to ignore the fact that the WHO Health Criteria 203 of 1998 (point 3 of Recommendations and Conclusions) have confirmed that:

“Exposure to chrysotile asbestos poses increased risks for asbestosis, lung cancer and mesothelioma in a dose dependent manner. No threshold has been identified for carcinogenic risks”…“Where safer substitute materials are available for chrysotile, they should be considered for use”…“Some asbestos containing products pose particular concern and chrysotile use in those circumstances is not recommended. These uses include friable products with high exposure potential. Construction materials are of particular concern for several reasons. The construction industry work force is large and measures to control asbestos are difficult to institute. In-place building materials may also pose risk control to those doing alterations, maintenance and demolition. Minerals in-place have the potential to deteriorate and create exposures”.

349. It follows that the WHO, like so many other scientific publications cited in the report of INSERM, consider that chrysotile asbestos poses similar risks and the same type of problems as regards controlled use to those posed by amphiboles asbestos. So, all the references in the scientific literature in the INSERM report remain valid and pertinent.

Question 8: Does the EC not recognize that amphiboles is more toxic than chrysotile? If so, shouldn’t the EC focus only on studies that at least attempt to distinguish the two? If not, how does the EC explain its position in light of the conclusions of relevant studies, including the INSERM report itself, concluding that amphiboles is more toxic?

350. Lung cancer is caused, with a comparable carcinogenic effect, by chrysotile asbestos and by amphiboles asbestos. Chrysotile asbestos, on the other hand, presents a lesser risk than amphiboles as regards mesothelioma. In both cases these diseases are currently untreatable and fatal, and it is no less serious to die of lung cancer than to die of mesothelioma.
ANNEX III

Publications and Documents Referred to by the Experts (Section V)

Dr. Henderson

Three key references on chrysotile asbestos – published in 1998 and 1999 respectively – are quoted or cited frequently throughout Dr. Henderson’s report in abbreviated form:


1. Documents referred to in Introductory Comments and Comments to the Questions by the Panel (Section V.C.1-2)


2. Documents referred to in the Endnote (Section V.C.3)


3. Documents referred to in Supplementary Comments (Section V.F)


Dr. Infante:


Stauder B. et al. X-ray results in roofers after exposure to dust from working with asbestos-cement for many years. trans. 1982.


ANNEX IV

Canada’s Comments on the Experts’ Responses to the Questions from the Panel

APPENDIX A

EXAMPLE OF THE APPLICATION OF A CONTROLLED USE POLICY
IN THE FRICTION INDUSTRY

The following example applies to friction product manufacture and use. In fact, for this sector, data indicate that even with past work practices, the risk, if any, for friction product manufacturing workers and mechanics has been extremely low. Canada presents it as a manner of achieving France’s desired level of safety, which is less trade restrictive than the ban. Company M wishes to manufacture brake linings, brake disc pads and dry clutches using chrysotile asbestos. The asbestos will be purchased from a producer of chrysotile P. Manufacturer will sell friction products through a distributor D to automobile manufacturers and to automobile service centres G.

The steps that would be followed under the proposed controlled use programme are as follows:

1. Company M requests a permit from the competent government authority to import chrysotile asbestos.

   The government authority grants permit only if:

   (a) Company M has in place the equipment, training programmes and work practices to protect workers from chrysotile exposure throughout manufacture and disposal of any waste materials.

   (b) Producer P would inspect the plant to ensure that all the regulated fibre-handling processes are in place to eliminate/minimize any potential for exposure.

   (c) Company M would provide the results of periodic measurements of the exposure of workers to the producer.

2. Once Company M has the import permit, the chrysotile producer will supply chrysotile to it, on the understanding that shipments will cease immediately if Company M fails to meet or exceed all applicable standards.

3. The chrysotile will be shipped in sealed containers to preclude exposure of workers, transportation personnel or the public during chrysotile shipment.

4. On arrival at the plant, the transfer of fibres to the process will be such as to eliminate workers opening bags of asbestos [e.g. automatic, sealed bag handling].

Risks of Lung Cancer and Mesothelioma for Friction Manufacturing Workers

A study of some 13,000 friction manufacturing workers in the United Kingdom in which lifetime estimated exposure to chrysotile ranged up to 356 f/ml-years [i.e. equivalent to 40 years exposure at just below 9f/ml] found no chrysotile-related increased risk of lung cancer or mesothelioma.

In this study the authors concluded: “with good environment control, chrysotile asbestos may be used in manufacture without causing excess mortality” [Berry & Neuhouse 1983, Neuhouse & Sullivan 1989, Berry 1994]. This study involved workers exposed up to 50-60 years ago, so controls were poor relative to present day standards.
In the USA, another study found no mesothelioma among 1630 deaths in persons manufacturing friction products [McDonald et al 1984].

Exposure Levels

The concentrations to which persons in the cohort studied by Berry and Newhouse were exposed were considerably higher than those reported in the Australian friction product plant by Dr. Henderson, even including peaks above the standard.

Control Feasibility

The technology and work practices to control exposures during manufacture exist and the experts appear to agree that exposures during product manufacture can be well controlled.

5. After manufacture under controlled conditions, the products will be shipped to the distributor in sealed packages. The manufacturer will ensure that all distributors have the knowledge and have a proper place to store the products without removing them from the original packaging.

Exposure Levels

As the product is composed of chrysotile in a high-density matrix and as it is sent in sealed containers [e.g. boxes], no persons at the distributor have potential for exposure.

6. On request, the distributor will deliver the products to the automobile manufacturers in the sealed containers. The manufactured products will be ready-mounted linings and pads and clutches which require no modification by the installer.

Exposure Levels

The brake linings and brake disc pads consists of chrysotile embedded in resin. Measurements have shown that exposure from the handling of these products is, at most, minuscule.

7. On request, the distributor will deliver the products in the sealed containers to automotive service centres. The range of sizes of brake linings (e.g. oversize) already exists so that when brake drums are turned (to eliminate scoring etc.) the lining of appropriate thickness is available for installation without modification. If it becomes necessary to have linings “ground to size” this would only be permitted at “authorized centres”, equipped with the appropriate exhaust ventilation systems. These centres would be identified to the manufacturer and competent authority. They would be at the same locations as those undertaking the turning of brake drums so that the linings could be fitted under controlled conditions and it would not be necessary to modify them when they are returned to the automotive repair centre.

Exposure Levels

The technology exists to do this work with virtually no exposure. [See NIOSH Report]

8. The distributor will provide the manufacturer and competent authority with a list of the names and addresses of purchasers. The purchaser will be informed that this list has been provided to the manufacturer and competent authority. In this way, the competent authority can readily target those locations where chrysotile-containing products are used. If the distributor, manufacturer or competent authority has reason to believe that safe work practices are not being followed, the supply of friction products would be discontinued.

9. The removal and installation of brake shoes, brake disc pads and clutches will be carried out according to precise codes of practice. This would include clean up and disposal requirements.
Risks of Lung Cancer and Mesothelioma for Brake/Clutch Repair Mechanics

The requirements for controlled use here go well beyond those necessary to protect worker's health.

The following studies demonstrate that workers carrying out brake repair work are not at an increased risk of lung cancer or mesothelioma.


The fact that there is no increased risk of lung cancer during manufacture of friction products shows that at exposure levels well above those of brake mechanics, there is no chrysotile related increased risk of lung cancer or mesothelioma.

Exposure Levels

The requirement and work practices exist and have been shown under field use conditions to reduce workers’ exposure during brake repair work to well below 0.01 f/ml. [See NIOSH reports].

1040
In the 1980s the average concentrations to which brake repair mechanics were reported to be exposed in Finland were less than 0.05 f/ml for automobile brake mechanics and less than 0.1 f/ml for truck and bus brake mechanics. [Kauppinnen & Korhonen]. Similar results were found in Germany where the lifetime exposure of brake mechanics after more than 20 years of full time brake work was less than 14 f/ml-years. These exposures took into account grinding, bevelling, sanding and otherwise modifying the brake linings as well as using compressed air to remove brake wear debris from brake drums.

The exposure of workers from work on clutches in the past was even lower than that associated with brakes [Lynch (1968), Kauppinnen & Korhonen (1987), Jacko & Ducharme 1973].

10. On removal of brake shoes, brake discs and clutches from the vehicles, these will be placed in containers provided by the distributor and returned through the distributor to the manufacturer.

11. As the worn brake shoes are returned to the manufacturer, any re-lining by unauthorised companies/persons is precluded. Any re-lining of brakes will be done as subcontracts by the brake lining manufacturer and with equipment and work practices that are no less stringent than those required of the manufacturer. There will be no brake lining material sold to other "re-lining companies”.

12. Disposal of any used brake lining, clutch facing or brake disc pad will be done according to jurisdictional requirements.

Environmental Releases and Public Health Risks

Data show that during braking or use as a friction product, chrysotile is altered to non-asbestos mineral or amorphous silicates. Thus the bulk of the material to which workers are exposed from used brakes is not asbestos as mentioned by one of the experts. Also, almost all residual fibres are very short [e.g. > 80% of fibres are less than 0.4um in length].

Because of the mineralogical and particle size alterations, the environmental release of chrysotile fibres greater than 5 µm from the use of chrysotile containing friction products is extremely low in the case of brakes and essentially nil in the case of clutches. [Lynch, JR (1968) Brake Lining Decomposition Products. J. Air Pollution Control Assoc. 18: 824-826]. Concentrations of chrysotile fibres measured at street level have also been very low. The data obtained in the United Kingdom indicate that the use of asbestos in brake linings does not measurably contribute to atmospheric asbestos concentrations in the urban environment. Even at two heavily used intersections in the London metropolitan area, concentrations vary from 0.0002 to 0.0004 f/ml. Jaffrey, S (1990) Environmental Asbestos Fibre Release from Brake and Clutch Linings in Vehicular Traffic. Ann. Occup. Hg. 34:529-534.

As there is no indication of an increased risk of lung cancer or mesothelioma in friction product workers or brake mechanics exposed at many orders of magnitude above the general public, the actual risk for the public at their levels of exposure will be epidemiologically undetectable.
APPENDIX B

EXAMPLE OF THE APPLICATION OF A CONTROLLED USE POLICY
IN THE ASBESTOS-CMENT INDUSTRY

The major portion of current chrysotile cement products is for outdoor applications, such as roofing, exterior wall cladding, rain gutters, pipes, etc. Chrysotile fibres are transported from fibre suppliers to asbestos cement plants, packed in sealed 50 kg plastic bags piled and “stretched-wrapped” on pallets, and are delivered to the plant premises in closed containers. Thus the possibility of dust emissions during transport is practically nil. Fibres are delivered to asbestos cement plants that comply with a “Controlled use” code of practices. This includes prohibition of reselling of unused fibre inventories to third parties by the asbestos cement plant manufacturers. Suppliers of chrysotile asbestos from Canada, Brazil, Zimbabwe and Swaziland have signed and endorsed a Memorandum of Understanding on Responsible-Use of Chrysotile Asbestos, whereby the signatories agree in particular that they will “provide a written commitment to appropriate national authorities indicating that chrysotile asbestos will be supplied directly to chrysotile asbestos-product manufacturing facilities on condition that chrysotile asbestos not be resold upon delivery ...”

Typical flow of the main steps from manufacture to disposal

1. In-plant handling of chrysotile fibres through the different steps (wet process) leading to the finished product:

   (a) Bag opening inside hoods under negative pressure. Operators must wear protective equipment;

   (b) wet-processing of the fibre cement mix, shaping of the product, wet-curing as the case may be, and wet practices for final shaping and cutting of the various products.

Comment

All work must be carried out according to safe work practices such as those described in the ILO Code of Practice “Safety in the Use of Asbestos”, chapter 13, and under engineering controls that have been shown to reduce occupational air concentrations to levels presenting a negligible, undetectably low health risk, as shown by the following published data:


In an asbestos cement factory using chrysotile only, 1,970 workers were traced, and their mortality experience was examined. There was no appreciably raised standardised mortality ratio (SMR) for the causes of death investigated, including all causes, all neoplasms, cancer of the lung and pleura, and cancers of the gastrointestinal tract. The authors indicate: “Thus the general results of this mortality survey suggest that the population of the chrysotile asbestos cement factory studied are not at any excess risk in terms of total mortality, all cancer mortality, cancers of the lung and bronchus, or gastrointestinal cancers.”


An investigation of 5,645 asbestos cement manufacturing workers, showing no raised mortality resulting from exposure for 20 years to chrysotile asbestos at exposure levels equal to or less than 100 MPPC.years (corresponding to approximately 15 fibres/ml.years). The authors state:
“… However, the demonstration that low cumulative and short-term exposures did not produce a detectable excess risk for respiratory malignancy may be of assistance in the development of regulatory policy, because a scientifically defensible position based on these data is that there are low degrees of exposure not associated with a demonstrable excess risk”.


A cohort study of 1,176 asbestos cement workers in a Swedish plant using chrysotile asbestos showing no excess related mortality at exposures of about 10-20 fibres/ml.years.


A cohort study carried out on 2,167 subjects employed between 1941 and 1983. No excess of lung cancers or other asbestos-related excess death is reported, at mean fibre concentrations below 1 f/ml, although higher levels had probably occurred in certain areas of the asbestos cement factory.

2. Delivery of pre-cut, pre-drilled chrysotile cement products (according to client’s specifications) to licensed contractors, with notification to government authorities.

3. Installation on the work site of the pre-sized, pre-drilled chrysotile cement product must be performed by workers who have received an approved training programme, which includes mandated work practices and working tools, such as those described in the above-mentioned ILO Code of Practice, chapter 13.4, and also in “Catalogue of Tools for Working with Asbestos Cement Products on Site” (AIA Recommended Control Procedure No. 2A). This will ensure that fibre emissions are kept at levels where measurable health risk is unlikely, and undetectably low.

Comment

Evidence published in 1980 by Rödelsperger et al (IARC Sci. Pub. No., 30, pp. 845-853) shows high levels of exposure of up to 100 f/ml peak exposure for workers installing roofing shingles when using high-speed grinding power tools. This is clearly not a situation where “controlled use” was observed. However, more recently available data show that installation of asbestos cement roofing shingles will not result in workers’ personal exposures to levels associated with detectable risk. All sample results of measurements are below 0.1 f/ml.


The same observations (below 0.1 f/ml) were made during removal of old roofing asbestos cement shingles. Complete data is found in:

Bonacci et al (1998) “Report of Industrial Hygiene Survey at 10233 Norton Road, Potomac, MD”. SSM Analytical Laboratory, Reading, PA, USA.

Similar low exposure levels have also been measured during various operations on old weathered asbestos cement sheets (water jet cleaning or painting, demolition by removal of whole sheets from roofs and walls). This data can be found in:


In case of asbestos cement pipes, when improper tools and work practices, such as using high speed, abrasive disks for cutting pipes for instance, exposure as high as 35 f/ml may result. However, operations such as using a manual or power lathe for cutting sewer pipes, and using the
adequate power hole cutter (about 15-20 min duration) will result in exposures in the 0.1-0.2 f/ml range. Such data can be found in:


As mentioned earlier, chrysotile cement products are essentially found in outdoor applications or in underground pipes, thereby not very likely to be subject to interventions by tradesmen such as plumbers or electricians, after their installation. If interventions are required, use of appropriate tools and work practices as described in many international standards (ref. previous comments on question 5(a)) will prove sufficient to manage any potential risk, if any.

4. During the normal service life of the asbestos cement product, emissions from in-place asbestos cement do not result in measurable increases above the average, naturally occurring environmental air concentrations.

Comment

Evidence supporting this can be found in the following published data:


…“A comparison of the asbestos fibre concentrations in those areas with and without A/C roofing … lead to the conclusion that there is no statistically significant connection between the use of asbestos cement materials and the asbestos fibre concentrations found in the various measurement areas”.


…“The study of emission conducted on coated and uncoated roofing materials revealed low asbestos fibre concentrations, even though severe corrosion was observed on uncoated asbestos cement roofs and a considerable quantity of material containing asbestos could be removed by blowing or suction. The asbestos fibre concentrations that were measured in populated areas are well below the level considered acceptable by the Health Authorities of the Federal Republic of Germany, i.e. clearly below 1000 f/m³ (length = 5 µm)”. (translation)

5. Disposal of asbestos cement plant and demolition waste must be carried out according to well-known waste management practices approved by national authorities.

Comment

Proper disposal site management practices have shown that there is no measurable additional burden to the naturally occurring environmental fibre concentrations, as is illustrated in the following example:


This report is about a survey of air concentrations at disposal sites in Germany, showing the following data:

- directly over disposal sites: 0.0005 to 0.003 f/ml
- vicinity of disposal sites: 0.0001 to 0.0009 f/ml
## ANNEX V

Comments of the European Communities on the Replies by the Scientific Experts to the Questions from the Panel

### SUMMARY OF REPLIES BY THE EXPERTS

<table>
<thead>
<tr>
<th>QUESTION NO.</th>
<th>P. INFANTE</th>
<th>N. H. DE KLERK</th>
<th>D. W. HENDERSON</th>
<th>A. W. MUSK</th>
</tr>
</thead>
<tbody>
<tr>
<td>1(a) Main categories of workers at risk</td>
<td>Principally secondary users (building industry, intervention, maintenance, …)</td>
<td>Principally secondary users (building industry, intervention, maintenance, …)</td>
<td>Principally secondary users (building industry, intervention, maintenance, …)</td>
<td>Principally secondary users (building industry, intervention, maintenance, …)</td>
</tr>
<tr>
<td>1(b) Mainly an occupational or an environmental risk</td>
<td>Mainly an occupational or occupation-related risk</td>
<td>Mainly an occupational or occupation-related risk, but also environmental</td>
<td>Mainly an occupational or occupation-related risk</td>
<td>Mainly an occupational or occupation-related risk</td>
</tr>
<tr>
<td>1(c) Release of fibres through degradation of asbestos cement</td>
<td>Fibre release; small, non-quantifiable risk</td>
<td>Fibre release; small, non-quantifiable risk</td>
<td>Fibre release, small, non-quantifiable risk</td>
<td>Fibre release, small, non-quantifiable risk</td>
</tr>
<tr>
<td>1(d) Fibre release during intervention on asbestos cement</td>
<td>Release of large quantities of fibre, established risk</td>
<td>Release of large quantities of fibre, established risk</td>
<td>Release of large quantities of fibre, established risk</td>
<td>Release of large quantities of fibre, established risk</td>
</tr>
<tr>
<td>1(e) Release of fibres during intervention on non-friable products containing chrysotile</td>
<td>Release of fibres; established risk for workers and handymen</td>
<td>Release of fibres, established risk for workers and handymen</td>
<td>Release of fibres; established (non-quantifiable) risk for workers and handymen</td>
<td>Release of fibres; established risk for workers and handymen</td>
</tr>
<tr>
<td>1(f) Danger of the fibres released by asbestos cement</td>
<td>Fibre release, small, non-quantifiable risk</td>
<td>Fibre release, small, non-quantifiable risk</td>
<td>Fibre release, small, non-quantifiable risk</td>
<td>Fibre release, small, non-quantifiable risk</td>
</tr>
<tr>
<td>1(g) Risk during demolition and removal of asbestos</td>
<td>Established risk</td>
<td>Established risk</td>
<td>Established risk</td>
<td>Established risk</td>
</tr>
<tr>
<td>1(h) Risk of wastes</td>
<td>Theoretical risk, probably low</td>
<td>Theoretical risk (if not handled properly)</td>
<td>Theoretical risk (if not handled properly)</td>
<td>Theoretical risk (if not handled properly)</td>
</tr>
<tr>
<td>2. Risk associated with other applications of chrysotile</td>
<td>Established risk</td>
<td>Established risk</td>
<td>Established risk</td>
<td>Established risk</td>
</tr>
<tr>
<td>3(a)(b)(c): Relative pathogenicity of chrysotile/amphibole</td>
<td>Chrysotile and amphiboles are carcinogens for lung cancer and mesothelioma</td>
<td>Chrysotile and amphiboles are carcinogens for lung cancer and mesothelioma</td>
<td>Chrysotile and amphiboles are carcinogens for lung cancer and mesothelioma</td>
<td>Chrysotile and amphiboles are carcinogens for lung cancer and mesothelioma</td>
</tr>
<tr>
<td>4(a): Epidemiological data for low levels of exposure to chrysotile</td>
<td>Established risk for many professions</td>
<td>Some studies do not show a high risk</td>
<td>No data quantifying the exposure-effect relationship</td>
<td>Some studies do not show a high risk</td>
</tr>
<tr>
<td>4(b): Safety threshold</td>
<td>No threshold for any disease</td>
<td>Impossible to demonstrate the existence of a threshold</td>
<td>No threshold for any disease (except for asbestosis)</td>
<td>No threshold for any disease</td>
</tr>
<tr>
<td>QUESTION NO.</td>
<td>P. INFANTE</td>
<td>N. H. DE KLERK</td>
<td>D. W. HENDERSON</td>
<td>A. W. MUSK</td>
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<td>4(c)(d): Linear model</td>
<td>The linear model is the most appropriate; no credible alternative model</td>
<td>The linear model is the most appropriate; no credible alternative model</td>
<td>The linear model is the most appropriate; no credible alternative model</td>
<td></td>
</tr>
<tr>
<td>4(e): Concentration/duration of exposure to chrysotile</td>
<td>No lower level of exposure without risk</td>
<td>No lower level of exposure without risk</td>
<td>No lower level of exposure without risk</td>
<td>No lower level of exposure without risk</td>
</tr>
<tr>
<td>5(a)(b)(c)(d)(e): &quot;Controlled&quot; use</td>
<td>Impossible in practice in the vast majority of situations</td>
<td>Impossible in practice in the vast majority of situations</td>
<td>Impossible in practice in the vast majority of situations</td>
<td>Impossible in practice in the vast majority of situations</td>
</tr>
<tr>
<td>6(a): Risks of non-fibrous substitutes</td>
<td>Not carcinogenic</td>
<td>No direct reply: only fibrous substitutes should be taken into account for cancer risk</td>
<td>No direct reply: only fibrous substitutes should be taken into account for cancer risk</td>
<td>No-direct reply: only fibrous substitutes should be taken into account for cancer risk</td>
</tr>
<tr>
<td>6(b)(c): Physical and chemical characteristics of substitute fibres; relative risk in comparison with chrysotile</td>
<td>Dimensions and form (whether or not they can be inhaled) and durability are associated with toxicity</td>
<td>Dimensions and form (whether or not they can be inhaled) and durability are associated with toxicity</td>
<td>Dimensions and form (whether or not they can be inhaled) and durability are associated with toxicity</td>
<td>Dimensions and form (whether or not they can be inhaled) and durability are associated with toxicity</td>
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<tr>
<td></td>
<td>All substitute fibres are less dangerous than chrysotile</td>
<td>All substitute fibres are less dangerous than chrysotile</td>
<td>All substitute fibres are less dangerous than chrysotile (doubts concerning refractory ceramic fibres)</td>
<td>All substitute fibres are less dangerous than chrysotile</td>
</tr>
</tbody>
</table>
ANNEX VI

Meeting with Experts – 17 January 2000

Transcript

Chairman

1. I would like to welcome the four scientific experts and the delegations of Canada and the European Communities. I should like to introduce the Panel members and the Secretariat staff especially for the benefit of anyone who wasn’t at previous meetings. My name is Adrian Macey, on my right is Mr. Lindén and his right Mr. William Ehlers. In the Secretariat staff, the Secretary to the Panel is Ms. Mireille Cosy and Assistant Secretary Ms. Doaa Abdel-Motaal. The Legal Officers are Mr. Yves Renouf and Ms. Kerry Allbeury. I would like to remind everybody that we have simultaneous interpretation in French and English. Secondly, the proceedings will be recorded and subsequently transcribed. The verbatim transcript will become an integral part of the final report. I would like now to invite the experts to introduce themselves, going in alphabetical order.

Dr. de Klerk

2. My name is Nick de Klerk, I work as an epidemiologist in asbestos-related diseases in Western Australia.

Dr. Henderson

3. My name is Douglas Henderson. I am the Professor of Pathology at the Flinders University of South Australia and the Flinders Medical Centre. I have been pursuing an interest in asbestos-related diseases for some 32 years.

Dr. Infante

4. Peter Infante. I am an epidemiologist and I am with the United States Occupational Safety and Health Administration.

Dr. Musk

5. I am Bill Musk, a clinical professor of medicine and public health of the University of Western Australia.

Chairman

6. We have received lists of the two delegations, Canada and the European Communities. Could we ask the delegations’ leaders to indicate who is who amongst your delegations? Can I ask firstly Canada to introduce themselves?

Mr. Hankey (Canada)

7. Thank you, Chairman. I am Blair Hankey, Associate General Counsel at the Department of Foreign Affairs and International Trade. I have on my right Maitre Thomas-Louis Fortin who is Legal Counsel at the Ministry, Eric Wildhaber, who is also Legal Counsel and Sebastien Beaulieu, also Legal Counsel at the Department of Foreign Affairs and International Trade. Also opposite me is André Dulude, who is Director of the Regulation and Technical Barriers Division at the Department, and Pierre Desmarais from the same Division. Behind me, I have Louis Perron from the Canadian Ministry of Natural Resources, and on his left, Gilles Mahoney, who is Director of the mineral industry for the “Ministère des ressources naturelles” of the Government du Québec. Then, on my left, I have Professor Corbett McDonald, as scientific adviser to our delegation, and Professor Alison McDonald. On my right, I have Dr. Graham Gibbs, who is also an expert, and behind me,
Dr. Jacques Dunnigan and Dr. Michel Camus, also experts. I would also like to add that the Professors McDonald are serving as honorary members of the delegation and have declined to accept any compensation from Her Majesty in order that both their independence and the appearance thereof may be guarded.

Chairman

8. Can I now ask the representative of the European Communities to briefly introduce their delegation?

Mr. Christoforou (European Communities)

9. Thank you, Mr. Chairman. My name is Theofanis Christoforou, and I am a Legal Adviser of the European Commission in Brussels. We have a big delegation, composed partly of Commission officials and French representatives. We have scientific experts and members of the Member states' delegations which are based here in Geneva. As the presence is quite long, I would rather leave it to each member to present himself or herself shortly.

10. Jean-Jacques Bouflet, Legal Adviser of the Delegation in Geneva of the Commission; Hubert van Vliet, member of the Legal Service of the Commission in Brussels; Dr. A. Tossavinen, Scientific Adviser; Marcel Goldberg, Scientific Adviser; Maud Valat-Taddei, responsable de la réglementation concernant l'amiante, Ministry of Employment and Solidarity, France; Sophie Chaillat, Ministry of Health, France; Marie-Christine Poncin, Ministry of Economy, Finance and Industry, France; Pierre Monnier, Legal Adviser, Permanent Delegation of France to the WTO, Geneva; Christian Forwick, Permanent Mission of Germany in Geneva; Mr. H. Rieck, Permanent Mission of Germany in Geneva; Mr. M. Nielsen, Permanent Mission of Denmark in Geneva; Sergio da Gama, Legal Adviser, Portuguese Mission in Geneva; Jacques Bourrinet, Professor à l’Université d’Aix-Marseille; Mrs. A. Bensch, DG Trade, EC Commission, Brussels; Dr. B. Terracini, Professor, Scientific Adviser; Dr. P. Huré, Scientific Expert; Mr. B. Castleman, Scientific Adviser; Mrs. Mchanetzki, Ministry of Economy, Finance and Industry, France.

Chairman

11. Thank you very much. I would like to explain how the Panel intends to organize its work for today. I would like to thank the four experts for having agreed to serve as advisers on the Panel and for the very hard work that they have managed to perform over such a short period of time. We do operate under significant time constraints in the WTO dispute settlement system and we have to produce reports within certain deadlines. This does put pressure on everybody involved. The purpose of this meeting is essentially to allow the experts to expand on the written responses that they have already given us to the Panel's questions. Obviously, these documents are substantial; we have all received them and it is not a matter of repeating what is already before us. The experts will initially be given an opportunity to make any general comments they may have, and reactions to their own colleagues' reports, as well as on the written comments received from the parties. The parties will be given the opportunity during the meeting to seek clarification on the experts' reports, to express their views on them. The focus, I would stress, is on the experts and on questions to them. So we would ask the parties to limit their interventions to questions and comments directly related to the issues that the experts have raised, either in the existing written comments or orally during the meeting. The experts will, of course, have the invitation to react as they wish to what is being said by the parties. Let us come back again to the point that the priority of the meeting which is to hear the experts. I hope that the meeting will give us a full opportunity for an exchange of views between the experts, the parties and the Panel, so that, at the end of the day, the Panel can be as fully informed as possible about the scientific and technical issues involved in the case. It is not the purpose of the meeting to hear new evidence which the parties have not previously submitted and we will have to reserve our right to disregard any argument or evidence which is not directly related to statements made by the experts or the other party.
12. Firstly, I need to remind everyone that the proceedings, according to the rules of dispute settlement, are confidential. We will handle the meeting as follows. Firstly, we would like to invite opening statements or comments by each of the experts, which we will do in alphabetical order. It may be that following their brief introductory comments, one or other or more of the experts might like to take up additional points, which they are welcome to do, but when this initial introduction is concluded, it is over to the parties to put their questions and comments. We would like to focus this main part of the meeting question by question, i.e. the questions that we sent to the experts fall under six main headings – there are essentially six main questions with a number of sub-questions under each one. For the clarity and the good order of the meeting, we will try to focus our discussion question by question. So that means that after any opening comments by the parties, we will, in the case of each question, invite first Canada and then the European Communities to raise comments and questions that they have under the first question, and then we will go on to the second question until we get to question six. I would like to invite the parties to be selective in the items that they comment or ask questions on, so that they can be sure that the key points they see as relevant to the dispute can be covered in the time available. We will of course be flexible as the meeting develops, and the Panel will do its best to make sure that we assist you in a smooth progression of the discussion and ensure that we don’t leave any important issues aside.

13. There is limited time available, so we can now proceed to the first stage of our meeting, which is to invite each of the experts to make any opening comments that they might wish to do, beginning with Dr. de Klerk. When we have had each of your opening comments, I might return to the experts again and ask if there are any points which you may want to follow up.

Dr. de Klerk

14. I think the six questions fall into three main ones. The first one is whether chrysotile is dangerous at all, and I think the generally accepted view is that it is. A subset of that question is which dose-response relationships are appropriate and whether you should use linear extrapolation or not. A corollary of that is which equation one actually uses, which dose-response, which group of studies one uses to do that, and that seems to be open to a bit of argument. The second question, and perhaps the more easy to answer, is the question about controlled use. It is clear, just by observation in Australia, that you can’t control the use of a dangerous product like chrysotile all the way down the chain of use. It is possible that you can control in the factories that produce it, but certainly down towards end-users it seems almost impossible, and in fact most cases of asbestos-related disease have arisen through such downstream use, probably mainly of other forms of asbestos, but certainly that sort of use. The third question is whether substitutes for asbestos are known to be safer – I would say that the evidence at present is that they are. I think that is a summary of how I see the issues.

Dr. Henderson

15. At the outset, I would express my great appreciation to the WTO and to the Panel and the Chairman, Dr. Macey, for asking me to act in my capacity as an adviser to the Panel. I would also like to thank the Secretariat and in particular Ms. Mireille Cossy for her consistent helpfulness at all times in responding to my questions and requests. My introductory remarks can be given quite briefly. My full opinion is set out in quite extensive reports already submitted to the WTO, both my original report of November last year with an attached Endnote and, following submission of new information, I did prepare a 26-page summary of additional Remarks which clarify and amplify some of my opinions and conclusions. Broadly speaking, I see the issues in a way which is very similar to that of my colleague, Dr. de Klerk. The three key issues as I see them are: firstly, is chrysotile carcinogenic for the lung and for the mesothelium? My answer to this is that the evidence is strongly in favour of the fact that it is, and that it is capable of inducing both lung cancer and mesothelioma at reasonably low levels of exposure; for example, low levels of exposure, such as occurred in the South Carolina asbestos textile workers in the studies carried out.
by Dr. Dement and his colleagues, led to a greater than two-fold increase in the standardized mortality ratio at quite low levels of exposure for white males, in the order of 2.7 to 6.8 fibre-year. And the additional information submitted in this case concerning the possible significance of amphiboles in the lung tissue of those workers does not, in my opinion, detract from the significance of that observation, and the reasons why are set out in my Supplementary Remarks to the Panel.

16. The second point, which I think is a crucial point, is whether or not the use of chrysotile can be controlled at all points of use. Again, I would be in close agreement with my colleague, Dr. de Klerk, that it cannot. My own series of mesotheliomas, amounting to in excess of 2,000 cases, indicates that by far the greatest number of mesotheliomas that I see occur – not in miners and millers nor in products manufacture – but they occur in those exposed to asbestos at the multiple points of end-use. In the Australian Mesothelioma Register, which represents a systematic compilation of all mesotheliomas found in Australia, there is good evidence that the greatest number of mesotheliomas that we see occurs among carpenters, builders’ labourers, plumbers, plasterers, painters and all others involved in building construction in particular. In Australia, this group represents a very large workforce; it is usually employed by small business, or the workers are self-employed. In the past, it has not been possible to extend controlled use of asbestos to this group of workers, and to the best of my knowledge, this situation continues today, although the use of chrysotile in building construction materials in Australia was phased out in 1987 or 1989.

17. The third issue that I believe is of significance here, is whether alternative substitute materials for chrysotile are safer than chrysotile. Again from my survey of the literature, I would be in close agreement with my colleague Dr. de Klerk that the evidence available to me indicates that substitute fibres are – according to national and international health authorities – safer for end-use than the use of chrysotile. And these are, I believe, the three key issues for resolution by this Panel.

Chairman

18. Thank you. Dr. Infante, please.

Dr. Infante

19. Thank you for asking me to participate. First, I would like to state what it is that I feel that all of us experts agree upon. That is that chrysotile presents a high risk of cancer to society, to exposed individuals. It is unlikely to ever be controlled enough to use safely. Substitutes appear available, and there is no evidence that they are as harmful as chrysotile asbestos. Regarding some particular studies - I did express this in writing, I want to reiterate it - the Dement study which has been reviewed, analysed and critiqued, the study of chrysotile textile workers, shows one of the highest risks of lung cancers ever observed among any asbestos-exposed population on a fibre-per-fibre basis. The increase in the relative risk from this study is 2 to 3 per cent per fibre per c.c. year. There are two additional studies of chrysotile textile workers, the Rochdale chrysotile workers which shows a risk of 0.5 to 1.5 per cent. There is a risk assessment based on a study by McDonald et al. which shows a relative risk of about 1.25 per cent increase per fibre per c.c. year. There has been a lot of discussion about the McDonald study of miners and millers; this study shows a significant excess of lung cancer but the dose response is about 30 times lower than the 2 per cent relative risk from the Dement study and about 16 times lower, if one assumes that the relative risk is 1 per cent per fibre per c.c. year of exposure. I suspect, in this study, that there is a fair amount of misclassification of exposure. Because, when you have misclassification of exposure, you are going to dampen the dose response and be biased towards a flat dose response curve. For mesothelioma, I think the recent analysis by Landrigan and Nicholson et al., which concludes that chrysotile is only one half to one quarter as potent for causing mesothelioma as crocidolite asbestos, I think that is a reasonable analysis. There may be certainly other reasonable analysis as well. But, even if chrysotile asbestos did not cause mesothelioma, which in my opinion it does, there is still enough risk from lung cancer alone, that there should be intervention to substitute for chrysotile
asbestos. There is a recent paper that was sent out after we completed our initial reports from Case and Dufresne. It was stated that I might change my opinion regarding the Dement study after I had reviewed this study, so I want to comment on that. I would add that this is an unpublished study and that the authors are much more restrained in their interpretation of the study than is Canada’s submission about this study. The authors state that they can’t determine to what degree the findings of fibre content in the lungs examined are representative of the entire cohort; their lung tissue fibre analysis only represents what is retained in the lungs by the time of death and there is a tremendous difference here between the miners and the millers and the textile workers between the time from cessation of exposure to death. Therefore you would expect a lot more chrysotile clearance from the textile workers’ lungs. Dr. Henderson has done an analysis based on assuming various half-lives of chrysotile fibres in the lungs which I think is a reasonable analysis, which indicates that there would be much more chrysotile in the lungs of the textile workers. But also on the basis of this new report, or furthermore, if the lung cancer in the Dement cohort study of textile workers was related to amphibole exposure, one would expect more than two mesotheliomas in this cohort. So I think that is striking also. Furthermore, Dr. Dement has done an analysis in response to this paper, which he has provided to me and which I would be happy to share with the Panel. What this study shows is that, regardless of when you analyse your data, because Green had found that only one of 39 workers hired in the 1940s or later had significant amphiboles in their lungs – these were the chrysotile textile workers – only one of 39 that were hired after 1940 had significant amphiboles in their lungs, Dement did a new analysis where he looked at the entire group of 126 lung cancers in his study and he gets the same dose response whether he looks at total employment or employees who were first employed after 1940, or who were employed before 1940 or 1950. So what it shows is the same dose response accounting for different periods of employment. So I thought that was impressive.

20. Regarding controlled use, it is my opinion that it may be theoretically possible, but is highly unlikely that chrysotile can be controlled in commerce. My point from my written submission was, that while it may be possible, in the United States alone, we have had over 4,000 violations of our asbestos standard in the last three years. In the United States there are monetary penalties that go along with these violations and yet, if we have this large amount of non-compliance in the United States in the presence of monetary penalties, and also in some cases there can be criminal penalties, then what does this bode for other countries that might not have this stringent requirement or penalties. Canada’s document was criticising, I believe, that I didn’t understand their controlled use programme. It seems to me that from recent articles that I have seen in countries where Canada appears to, or is importing, its chrysotile asbestos, in Morocco, Brazil and India, recent reports just came out indicating that asbestos is not controlled according to its controlled use programme. Therefore, in my opinion, the programme has little credibility to me. My point is that if it can’t be handled in the United States, I suspect that it is going to be even more difficult to control its use in other countries. Regarding substitutes, I feel that the substitutes do not present the cancer risk that chrysotile asbestos does. Three have been studied experimentally, two of the substitute fibres have been negative in animal cancer studies; fibreglass has been positive. I did not mention refractory ceramic fibres because the question was not specifically asked about refractory ceramic fibres. Refractory ceramic fibres are carcinogenic in experimental animals. I definitely think that there should be various serious concerns to humans exposed, but these fibres are limited to special high heat applications and I don’t believe that these fibres would be substitutes for chrysotile in most current applications of chrysotile asbestos.

21. It was commented in the last submission by Canada that I had a different opinion about the carcinogenicity of fibreglass compared to asbestos than what I had published in 1994. Looking further at data, I feel that there is not sufficient evidence in humans that fibreglass is carcinogenic, but I think that one should presume that these glass fibres are carcinogenic to humans; that doesn’t mean that it is proven, but I think that there is enough evidence that we should be concerned about that. But I don’t feel that they are as potent as chrysotile asbestos. As I indicated, I recently spoke with several workers who are employees of the fibreglass manufacturing facility that showed a two-fold risk of lung cancer. Those
workers explained to me that there were other known human carcinogens to which they were exposed at that facility which had not been mentioned in the report, namely, they were exposed to asbestos and to crystalline silica, along with several others which I mentioned in my report. Because of that, I feel that one cannot look at that study in terms of the fibreglass/fibre count in relation to the elevated risk of lung cancer, there is confounding from other known carcinogens or highly suspected human carcinogens in that population that are not accounted for.

22. On page 49 of the Canadian response, it states that there are three studies in which cellulose exposures have been investigated but that I did not identify them. Cellulose has not been studied for carcinogenicity in experimental animals. What I had indicated was that there are three industries where there is cellulose exposure, namely the paper industry and this study in this industry does not indicate any elevated risk of lung cancer or mesothelioma. I didn’t identify the literature, there is an entire IARC monograph on the paper industry. The same with wood dust. I didn’t cite any particular specific studies of workers exposed to wood dust which contains cellulose. IARC has an entire monograph on the furniture manufacturing industry and a more recent monograph on wood dust. There is no indication of any excessive risk of lung cancer or mesothelioma. Cotton dust, I didn’t cite any particular study but there is a tremendous literature on workers exposed to cotton dust. The Occupational Safety and Health Administration in the United States issued a new regulation for cotton dust a number of years ago, and cancer was never an issue that was raised as a health concern; it was byssinosis from workers exposed to cotton dust. Regarding the cotton dust exposure and the byssinosis, it was never proved whether it was the cotton fibres per se or the contaminants that were related to the byssinosis. In any event there is no indication of lung cancer or mesothelioma from cotton dust exposure. So while I didn’t cite those, there is a tremendous literature on those.

23. Finally, I would conclude by saying that, once it is known that these fibres are carcinogenic, one should not need to demonstrate their carcinogenicity in ever sector where blue-collar workers come in contact. Once you have identified the hazard, it is not convincing to say a particular study does not show an excess, it is the exposure that we are concerned about. We already know that exposure to these fibres are dangerous. This is an industrial health problem of abating the hazard, not continuing to identify the hazard in new populations that have not heretofore been studied. There have been epidemiological studies, that is, not of controlled environments like the laboratory setting, there are always errors and misdiagnosis of disease, there is incorrect recording of disease on death certificates, there is misclassification of exposure. All of these factors, particularly the misclassification of exposure, lead to flattening the dose response, so that you don’t find any dose response. So I think that we have identified the hazard, and it is my opinion that since there are substitutes available for it, I would recommend a substitute for asbestos. Thank you.

Chairman

24. Thank you very much. Dr. Musk, would you like to make some introductory comments?

Dr. Musk

25. Thank you. I would like to echo my fellow panelists’ gratitude for inviting me. My analysis and distillation of evidence from my own work and from the literature is that all forms of asbestos may cause disease. The main diseases being well known are: asbestosis, lung cancer, malignant mesothelioma and pleural plaques. The issues of main concern are the malignant diseases that may arise from chrysotile. The approximate relative potency of the different forms of asbestos to produce the different outcomes is summarized and tabulated in my submission. I want to stress that it is obvious that these are very ballpark estimates. I hope that we don’t argue too much about the numbers themselves. The outcomes from exposure to asbestos appear to be determined by the dose of exposure, the dimensions of the particles, their durability and chemical properties – least well understood.
Not all the data is consistent, particularly regarding the effects of chrysotile and mesothelioma, thresholds of exposure may exist, especially for asbestosis, but there is no direct evidence for them that I am aware of and I believe that it is unlikely that there are thresholds for carcinogens. Control of asbestos-related diseases is dependent upon control of exposure in the people who handle it from mining onwards or are otherwise exposed to it in the chain of events to ultimate or penultimate disposal. Controlled use of asbestos in production and manufacturing may be feasible. But controlled use does not seem feasible when extended to subsequent use or handling or incidental exposures. Finally, substitutes are probably safer, in my assessment, than chrysotile.

Chairman

26. Do any of the experts, having heard what their colleagues have just said, want to comment further? No. Well, we will move into the part of the meeting that gives parties and the Panel, of course, the opportunity to ask any additional questions. Before I give the floor to Canada, I think it would be helpful if delegations could limit general statements they may wish to make, if they could be quite concise. I would like to give each delegation the opportunity, should they wish, to begin with any general comment they might have. Could I ask Canada first if you would like to begin with the general comments, or whether you would like to go straight into addressing question to the experts?

Mr. Hankey (Canada)

27. I would like to thank the experts for the hard work they have done. We look forward to discussing their views with them today with the object of shedding more light on this complicated scientific question, insofar as it is relevant to the issues before the tribunal. In its response to comments by the experts, Canada referred without citation to an unpublished study by Case, Dufresne, Sebastien, and two distinguished members of our delegation, Professors A. D. and J. C. McDonald. In the response to the comments by the experts we indicated that this study had been the object of a presentation at a conference held in Maastricht this last fall. When asked by the Panel to provide a text of this study, we submitted it to the Panel’s Secretary in its draft form which was found on the Internet. This Internet version was marked “Not for citation.” It should be pointed out that this version of the study had yet to be seen or approved by either Dr. A. D. or Dr. J. C. McDonald. They acknowledge that there were statistical errors in this text which have been pointed out to Dr. Case both by the McDonalds and by Dr. Henderson in his Supplementary Remarks. These errors have been corrected in the version which is to be submitted for publication. Finally, I note that the statistical errors noted by Dr. Henderson in his Supplementary Comments do not detract from the essential findings of this study including that amphiboles are present in textile workers’ lungs. I have one other question. Mr. Infante has just made reference to a text by Mr. Dement which I believe has not been filed before the tribunal. I wonder, you had said that it was not your intention to allow the admission of new evidence at this time. Does that apply only to the parties and not to the experts? Because we have not seen that study.

Chairman

28. Could I come back to that point later, and ask the European Communities whether they would wish to make any introductory comment.

Mr. Christoforou (European Communities)

29. Can I join you and my Canadian colleagues in thanking the four experts for the hard work they have done and the time they have taken to provide so detailed and pertinent replies to the questions we are facing here. I would request the four experts, when they reply to the questions by Canada – this is not a polemic comment – to try to identify each time what their views are. We say this because we have the feeling that in Canada’s comments, sent to the Panel on 13 December, there is an underlying attempt by Canada to somehow confuse by grouping all the scientists together – saying “the four scientists” or...
the scientists - whereas in some of the cases probably only one of the scientists had said something. So I would appreciate it if the experts tried each time to identify what their personal views are on the specific questions. It is very important for the record to show what the views of each of the experts are and not leave with generalizations. One procedural issue, Mr. Chairman: I did not hear anything about the time that is allowed to the parties, especially I did not hear the word “equal time” to the parties. So I would assume since we are second under … [END OF TAPE] … otherwise we may end up here today with us having no time to ask questions to the experts.

Chairman

30. Thank you. To take the latter point first. I said that the Panel would make sure that we are able to do our work as efficiently as we can and I think that we will have to see how the meeting goes, but we are not yet at the point of having to allocate time to each party as we are not yet at a point when we are under any pressure for time. But we will need to see how the discussion goes but obviously, yes, we will make sure that both parties have a fair opportunity to have their points heard.

31. On the other point of evidence, we did not obviously limit the evidence that the experts might supply for the amplification of their views. And the parties today have the chance to comment on that evidence. I think that the point was in terms of the due process of the Panel’s case itself that we were not, on the part of the parties, we were not in a position to be able to look at completely new issues that had not been raised before. We think we are at the point where we can begin with the questions to the experts. I would invite you as far as possible to follow the path we suggested to take the issues question by question. Obviously there may be some overlap in the questions, especially in questions 1 to 4 which are all concerning chrysotile, question 5 was concerning controlled use and question 6 concerning substitute fibres. For the good order of the meeting and to be able to monitor our progress through the issues as we go, it would be helpful if the parties can as much as possible try to address questions to the experts under each of the headings of our questions 1-6. I give the floor to Canada on issues concerning question 1. Are you ready to begin or do you want a few minutes to consult among yourselves, in the light of what the experts said in their introductory comments?

Mr. Hankey (Canada)

32. No, Sir. I am ready to begin. I just want to come back to that procedural issue I raised at the beginning, that is to say the admission of evidence. I take it that inasmuch as the experts have admitted or led new evidence either in their submissions today or for example in the extra procedural filing by Mr. Henderson a week ago, that we would have the opportunity to lead evidence that we may need to rebut or to respond to the new evidence brought forward by the experts.

Chairman

33. I think that it was made clear that we have at our disposal a limited amount of time for this expert phase of the Panel process. That phase essentially concludes at the end
of today. I think that rather than delay our proceedings in any further discussion of a procedural nature, it would be very helpful if we could begin straightaway on the questions themselves. So I would invite Canada to begin. What I would suggest is that we can alternate questions between Canada and the EU. Canada, would you present your question on question 1 first, and then we can follow with the EU.

Mr. Hankey (Canada)

36. I do just want to signal to you at this time that we agree to proceed with this part of the process as you suggest. But on Thursday I will be raising what we consider to be serious procedural problems with the way the expert consultations have taken place. But let’s not bother with that now.

37. This question is directed to all of the experts. A majority of you have identified construction workers as being the population at greatest risk. Who do you include in the definition of construction workers? Do you include, for example, skilled workers such as electricians and plumbers?

Chairman

38. The parties are free to ask their questions either to an individual expert or to the experts as a group and in cases such as this one, where questions are being asked to the experts as a group, we will leave it to the experts themselves as to which question they wish to respond. I would just like to give the floor briefly to Mr. Christoforou.

Mr. Christoforou (European Communities)

39. I really regret having to intervene, but I would suggest – Canada is free of course to ask and to term the question the way it wishes – but I would make a second plea to avoid words like the “majority” without knowing who of the four scientists had said what. I would request Canada to identify which of the scientists had said what, words like the “majority” or “most of you”, it is our suggestion that they should be avoided. We need to know who said what instead of referring to the majority of the scientists. Thank you.

Chairman

40. Thank you. Take note of that, please.

Mr. Hankey (Canada)

41. Thank you, Mr. Christoforou. I could rephrase the question if it is helpful, either to say “some of you have identified construction workers etc. etc.” And those of you who wish to respond may do so. I don’t insist, I am not in a position to insist, that anyone responds who doesn’t think the question pertinent.

Chairman

42. I pass the floor to whoever wants to respond to that question. Mr. Hankey, would you mind repeating the question?

Mr. Hankey (Canada)

43. Some of you have identified construction workers as being the population at greatest risk. I suppose I can address the question to those of you who have done so. Perhaps you haven’t all done so and perhaps the majority of you haven’t done so, and perhaps we don’t count so well. Who do you include in the definition of “construction workers”? Do you, for example, include skilled workers, such as electricians and plumbers?
Dr. de Klerk

44. Speaking for myself, I was talking about people in the construction industry, so that would include electricians, plumbers, carpenters, laggers, boiler makers, anyone in any form of construction. It’s basically the group of workers who form the largest part of people who come down with mesothelioma. And where regulations are going to be hardest to police.

Chairman

45. Dr. Henderson was going to make a point.

Dr. Henderson

46. My inclusion amongst construction workers would include a large and disparate workforce which includes both skilled and unskilled workers involved largely in building construction and building maintenance and so forth. If one looks at mesothelioma as an index tumour for asbestos exposure and you go to the attachment I gave to my first report of the professions or workers included in the Australian Mesothelioma Register, they do include, going down them alphabetically: people who carry out maintenance on asbestos dwellings, fences, they include builders, brickworkers, builders’ labourers, carpenters, joiners, construction workers, civil engineer, demolition worker, electrical engineer, electrical fitter, electrical mechanic, electrician. Going further down the list, labourer, locksmiths, machine fitters, maintenance carpenters, maintenance electricians, maintenance fitters, mechanics (they’re not involved in building construction, of course, they are a different group). They do include painters, plasterers, plumbers. Together I think it adds up to a fairly large and disparate workforce which is very poorly regulated in Australia.

Chairman

47. Thank you. Any expert wishes to add anything?

Dr. Infante

48. I would agree with that. It is both skilled and unskilled in the rubric of construction workers.

Dr. Musk

49. That would fit in with my ideas. We might argue whether construction and demolition aren’t opposite processes, but there is so much overlap in the sort of tasks that people in the construction industry undertake, that we could probably include demolition with construction.

Chairman

50. May I invite any further comments or issues that parties might like to raise in connection with this question? No, in that case can we turn to the European Communities for their first question or comment.

Mr. Christoforou (European Communities)

51. This question is addressed to all the scientists, in particular, to Dr. Infante and Dr. Henderson. In your reply to question 1(e) of the Panel, where you are discussing occasional interventions on asbestos, (for example Dr. Infante states “mesothelioma has been identified from these exposure situations because it is a marker cancer related to asbestos exposure”). We would appreciate it if you could expand on this, and whether you think there are data from the mesothelioma registers which support this, and what is the part of the population which is at most risk. And therefore the question of public health concern.
Dealing with this group of workers, and in particular, the occasional workers, I think that it is fair to say that the risk of mesothelioma and of lung cancer will be related to the frequency and to the cumulative exposures that these individuals sustain, because professional workers, for example professional carpenters, will be working most consistently and regularly with asbestos-containing building materials. It is they who will sustain the highest cumulative exposures, and therefore suffer the greatest risk of both mesothelioma and lung cancer. For the occasional worker, the risks will be substantially less because the cumulative exposure will be less. But in my own series of mesotheliomas in Australia, I have a number of cases of individuals who simply dwelt in asbestos cement houses and who carried out maintenance and renovation on the houses. It so happens that most of those individuals would also have sustained exposure to the amphiboles. Given the relative potency differential between the amphiboles and chrysotile, I would expect the risks of the occasional worker with pure chrysotile cement materials to be substantially less than those exposed to mixed asbestos cement materials. However, I would also point out that in Australia there are individuals who style themselves as “home handymen” and they make a career of buying dilapidated houses, often asbestos cement houses, and they live in them for a year while carrying out extensive renovations and maintenance work. They then sell these houses a year later and because they have dwelt in the house for a year, the profit that they make is not subject to taxation. These individuals call themselves “home handymen”. The houses they buy are often called “handyman specials”, because they require maintenance and renovation. These individuals will move through a succession of houses at yearly intervals. Now, it so happens that if you look at their cumulative exposure, they may approach the types of cumulative exposure one would expect for a professional carpenter. So I would have to say that the risks would be related to the frequency and the duration of the exposure, and its intensity, and therefore to the total cumulative exposure.

In these situations there is not good information about exposure but rather as the scenario type of exposures. It’s this intermittent exposure that you really don’t know how much fibre these individuals are exposed to. But the fact that you see some mesotheliomas, it indicates that without being able to add up the cumulative dose of their fibre exposure, it indicates that it was enough in some situations to induce mesothelioma. We don’t have specific information on dose-response from these operations because we don’t have information directly on dose or fibre counts over time. The point I was trying to make was that if you identify mesothelioma from these types of exposure, there will also be the unidentified risk from lung cancer from those types of exposures. Lung cancer is more difficult to identify because it has got a high background level and there are other factors that relate to lung cancer in addition to asbestos. So it is difficult to identify the lung cancer cases.
as a co-factor, usually with tobacco smoke, but the estimates range from less than 3 per cent to 20 per cent in different countries. In Australia, when we look at the Mesothelioma Register and the Dust Diseases Register for New South Wales, the data for mesothelioma are very good. But the data for lung cancer are very poor. It has been suggested that normally one sees about somewhere between one lung cancer for every mesothelioma to up to 10 lung cancers for every mesothelioma. In New South Wales, despite the adequacy of mesothelioma data, the data we have for lung cancer are very poor, so that if you look at compensated cases for lung cancer in New South Wales, we see a reversal of the lung cancer to mesothelioma ratio. So, we see ten mesotheliomas compensated for every lung cancer. The situation probably is that most of these asbestos-related lung cancers are passing unrecognized by the medical attendants, because the patient is a cigarette smoker – that is explanation enough and no further explanation is sought. And even the cases that come before the Register, a large number of them are rejected on the basis that the data on exposure don’t suggest that there is sufficient exposure for the individual case to be compensated. But if you approach this on a population basis it seems that a large number of our lung cancers have a contribution from asbestos exposure, passing unnoticed by the national health authorities and regulatory authorities.

Chairman

57. Any additional comments on this point? Canada.

Mr. Hankey (Canada)

58. Dr. Henderson, I wonder if we may get back to the question that was actually asked, which was I believe about your Register study of mesotheliomas. How many of the mesotheliomas deaths do you consider to be attributable to chrysotile only?

Dr. Henderson

59. That is a very difficult question to answer. I am afraid I cannot give a precise answer because many of the individuals will have sustained mixed exposures or they have sustained exposures for which we have no precise data as to fibre type. However, if you look amongst the Australian Mesothelioma Register data, there is a figure of 58 mesotheliomas among automobile and brake mechanics whose only exposure was to brake linings, and brake blocks themselves. For decades in Australia, brake blocks and brake linings have only contained Canadian chrysotile in a bonding matrix, so there have been no amphiboles in that material for some decades.

Mr. Hankey (Canada)

60. What sort of controls did you have in place for this study?

Dr. Henderson

61. This is not a study. These are figures from the National Mesothelioma Register where the occupational histories are quite good. Again, it is one of those things that one is reliant upon the data supplied to the Register. But the increase in incidence has been worked out by comparison with the Australian census figures for the total number of automobile mechanics, including all types of mechanics, and the number of mesotheliomas occurring among them over a particular period of time. The same type of figure is also given in the NICNAS document which I submitted to the Panel as an annexure to my original report.

Mr. Hankey (Canada)

62. Thank you. Dr. de Klerk, what value would you attribute to a register study that is conducted without controls? What probative value would you consider that it has?
Dr. de Klerk

63. Mesothelioma register studies are widely used. Basically, the problem being that you cannot compare, generally, the rates with other groups if you haven’t got any population basis. But I think Professor Henderson said that somebody actually looked at this in relation to the population in that occupational group. So obviously if the occupational group, you know you have the total population, you can ascribe a rate of disease in that group and you can compare that with the overall rate in the population. In terms of the proof scenario, people always put case series down at the bottom of the list but it is often in medical history that the case series come up with the sort of proof, well not the proof, but come up with the first idea as to something being a risk factor. You only have to look at all the nickel workers in Wales, the chimney sweeps and all those kind of things. They were all first observed purely through case series. So I think the case series is a very valuable epidemiological tool.

Mr. Hankey (Canada)

64. So, do I understand you to say that in terms of probative value in scientific rigour, you would place it near the bottom or at the bottom of methodologies that are used to determine relative rates of disease in one occupational group as opposed to another?

Dr. de Klerk

65. In standard epidemiology texts, they always start off by saying that the best thing for showing an effect is the randomized control trial. You can’t really do a randomized control trial in this situation. So the next thing you do is a cohort study and because these are all people in disparate industries, you can’t do a cohort study. Then you could do a case control study, but the exposure is fairly rare, so it is not very good to do a case-control study. So you end up with the case series. I can see your point, but at the same time, if you have got this number of cases with only this exposure, it has got to carry a fair amount of weight in terms of if you choose to stand next to somebody blowing out dust from their brake drums, if you see what I mean?

Mr. Hankey (Canada)

66. You say you attribute a fair bit of weight, to what? What conclusions would you draw from the study?

Dr. de Klerk

67. Well there are a lot more brake mechanics getting mesothelioma, I mean the rate in brake mechanics is a lot higher than the rate in other groups of the population. Therefore one would attribute a fair amount of weight to that study.

Mr. Hankey (Canada)

68. Are you aware that there have been four case controlled studies of garage mechanics, two in the United States. (McDonald and McDonald, Teta et al.), one in Canada (Teschke) and another in Germany (Woitowitz and Rödelsperger). They have all shown no increased risk of mesothelioma for garage and brake mechanics. Do you accept these data?

Dr. de Klerk

69. If those studies are there and that is what they show.

Mr. Hankey (Canada)

70. And what would you consider to be the more scientifically rigorous methodology and which would have in a court of law the greatest probative value: the register analysis
that has been done by Dr. Henderson or these kinds of case control studies?

Dr. de Klerk

71. I think that you would have to look at them on an individual basis. The problem with case control studies is that it is very easy to do a bad case control study, where you have a sort of register in place that is sort of collecting data as fully as it possibly can, one might make the point that the register might be better. At the same time, in the case control study, there is a problem with sample size: I mean to show no increase in risk is not the same as showing that there is no risk. It is just showing that the study doesn’t have sufficient power to detect an increase if it is there, and I make that point somewhere else in my document, the standard case control case is bedevilled by small sample size problems. I wouldn’t like to generalize too far, there may be heterogeneity in the cases in the study, there may be different work practices in the different countries. It is just that certainly in Australia, there seems to be good evidence that the brake mechanics do have an increased risk of mesothelioma.

Mr. Hankey (Canada)

72. I don’t think that you got the point of my question. I am not asking you to attack the method by which Dr. Henderson conducted his study, as related to the rigour of the four case control studies on garage mechanics to which I have referred. But rather I am asking you: *grosso modo*, as a form of analysis, as a form of enquiry, which is generally considered to be the more reliable in terms of producing hard results?

Dr. de Klerk

73. Well, the case control study.

Mr. Hankey (Canada)

74. Yes, all right, good. Now, Dr. Henderson, are you aware that a proportional mortality study of mesotheliomas in England and Wales covering the period 1979-1980 and 1982 to 1990 showed no evidence, and I repeat no evidence, of an increased risk of a mesothelioma in motor vehicles? That is the study by Hodgson et al.

Dr. Henderson

75. Yes, I am aware of the studies that have shown negative findings with no detectable increase in risk. I would simply amplify the comments that Dr. de Klerk has already made. (And I don’t regard myself as an expert epidemiologist). I would simply say that if you are looking at a small effect in a small population, you may not detect an effect. When you deal with national populations, yes, the quality of the information and the controls may diminish, but you are not looking at the same issue in many respects, and we are not looking here – when we look at the Australian incidence of mesothelioma among automobile mechanics – to provide proof in a court of law. We are looking for an indication of an effect that might be used for the formulation of national occupational health and safety policy, which I think is an entirely different exercise. But yes, I am aware of those negative studies and I have to counterbalance those with the indications that we have – not only from my own looking at the Register – but from the National Occupational Health and Safety Commission looking at the Register, to say that there is an indication of an increased frequency of mesothelioma amongst brake mechanics in Australia and that the increase is in the order of 1 to 2 per cent increase per year, which is roughly comparable to the overall growth of mesotheliomas in the Australian population. But when we look at that type of effect, we need to ask what group are we going to compare them against. And if you look at the background rate of so-called spontaneous mesothelioma of 1 to 2 cases per million of the population per year, we do have an indication of an increased effect in making that comparison.
Mr. Hankey (Canada)

76. I just want to point out that, although you pointed out a difference in the kind of evidence that you seem to think is appropriate for setting policy and the kind that is appropriate in the court of law, but I think that, in both circumstances you would admit that what is important is the probative value of the evidence studied. In both cases, people are trying to draw conclusions to complex and difficult questions. But I just want to conclude by saying, of the four studies that I cited earlier in my question to Dr. de Klerk, these studies of garage mechanics by McDonald and McDonald, Teta et al., and Teschke, Woitowitz and Rödelsperger, of these four studies, one of them was a strictly controlled series of mesothelioma study by McDonald in 1980. The mesothelioma study considered all 344 cases of mesothelioma reported by pathologists throughout North America during the reference period. These were compared with 344 strictly matched controls. Of these 344 cases, 12 cases had been garage workers. This perfectly matched the 12 controls who were garage workers, indicating that the rate of mesothelioma among garage workers is the same as in the general population. Do you accept these data?

Dr. Henderson

77. Well, you are going into some highly specific details, amongst thousands and thousands of pages of information that I have tried to digest in preparation for this meeting. But yes, I agree with the general conclusions, and the simple fact that I would draw attention to is that with so many studies on asbestos-related diseases, one is dealing with contradictory sets of data. The question arises as to what weightings one places upon one set of data as opposed to another and what significance one gives to a particular set of data when trying to set national occupational health and safety policy.

[Coffee break]

Chairman

78. … [Not recorded] Dr. Infante said that he wished to intervene on the previous question we were discussing.

Dr. Infante

79. My comment relates to which is a better study, a case control study or using the mesothelioma registry in Australia to estimate the risk of mesothelioma. In a case control study you are sampling your controls, hoping that they represent the universe. The extent to which they do or not, you don’t know, but you use certain matching criteria and hope that they do. The extent to which they do, may affect your findings. On the other hand, looking at the mesothelioma registry for the entire country of Australia, you don’t need to sample the universe, because the denominator is already the universe. So you don’t have any sampling error that you have to be concerned about. Then Dr. Henderson estimated then what the incidence of mesothelioma would be in the general population of Australia, based on the cases that were reported to the registry. In my opinion, he overestimated the denominator, by making certain assumptions. But nevertheless, he had quite a high incidence of mesothelioma per million population from his analysis. So, in my opinion, in this particular case, I feel that the registry is a very good source and in fact may be superior to using a case control study where you are trying to estimate what the incidence is and the relative risk compared to the universe which you are presuming from your controls. And also it is like we are talking about asbestos exposure in mesothelioma here, it’s not that we are looking for some new disease related to asbestos. It is a disease that has already been indicated as being associated with asbestos. So, I feel that using the registry, where we have the entire data based on the entire country, may in fact be preferable to a case control study where you are trying to sample or estimate what the frequency is and the comparison population.
Chairman

80. Thank you. Canada wishes to further comment on this question?

Mr. Hankey (Canada)

81. I would like to ask Professor Corbett McDonald, who conducted the largest cohort of asbestos workers for the longest period of time of any study that has ever been conducted. I would like him to comment on the relative merits of the various forms of studies that are being discussed here.

Dr. C. McDonald (Canada)

82. I suppose my question would be to Dr. Infante, as an epidemiologist, to ask whether you really are in a controlled study trying to sample the universe. Are you not trying to sample the part of the universe that is comparable to your cases? That is people from the same place, of the same age and sex, with questions about their occupation which are comparable and which are analysed blind, without any preconceived idea of an association. Is not that the object of a properly designed case control study? And secondly, is it not true that in a register study, of which I have done many, the issue of asking information about occupations is almost certainly biased by concepts of what you believe or what the people believe to be the truth? Is not in Australia, the biggest producer at one time of crocidolite, liable to get questions which suggest that lots of occupations may be due to mesothelioma? Is not the object of an objective, scientific epidemiological study to remove sources of sampling bias and information bias?

Dr. Infante

83. Yes, when you are sampling, you are trying to eliminate the bias in your study design and the extent to which you do that depends on the success of selecting your controls. If you do a study and you are looking at mesothelioma in North America then you are really trying to sample, as you said, the individuals that match closely the cases, and the cases are coming from the entire North America. So in my opinion, you are still trying to estimate the universe in that particular type of a case control study. In terms of bias from a registry source, you can have bias from a registry study, you can have bias from the registry. It depends on how the questions are asked.

Chairman

84. Canada?

Mr. Hankey (Canada)

85. I have one final point. It is to Dr. Henderson and Dr. Infante and it is simply this. If we have a registry study, such as that conducted by Dr. Henderson and we have ...., I forget what number of garage mechanics he found, but let’s say that it was 50, how does he determine, without controls, whether that is a large number or a small number, or a number that is more or less in proportion to what you would find in the general population. What is the significance of that number? What does it tell us if there are 50 garage mechanics, if among mesothelioma victims there are 50 who are garage mechanics?

Dr. Henderson

86. The figure in the 1999 Register was 58 mesotheliomas in individuals designated as brake mechanics for which there was only exposure to asbestos derived from brake lining and brake block materials. This was a group separate from the individuals who had multiple other exposures to asbestos. In this respect, most of the history data for the Australian Mesothelioma Register, are fairly accurate, as much as one can ever achieve with a population-based set of statistics, in that the occupational histories in Western
Australia and New South Wales are taken by professionals that are asking histories, for example, the New South Wales Dust Diseases Board. We don’t know exactly how many brake mechanics there are in Australia, but the 1996 Australian census figures came up with a figure of approximately 87,000 male automobile mechanics, that is mechanics of all descriptions [among whom brake mechanics would constitute a smaller] … [END OF TAPE] … number of individuals; what I then did in calculating the statistics was to round it off at 100,000 or 200,000 to take into account the number of mechanics who might have left the industry and retired. I compared that simply with the estimated background rate of mesothelioma for the general population as being one to two cases per million per year. Using the upper figure of two cases per million per year, I still came up with an increased number of mesotheliomas above what I would have expected for purely spontaneous or background mesotheliomas. Now, this was no systematic study on my part, it was a set of calculations. But the interesting thing was that the figure I came up with was roughly comparable to the figures given from the National Occupational Health and Safety Commission in Australia, which had found an increased incidence of mesotheliomas, and that the rate of increase is roughly proportional to the rate of increase of mesotheliomas amongst the rest of the population. I don’t pretend that the statistics are anything more than that, but to me they are an indicator in terms of approaching a problem at a national occupational health and safety level of indicating a possible effect and therefore the need for a cautious and prudent approach.

Mr. Hankey (Canada)

87. Sir, it sounds to me like you are saying that your study, the probative value of your study is your intuition as to what the percentage of garage mechanics is in Australia because the kind of calculations you have just suggested do not strike me as the kind of scientific rigour that would be required in order to produce a study which would have probative value in any court of law.

88. But I have another question, Dr. Henderson, well this is for Dr. Henderson, yes, again. Are you aware that in about 1990, Dr. Woitowitz and Dr. Rödelsperger published a short report noting that they had collected a number of cases of mesothelioma in men who had been automotive mechanics and they were concerned that, because these men had low exposure to chrysotile during brake work that their mesotheliomas would indicate that low exposures to chrysotile were causing mesothelioma. In 1994, they reported that they had carried out a case-control study with two different types of controls and that both showed that there was no association between work as a mechanic or brake repair work and mesothelioma. Now, in view of these findings and the overwhelming evidence for no association between friction products and mesothelioma, wouldn’t you agree that no conclusion should be drawn from the Australian Registry regarding the relationship between mesothelioma and brake repair work as no controlled study has been done?

Dr. Henderson

89. Well, I am no epidemiologist, but I would not draw the conclusion that no conclusion can be drawn from the Register figures. We have figures for a number of different occupations, from the Register, who have mesothelioma and the fact that we don’t have precise data on all of those other occupations represented on the Register, does not mean that we can draw no conclusions from them. I am aware of the study carried out by Dr. Woitowitz, (in fact, if you read my report initially submitted to the WTO, you’ll find that I did discuss that report), but I will also point out that at the conclusion of their revised report, Dr. Woitowitz pointed out that some of their cases had in fact sustained amphibole exposure as well, and that, taking that into account, they could not identify an excess risk. But they also indicated that their study had low power to detect small risks and they indicated that, if the risk of mesothelioma was small, their study would not have detected it. Now, all I am saying is that when we are dealing with a national population of 18 or 19 million individuals and we look at all of the mechanics amongst that population, we are dealing with a larger population, and although it may not have the precise rigour of a case-control analysis, the figures nonetheless do indicate that there is an increased risk of
mesothelioma among brake mechanics who are exposed only to chrysotile asbestos from grinding brake blocks. The other point that I would make is – when I looked at the figure for the mechanics across Australia and I compared it to the figures also given for North America – the figures were that the number of estimated brake mechanics for the two populations were surprisingly similar. So one of the points that I would emphasize is that if my figures are inaccurate – and they may well be inaccurate – they are inaccurate on the side of conservatism, in that I have overestimated the total number of brake mechanics and, therefore, perhaps underestimated the effect.

Chairman

90. I think we are keen to move on through the list of questions and I have just consulted briefly with my colleagues, and we feel that on this present question plus set of supplementary questions, we have gained a great deal of clarification of the experts’ views. So I would invite Canada maybe to make one last comment or raise one last issue concerning this group of questions before we move on. Thank you. Before that, we invite Mr. Christoforou to take the floor who has been seeking to make a comment for some time.

Mr. Christoforou (European Communities)

91. Thank you Mr. Chairman. I would like to ask a question on this because I’m afraid that the way we proceed – it is up to you Mr. Chairman - but it will take us quite the entire day probably. We will not finish, and on this point we run the risk of trying to see an individual tree and would lose the entire picture of the entire wood in this case. So I would like to come back to this question with one and then I still have another question to ask, as I pointed out.

Chairman

92. I am sorry I didn’t quite understand what precisely you wanted to come back on.

Mr. Christoforou (European Communities)

93. I want to ask a subquestion on this point and then ask the other question I have. I announced two questions on this first point.

Chairman

94. OK. I did invite Mr. Hankey to make a final comment on this particular set of issues. Please go ahead.

Mr. Hankey (Canada)

95. My question is: taking account of the definition of construction workers, of those categories of workers that you each identified as construction workers in response to my first question, over a one-year period, is a construction worker at greater risk from exposure to low-density asbestos products in place, or from exposure to products at issue in this case, that is, high-density chrysotile-cement or friction products? That question is to each of the experts, thank you.

Chairman

96. Would you mind repeating the question, I think the experts are not clear exactly what the question was?

Mr. Hankey (Canada)

97. Of course, Sir. I said that, taking into account the definitions you gave earlier, or rather the list of workers, the universe of workers, that you consider to fall under the general
rubric “construction workers”, over a one-year period, is a construction worker at greater risk from exposure to low-density in place asbestos products or from exposure to the products at issue in this case, that is to say high-density chrysotile-cement or friction products?

Dr. Henderson

98. The question is a little bit like asking how long is a piece of string. It depends on so many different variables that the answer will vary according to those variables. It depends on what the worker is doing with low-density asbestos-containing insulation materials, how often he or she is doing it, the frequency with which this is happening and so forth. I would believe that for an individual who works consistently with low-density friable insulation materials, either applying them or removing them, the exposure levels are likely to be consistently high and that person would be at higher risk for both mesothelioma and lung cancer. When one is dealing with a high-density product such as asbestos-cement, it is very difficult to make direct comparisons. But if you look at individuals cutting asbestos-cement building products with a power saw for example, that can generate very high airborne fibre concentrations and again the effect in terms of mesothelioma and lung cancer induction will depend on the levels, the frequency and duration of the exposures and therefore the total cumulative dose. But even if one takes into account the fact that, or one concludes that, the worker dealing consistently with friable insulation materials is at greater risk, the point that I would make is that in Australia, most of the building construction workers we come across give a history of consistently working with high-density asbestos-cement building products. Although their risks might be less than the corresponding insulation worker, there are many more such individuals engaged in that type of activity, manipulating high-density products, and therefore a lower risk needs to be multiplied against a greater number of workers, so that the total effect we see in terms of mesothelioma incidence is greatest for example, among carpenters in the Australian building industry, who consistently cut high-density asbestos-cement building products.

Chairman

99. Thank you. I would now like to give Mr. Christoforou the opportunity to make his comment and also to ask the second aspect of the original question.

Mr. Christoforou (European Communities)

100. Yes, Mr. Chairman, thank you. Before I ask the question of clarification. Canada says the products in this case are high-density cement asbestos products, but that is not the case. Here, Canada exports chrysotile, period. We are not dealing here only with asbestos-cement containing products. I don’t see what is this limitation referred to by Canada. Canada exports chrysotile. A substantial part of it may go to the production of asbestos-cement containing products, but this is not the only use, so we cannot really reduce the entire issue to this product and try to probably confuse everybody around this table. My question, or rather subquestion, I wanted to ask to the previous issue we have been discussing, was the following: I guess everybody agrees - and the scientists here have already said it so - international institutions like the International Agency for Research of Cancer, have since a long time classified all forms of asbestos, including chrysotile, as a proven human carcinogen. I guess there are good scientific reasons before an international institution does so. The scientists here have all agreed on this. Now, we also – given the previous definition of the large, wide category of skilled and unskilled workers which are involved in the everyday handling in their jobs of these substances, and also outside non-skilled, non-workers, like the handyman-type situation, which we also discussed in the papers – given this entire category of people that come in contact with chrysotile, how reasonable it is, or does one really think one can explain only by the data referred to by Canada on brake mechanics, that we can cast doubt on the evidence we hear from so many other sources? Mr. Henderson has referred to inputs from a number of countries in his reply, first set of replies. I refer to pages 35 and 37 of his report on inputs from Russia, the former German Democratic Republic, Italy, from China, and so on, which involved only

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inputs of chrysotile exclusively, nearly exclusively.\textsuperscript{232} And we also know that France has been importing over the last 50 years, exclusively chrysotile, 95 per cent or even more. And still we see so many cases, more than 1,000 cases per year in France of mesothelioma lung cancer cases. Do you really think it is possible to attribute all these cases of asbestos-related diseases which we see by the small percentage, infinite percentage of other types of asbestos like amphiboles, crocidolite or everything else in explaining these cases we see. Can we really use the example of [ ... ], if there is any doubt - we think that there is not - from the brake mechanics to question the entire evidence which has lead international institutions to classify all forms of asbestos as a proven human carcinogen? It is addressed to all scientists, in particular, Dr. Henderson. Thank you.

Dr. Henderson

101. Well, I would be in broad agreement. I think the argument would be made that most of the mesotheliomas – and some would argue that the lung cancers that we see – are not so much due to the chrysotile, but due to co-existent amphiboles in place and encountered by building construction workers. Certainly most of the mesotheliomas, but not all, that I see occur among workers who have had a history of mixed exposure to asbestos cement building products that contained chrysotile and varying amounts of amosite and crocidolite, or both at different times. However, as I have indicated, I have seen mesotheliomas among brake mechanics who only had exposure to chrysotile. So I think that this becomes an argument, as to whether one says that the chrysotile has no effect whatsoever – and that all the effects we are seeing are due to the amphibole content – or that one is looking at a mixed response to amphiboles plus the biological effects of chrysotile. I suppose that one of the concerns I have about the continued use of chrysotile, particularly in situations where it cannot be controlled, is that many of the workers who will be handling that type of material may have a pre-existent amphibole and chrysotile content in their lung tissue and we have few, if any data, on the additive or multiplicative superimpositional effect of extra chrysotile exposure on top of a pre-existing amphibole burden. Although I can’t quantify the effect, one suspects that it would not be a negative effect and that it would contribute both to mesothelioma and lung cancer incidence. But perhaps the others might prefer to elaborate upon that.

Dr. Infante

102. Yes, I think if I understood your question, how I interpreted your question was, that if you have an individual who is diagnosed with mesothelioma and they have been exposed to amphibole and chrysotile, can you dismiss the component of the chrysotile exposure as contributing to that mesothelioma? My answer to that question is no. We know that chrysotile is capable of inducing mesothelioma, so just because individuals have mixed exposure to amphiboles and to chrysotile, you can’t exclude that individual’s chrysotile exposure as contributing to the development of mesothelioma.

Chairman

103. Would either Dr. de Klerk or Dr. Musk like to add anything to what has just been said? Canada, please.

Mr. Hankey (Canada)

104. Sir, I have a follow-up question to that question. This is directed to Dr. Infante. Dr. Infante, can you identify for us any controlled studies of cement or friction product workers showing that the risk of lung cancer to chrysotile cement or friction product workers is as great as the risk of lung cancer to amphibole workers?

Dr. Infante

105. Yes, the 1987 study by Hughes et al. They analyse their data by workers exposed to chrysotile only versus workers exposed to chrysotile and crocidolite. The dose response for lung cancer in that study is similar.
Mr. Hankey (Canada)

106. Could I ask Professor McDonald to comment on that please?

Dr. C. McDonald (Canada)

107. Dr. Infante, I haven't got the paper of Hughes in front of me, but I am very familiar with it, and in fact are you correct? I am fairly certain that the slope was appreciably higher in the factory which had the amphibole workers. Moreover, they had more mesotheliomas.

Dr. Infante

108. I was answering the question related to lung cancer and I believe that if you look at table 10 in that report, they did an analysis, for lung cancer, looking at individuals exposed to chrysotile only and then they did an analysis looking at individuals manufacturing cement products exposed to chrysotile plus crocidolite, and the dose response is similar for lung cancer.

Mr. Christoforou (European Communities)

109. Mr. Chairman, on this question, we know the Environmental Health Criteria 203, which has been cited by Dr. Henderson with his reply to question 1(e) on page 59, (he quotes a long passage from the WHO Environmental Health Criteria Report, it is already filed with the Panel). I interpret this citation as Dr. Henderson agrees with this study which is cited there, where even in Canada there was a study, where exposure to chrysotile and amphibole was separated and there was exposure to several types of situations. The study here has identified a double increase almost 2 per cent increase of chrysotile, of mesothelioma and lung cancer. Would you think that this is one of the studies which is searched after and Canada would like you to point out because it is used already by the Environmental Health Criteria Report?

Dr. Henderson

110. Well yes. I am familiar with that passage. I took it from the WHO book, Environmental Health Criteria 203, and as far as I can see, that's an accurate quotation. I really cannot elaborate further.

Chairman

111. Canada, please?

Mr. Hankey (Canada)

112. I would like to come back to that later Sir, because the name of the study wasn't identified. But I just want to ask a follow-up to Dr. Infante. Dr. Infante, isn't it the case that the Hughes study contained large numbers of temporary workers and that, when those temporary workers were abstracted out of the study and the figures were calculated only for the permanent workers, that permanent workers exposed only to amphiboles had 25 times more cancers than permanent workers exposed only to chrysotile at the same dose.

Dr. Infante

113. Are you referring to lung cancer or are you referring to mesotheliomas?

Mr. Hankey (Canada)

114. I meant to say lung cancer.
Dr. Infante

115. I can’t recall the particulars of that study right now, but I do recall table 10, like I said, which showed a similar dose response. If your question to me is that, well, if they then removed the short-term workers, were the results different? I don’t recall the data that well to answer that question at the moment. But the authors in the study pointed out that the dose response was similar for the two groups as I had mentioned earlier. And besides if there is some difference because of short-term workers in a study and if in a particular study short-term workers demonstrate an excess of cancer than other workers, or even a greater risk than other workers, you have to find out in that study what the particulars are among those short-term workers. Were they exposed to – do they have higher levels of exposure? You have to explore it. Quite often, short-term workers have the dirtiest jobs and so they have the highest levels of exposure, but I can’t recall the particulars from the Hughes study.

Mr. Hankey (Canada)

116. We have them here. I’d like Dr. McDonald to read them into the record and we will, now that the study has been cited, if it is not already annexed, we will annex it. Thank you.

Dr. C. McDonald (Canada)

117. I have the slopes published, the exposure response slopes for lung cancer from this study. And, in plant 1 which was the one of chrysotile only, the slope of risk per fibre million litre year was 0.0003. In the plant 2, which also entailed exposure to amphiboles, the slope was 0.0076. On the point of short-term workers, again, I am sure that Dr. Infante, as an experienced epidemiologist, will be aware that virtually every cohort study of any material shows high risk of lung cancer in short-term workers. Sorry, do I make it clear? A higher risk of lung cancer in short-term workers, whatever the material.

Chairman

118. Thank you. Is there any further comment from the experts on this point? If not, by my recollection, all that previous discussion was dealing with the two issues raised by the European Commission, so if we could return to Canada to ask its next question.

Mr. Hankey (Canada)

119. I’d now like to turn to asbestos products, I’m sorry, chrysotile-cement products because, as we know, most Canadian chrysotile is exported to France and to other places, the great majority of it is used for chrysotile-cement products. Gentlemen, Canada wishes you to consider available scientific and properly controlled evidence on the probable risks associated with the manufacture and use of chrysotile cement products. For chrysotile-cement products, we know of four cohorts: Thomas, (1982); Ohlson, (1985); Gardner, (1986) and Hughes, (1986). In total 6,843 men were studied with 1,432 deaths. 118 of the deaths were from lung cancer, that’s an SMR of greater than one, that is to say the mortality rate is greater than one relative to the general population. I’m sorry, mortality rate less than one, did I say greater? So the mortality rates overall were less than those in the general population from lung cancer. This means that overall there were fewer deaths from lung cancer than expected in the general population and these are cohort studies of 6,843 men. I would like you to comment on those deaths, Sir. This question is addressed to all four, but perhaps I might address it first to Dr. Musk.

Dr. Musk

120. I’m afraid I’d need those studies in front of me to address that. I’m not familiar enough with them, although I have read them.
Dr. de Klerk

121. I think, because we are looking at asbestos-cement and, therefore, because Canada is saying that most of the products that they want to export to the European Union, presumably are used in asbestos cement, that therefore one should ignore all the other evidence about chrysotile apart from that from asbestos-cement workers, I think that it is a bit of a ..., I can't think of the word, but anyway I think you know what I mean.

Mr. Hankey (Canada)

122. I don't know what you mean. I would be grateful if you would elaborate on that.

Dr. de Klerk

123. Well it's a ..., I can't think of the word, I will explain what I mean. It's, you are sort of ignoring a lot of the fact that chrysotile will be completely different in its actions and effects; because it's in a cement product, ignores the facts about the fibre clouds produced by asbestos-cement products and the evidence from other forms of use of chrysotile. You know, even when you add these four studies together, they are in fact quite small, if you look at the numbers of deaths and to show that they show no effect is, I doubt very much whether you could actually rule out what could be quite an appreciable effect. So I think that you have to look at all the studies about chrysotile together, rather than just concentrating on asbestos-cement products. I think it is a bit of a ... it will come to me.

Chairman

124. We will let Dr. de Klerk come back on that when he finds the exact word. Thank you.

Mr. Hankey (Canada)

125. I have a follow-up question to Dr. de Klerk, but would you like Sir, that the other Doctors ...

Chairman

126. Maybe give the other experts an opportunity to speak first.

Dr. Infante

127. I am looking at the summary in the document 203 of the studies of asbestos-cement production. As I had mentioned earlier, we already know the toxicity of chrysotile-asbestos and the toxicity of these fibres does not have to be demonstrated in every occupational situation where it occurs. What we need to do now is to control the hazard that's recognized. Having said that, when you look at the four studies, the numbers of the largest study, the Hughes study for chrysotile only demonstrates a significant excess of lung cancer. The Hughes study of chrysotile had included crocidolite and amosite exposure, there is a 17 per cent excess, that's not statistically significant. The Gardner study, the SMR is 92, that's not significant elevation or deficit. In the Hogstedt and Ohlson study, there is a 58 per cent excess and that's not statistically significant, and in the Thomas study it isn't.

But we already know the hazards of these fibres and, also, if you look at the upper 95 per cent confidence limit, some of these studies were only 95 per cent confident that the risk, for example in the Ohlson study is not as high as three-fold for lung cancer. So I think that you have to look at not only the SMRs, but the confidence intervals around these studies.

Chairman

128. Professor Henderson, would you like to add anything on this point?
Dr. Henderson

129. I couldn’t add anything unless I have the particular reference in front of me. It is part of a large volume of material and I can’t remember the precise details. In general, though, I would point out that certainly in the manufacture of high-density chrysotile products, at least in Australia, where it is almost a totally closed operation and the airborne fibre concentrations are extremely low with a predictably low risk for that particular cohort. My major concerns about the use of these products is in the end-users who manipulate, saw, drill, rasp, grind or otherwise handle these materials and that one knows that some of the fibres released from these operations will produce elevated airborne concentrations of fibres which are in the dimension range known to be associated with carcinogenicity, even though in some circumstances it might be a relatively small proportion of the total fibres released.

Chairman

130. Thank you. Canada, please.

Mr. Hankey (Canada)

131. I think we have really gotten to the heart of the matter, because it is incontestable that in these four studies - which are the only cohort studies of persons working with chrysotile cement, the only control studies - they all show together, collectively, that there are fewer deaths from lung cancer than expected in the general population. I don’t think those data, Sir, can be swept under the carpet. Dr. de Klerk, I think really does identify what is the issue here. He says that we should look at data for other industries and apply it to the cement industry or the friction industry, because asbestos is a known carcinogen. So it seems to me that Dr. de Klerk proposes that we compare apples with oranges. Now, in my business, Sir, as a lawyer, when we deal with evidence it is always a requirement that we compare like to like. There are many, many rules of jurisprudence that require that. So I would like to ask, now, this question generally because it is on the very same point to each of the experts who wish to take it up: given that each chrysotile industry sector has its own particularities, that is to say, wet or dry processes, open door and closed processes, different fibre lengths and the possibility for oil treatments, doesn’t it make sense to base risk assessments as much as possible in one given sector on the particular experiences of the workers in that sector and not in workers in a completely different sector?

Dr. de Klerk

132. Well, it would if there were data available. But I think, as Professor Henderson has pointed out and we all agreed on in our reports, I think the main risks that people are worried about are in the downstream user, the builder, the construction workers etc., and there aren’t any data for those. So what one does is extrapolate from other studies where fibres of the relevant size, shape and density are available so that, as Professor Henderson said, the asbestos cement industry itself has been well controlled for a long time. The worry is not the asbestos-cement industry, it is the asbestos-cement user, I think.

Chairman

133. Does any other expert wish to comment?

Dr. Henderson

134. Really I can’t elaborate beyond what my colleague, Dr. de Klerk, has said. One of the problems we are dealing with trying to assess the risks to different groups is that we’re faced sometimes with conflicting or contradictory data for areas for which we have no direct observational data. Therefore we need to proceed in an area where there is some uncertainty as to the exact risks for a particular population. This is one of the reasons why we tend to use extrapolation models as Dr. de Klerk has said, and also use other
investigations. For example, the South Carolina chrysotile-textile workers [are used] almost as a worst-case scenario in order to formulate prudent approaches to population safety.

Mr. Hankey (Canada)

135. I’m still following up on this question. This question I could direct to both Dr. Henderson and Dr. Infante because Dr. Infante is obviously very keen on Charleston, as am I, it is a wonderful city. My question is, if indeed there are no applicable studies to the use of asbestos cement in construction, then evidently we have to find a surrogate study, something that is closest to it and it seems to us, and that is why we put the question, that the other sector where asbestos cement is being handled and being used, that is to say in its manufacture, that that would constitute the best surrogate. The results of those studies incontestably demonstrate that there is no increased risk of mesothelioma or lung cancer, there is no dispute about that. So instead, the surrogate that Dr. Infante takes us to, and I gather Dr. Henderson, is the textile industry in Charleston. Now, I must say that it is a little difficult for me to figure out exactly what the relevance is of textile workers to asbestos cement workers. First of all, Canadian asbestos is not used, cannot be used these days in the manufacture of textiles, because nowhere do we know are these textiles being now manufactured. Certainly not in the European Union or in North America. It is our view that the Charleston data are very unreliable for a number of reasons. Some suggest that data from the Charleston textile cohort is relevant to this proceeding. Unlike asbestos cement in friction material processes, friction material production processes, the processes used in producing textiles differ enormously from those used to produce chrysotile cement and friction products. For example, the Charleston textile cohort was exposed to crocidolite and amosite amphiboles. Chrysotile during carding, a dry process in which the fibres are split and torn apart into numerous fibrils without substantial controls. (I know people who have visited that factory and the stuff was spinning around and it was hanging from the ceiling like cobwebs) This is totally unlike the conditions in a chrysotile-cement factory, or on a construction site, using chrysotile-cement. Finally, carcinogenic oil was sprayed on the fibres at the start of the production processes and it is doubtful that a valuable allowance can be made for oil fibre or oil-smoking interactions. Given these quite substantial and different exposures, why do you think it makes sense to extrapolate from textile cohorts to asbestos cement and friction product cohorts?

Chairman

136. Thank you. I guess the question was addressed to Dr. Infante and to Professor Henderson, so I will give them both the opportunity to respond to that. Then Mr. Christoforou also wanted to make a comment.

Dr. Infante

137. Thank you very much. First of all, in your question, you had an incorrect factual statement. That is if you look at asbestos cement production, the Hughes study, which shows a statistically significant excess, furthermore demonstrates a dose response from exposure to chrysotile-cement production only and lung cancer. In fact the potency estimate is 0.7 per cent which is just a little bit less than the 1 per cent estimate per fibre per c. c. year which has been identified in two other studies of workers exposed to chrysotile textiles and a little bit lower than the estimate from the Dement study. But there is really not a great deal of difference in my opinion, between 0.7, 1 or 2 per cent. So in fact you do have dose response and when you demonstrate dose response in an epidemiological study, that is a very powerful tool. So there is evidence, even though some of the other studies don’t show an excess of risk, they didn’t find it for whatever reason, but I would submit that the Hughes study does demonstrate an excess and does demonstrate dose response. In terms of the abuse of the Dement study to estimate risk, it’s the study that has been the most thoroughly evaluated, the exposure estimates in that study, in my opinion, are superior to any other study because they took simultaneous samples doing particle counts and counting fibres. They used different correction factors depending on which operation was being used in that study. So the Dement study is very good in terms of characterising exposure.
There was some amphibole exposure to that cohort, but when you bring in some amphibole exposure to a cohort for a certain period of time and then it is no longer there, that in fact may dampen your dose response, when you are looking at fibre exposure, if your lung cancer would only be related to the amphibole. So, and furthermore, Dement has done analyses, which I had mentioned earlier, where he identified his cohort, depending on whether it was the entire group, his case control analysis for a dose response, or people who began after 1940, or began before 1940 or 1950 and he has the same dose response. So, in my opinion, that is a very strong study and it should be used to estimate risk of lung cancer from exposure to chrysotile asbestos. And it is not dissimilar from the dose response from the Rochdale chrysotile workers or the Pennsylvania plant in the US.

Chairman

138. Thank you. Professor Henderson, please.

Dr. Henderson

139. I basically agree with the comments made by my colleague Dr. Infante. Yes, on the face of it, the South Carolina, the Charleston chrysotile-textile workers appear to be different from asbestos-cement manufacture. I think the differences are partly explicable by the fact that, as I have said, in Australia asbestos-cement manufacture is a completely closed operation so that there are very low airborne fibre concentrations. But one knows that for downstream users, the operations carried out on those high-density products will produce elevated levels of airborne respirable asbestos fibres. Some of those fibres will have the dimensions that are known to be associated with carcinogenicity.

140. The point about the asbestos textile industry - and I agree with Dr. Infante that this is one study which has been regarded as a classical study and a very rigorous one in its methodology - is that for white males in the Charleston factory, there was a greater than two-fold increase in the standardized mortality ratio for lung cancer at quite low airborne fibre concentrations. That was quite different from the Quebec chrysotile miners and millers. But for many other studies we don’t have direct observational data and one needs to try and take into account the fact that some of the downstream uses of asbestos-cement products may generate airborne fibre concentrations which at a cumulative inhaled dose may approach the levels of carcinogenic fibres that have been reported for the Charleston cohort. It might be argued that we don’t know that the response from those workers will be the same. But equally we don’t know that it will not be the same, because we don’t have any data. In relation to the Charleston cohort, it was claimed that these workers had crocidolite and amosite in their lung tissue. I guess some of them did, but one of the problems about this and particularly the Case-study, is that those fibre concentrations were not linked to lung cancer as an outcome. It was simply a study done on this particular group of workers and in fact, when you look at the lung cancers that were studied in this group, and particularly the lung cancers amongst the Quebec chrysotile workers, there are substantial differences which indicate that the two groups simply are not comparable. For example, the interval following cessation of exposure until death, when the fibre burden analysis was carried out. But there is another factor which needs to be taken in to account, is that it was argued that the Charleston workers had [commercial amphiboles, crocidolite and amosite, in their lung tissues, but if one looks at the total amphibole content]... [END OF TAPE]... that is tremolite plus amosite plus crocidolite, it was higher for the Quebec group. So that if the lung cancer effect is related to amphiboles, why aren’t there more lung cancers and a higher lung cancer risk and mortality rate amongst the Quebec workers? When you look through the evidence, I don’t think that this idea of commercial amphiboles being present withstands serious analysis. The other point that was raised was that the Charleston workers may have used carcinogenic oils. Again, I think that there is no evidence for this proposition. All of the reasons for the difference between the Quebec chrysotile workers and the Charleston workers have been explored and, at the moment, there is no convincing explanation for the many-fold difference in lung cancer risk between the two groups. When there is no obvious explanation, my policy is to proceed on a approach of caution, to make sure that people are not exposed to a substantial lung cancer risk. The only other thing that
I would say is that it was said that we can’t use the Charleston chrysotile workers as a paradigm for lung cancer risk assessment. But this is exactly what Drs. Case and others said in the Abstract that they submitted to the Maastricht meeting. In the Abstract they said that risk assessment for asbestos exposure is based on lung cancer risk for textile workers rather than the miners and millers.

Chairman

141. Thank you. Mr. Christoforou asked for the floor a while ago. Would you like to make a comment now?

Mr. Christoforou (European Communities)

142. In terms of the population that is potentially at risk, as we have identified the people we are concerned, especially the case was of France, when it adopted the Decree in question. You have identified a large segment of the population, both skilled and unskilled and you have said that all these people are at risk. Canada is asking the question and trying to limit the issue on high-density chrysotile-containing cement products and their manufacture. But they export chrysotile as a product. Now, if I may ask, especially Dr. Infante for your experience in the regulatory approach to these questions: do you think it is reasonable, knowing the potential large segment of the population that deals with not only the manufacturing but subsequently during the lifetime of the product that remains in the place, is it reasonable, knowing the evidence we have, to take measures that prohibit the use of asbestos rather than leave it and allow it to be imported and be faced with all its potential effects on all these categories of people. Thank you.

Chairman

143. I’ll give the floor to Canada next, if you want to follow up any more specific aspects on the responses just given, and then invite the experts to respond to the point made by Mr. Christoforou.

Mr. Hankey (Canada)

144. I would like Dr. Henderson to recite again, because I didn’t perfectly understand, why he feels that the Charleston textile study is a better paradigm or a better surrogate for exposures and risks in the chrysotile-cement construction industry than is the chrysotile-cement manufacturing industry or the friction manufacturing industry or brake mechanics who are exposed to chrysotile. Sir, I think it is incontestable that the Charleston study is an outlying study, it’s a study whose results do not conform to nearly all the other studies we have on asbestos. I am not aware, but there may be and I am evidently a layman, I’m not aware of any other study that could be said to corroborate the results of the Charleston study. Yet we have cited many today even, and in our pleadings, many studies that show no increased risk of cancer from exposures to chrysotile cement and to friction products. None of the experts have contested these assertions. Yet you picked this study from Charleston which has figures which are radically at odds with all the other studies and I am still ... knowing what I do know about the conditions in that plant which are well documented. By the way, it is not factually correct to say that it is not known whether oil was used; oil was used and that is recorded in the studies and used frequently and used consistently, so we know that oil was used and we know that oil is a carcinogenic product. It seems to me that one cannot at all exclude that as a valid hypothesis for the difference. Even apart from the oil, how and why would you justify using that textile mill, where conditions were so different to those in the chrysotile-cement industry, a plant that was clearly negligently controlled by the South Carolina authorities, why would you use such a surrogate for the modern up-to-date chrysotile-cement industry?

Dr. Henderson

145. Well if the question is asked why do I use it? The answer is many others do also. In relation to the comment about oil, I may not have expressed myself with clarity. I was
not saying that oil was not used – what I am saying is that studies on the carcinogenicity of the oil have yielded negative results and cannot explain the difference between the two worker cohorts. I think this point was made by Professor McDonald himself in a recent editorial on this issue, that really the difference for the dose-response line for the two groups for lung cancer remains basically unexplained. Because we do not know the explanation, it is difficult then to control for whatever that unknown factor is. In approaching national policy for occupational health and safety, one often adopts a prudent approach, using a conservative or worst-case scenario on the “first do no harm” principle. When it comes to the fact that the two groups are different from, for example, asbestos-cement manufacture, I don’t dispute for a moment that asbestos and friction product manufacture nowadays carries a very low risk because there is a low airborne fibre concentration with low cumulative exposures. The reasons why I use the Charleston group as an approach is that it identified a high lung cancer risk at low exposure and that the types of fibre released during that operation can be released during end-work, that is machining asbestos-cement products. If this can produce comparable cumulative exposures, one needs to assume that we have not proven that those exposures will have no effect. Therefore it is basically an approach of safety and prudence for the formulation of the national health policy.

146. The other thing is, I’ll just reiterate, that if you go to the Abstract for the paper by Drs. Case and others, they themselves said that the risk assessment for asbestos exposure for lung cancer is based on lung cancer risk for the textile workers rather than the miners and the millers. The question is which is the outlier – do you take the Charleston textile group as the outlier and ignore it or do you say, well, perhaps there is something peculiar about the Quebec chrysotile miners and millers and, in terms of their exposure and dose response, then they are an outlier in that they showed a very low slope to the lung cancer dose-response line. Other studies have showed an intermediate risk. So the question is which one do you adopt for formulation of national health policy?

Chairman

147. As we conclude the morning sessions, could I perhaps make a couple of comments which might help us into the afternoon? Just looking at the questions and the discussion so far, it seems to me that we have had some quite good coverage of the first broad subject which is chrysotile itself. From the Panel’s point of view, it will be important that we also have some time to discuss the two other broad topics of controlled use and substitute fibres. So I’d reiterate my request or my suggestion to the parties to be selective in their questions and comments. It obviously is not going to be possible to deal with all the issues in an exhaustive manner. So what we will do, I think, is see how we progress in the afternoon. The Panel will want to make sure that there is time for those other two issues I mentioned and we may have to make time for those issues and then come back to the first set of issues concerning chrysotile at the end if we have time. I’d also ask the parties, when you are asking your questions, could you please as far as possible refer directly to either the Panel’s original questions or to the parts in the experts’ reports where they have addressed these issues so that it is easier for the experts to respond and see exactly what you are each referring to, especially when we have cases of some of the studies that have been cited. They are usually there in the material somewhere, but perhaps sometimes a page reference would be helpful to enable the experts to respond more quickly. We will reconvene at 3 p.m. Thank you very much.

[Lunch break]

17 January 2000, p.m.

Chairman

148. We were part way through the discussion of one question. I just checked with Mr. Hankey as to how far through the list of those questions relating to the first four questions submitted by the Panel Canada was and I understand that they’ve made very good progress through them. So that being the case, I think we will continue on with this current question...
where there may be one or two comments still to be heard, then proceed on through the
remaining one or two questions concerning the first broad heading of chrysotile asbestos
itself. That would then give us adequate time to work through controlled use and the
question of substitute fibres. I would really express the hope that by 3.30 we should be
beginning our discussion of controlled use. If that is acceptable, we now open the floor.
We were in the midst of a discussion, there was a point that had been made by
Mr. Christoforou which had not yet been responded to. I think Canada also had one or
two additional elements that they wanted to mention. So unless the experts feel that they
need to add any points to the responses they have given so far, I could pass the floor back to
Canada if you were following up on the question currently under discussion. Then we can
ask the experts to respond to Mr. Christoforou’s point.

Mr. Hankey (Canada)

149. Thank you Chairman. Our follow-up just has to do with this issue that in the
absence of much direct data on the use of chrysotile-cement products in the construction
industry, what would constitute a good paradigm or surrogate among the various studies
that do exist. We know that there are close to sixty studies about the use of chrysotile
asbestos and we have had a lot of reference to the Charleston study: Charleston is wonderful
for the jazz festival but I am not so sure it is very relevant for the issue before the tribunal.
So Dr. McDonald would address the issue of what he thinks might be a more appropriate
paradigm or surrogate to examine the issue of risk exposures in the use of chrysotile-cement.
Thank you, Dr. McDonald.

Dr. McDonald (Canada)

150. I’ll try to be as brief as I can. The first point being that, of course, the Charleston
cohort of textile workers is not unique in textile cohorts. There have in fact been three:
one, Charleston, is almost entirely chrysotile but there are two others which have been
mentioned briefly, in which there were substantial amounts of crocidolite used. I only
want to make the point that all three of these textile cohorts show this anomalous high
level risk of lung cancer, whether it was chrysotile or whether there were amphiboles,
whereas so far as mesotheliomas are concerned, the presence of crocidolite clearly correlated
with the incidence of mesothelioma. In other words, there was no excess of mesothelioma
in the Charleston cohort, any more than there is in asbestos mining and milling. Whereas
in the other two textile cohorts, there were substantial numbers of mesotheliomas. Therefore,
those of us who have been trying to understand why the textiles are different conclude, I
think, that there is something funny about textiles. It is precisely that point that makes me
say I would be personally wondering why you would choose something anomalous rather
than something that is in line with the rest. I refer now to the fact that by far the biggest
scientific study of chrysotile workers is the one of chrysotile miners and millers in Quebec
which has been of some 11,000 men studied continuously for 35 years and now 80 per cent
of them are dead. So we have one of the most complete pictures of mortality in chrysotile
workers almost ... than you can imagine. There is nothing comparable. These men were
exposed in the 1930s and 1940s to astronomically high levels of chrysotile exposure. Now
the issue of exposure levels has been questioned, if you like, by Dr. Infante who said that
the methods used in the Quebec cohort were different, if you like, by Dr. Infante who said that
the methods used in the Quebec cohort were different from those in Charleston. I would
like to point out that it is not so. We also estimated exposures individually in relation to
fibre conversion and in fact published a detailed report in 1980, showing that the risk estimate
based on individual estimations by fibre gave us exactly the same estimate risk as using
the average. I can give the reference for that: it was published in an international meeting
in Lyon by the IARC and I am sure Dr. Infante is familiar with it. I hope that that would
reassure him that there is no reason to think that the exposure estimates in Quebec were
any better or any worse, shall I say, than in Charleston. In fact, they were based on a very
much larger amount of data, a very much larger, with parallel counts by fibre and dust, as
in Charleston. There was no difference. So we are left with the fact that Charleston is an
anomaly. Now I don’t want to go into the Quebec result in detail but we have 8,000 deaths
which is, I suppose, as big as there is in any study. What everybody, I think, is familiar
with, is the fact that it showed a very modest risk of lung cancer, except at quite high levels.
Indeed, the cohort mentioned as asbestos cement workers in that part by Hughes, in the part of the study which was chrysotile only, gave a slope almost identical with the miners and millers. I would also point out that the type of work in mining and milling, in which you sort out the fibres, is very similar to that in cement workers and in friction product workers, very similar and quite different from that in textile workers. And what that showed, as I say, was a very modest increase at low levels, a substantial increase in lung cancer at high levels, a substantial one, but at levels below about 25 fibres per c.c. for forty years' work, we could not detect an increase in lung cancer. It doesn't mean there wasn't one. We're quite prepared to accept the concept of a linear relationship but the fact remains that for people below that level, we couldn't detect any increase. This is a very big cohort. Equally, yes, there were mesotheliomas in this cohort but not very many in relation. There were 33 deaths from mesothelioma in miners and millers: not a single one in a man who had worked for less than two years and only one in a man who had worked for less than twenty years. Surely that suggests that the risk of exposure in that very large cohort was quite modest and that exposures at modern levels of say one fibre per c.c. wouldn't possibly be detectable. But I would submit to you that surely this experience is much more in line with asbestos cement and friction product workers than textile workers, the explanation of which we really don't know.

Chairman

151. Thank you, Dr. Infante.

Dr. Infante

152. Dr. McDonald, you made some comment that the Hughes study gave a slope identical to … I didn't understand then which group you were referring to?

Dr. McDonald (Canada)

153. You will recall there were two plants in the Hughes study, one of which was thought to be essentially chrysotile only and gave a slope of 0.0003 if I remember, which is when I say identical, almost identical with the slope in Quebec. That was one. Not the crocidolite one which was something like 25 times higher.

Chairman

154. Thank you. Perhaps we should now pass to the question or element of question which Mr. Christoforou asked us shortly before we concluded. I think it might be helpful if we asked Mr. Christoforou to repeat the question so that the experts can respond. Thank you.

Mr. Christoforou (European Communities)

155. Thank you, Mr. Chairman. What I said was, following up on what Canada was saying about high density chrysotile-containing cement products and I was confining the argument about the manufacture of such products - and the argument was whether there was any evidence that these may have effect and what was the level, whether there were any worries about the level of exposure and the consequent asbestos-related diseases. I said we want to somehow reposition this argument and request Dr. Infante from his experience and also Professor Henderson: given the fact that even Canada does not dispute that all forms of chrysotile have been classified by international agencies, like the International Agency for Research on Cancer. There is a proven human carcinogen - I don't think anyone in this room would dispute it. And given the fact that the four scientists have defined very broadly the population most at risk to include both skilled and non-skilled workers, not only those dealing in the manufacture of cement, high-density cement and products containing asbestos. The question was then addressed to the experts was from the regulatory point of view, and Dr. Infante has such experience, is it really reasonable to believe that a country like France, which has been importing for the last fifty years, more
than 95 per cent of chrysotile asbestos, and we see so many cases of asbestos-related diseases, is it really reasonable to attribute these cases to chrysotile, is it reasonable to confine the argument to cement products, which anyhow Canada does not export - Canada exports asbestos as a product.

Chairman

156. Thank you. I pass the floor to the experts. Anyone who may wish to respond?

Mr. Christoforou (European Communities)

157. Mr. Chairman, if you wish, because there is an element of controlled use here which can probably lead us to the next question. If we speak about cement asbestos, high density cement products containing chrysotile, and we know from the comments Canada has sent on 13 December, they speak about pre-sized, ready-made, ready-tailored cement products which are delivered to the construction industry and the question we would like to ask Dr. Infante, because he does make comments on this aspect in his replies: is it really reasonable to believe that even in the construction industry, these cement-containing products will never be required to be modified, cut and changed so as to fit for the purpose of construction? Can we really compare only this type of situation with the other possible parts of the workers that are exposed to chrysotile products, just to make the comparison between strictly cement construction with the rest of the population who later on will come into contact with chrysotile in the different types of activities they are involved: plumbers, electricians, insulation workers and so forth?

Chairman

158. Thank you. Does that clarify the question for you? Dr. Infante?

Dr. Infante

159. Let me see if I understand the question. Is the question: is it possible to control exposure to chrysotile in the construction sector of industry, outside of, talking about manufacturing? Is that essentially the question?

Mr. Christoforou (European Communities)

160. Yes, yes. I can give you an exact reference. It is on page 19 of your replies22 where you talk about pre-sized … and the need to modify these products and whether it is realistic to argue, as Canada does, that these high-density cement products will never be changed so that the risk will be, as Canada argues, in this type of situation very low levels of exposure.

Dr. Infante

161. It’s my opinion, that I stated here, that I don’t think you can have chrysotile asbestos cement products in commerce without presenting risks to individuals who may need to manipulate those products. Even if they are pre-sized, they periodically have to be cut, those that are in place sometimes have to be cut into to get into the contents inside pipes that are carrying whatever they happen to be carrying. For example, I know, in the United States, if you take chrysotile-cement to a dump, you are charged by the dump for the volume that you take to that dump site. So, for example, if you were to take a large chrysotile pipe to the dump site, you are charged for the entire volume. So it’s beneficial to the construction worker to chop the cement up into pieces which then adds to the fibre exposure because, one, it’s easier to remove it in pieces and two, it’s cheaper when you deliver it to the dump site. I don’t know what policies are in other countries or how they do business but that’s how it is in the United States and that creates exposure. I feel that, and I think I’ve said, if you cannot control exposure in the occupational setting, in the United States particularly where even in manufacturing you can’t control it, how are you
going to control it in the construction sector? There are just too many variables that you can't control. People don't get educated well enough, people don't wear the appropriate respirators, there just are not programmes that can extend that far in my opinion to protect those workers. Even in the manufacturing sector, just this past October, we fined an asbestos brake manufacturer $125,000 for being over the permissible exposure limit, for not providing respirators, for doing dry sweeping. That's in the United States where we've had an asbestos standard in place for a number of years. So, my point is that it may be theoretically possible but it's not practical to think that you can control exposure to asbestos even in the example I gave in manufacturing and it's certainly less practical to begin to control it in construction.

Chairman

162. Thank you. We appear to have made a seamless transition to controlled use at the moment. I invite new responses by Canada on that.

Mr. Hankey (Canada)

163. Before you get too excited about controlled use, I'd like to bring us back to the question … There was a premise in Dr. Henderson's answer which I think needs to be examined. He said “even if we can't control exposure to asbestos in the manufacturing industry … “. But Sir, the only evidence you've cited or any of the experts here or the European Communities have cited that indicates maybe we can't control it in the manufacturing sector, if I'm not mistaken, relates to textiles. As we have demonstrated, or at least argued I think quite coherently, this is an entirely different sector and one in which asbestos is not used and has not been used for many years in the European Union, certainly not in France. We have data relating to some fifty studies of the use of asbestos in the manufacturing of cement and friction products. We know of no instance that indicates that, in these current manufacturing facilities, there are levels of exposure, cumulative levels of exposure, to asbestos that cause danger to human health. If you or your colleagues or the European Union can put evidence on the table that indicates otherwise, I'd be happy to see it but it seems to me that the premise you base your conclusions about the use of asbestos in the construction industry is simply not viable.

Dr. Infante

164. I gave one example in manufacturing that was surprising to me because one would think, in manufacturing, you can control and you should control. It was just a shock to me to see this company clearly manufacturing asbestos brakes in the United States last fall. They were way above the exposure limit and not doing anything about it. The basis for my opinion in the construction sector is, what I indicated in my written response, that in the last three-year period there have been over 3,000 violations to our standard. A large portion of those are in the construction sector.

Mr. Hankey (Canada)

165. Tell me, Mr. Infante, how many of those violations concern excess exposure limits?

Dr. Infante

166. No, I can't do that because we don't have the data to break out that way. I'm not talking about exposures above the permissible exposure limit because quite often in construction we don't take atmospheric samples. The reason we don't take atmospheric samples in construction is that by the time you would get the sample result back, they're onto the next job. So, rather than taking atmospheric samples in construction what we do is look for other violations of the standards and proper use of respirators or no respirators, improper hazard communication to the workers that are involved, not having a person who has expertise in the hazards of asbestos responsible for the job. It's those kinds of violations that I'm referring to. I was not referring to levels over the permissible exposure limit because in construction we don't take a lot of samples.
Mr. Hankey (Canada)

167. Sir, if you would recall, the question I brought you to, or sought to bring you to, because you do keep on moving around - you’re very agile - was your premise that even if we can’t control exposures in the manufacturing industry, so let’s forget about construction sites for a minute. As I understand it, the example you referred to was not a study. You simply found a violation, that is to say an excess of exposure limits in a single manufacturing facility. Do you have any notion or any data concerning the health effects of that high exposure?

Dr. Infante

168. Do you mean that particular exposure?

Mr. Hankey (Canada)

169. Yes, that particular exposure.

Dr. Infante

170. I don’t think anyone could answer that particular question. They were above a permissible exposure limit that is already considered by the United States to present a significant risk of health hazard. The day that the compliance officer was there, they were exposed above the permissible limit which is 0.1 fibre per c.c. and you’re asking me what are the health consequences of that day of exposure. Well, I don’t think anyone can answer the question to that. Canada is arguing controlled use and my point is that that’s something to aim for, but because you aim for, or have a policy, doesn’t mean that it gets implemented. I’m giving that as one example.

Chairman

171. I’ll just interrupt you for a moment. As we’re taking an ad verbatim transcript here, it’s probably better if we go through the Chair so that I can then announce clearly who is speaking each time. So I give the floor to Mr. Hankey.

Mr. Hankey (Canada)

172. Thank you Sir. Insofar as the risks of exposures in the friction products industry, I cited you considerable data earlier which you did not contest. These data indicate no excess risk of lung cancer or mesothelioma to workers in the friction manufacture industry as compared to the general population. Speaking of, for example, Berry and Newhouse, McDonald, Teta, Teschke, ....

Dr. Infante

173. Can I respond? I thought that our discussion earlier had to do with asbestos-cement production, not friction products.

Mr. Hankey (Canada)

174. Am I mistaken? Did you not raise the issue of exposure levels in a facility that was manufacturing friction products. Am I mistaken?

Dr. Infante

175. Just now I did, yes. But earlier you said I didn’t challenge something you had on friction products and my point is I was responding earlier to your comments on asbestos-cement production and that’s why I cited the Hughes study. That’s a study in asbestos-cement production. I wasn’t talking about friction products earlier. I just now gave that as
an example where an inspection was made and the company was above the permissible exposure limit and had other violations as well. I commented on friction materials studies.

Mr. Hankey (Canada)

176. Perhaps could I then go through the friction material studies, Mr. Chairman, with Dr. Infante. Because I did recite them all before lunch but perhaps he was not engaged in that particular discussion, I don’t recall. I could certainly lead him through the evidence and we could see whether he’s familiar with it and concurs with it or differs with it because he has raised, Sir, the issue of manufacturing of friction products. I thought we had demonstrated conclusively that there is no excess risk of disease from the manufacture of chrysotile-containing friction products.

Chairman

177. I think I would, on that particular point just raised, give the opportunity to any of the experts who wish to make a brief comment. It does seem to me that we’ve spent a considerable amount of time on this particular point. It would be, from the Panel’s point of view, useful to move as much as possible toward the various issues concerning controlled use. But let us perhaps invite a brief comment from the experts and if Canada still wants to come back on that one you may. Dr. Infante, please.

Dr. Infante

178. If I look at the document 203, on page 109 and table 23, they list several studies on friction materials production. The study overall by Newhouse and Sullivan does not show any excess like the SMR is 93, the study by McDonald et al. 94, (we’re talking lung cancer now), shows a statistically significant excess. Then there are mixed products in friction materials, several of those, in fact all of them, show a significant excess of lung cancer. Granted that these are mixed products, but nevertheless they show an excess and you can’t totally discount, in my opinion, the chrysotile contribution. The study by McDonald et al. in 1984 shows a significant excess in lung cancer and the majority of that excess, not all, was in short-term workers. That’s noteworthy in that study. So you say, well what does that relate to? I think you have to know something about the short-term workers to know why you have the excess in short-term workers, it’s not the first study: workers exposed to beryllium, they were exposed for a short term, show a significant excess of lung cancer and it was all initially in the short-term workers. We know that beryllium is a human lung carcinogen. What the study shows in terms of dose-response is that there is not much potency. One of the problems in doing dose-response in the study is that you have this excess in the short-term workers. So you are not going to expect in that study to find a dose-response because presumably the short-term workers had low exposure and that was the majority of individuals in the study. You are not going to expect to be able to find a dose-response and there is not a lot of statistical power in that study when you go beyond the short-term workers. You can have a U-shaped curve in terms of the dose-response; you have a high risk in the low-exposed group, you have a slightly lower risk in the medium and you have a high risk in the highest exposed group. I don’t know what you can say about dose-response in that study given that you have got some kind of observation that you need to try and understand, in my opinion, before you do dose-response.

Chairman

179. Would any of the other experts want to add anything to the point made by Dr. Infante? Dr. de Klerk.

Dr. de Klerk

180. Can I just semi-respond to Corbett McDonald’s points earlier on. I just think that they need some kind of reponse because his basic conclusion, I thought, was that
because the textile industries were different, because they had higher risks of lung cancer, then we should ignore them in terms of setting health standards. I suggest that’s not really the way you should go about setting health standards. The one thing they all have in common is they’re textiles but they are also chrysotile. Therefore, as Professor Henderson has been saying, in terms of setting prudent health policies, if you’ve got some evidence that a substance is dangerous and then it’s going to be used by a lot of people where the properties are unknown and, I thought we’d agreed earlier on that the majority of people we are concerned about are not friction product manufacturers, asbestos-cement manufacturers, we’re worried about the people using the products later on and we don’t know what characterizes their exposure, only that they will be exposed to chrysotile. We have some evidence that chrysotile is dangerous. We have a lot of evidence that it’s dangerous and that we are not in a position to control that exposure. So, to say that we should ignore evidence that it is dangerous, I think is imprudent at best.

Chairman

181. Thank you. Professor Henderson.

Dr. Henderson

182. In relation to my colleague, Dr. de Klerk’s, observations, I would have to agree with him. I was struck in Professor McDonald’s comments that he pointed to the consistency of the high lung cancer risk among textile cohorts. He also indicated that the explanation for this difference between the textile workers and other groups of workers still awaits elucidation. We have no clear explanation for this difference. In the absence of something which we cannot explain and therefore take measures to control, prudence should lead us to take the position of maximal caution because we don’t know that the extremely low risk of lung cancer found in the Quebec chrysotile miners and millers will be translated across other cohorts. In this respect, it’s what I said in one of my earlier reports that when in doubt, or there are uncertainties or lack of observational data in comparison with cohorts, one adopts a principle of “first do no harm” or when in doubt play it safe for the setting of national occupational health policy. I was also heartened to hear Professor McDonald basically say that there is a modest risk of lung cancer at low levels, that he did endorse the linear relationship model and he did state that the explanation for these differences is not clearly known. Because of these uncertainties concerning risk, I would adopt the same policy as Dr. de Klerk and argue that one takes a conservative scenario in order to avoid a risk of harm – here we’re talking about cancers with close to a 100 per cent mortality rate – for the benefits of the average population.

Chairman

183. Thank you. Dr. Infante wanted to come back on a point.

Dr. Infante

184. I wanted to comment on what Dr. McDonald had said earlier. I think his point was that why would one rely on Dement’s study or the other studies of chrysotile-textile workers when the results seem so different from the results from the miners and millers study that he conducted. He also indicated that the Hughes study of asbestos-cement production workers gave a slope closer to the slope of the miners and millers study. Is that . . . , you are shaking your head? Yes, that’s right. But as I look at the data from the Hughes study, it gives a slope closer to the textile workers study and he just said that the cement production would be closer to miners and millers. When you look at the slope from the Hughes study, if you look on page 168 of that study, they indicate that for the chrysotile group only the slope, this is per unit of fibre, the slope is 0.01 for the chrysotile group and 0.016 for the mixed fibre groups. So it appears to me that that slope is closer; that’s close to what the slope is from the study of the textile workers based on McDonald’s Pennsylvania cohort and the Rochdale study done by Peto which is about 1 per cent, and it’s a little lower than that based on the Dement study which shows 2 to 3 per cent.
185. Professor McDonald.

Dr. McDonald (Canada)

I would like to say that the slope in the textiles is of the order of 0.1. The slope in the Hughes chrysotile plant was 0.0003. That is indeed approximately similar to the Quebec chrysotile miners and millers. We agree entirely that the textile plants are out of line with that by an approximately fifty-fold difference. All I can say is that the Quebec plant is not isolated. The thing that is isolated are the textile workers. The Quebec miners and millers are similar to the cohorts of the chrysotile cement workers and similar to the cohorts of friction product workers. Indeed, there are only something like eight studies where anybody has measured the exposure at all. And seven of the eight agree with the miners and millers and only the textile workers don’t. I would agree that if we had to decide about the continuation of textile work, we would be absolutely right to say let’s be careful about it. But that’s seems to me a rather historical question.

Dr. Infante

I just have one point of clarification. The risk of 0.0003 that you indicated for asbestos-cement production according to the document 203, that is the potency estimate for use at plant 1 which was chrysotile, crocidolite and amosite. The risk level for plant 2, which I understand is chrysotile only, was 0.007 and so that’s 0.7 per cent.

Dr. McDonald (Canada)

It’s the other way round, but I’d say we really ought to discuss this somewhere else.

Chairman

Well, could I suggest that we should now try to focus ourselves solidly on controlled use, given that most of the discussion so far in this afternoon session has tended to follow on from really the same issues as the morning session. As I say, if we have time at the end of our meeting after we’ve managed to deal with both controlled use and some aspects of substitute fibres, maybe we can come back and continue some of the discussion covering the first four broad questions of the Panel. Are the parties ready now to address issues specifically concerning controlled-use? I think probably the floor is to Canada for the next main question.

Mr. Hankey (Canada)

Sir, my first question has to do with the construction industry. You may regard it as preambular to the issue of controlled use because it really, I think, gives rise to what kind of a controlled-use may be appropriate in that industry. I refer to the 1980 paper by Rödelsperger et al., entitled Estimation of Exposure to Asbestos Cement Dust on Building Sites. In that paper the office observed that for past uncontrolled use/uncontrolled conditions, exposure levels reached 10 fibres per mm. during the sawing, cutting and grinding of chrysotile cement sheets and calculated time-weighted average exposure levels of 0.6 to 1.2 fibres per millilitre for the installation operation. Because such operations occurred only one day out of six the resulting average exposures were 0.1 to 0.2 fibres per millimetre, which is one or two orders of magnitude, that means up to one hundred times, lower than the exposure levels for past mining, milling, asbestos-cement and friction product workers. I’m wondering what you make of this data and its relevance to the subject before us since
it is data from the very sector we’re talking about i.e. the use of cement products in the construction industry.

Chairman

192. I must say it’s seems to be a question of exposure rather than controlled use, but given that we have had a question already from the European Communities on controlled use, perhaps we could ask the experts to give us a brief response after which I will again pass the floor to the European Communities.

Dr. de Klerk

193. I thought that in that paper the levels went up to as high as a 100 and 120, as I recall, and not 10. The averages are based on specific jobs and if you are getting levels of a 120 next to somebody cutting an asbestos sheet, it depends on the structure of your job how much you would get over a week and over a year, it just happened that the average was over that particular job. There would be other jobs where you would be doing that all day long, I would have thought.

Chairman

194. Canada please, Mr. Hankey.

Mr. Hankey (Canada)

195. Perhaps I’m in error but I have the impression that, generally speaking, in the construction industry exposures would tend to be intermittent and therefore it was the accumulative factor of peak exposures which was the relevant measure of what would constitute risk. Perhaps I’m not right about that.

Dr. de Klerk

196. If you’re putting up asbestos-cement fences, you would be exposed to that kind of level all the time.

Mr. Hankey (Canada)

197. If you’re putting up asbestos-cement fences as a sort of a full-time occupation, wouldn’t that be a little like working in a nuclear field? If indeed you understand it to be a highly dangerous job, wouldn’t it be the sort of job in which presumably controlled-use should, ought to be, I would hope, is enforced and properly administered.

Dr. de Klerk

198. Yes, you would hope so in theory, but it’s the kind of thing that doesn’t happen in practice and that’s I think one of the crucial issues: there’s been asbestos regulations in place for over a hundred years and there is ample evidence that in very few places have those regulations ever been adhered to by the people using it.

Chairman

199. Any additional comments from the experts? Professor Henderson.

Dr. Henderson

200. Again I noticed the estimate of the peak airborne fibre concentration cited from the Rödelsperger paper and again my recollection was that the peak concentrations were up to 100 fibres per millilitre of air, so it was a stated underestimate. Yes, one would hope that the use of these products in the building construction industry, in particular [could be “controlled” by best work practices or, alternatively, the use of chrysotile restricted to a
few special applications, analogous to nuclear fuels, but even in this latter situation] … [END OF TAPE] … a recent episode in Tokaimura, Japan, indicated that not even that is achievable. But the problem we have in Australia, in particular, with asbestos-cement building products is that they are so widely distributed in dwellings and buildings throughout the country, so that the largest group of mesotheliomas and lung cancers that I see related to asbestos comes not from the Wittenoom cohort - which although the exposures were high and the risks of mesotheliomas were high, was a relatively small workforce - the greatest number of mesotheliomas that I see comes from carpenters who give a history that day in and day out they cut asbestos-cement building products with handsaws, power saws, they used power sanders, they used angle grinders, electric drills and the like. We know that all of those operations can produce substantial elevations of the airborne fibre concentrations. If we are going to use mesothelioma as an index of exposure, the fact that we have such a large number of mesotheliomas among carpenters and building construction workers indicates that exposure did occur. Now, certainly many of those workers, perhaps the majority of them, also sustained exposure to the amphiboles. But here I’m using mesothelioma simply as an index to a marker for the fact that significant exposure did occur. The simple fact is that, among the many, many cases of mesothelioma that I see, a consistent theme amongst the workers is that they were not told by the employers that the materials they were dealing with were dangerous, there were never any airborne fibre concentrations measured in their working environment, only late in history were they provided with face masks, usually in the form of a surgical paper mask or a plastic mask, and we know that even more substantial respiratory protections are sometimes ineffective. So that, from my perspective in Australia, historically, we have never seen controlled use of asbestos and the very fact that no measurements or estimates of the risk were carried out indicates that controlled use has not been in place historically in Australia and so far as I am aware, it still isn’t. In fact, it was dealt with by phasing out chrysotile from asbestos-cement building products in 1987 or 1989 so that they are no longer used in this particular application. In this respect I’d have to harp back to the WHO document Environmental Health Criteria 203, which indicated that construction workers pose particular concerns because of the large and diverse nature of the workforce so that it is very difficult to disseminate information to all the individuals concerned in these types of operation. That document indicated that chrysotile use in that situation is not recommended.

Chairman

201. Thank you. Dr. Musk wanted to comment.

Dr. Musk

202. I’d just like to reinforce that. We’ve been arguing about which is the best sort of model of exposure in industry where it has been measured what the exposures are. But in the construction industry it hasn’t been measured and can’t be measured regularly, so it isn’t really controllable.

Chairman

203. Thank you. I give the floor now to the European Communities.

Mr. Christoforou (European Communities)

204. Thank you Mr. Chairman. I would request the four experts, if they can take a minute, to have a look at page 28 and page 29 of Canada’s comments of 13 December. Page 28 please.234 This is a document dated December 13th, called “Canada’s Comments on the Experts’ Responses to the Questions from the Panel”.

Mr. Christoforou (European Communities)

205. Page 28, and especially paragraph 6, where there are four bullet points which go over to page 29, where Canada describes what is in its view the so-called controlled use.
Canada, I would like to remind you in case you have not read all the documentation, has been changing position constantly since we started this dispute about what is controlled use and progressively moves and tries to restrict more and more what in its view is controlled use it has in mind. Now, I would like to request you to read these four bullet points and would appreciate if you could tell me if this type of situation described here, that is: to distribute products only to companies licensed to purchase these products; those companies must have workers trained and licensed to install products and must be in compliance with regulations; approved users shall not resell to third parties and any unused material must be returned to the manufacturer; to provide a list of users of products to the responsible government agency; to provide products cut to specification at established centres equipped to cut the products to size and where persons cutting the products are trained and are licensed to work with asbestos; and, fourth point, to police the downstream users in cooperation with the government; the product manufacturer visits, monitors and reports on the performance of the downstream users at regular intervals. There are penalties for failing to provide this product stewardship. The question is, from your own experience in dealing with these questions in your profession, do you think this is a feasible and realistic scenario taking into account the type of population exposed as you have defined it previously? Thank you.

Chairman

206. Thank you. Let’s give the experts a moment to decide who might want to respond first on that point or whether you want to take up aspects of it, as it’s quite a broad issue, individually. Dr. Infante.

Dr. Infante

207. I feel that this stewardship programme, when I read this, I feel that it’s not a reality; it’s a possibility but it’s unlikely and definitely not likely to occur in construction. With regard to point 6 about controlled use, that “this permit will be withdrawn if the company does not meet the following commitments”, what went through my mind when I read that was: withdrawn by whom? Who enforces this? The first bullet point about “those companies must have workers trained and licensed to install the product”, well who oversees that training? It’s not clear to me who would do that in countries that would be working with the asbestos? And bullet point 3: “to provide products cut to specification”. I think that’s good to do that but then there are always adjustments that have to be made, so even though products may be cut to specification, there are places where you have to trim or the pipe or something is too long and you have to make some adjustments, and the concern is when those adjustments are made, that proper precautions aren’t taken. Then, in the last bullet there are penalties for failing to provide this product’s stewardship. As I read this, I wondered what are these penalties and how many have been issued to date. This, to me, seems good in theory but it doesn’t seem real to me. Then when I just recently read an article about asbestos, chrysotile-asbestos exposure in Morocco which imports Canadian chrysotile and I see these photographs in this article just published this year – I have a copy of the article – and it shows that asbestos is just all over the place. So I’m wondering if the Canadian Government, if it has this partnership for a sustainable development, why are there countries like Morocco, Brazil and India that seem not to be following what’s required by this stewardship and the controlled use?

Chairman

208. Thank you. I think perhaps any further comments from the experts before we get into discussion on this item. Dr. Musk, please.

Dr. Musk

209. This sort of regulation would require a new system for enforcement which hasn’t previously existed anywhere that I know of. Secondly, it doesn’t take into account people working with products that are already installed, modifying and installing pipes, electricians, plumbers and the like. So it certainly wouldn’t cover all the opportunities for exposure.
Chairman

210. Professor Henderson, please.

Dr. Henderson

211. I’d have to agree with my two Panel colleagues that, as I’ve indicated, so far as I’m aware, controlled use for the stewardship-type of arrangement has never been used in Australia in relation to asbestos products of any type. As I’ve also indicated we don’t really have detailed dust measurements in almost all workplaces including asbestos manufacture; or where they have been done, their count seems to be artificially low in comparison to the fibre count seen in the lung tissue of the workers. So that historically in Australia I cannot see that this has ever been applied and as Dr. Musk said I don’t think that it is enforceable in law. It would require a whole new infrastructure in industry and legislation to bring into effect. Just as a common sense observation, as far as I can see for a products manufacturer to police the after-sales uses of its products would introduce a new dimension in Australia. I’m mindful, for example, of the fact that automobile manufacturers who sell cars, yes, they may sell them only to people who hold a driver’s license, and the government authorities do have a list of the license holders and the registration numbers; but for an automobile manufacturer to try and police dangerous driving, excessive speed or driving under the influence of alcohol, to monitor drivers on the road and then report them to the police, would produce an entire new dimension into Australian society at least. It’s one that I would think would create an immediate conflict of interest between sales and profitability on the one hand, and the policing and regulatory function on the other. But I think it’s fine in principle, but I suspect that’s it’s unworkable in practice in Australia, at least unenforceable at law.

Chairman

212. Dr. de Klerk, do you want to add anything?

Dr. de Klerk

213. I was just curious as to whether there was any sort of precedent for the system that they put into that document. I can’t imagine anything like that working anywhere with anything. But presumably there may be some precedent somewhere for that kind of system?

Chairman

214. Thank you. Does either party or Panel members want to comment on those responses? OK. If that’s not the case, then perhaps I could give the floor to Canada for their next question.

Mr. Hankey (Canada)

215. You all obviously have some doubts about the efficacy of controlled use: I guess it’s chiefly among construction workers, although perhaps your remarks aren’t entirely limited to that sector. But, I wonder which of the following aspects of control do you consider key to safeguarding health of construction workers using high-density chrysotile products? There are chrysotile products, low density chrysotile products in place, so evidently a certain amount of due care is required by people in the construction industry. I’m wondering what measures you would consider to be particularly necessary. I’ll list a number and perhaps you could indicate whether you think they are key, whether you think they are useful to making controls work or work better. We could start with hazard risk and risk assessment. Do you think that proper hazard and risk assessment assist in safeguarding health of construction workers using high-density chrysotile products?
Chairman

216. Would it help if you listed all of them?

Mr. Hankey (Canada)

217. I’m quite happy to do that. I’ll go slowly though because usually what happens when I read one of these long sentences, I’m asked to repeat it so I was going to take it one by one: Hazard/risk assessment; information; education; training of workers; registration of tradesmen; hazard control; personal protection; licensing for specific potential dangerous risks; sale of products to registered users only; and finally the point which has, I think, just been mentioned: removal of license to purchase chrysotile and chrysotile products if users are not in compliance with regulations. I don’t know which of you feel that you have expertise in this area of industrial hygiene, please feel free to answer the question.

Chairman

218. Let’s give the experts a moment or two to think about that and let’s decide who might want to answer.

Dr. Infante

219. Could you just quickly go over. I missed a couple of them as I was writing them down.

Mr. Hankey (Canada)

220. Sorry, I have hazard/risk assessment; information, education, training; registration of tradesmen; hazard control; personal protection; licensing for specific potential dangerous tasks; sale of products to registered users only; and finally, removal of license to purchase chrysotile or chrysotile products if not in compliance with regulations. And my question is, which of these following aspects of control do you consider key to safeguarding the health of construction workers using high-density chrysotile products? Perhaps I should also frame it slightly differently, that would make a material difference, a significant difference.

Chairman

221. Dr. de Klerk?

Dr. de Klerk

222. I can make a few points. It’s probably outside my area of expertise most of the latter ones. The only thing that I would consider myself vaguely expert in is in terms of risk assessment. It’s already been mentioned that, in fact, we don’t have risk assessment information for most of the downstream users, so obviously that’s important. I’ve been involved in studies where we’ve tried, using information in education and training - this is outside the asbestos area, this is to prevent accidents in industry and sometimes it works and sometimes it doesn’t. The fourth one, hazard control, obviously if you reduce the exposure you reduce the risk of disease, I assume that’s what that means. The thing that strikes me about all of them is the fact that, in essence, all of these steps were part of the asbestos regulations, certainly in force in Australia, at say, for example, the Wittenoom mine and mill and they weren’t really of any help at all in preventing disease occurring from there. There was risk assessment in the sense that people knew that heavy exposures caused asbestosis but those levels weren’t kept. There was information provided on the notice board that the mine was a registered mine, there were attempts made to reduce the dust but they didn’t reduce it, they just spread it around; people were encouraged to use face masks but in the heat they couldn’t wear them; the mine had a license which the government was supposed to supervise and it didn’t. When they broke the rules it didn’t
remove the license. So it’s an example of where, although you’ve got something in theory that should work, in practice it won’t.

Chairman

223. Do other experts want to add to that response? Dr. Infante, please.

Dr. Infante

224. I would agree that they would all be helpful, assuming that the hazards/risk assessments have already been done or we wouldn’t be here today. As far as information, education and training, yes, that’s important; registration of tradesmen, that’s important; hazard control, of course these are all important; personal protective equipment is important. They are all important but some of the problems are, you have personal protective equipment, what does that mean? Let’s take respirators, for example, when do you wear a respirator? Our standard requires a competent person who has to know about where asbestos may be, whether or not the product may contain asbestos. It’s not simply having a respirator available, but do you have a respirator fit-testing programme to assure that the worker who wears a respirator is getting the protection they should have; do you have a programme that cleans the respirator? Do you have different types of respirators that are available depending on what the exposures might be? So a respirator programme requires a fair amount of training in itself and knowledge on the part of a competent person. Then, one of the problems is that in the United States there is a tendency not to train short-term workers in the construction sector because it costs to train workers and you know they’re only going to be there short-term and they’re going to be moving on to another job where there isn’t asbestos exposure. Since they’re going to be gone shortly there’s a tendency to try to save money and not to train workers that would be there for a short period of time. So, all of these are good: the problem is implementing such a programme in reality, I think is difficult.

Dr. Henderson

225. Again, I would reinforce the comments from my two colleagues. In Australia, the use of respirators in the building construction or any other industry poses particular problems, despite penalties, in the form of fines, and even three breaches of the regulations and the worker is dismissed. The simple fact is that compliance is poor because in a hot, dry environment, where temperatures regularly go over 30°C and sometimes above 40°C, the thermal consequences of wearing a respirator create such discomfort to the worker that they will often discard the respirator, irrespective of penalties of work without them and so will their fellow work mates. In relation to regulation of the various practices outlined, the simple fact is that government regulatory agencies in Australia, increasingly have a diminishing capacity to regulate work hazards. For example, after the Conservative Government was elected in Australia, the National Occupational Health and Safety Commission was downsized and approximately one half of its workforce was made redundant, so they no longer have the capacity to supervise all points of end-use of asbestos or any other product at all times. As I’ve said, the building construction industry is of particular concern, simply because of the spectrum of different occupations represented in that group who have asbestos-related diseases and many of these individuals simply go straight into the building construction industry with minimal training or no training: they simply leave school and suddenly appear as an unskilled worker in the building industry and acquire their training on-site. Those who employ them are often individuals or very small businesses which do not themselves have a background and depth to provide training in the correct application of safe work practices. So, we’re dealing basically with a very large, diverse, often poorly trained workforce who have a very poor appreciation of the risks to which they’re exposed and a common theme amongst the cases I see is that the worker didn’t really know that it was asbestos or if he did, he didn’t know it was dangerous, didn’t know that the operations he was carrying out would in fact generate dangerous levels of airborne dust, and therefore the individuals are unaware of the risks they’ve been running. In many of the cases I see, for example mesothelioma, we really have to use the
tumour as an index of exposure in order to uncover some pattern of asbestos-exposure which even the worker has been unaware of and I cited a couple of examples in the supplementary remarks to my report. The other point I would emphasize is that, from my perspective, controls are most certain when there is a minimization of the total amount of asbestos introduced into society, into the workplace and into the general environment. And if you don’t introduce any more, then hopefully, provided you can try and implement reasonably safe work practices, you can minimize exposures to those products that remain, but their total amount will diminish over time. But one must recognize that because of the diversity of this group, that the training programmes will not always be followed, there may be poor worker compliance, and that many of the programmes are not always effective in any case.

Chairman

226. Thank you. Dr. Musk, would you like to add something?

Dr. Musk

227. Once again, this sort of programme would ignore the people handling asbestos that is already in situ. I think it would certainly act as a deterrent to using asbestos at all because it would be pretty unwieldy to implement, if it was implemented properly, so people would probably look for other products to substitute but it does ignore the asbestos that’s already there.

Chairman

228. European Communities, please.

Mr. Christoforou (European Communities)

229. If I can continue with a follow-up question on this point. One can then legitimately pause for a moment and ask the question: all these requirements indicated by Canada, all these steps one has to go through, where do they come from? How did Canada come up with this list of steps to go through before applying controlled use? And I would like to ask the experts, because I see Dr Infante says on page 17 of his replies that he is not aware of any international standard that would prescribe a controlled use, let alone a controlled use in the sense of containing all the steps indicated by Canada. So where do they come, all these requirements? Is there any international standard that would require such steps? Thank you.

Chairman

230. Thank you. On the point of the existence or otherwise of an international standard, Dr. de Klerk, first.

Dr. de Klerk

231. If I have understood correctly, and if this is the same question I asked earlier about whether there was any precedent for such a system, is that what you’re saying. Because I don’t know of one and that’s why I was asking the Canadians.

Chairman

232. Do other experts wish to address the question just raised by Mr. Christoforou? Dr. Musk, please.

Dr. Musk

233. I’d be interested to hear Canada’s response to the question because I don’t know where they came from and I’m not aware of them existing elsewhere.
Chairman
234. Does Canada wish to comment on that?

Mr. Hankey (Canada)
235. No. I have a question for Dr. Musk actually.

Chairman
236. I think perhaps Mr. Christoforou would like to clarify his question.

Mr. Christoforou (European Communities)
237. Mr. Chairman, this is partly scientific in the strict sense and partly relevant for the discussions we’ll have later on. But I ask this question because of the express sentence in the replies of Dr. Infante on page 17 in the fifth middle paragraph. Just to indicate that Canada has been portraying until now that one ILO convention, International Labor Organization Convention 162, prescribes something which can be applied and achieve controlled use, which will achieve a level of exposure below a level threshold that is not dangerous. Just to say that the type of controlled use, just now indicated by Canada, exists nowhere and I am glad to see that the scientists confirm they have no knowledge whatsoever of any of this type of controlled use applied anywhere.

Chairman
238. I’ll give the floor to Mr. Hankey to ask the question you were going to ask Dr. Musk.

Mr. Hankey (Canada)
239. I just want to clarify your last statement but one. You said something about, in response to my question about which of these aspects of control would be key to safeguarding the health of construction workers, you said something about in-place asbestos. Could you just repeat that? I want to fully understand the import of your remark.

Dr. Musk
240. I was saying, I don’t think any of the measures addresses the handling of asbestos that’s already in place to the extent that, if the measures were put into place, there would still be asbestos going into construction. That asbestos would then be there for plumbers, electricians and anyone else coming along later. It doesn’t address their exposure.

Mr. Hankey (Canada)
241. What, Sir, would you propose to do about asbestos already in place?

Dr. Musk
242. I think, as a general principle, one needs to minimize the exposure to it and work practices are important in that area. Once it’s there, as far as I’m concerned, it ought to stay there until there’s a good reason to remove it. And then when one does remove it, it should be removed with due care.

Mr. Hankey
243. If I understand you correctly, you would say that asbestos which is already in place should be left there and that due care should be taken with its use. By due care, could controlled use be another way of expressing the same idea or not?
Dr. Musk

244. The notion of controlled use I interpret from these measures relates to providing new asbestos products for the construction industry, not to protecting workers against asbestos that is already in place.

Mr. Hankey (Canada)

245. What measures would you propose? What kind of due care measures, perhaps you wouldn’t call them controlled use, I don’t know what appellation you would give them, but what kind of measures would you propose to deal with asbestos already in place? Because Sir, I don’t know what the situation is in Australia, but I do assure you that in France, which is the country that’s at issue here, there are vast amounts of asbestos in place, including very great amounts of low-density asbestos, much of it of mixed fibre, although we have some dispute with the European Union as to how much of it is mixed fibre, certainly a substantial amount of it is. It’s incontestable, as a matter of social history, that the very issues that gave rise to the ban that’s currently in place, which is the very subject of this dispute, is in-place asbestos, old uses, high density, low density asbestos in places like Jussieu and there are vast amounts of in place in France, so the issue of how to deal with that in-place asbestos strikes me as extremely relevant. So I would like to know, Sir, you seem to think that these measures that I have proposed or put out for comment are not applicable to in-place … and you also propose that it should be not removed. Now I think everyone would know that it’s a real and present danger, these old uses of asbestos, vast amounts of which still exist in France, how would you propose to deal with it if you were a policy maker?

Dr. Musk

246. I’m not a policy maker and this isn’t my area of expertise, but I would say that when the time comes that it’s required to be removed in buildings where it’s past its use-by date, or the insulation is deteriorating or the asbestos-cement products are cracked and broken, the roofs and there’s a lot of asbestos-cement roofs, where I come from, have deteriorated to the extent they’re not doing their job, then the people permitted to remove them need to be policed to use methods for removal that will not expose the worker. There are in Australia licensed asbestos removers and they are required to have air-supply respirators and they do the major jobs for removing asbestos from buildings. But the most exposed people are the small businesses or the handyman who does it himself and nobody gets to know that it’s happened till it’s passed. So it’s relatively unregulated.

Chairman

247. Thank you. That’s a last point before we break for coffee.

Mr. Hankey (Canada)

248. I am very glad to hear that in Australia you are able to exercise control, it seems, when necessary to remove this stuff. I’m really, though, very interested in what goes on when the stuff is there because it may not be removed, I don’t know, for twenty, thirty or forty years, you haven’t given me any indication but you say you’re not involved in the business of policy-making, but fortunately we have at least a couple of people on the Panel of experts who are … Mr. Henderson, for example, in his paper, in his summary of conclusions, prescribes indeed the remedy which this Panel should provide in this case, so he is clearly in the business of making policy, or at least he has a very great interest in it. And I wonder, Sir, what remedy you might propose, relative to the vast amounts of asbestos, including very much low density products of asbestos in place in France, much of it containing mixed fibres. What would you do about it, Sir?
Chairman

249. I’ll give Professor Henderson the opportunity to respond to this point, and then we will have a coffee break of 15 minutes. Professor Henderson.

Dr. Henderson

250. The question is based on a false premise. I’m not involved in setting public policy on this issue: this is done by others, and particularly, the National Occupational Health and Safety Commission. My comments on the disposal of existing asbestos products in place are similar to those of Dr. Musk. I think that some of the procedures that you’ve outlined should be implemented, as a matter of common sense, to try and minimize exposures to existing products. As Dr. Musk says, there are licensed asbestos removal organizations in Australia, which are meant to carry out these operations under controlled conditions and at minimal risk to the asbestos-removal workers and to the general public. However, just in the last six months, I’ve come across two mesotheliomas that have been a direct consequence of asbestos-removal programmes because it appears that those procedures were not followed. One of them was a fireman who was regularly called to buildings which had been incinerated by fire and where fire alarms were set off by high airborne dust fibre concentration as a result of asbestos-removal programmes. This fireman visited these buildings at least once a month to check them through and was, we believe, exposed to elevated airborne fibre concentrations. Another one concerned a university lecturer who for a period of weeks had to walk to and fro through a building where an asbestos-removal programme was being carried out. Although the removalist was supposed to encapsulate the material and seal it in polythene bags, it appears that they left it lying on the ground in an unprotected state and this person, the lecturer, walked past this asbestos material quite regularly over a period of some weeks. So, I agree that best work practices should be aimed at, in order to try and minimize exposures, but my concern is one of caution and prudence, to realize that not everybody is going to implement these procedures at maximal efficiency all the time and that exposures will occur. I’d agree with Dr. Musk that probably the best thing to do with existing asbestos in place is to encapsulate it until such time as the building is demolished or unless it can be shown that elevated airborne fibre concentrations exist in the building, and again I’ve got other mesotheliomas which have occurred simply from individuals who worked in department stores where there is friable asbestos insulation with elevated airborne fibre concentrations. So, I think you need to balance the risks of removal against the risks of the asbestos continuing in place until the time of demolition. But implementation of best work practices should minimize exposures but ultimately exposures will be best minimized when there is no new introduction of asbestos materials into the workplace where they can remain for 20, 30 or 40 years and be subject to periodic and sometimes regular maintenance and renovations.

Chairman

251. Thank you Professor Henderson. We’ll obviously be continuing the discussion on controlled use after the break, so we’ll have a coffee break now, and return at 16h50.

[Coffee break]

Chairman

252. We left for coffee just after Dr. Henderson has responded to a question from Canada and that in itself was part of the discussion on the question Mr. Hankey initially raised regarding the various precautionary measures to take in the case of prevention of exposure to asbestos. Could I ask if there are further comments or follow-up questions on the remarks by Professor Henderson? Canada, please.

Mr. Hankey (Canada)

253. In the overriding interest of not missing my evening cocktail, I was going to desist from further discussion on controlled use. So, if we are moving on the substitutes, I would
have no more to say. But if we are going to be staying with controlled use, then, yes, I would have ...

Chairman

254. The Panel’s view was that we could perhaps continue for another ten to fifteen minutes on controlled use, if you wish, and, by 5.10-15, we should be into the discussion on substitute fibres. Mr. Christoforou, you would like the floor as well?

Mr. Christoforou (European Communities)

255. Yes, I would like to ask one more question on controlled use.

Chairman

256. Yes, please. We’ll continue the discussion for another ten to fifteen minutes, and then we’ll move to substitute fibres.

Mr. Christoforou (European Communities)

257. Thank you. The question is addressed to all scientists. It is prompted by a comment made by Drs. Infante and Henderson, that controlled use is even much more difficult to be applied in non-occupational circumstances. I think it is an obvious statement, but I would like you to comment on this and say if you know whether the equipment suggested to be used (the mask and all the other equipment) and the proceedings to follow, will always, constantly, achieve, in occupational and non-occupational circumstances, a level of exposure which is below 0.1 per cent of fibre per mml. Do you think that it will always be achieved below that threshold? Thank you.

Chairman

258. Dr. Infante, please.

Dr. Infante

259. The point of my written comments was that, I don’t think controlled use is likely to occur in the occupational setting and so that in the non-occupational circumstances, it would be even much more difficult, because there is no…, you don’t really have the potential here for training that you do with the occupational setting, or even the construction sector, where training quite often doesn’t take place nor any of the other programmes related to controlled use and exposure to asbestos. If you are asking, well, in non-occupational circumstances, could fibres exposure exceed 0.1 fibre per ml, it depends on what the individual would be doing. I would says, yes, that’s possible to occur. For example, I know that with using a Transite, which is an asbestos-cement product, when individuals tear this off a wall, it usually breaks apart, because it’s either nailed or screwed on, and it’s much faster to just simply pull it off the wall and quite often, it will pull the nails out with it. So, it is a lot faster to remove it that way. But quite often, as is usually the case, it breaks into pieces and when that occurs, you can generate fibre levels above 0.1. That’s one example. It just depends on what’s being done in a non-occupational circumstance and what product is being manipulated.

Chairman

260. If there is no further comment on that point, I’ll pass the floor to Mr. Hankey.

Mr. Hankey (Canada)

261. My question is in relation to the products that we are discussing, in particular those used in construction, that is to say chrysotile-cement products, do you consider the greatest
risk to be at the point of installation, or maintenance – that is to say interventions after it’s put in, by electricians, carpenters, plumbers and so on and so forth – or its demolition and removal? Where, at which point would you consider the risk to be greatest? And I wonder if each of you could answer that question. Thank you.

Chairman

262. Dr. de Klerk first.

Dr. de Klerk

263. It depends on the exposure, really. I know we said it before but it’s obviously where the greatest amount of dust is generated, it’s going to be the process that gives the greatest risk in terms of existing measurements made around operations that are available – obviously demolition and removal have the highest exposure levels. But in some ways the people in demolition and removal may experience less exposure because there is more of a likelihood that precautions will be taken. … [END OF TAPE] … But again, even within this sort of full-face respirators there is a measurable level of asbestos found. So obviously they are all at risk and it depends how well the operations are done. I don’t think you can be sort of make hard and fast rules about it, but I mean historically people installing have not taken precautions and have experienced great risks and as we can see from them being the group with very high levels of mesothelioma on all registers. Maintenance, you’ve got the people who again tend not to historically take any precautions, and again there are groups of people, plumbers, electricians who, using Professor Henderson’s point about using mesothelioma as an index of exposure, there are people there who have obviously been exposed. Obviously, though, I’d like to say that they are all at risk in one form or another and it depends on the level of exposure and the precautions that are taken.

Dr. Infante

264. I don’t think that, as a general statement, you can say that one has a greater risk than the other. I think they all carry, you know, a great risk depending on how the installation, or the maintenance or the demolition is carried out. That is what relates to the fibre exposure.

Dr. Henderson

265. Again I’d agree with the comments from my two colleagues. I see cases of mesothelioma related to all of these types of activity using mesothelioma as an index-marker of exposure. Again, it’s my belief that the risks will be dependent on the frequency of the operation, the types of operation carried out, the airborne fibre concentrations generated and the duration or the type of work. I see mesotheliomas resulting from all of these activities, for example, among carpenters, and for example the handyman who regularly carries out maintenance and renovations on houses, where he might use a power saw to cut a new doorway through an asbestos-cement clad wall, will generate fibre concentrations equivalent to the carpenter carrying out this type of work day after day. It’s just the frequency with which he does this type of operation, may be less. The same can also apply to demolition, particularly of small dwellings, if precautions are not carried out during building demolition and disposal of the asbestos-cement product. So I’d have to say that I couldn’t give a figure for the risks to each of these groups because they would vary according to the variables I’ve already mentioned, but I do see cases of mesothelioma resulting from all of these types of activity.

Chairman

266. Thank you. Dr. Musk, would you wish to add anything to those three comments?
Dr. Musk

267. I’d agree with the three previous speakers. I’d suggest that people involved with maintenance, being the least regulated group, and least easily regulated group may be at greater risk but like Dr. Henderson I see cases of mesothelioma from people involved in all those activities.

Chairman

268. Thank you. Mr. Hankey.

Mr. Hankey (Canada)

269. Thank you. If I could just try to make a synopsis of what I just heard. I think each of you said essentially, although Dr. Henderson’s answer was I think more complex than the others, but certainly each of you said really it all depends on what precautions are taken. Dr. de Klerk said precisely that, and as did Dr. Infante, Dr. Henderson did say that but along with a number of other things, and finally, Dr. Musk said exactly that and added to that he thought that perhaps maintenance was perhaps the biggest problem because it was the most unregulated. So, if you understand you correctly, then the issue at each point, that is to say installation, maintenance - and by maintenance I mean interventions once it is already there by tradesmen such as plumbers, carpenters and electricians and so on and so forth. And then the removal -you consider you can’t distinguish between these risks, you say it all depends on what precautions are taken at each point. That’s what each of you said. Now, I’m wondering still if we could come back to this problem about the asbestos in place because we all recognize - and I don’t think there is any issue about this – that the asbestos in place, if you like, fibre for fibre and man for man in terms of the exposure to it represents still the greatest risk. I concede that we don’t know what the risk will be perhaps 100 years or 200 years from now, that’s another question. But currently, I recall, Dr. Henderson said early this morning that, when I asked which he thought was the greatest risk, he indicated, if I understand correctly, that, yes indeed, the greatest risk from an exposure at a given level, or for the same amount of exposure time, I think was really the point, but you can correct me if I’ve got it wrong, to low-density products which may contain mixed fibres. You thought that would be greater - sort of intervention for intervention - than interventions in these high-density chrysotile-only products. You said you had difficulty calculating the overall risk because indeed, you felt there were more interventions; more people were perhaps coming into contact with chrysotile-cement products than with these old kinds of products. Is that a correct statement, Sir, of what you have said this morning? I haven’t finished my question, but I’m basing it partly on what you’ve already said. I want to make sure that I’ve got that right.

Dr. Henderson

270. Well that is not quite correct. What I was trying to say this morning is that the risks of lung cancer and mesothelioma will be dependent on the type of operation carried out, and therefore the airborne fibre concentration, the frequencies with which those operations are carried out, and their durations – that you are looking at a risk related to cumulative exposure levels; and the point that I was trying to make this morning was that, if you take a cohort, for example, the Wittenoom cohort in Western Australia, those individuals have a very high risk of mesothelioma and yet, the cohort, which numbered about 7000 individuals, was relatively small. Although, if you are looking then at a lower risk in a larger group of workers, for example, carpenters, because there are many, many more carpenters in Australian society than Wittenoom workers, then the total number of mesotheliomas you will see in this larger group at lower risk will be equivalent to those you see from the Wittenoom cohort or even larger in terms of absolute numbers. When I took that figure I took the figure for carpenters only, but if you add in plumbers, plasterers, other building workers, it adds up to a very large group, and probably one of the largest groups represented in the Australian Mesothelioma Register.
Mr. Hankey (Canada)

271. I’m a little sceptical about that thesis, because it is a bit like saying, if each of us here, if we have a keg of beer that is brought in and each of us gets a glass of it, that’s as much risk as if I drink the whole amount and then go out and drive my car. I rather expect the authorities who are regulating drinking and driving probably wouldn’t concur with that approach to the matter. In any case, let me proceed with the real point of my question. If it all depends really on what precautions you take and if the dangers are the same at installation and interventions when it is installed and then removal, I still don’t think I have received from any of you any kind of satisfactory answer to what we do about the in-place old uses of asbestos which, I still have to insist that if we look at the social commentary on why France introduced the ban, the ban was introduced precisely in order to remedy those problems. That, or at least I shouldn’t say precisely, is certainly what led to the political pressure to introduce the ban, and a study of the French media at the time definitely would prove that; so, if controlled use doesn’t work for these new products which, I must say I think most commentators would agree, are less, product per product, dangerous than the old ones and the low density ones that contain mixed fibres and amphiboles, … what is the political, the social remedy to that stuff in place, if indeed controls don’t work because when I ask you what happens to remove any type of product, whether it is the old one or new one, you tell me that really it all depends on what precautions you take, which seems to me that you are saying it all depends on what controlled-use mechanisms are put in place. So I am still at a loss as to what we are going to do about this huge danger facing society with the in-place old products.

Chairman

272. I’ll give the experts the opportunity to respond on this point and then I think we need to move on to substitute fibres and then immediately after one or other of the experts has responded on this current question, I’ll give the floor to the European Communities. Or did you want to make a point, Mr. Christoforou?

Mr. Christoforou (European Communities)

273. I would like to hear the follow-up question after I hear the replies of the experts on this point, Mr. Chairman, please.

Chairman

274. OK. Fair enough. You may do so. Professor Henderson first, please.

Dr. Henderson

275. Well, in reply to my comment about the workers at risk, I can only reiterate my comment, it is not so much on the controls in place, although hopefully by disseminating information one can try and implement best work practices to minimize exposures to those products that remain in place. When you disputed the estimates I gave for a lower risk among carpenters in comparison to the Wittenoom cohort producing a larger aggregate number of mesotheliomas, your doubts are not supported by the figures from the 1999 Report for the Australian Mesothelioma Register, which records, among carpenters and joiners, 187 mesotheliomas due to single exposures only, 33 additional mesotheliomas from workers with multiple exposures, making a total of 220 cases. Whereas the Wittenoom cohort accounted for 189 mesotheliomas (single exposure) and an additional 25 (multiple exposures), making 214 cases. So, although the risk of mesothelioma is high in the Wittenoom cohort and among non-smoking survivors, mesothelioma is now the most common cause of death, the numbers in aggregate are slightly less than the number of mesotheliomas in absolute numbers we see among carpenters, simply because – although the carpenters are at lower risk – there are many many more carpenters in Australian society than there were Wittenoom workers. So, that low risk needs to be multiplied against a larger population. That is the point that I was making.
276. As for the problem of asbestos in place, I agree entirely that this is a major problem. What do we do about the asbestos which is in place, and how do we minimize exposures? Some of the strategies that you’ve indicated, in terms of informing people, trying to implement these best work practices, will hopefully minimize the exposures but so far as I am concerned this is an ongoing problem for which we have no easy solution, taking into account that many of the people who carry out interventions on those products, by way of building maintenance and renovation, are almost completely unregulated. Although it is very regrettable, despite our best efforts, I believe that we are going to continue to see mesotheliomas from that type of exposure. But having pointed out the difficulties of minimizing exposure to asbestos in place, that does not by itself, from my perspective, represent a justification for the introduction of more asbestos into the environment whereby the total quantity will become greater and the scope for people to be exposed, even at lower levels, will be translated into an ongoing population over time.

Chairman

277. Thank you, Professor Henderson. I will give the floor briefly to Mr. Christoforou for the follow-up question he wanted to ask. Could I ask that you do make it brief and hopefully the reply could be brief so that we don’t lose any more time before getting on to the substitute fibre questions. Thank you.

Mr. Christoforou (European Communities)

278. Mr. Chairman, I renounce to ask the question because the reply of Dr. Henderson covered my point. Thank you

Chairman

279. Well, in that case, I would give the floor to the European Communities, if they wish to ask a question concerning substitute fibres.

Mr. Christoforou (European Communities)

280. Yes, Mr. Chairman, thank you. We would like to request all experts to elaborate on your replies concerning alternatives products which are non-fibrous and whether, in their knowledge and experience, such non-fibrous alternative products have been classified as proven human carcinogens, as is the case with chrysotile asbestos. I highlight the word non-fibrous alternative products.

Chairman

281. Yes. Dr. de Klerk.

Dr. de Klerk

282. I’d just like to answer fairly briefly. The question, as it was asked before, was really asking about alternative fibres but when you look at non-fibrous products, as far as I am aware, anyway, it is the fibre quality of asbestos that makes it dangerous and if you’ve got a product that isn’t fibrous then it doesn’t have those qualities and therefore is unlikely to be risky in that same kind of way.

Chairman

283. Thank you. Question six did concern substitute fibres. It was not specifically asked about non-fibrous substitutes. If there are no further comments on that point, could I now pass the floor to Canada on the fibrous substitutes issue.
Mr. Hankey (Canada)

284. You may indeed. I mean, I do have a comment about that question but perhaps if you rule the question at the border, perhaps I need not comment.

Chairman

285. Well I think, as I see it, the issue that was concerning the Panel was the question of fibrous substitutes particularly.

Mr. Hankey (Canada)

286. My question is to any of the experts who really cares to answer, but I’d perhaps suggest that I would like Dr. Infante, among others, to answer because I believe he has considerable expertise in this area. Basically my question is that “Do you agree that the information base regarding human exposure to substitutes is meagre compared to what we know about chrysotile?”

Chairman

287. Dr. Infante.

Dr. Infante

288. I think that compared to what we know about chrysotile asbestos, the data on most toxic substances is meagre in comparison.

Mr. Hankey (Canada)

289. I just wonder then if Dr. Henderson and Dr. de Klerk and Dr. Musk would agree with that statement.

Chairman

290. Dr. Henderson please.

Dr. Henderson

291. I would agree with that statement in broad terms. So far as I am aware, except for a couple of large cohort studies on man-made mineral fibres there are virtually no epidemiological investigations of human populations for the majority of the substitute fibrous materials. Evaluation of their effects is based basically on fibre characteristics and experimental models.

Chairman

292. Dr. Infante would like to add something.

Dr. Infante

293. I just want to elaborate. I think that for workers exposed to fibreglass, there has been considerable epidemiological study, but there has not been epidemiological study to my knowledge with the polyvinyl alcohol, the para-aramid fibres or refractory ceramic fibres. But there is experimental data for those substances and I think I mentioned earlier today what some of the findings were.

Chairman

294. Thank you. Any additional comments from the experts or further follow-up from the parties? Mr. Christoforou, please.
Mr. Christoforou (European Communities)

295. Mr. Chairman, this is a question that addresses partly the previous question on non-fibres and also on this question on fibres raised by Canada. I would like to request the experts whether, in their view, when asbestos-containing products are replaced in their use are they normally replaced in the majority, if not exclusively, or can they be replaced almost exclusively in their use by products which are non-fibrous? If you allow me to somehow rephrase the question: the European Communities have been arguing that there are - and I can give the example of cast-iron pipe, high-density polyurethane pipe, concrete pipe, metal roofing sheets, clay roofing tiles, plasterboards and so on, which can substitute asbestos contained in products in nearly all of its uses. Are you aware of this fact? Thank you.

Chairman

296. Could I just perhaps reiterate that we did not specifically ask the experts to address the questions of non-fibrous substitutes. The interest of the Panel in the scientific aspects of this were especially concerning the qualities, properties of fibrous substitutes. I could perhaps invite the parties and the experts to concentrate as far as possible on the specific issues that were asked under Question 6, which is really concerning the fibrous substitutes.

Mr. Christoforou (European Communities)

297. Mr. Chairman, with due respect, we don’t think this is the situation. Question 6 refers to both fibrous and non-fibrous and we would suggest that it is even more relevant, because, as we suggest here and as we have been making in our submissions, there are numerous non-fibrous products which can substitute asbestos for nearly all of its uses. So the question is very relevant to see the magnitude of the problem, of whether there is a problem posed by fibres, which will come later on.

Chairman

298. Having re-read the question very carefully, I can say there were one or two references to non-fibrous substitutes. I would invite the experts to respond on that point.

Dr. de Klerk

299. I’ll just chip in a couple of points. In terms of, in Australia anyway, I mean I haven’t really looked into this because I sort of assumed it was fibrous but, for asbestos-cement the main manufacturer uses cellulose instead of asbestos. I think in brakes it’s para-amid fibres, so that in fact, as a general rule, most of the substitutes are fibrous, well certainly in Australia. I would also like to add that most of the comments that I made in terms of this, because it is probably outside my area of expertise in a way, were based on a good review by Harrison et al., which I think everyone has probably read. I think that sort of summarizes the extent of knowledge at this time. I haven’t found anyone who disagreed with that at all.

Chairman

300. Thank you. Any amplification or further comment? Professor Henderson?

Dr. Henderson

301. Well again, like Dr. de Klerk, I focused on fibrous substitutes because so far as we know the agents implicated in the causation of mesothelioma are almost always fibrous materials namely, amphibole asbestos, chrysotile asbestos, or the naturally occurring mineral erionite. There are some concerns about refractory ceramic fibres; I am not aware that there are any data in humans but there are some experimental models which cause reason for concern. So when we are dealing with mesothelioma I think we are dealing with
substitute fibrous materials as opposed to non-fibrous materials. Of course, the non-fibrous materials may have toxic effects which are different, but so far as we know they are not implicated in mesothelioma induction; so I focused my answer on the substitute fibres like my colleague Dr. de Klerk.

Chairman

302. Thank you. Perhaps if there is no further follow-up on that one I could now give the floor to Canada for further question or comment on fibrous substitutes.

Mr. Hankey (Canada)

303. Yes. My next question is: do you believe that fibres used as substitutes for chrysotile in cement and friction products, for example, glass fibres, cellulose fibres, para-aramid fibres, PVA and RCFs, such as potassium octotininate should be used without controls? Perhaps Dr. Infante, you could start, and I would like the others to answer as well.

Dr. Infante

304. If you could perhaps refine your question. What do you mean if they can be used without controls. What do you mean by that?

Mr. Hankey (Canada)

305. Well, for example, would you suggest that workers who are installing them or removing materials made with, that contain these substances, any of them, should work without masks, for example, that they should saw it with high-speed saws. That would be two questions. I would have to really, I’m afraid, ask my experts to propose other answers, or help me formulate other questions. I suppose – I may be wrong – that for each of these they present somewhat different risks, and that therefore the measures you would impose would perhaps be different for each of them. Another thing might be exposure limits for example, would you say there would be a need for exposure limits for any of the materials I’ve indicated, and if so, which ones?

Chairman

306. Dr. Infante, are you able to answer on the basis of that?

Dr. Infante

307. I think, as a matter of industrial hygiene you should reduce exposures to the extent that you can in the occupational setting. Now by saying that that, doesn’t mean that these fibres carry the same risk as chrysotile. I don’t think that any of them do, but as a matter of proper industrial hygiene, we should try to reduce exposure levels or use good work practices. You can get some of these things perhaps in your eyes, from sawing them, so perhaps you would want to wear goggles, for example. I always think you should handle substances in the workplace appropriately. Should you be concerned about the same risk of exposure to these substitute fibres, as you should be concerned about asbestos fibres? I guess what I would say is that I don’t see the evidence that these fibres are as harmful; but yes, you should try to control them to the extent that you can.

308. You have to look at what some of the information is here. If you look at refractory ceramic fibres, for example, I think that they are hazardous, and that, if you are working with these fibres, yes, you should take precautions with them and you should wear appropriate protective equipment if you are exposed to these. But it is my understanding that the refractory ceramic fibres would not be a substitute for chrysotile on any large basis. That does not mean that they are not toxic. Is there evidence that they are carcinogenic
in humans? – No. But there is evidence in experimental animals, and on the basis of that I would take all of the precautions that I could. With the polyvinyl alcohol fibres, there have been some implantation studies that have been conducted on experimental animals and IARC concluded that there is insufficient evidence of carcinogenicity for those fibres. It is my understanding that their size is such that with a large diameter, it is unlikely that they would be respirable. So I think that it is good that that’s the case. So I don’t think there would be much biopersistence then if they are not able to get into the lungs. With the para-aramid fibres there has been, I believe, inhalation study and intra-peritoneal injection studies that IARC reviewed and they concluded that there was no evidence of carcinogenicity for the para-aramid fibrils. In terms of biopersistence, I think that I cited the study by Searl that indicated that these fibres greater than five microns in length are less biopersistent than chrysotile fibres greater than five microns in length. The para-aramid fibres, it is my understanding, are like somewhere between ten and twelve microns in diameter, so they would not be in the respirable range. However, there is the potential for some fibrils to break off from these fibres that are smaller. What hazard there is from these has not been studied in humans, but on the basis of the para-aramid fibrils from experimental studies, IARC concluded that there is no evidence of carcinogenicity. I am concerned about any exposure, but when you talk about what’s the potential disease risk I think you are dealing with a known factor with chrysotile asbestos; the studies that have been done don’t indicate any cancer response for these. So if it were me, I would, if I were in occupational health, I would prefer to see para-aramid fibres substituted for chrysotile in the appropriate applications. With cellulose fibres, they have not been studied in experimental animals or in humans. With glass fibres, in my opinion, there is evidence in experimental animals that these are carcinogenic. I think it is more likely than not that respirable glass fibres may be carcinogenic to humans, it is more likely than not. Does that mean it has been proven? – No it doesn’t. But I would exercise caution with those. And what I am talking about here is, you know, there may be a risk of lung cancer in humans. I have not seen any information that mesothelioma is associated with glass fibres – and I think I mentioned that in my report. As I had mentioned earlier, I, at one time, thought that this high risk in the Canadian study was specifically related to low fibres on the basis of the data that were available. But I have other information now about that, the exposure to those cohort members to other known human carcinogens, including asbestos. So I’ve qualified my comments about the potency of glass fibres compared to chrysotile asbestos. I mean at least we have standards for these in the United States, as nuisance dusts, which is like a fifteen mg. per cubit metre limit. So we do have some regulation of them, and I suppose as you have, if you have more information available that indicates that there should be better control of these and if they are not being controlled – and I think you need to implement that knowledge. But exposure to fibreglass in the United States, at least in manufacturing, has always been quite low. Going back to the 1940s, I think average exposures are 0.04 fibres per millilitre and that has always been low exposure. These fibres are used for blown-in construction and there fibres can get up to – I think the highest levels I ever saw was 7 fibres per ml, that’s the peak, the highest I’ve ever seen. Usually they are below, certainly below 1 fibre but I feel that these glass fibres aren’t as potent as chrysotile asbestos. And I have already commented on the ceramic fibres.

Chairman

309. Thank you Dr. Infante. Any point which the other three members would like to add?

Dr. Musk

310. I might just add to that that any particulate would be considered a nuisance dust unless it has specific properties and that is because, partly at least, of the possibility of occupational airway disease, so-called industrial bronchitis and airway narrowing. I mean that’s an entity that seems non-specific just related to particulate content irrespective of the nature of the particulate.
Chairman

311. Thank you. Are there any further comments to add. Yes, Professor Henderson.

Dr. Henderson

312. Again, my comments would closely mirror those of my colleagues. From the point of the potential carcinogenicity of substitute fibres, as indicated in my report and my supplementary remarks, the key factors seem to be the dimensions of the fibres, their persistence in lung tissue, and in various studies, their capacity to cause disease. Based on the review by Harrison and a number of the articles submitted, including annexes from Canada, and the recent press release from the Health and Safety Executive in the United Kingdom, there seems to be a growing body of opinion that the substitute fibres are safer in general, with the exceptions already indicated by Dr. Infante. Importantly that they are less biopersistent in lung tissue, so that presumably their capacity for carcinogenesis is proportionately less than chrysotile.

Chairman

313. Mr. Christoforou, please.

Mr. Christoforou (European Communities)

314. Will you allow me a follow-up question on this point?

Chairman

315. Yes. Go ahead.

Mr. Christoforou (European Communities)

316. The follow-up is what Dr. Henderson said, with a few exceptions mentioned by his colleague, and I think he referred to the statement by Dr. Infante. Dr. Infante has identified ceramic fibres and glass fibres as possible, probably, dangerous substitutes. The question I would like to ask is the following: I don’t know if you know of any country which has banned asbestos from use – all uses of asbestos – and it has substituted by glass fibres entirely all previous uses in which asbestos was used and employed. In other words, I wish Dr. Infante to expand on what he said on a large basis. Is it really true that these suspected – these two possibly suspected products – the glass fibres and the ceramic fibres, are a realistic substitute for all uses made of asbestos previously? Is there any country who has? Is there any knowledge about this? Can we really argue, as Canada is implying, that these are possibly dangerous and so because they are too dangerous, we should not ban asbestos? Thank you.

Chairman

317. Thank you. Dr. Infante.

Dr. Infante

318. No, I wasn’t implying that they were the substitutes. We were asked to address the toxicity of a number of fibres and I stated my view on the fact, I said that I didn’t – from what I understand it – you know ... refractory ceramic fibres are limited to very special high heat applications and it would not be a general substitute for chrysotile – certainly not with a chrysotile-cement products. It is my understanding that I didn’t realize that fibreglass was put into asbestos-cement, but I think you would have to ask someone else about that, it was more like the polyvinyl alcohol fibres that I believe that I had read in some submission might be one of the substitutes; and if that is the case, that would probably be good since they are of such dimension that they are unlikely to be respirable. Now knowing the toxicity
of them, as the data that are available for the polyvinyl alcohol fibres don’t indicate any carcinogenic response. But I’m saying that you don’t have to worry about that if they don’t get into the lungs at all, whether or not they might be.

Chairman

319. Thank you. Canada, please.

Mr. Hankey (Canada)

320. I have a follow-up question.

Chairman

321. Certainly, please.

Mr. Hankey (Canada)

322. Dr. Henderson, when you said it all depends on the characteristics of a fibre – and I think you were speaking of any fibre, whether it is a natural fibre like chrysotile or an artificial or man-made fibre. Would that be the case that these elements that you identify would be the same for any fibre as being the criteria by which you would determine its carcinogenicity?

Chairman

323. Professor Henderson.

Dr. Henderson

324. In broad terms the answer is yes, and the characteristic that I would focus on is the dose to which the individuals are exposed, the dimensions - are the fibres similar in character to those of chrysotile or amphibole fibres? - and the biopersistence of those fibres in tissues. Finally, in experimental systems, have those fibres shown a carcinogenic effect or not? They would be the four key parameters on which I would base assessment of substitute fibres.

Mr. Hankey (Canada)

325. I gather two of these factors have to do with the quality of the fibre itself, that’s to say dimension and biopersistence. They have something to do with the way the fibre is made, whether it is made naturally by nature or whether it is made by man. The other two - it was dose and you said - what was the fourth - I’m sorry?

Dr. Henderson

326. Dose, dimension, and durability.

Mr. Hankey (Canada)

327. Certainly, dimension and durability would be objective criteria by which any fibre could be measured. And presumably chrysotile is used for certain purposes precisely because it possesses certain characteristics of dimension and biopersistence. Or was biopersistence the correct idea? I mean certainly that it lasts - I presume - that it has a certain durability. Now if that is the case - perhaps it isn’t - but let me finish my question and you can demolish it at any point of the logic you see fit. If that’s the case, then won’t manufacturers be rather inclined to create artificial man-made fibres, substitute fibres as it were, that have similar characteristics to chrysotile; if indeed they tend to use it for the same purposes; in friction products and/or in cement products.
Chairman

328. Thank you. Professor Henderson will respond on that point.

Dr. Henderson

329. That really is an engineering question which is starting to fall outside my area of expertise. My understanding is that chrysotile fibres are chrysotile fibres, and that you can’t start engineering them to be of vastly different dimensions from what they already are, whereas for some of the substitute fibres they can be engineered or reproduced in such a way, that the fibres are either not respirable or that they do not have the dimensions that are normally associated with carcinogenesis from asbestos. That is the only point that I would make. I would suggest caution about any substitute fibre which had fibre characteristics which were very similar, for example, to those of the amphiboles and refractory ceramic fibres would be one example. I would treat refractory ceramic fibres with great caution but my understanding is that from the other features that I have mentioned, the fibres are either less biopersistent than chrysotile, or they have different fibre dimensions, or they have not been shown to cause cancers in experimental animals.

Chairman

330. Thank you. Mr. Hankey please.

Mr. Hankey (Canada)

331. You did say just now that artificial fibres can be engineered to produce more or less whatever characteristics the manufacturer wishes, including characteristics that would be similar to chrysotile. I think it is not a stretch to think that, if chrysotile is useful because it possesses such and such characteristics, dimension, length, endurance, there will be a certain incentive on the part of manufacturers to produce similar products, artificial products, to perform similar functions. That being the case, I wonder if you could tell me what controls are in place in Australia to ensure that any new fibres put on the market by manufacturers, or used within an entity, a manufacturing entity - you know, produced by one branch of a corporation and used in another branch of the same corporation - what controls are in place in Australia to ensure that any new fibres brought, created, engineered to replace chrysotile are not carcinogens and generally not as dangerous as chrysotile?

Chairman

332. Professor Henderson.

Dr. Henderson

333. Again, the point that I would make is that your concerns seem to be that the substitute fibres might be manufactured to have fibre properties similar to those of chrysotile. My perception is that there would be a pressure on a manufacturer to produce a substitute material with similar thermal characteristics and stability in the environment – but with fibre characteristics which are clearly dissimilar. It is a matter of deciding on - if you like - the properties of the material to be used versus the fibre characteristics. This is really an engineering question that falls outside my area of expertise and I really cannot comment on any of the engineering processes. In Australia, with the exception of fibreglass, as far as I am aware, the substitute materials are imported rather than manufactured on site; but certainly I do know that the National Health Authorities have recommended substitution of chrysotile in virtually all uses, provided that the alternative material is not more harmful and clearly is not as effective, and there may be certain restrictions for which chrysotile can still be used. For example, there is still some production of brake linings and asbestos-containing gaskets in Australia; but the simple fact is that in new automobiles, manufacturers have largely replaced the use of chrysotile by substitute materials. Now apart from this, I don’t know what particular controls have been implemented in Australia and what recommendations … [END OF TAPE]
Mr. Hankey (Canada)

334. I was just wondering, and if I could paraphrase your answer, it seems to be that you are not aware of any controls that are in place to ensure that substitute fibres are not carcinogenic, or otherwise dangerous to human health, but that you think that manufacturers would be decent enough not to produce them. That was more or less what I got from your answer. Would that be a fair restatement of it?

Chairman

335. Professor Henderson.

Dr. Henderson

336. It is not quite correct. The National Occupational Health and Safety Commission has concluded, largely on the basis of overseas investigations, that the substitute fibres are safer, and that there is some allowance for importation of small amounts of chrysotile into Australia, in the order of about a thousand tonnes per year, for the use of production of chrysotile-containing friction products materials and chrysotile-containing gaskets. But certainly, the Occupational Health and Safety Commission has recommended that there should be no introduction of any new material, that a material that has been already replaced by chrysotile should not in future be replaced by a chrysotile-containing material. Certainly the vehicle manufacturers in Australia, with one or two exceptions for older vehicles, have replaced the use of chrysotile in those vehicles. This is monitored to some extent by the National Occupational Health and Safety Commission, but the details of its controls, if there are controls in place, are not known to me.

Mr. Hankey (Canada)

337. Then if I understand, your answer is that the substitutes being used are believed to be more safe than chrysotile, based on experiments carried out and their use in foreign countries. I think that was what you said.

Chairman

338. Professor Henderson.

Dr. Henderson

339. Yes, this is contained in the document I refer to continually as NICNAS99, where the National Occupational Health and Safety Commission has concluded on the basis of overseas expert bodies that the substitute fibres are safer than chrysotile, and for that reason they have recommended that chrysotile be phased out over the shortest time interval possible, its use for the remaining periods of time be restricted only to a few applications, and that substitute fibres be implemented. I would add though, that if one is talking about, if you like, fibres released from brake linings of passing vehicles, apart from the use of those chrysotile-containing materials, to the best of my knowledge there are no controls in the general environment so that the substitute fibres are treated in that respect no differently from those that still contain chrysotile.

Chairman

340. Thank you. As time is moving on, perhaps I could just explain how we would tend to conclude the meeting. The meeting was supposed to finish at 6.00 p.m. It is possible that we can go on to 6.15 p.m. I do want to give the opportunity, but not the obligation, to the experts or any of the experts who wish to make any concluding comments. After they have done that, I would like to address one or two procedural questions, including the question raised by Canada earlier this morning about the two pages of comments submitted by Dr. Infante. So I think there would be time for perhaps one further point or question
from either of the parties. Or maybe one quick one from each of the parties if you have a burning need to ask one or two more questions. Mr. Christoforou.

Mr. Christoforou (European Communities)

341. Mr. Chairman, I would like, as a last point on the issue of substitutes ... I am sure the scientists know that for example the Environmental Health Criteria 203 has recommended the substitution of products of asbestos, all types of asbestos, by other products because they are safer. On the basis of the existing knowledge of which you are aware, with the possible exception of tile fibres and glass fibres, do you think the substitutes which are used are safer than asbestos-containing products?

Chairman

342. I think Dr. Infante will respond.

Dr. Infante

343. There is no evidence that they are harmful. We are talking about the polyvinyl alcohol, para-aramid fibres and fibrils, and cellulose fibres. There is no evidence that any of those potential substitutes are carcinogenic – there is no information at all on that. As scientists and as people involved in public health, we do exercise caution in using these fibrous materials. That is different than saying that they have met the same standard of toxicity as asbestos fibres because they haven’t. We just exercise caution. But I would, you know, recommend that, as the document says, chrysotile fibres certainly be substituted for.

Chairman

344. Thank you. It’s a good place to give the opportunity to Canada for a last comment or question.

Mr. Hankey (Canada)

345. I just have one short comment and then a question, and my comment is simply …, and it really relates to the debate I was having with Dr. Henderson. You kept referring to the “substitutes” for chrysotile as if those substitutes were a fixed universe. Now it is my impression, I may be wrong, but they are not in fact a fixed universe, that, given the fact that chrysotile has been banned rather recently in many jurisdictions, new products indeed, new substitutes are invented and come on to the market from time to time, and what I was really asking you was whether you were aware of any controls in place to assess and ensure the safety of these new products, before they were put on the market. I understood your answer to be “no”. Now, I’d like now to move on to my last question and it is simply this: a November 1999 report of an independent committee organized by INSERM238 states that “No significant excess risk of cancer has been ever detected from exposures to asbestos at the same exposure levels used to evaluate the carcinogenicity of substitutes.” Now if you like I could read from the original French but I believe that’s the best translation we can give to that. So would you wish that I read it in the French language or not? Well that is my question. My question is do you agree with this statement? And perhaps I might start with Dr. de Klerk.

Chairman

346. Well I think we will leave it to the experts as to who wants to answer first. But could I ask the experts to answer this one briefly so that we do have time to go on to any concluding comments you wish to make.
Mr. Hankey (Canada)

347. I would Sir, ask each of the experts to answer because each of the experts did indicate in one way or another that they thought that substitutes were probably safer than asbestos, than chrysotile. So I think it is a fair question.

Chairman

348. So we’ll have a first response and then leave it to the others as to how they indicate their views. Dr. Infante.

Dr. Infante

349. Excuse me. I just wondered, rather than repeating it in French could you just repeat it in English? I want to make sure that I understand the question.

Mr. Hankey (Canada)

350. Yes, of course. It says the INSERM Committee stated that no significant excess risk of cancer has ever been detected from exposures to asbestos at the same exposure levels used to evaluate the carcinogenicity of substitutes. That is the INSERM … A voluminous report they have just published on the issue of substitutes.

Chairman

351. Dr. Infante. Have you managed to absorb the question? Or maybe anyone else, I could give the floor to anyone.

Dr. Infante

352. Can you repeat it? No significant increased risk of cancer from exposure to asbestos …

Mr. Hankey (Canada)

353. … has ever been detected at the same exposure levels used to evaluate the carcinogenicity – I have a lot of problems with that word - of substitutes. Je peux le dire en français.

Chairman

354. While Dr. Infante is pondering his response, perhaps I could hear a comment from Mr. Christoforou. But we really do need to be brief here.

Mr. Christoforou (European Communities)

355. Yes, Mr. Chairman. Thank you. While the scientists are thinking, we have here the author of this report and he can probably put this citation into context because he has written this phrase and he can explain what it is meant and then the scientists give their reply.

Chairman

356. Well, I am happy for the gentleman to do that provided he too can be brief.

Mr. Christoforou (European Communities)

357. Yes, he will be very brief.
Chairman

358. Then we will ask a brief response from the experts.

Dr. Goldberg (European Communities)

359. Merci Monsieur le Président. Je suis Marcel Goldberg et je suis effectivement un des auteurs de ce rapport, et notamment, je suis le responsable de cette partie. Nous avons effectivement écrit la phrase qui a été citée, mais une fois de plus, je crois que la citation est extrait de son contexte. Il est vrai que nous avons écrit cela, mais c’est une discussion dans la partie qui traite uniquement des données épidémiologiques, et il faut rappeler que le rapport complet fait quelque chose comme 450 pages, et que nous avons pris en compte l’ensemble de toutes les données disponibles, y compris les données expérimentales, et que la conclusion de l’ensemble de tout nous a permis de conclure que, très vraisemblablement, le risque de cancer attaché à ce type de fibre était largement inférieur à celui du chrysotile. Merci.

Chairman

360. Thank you. I take it the translation has finished coming through. We will now ask the experts, do they wish to make any comment. Dr. de Klerk.

Dr. de Klerk

361. Does that mean, therefore, that the substitutes are at least as safe as chrysotile? Is that what you mean, is that why you asked the question? That therefore means that all the substitutes are at least as safe as chrysotile, is that what you are saying?

Mr. Hankey (Canada)

362. Yes, I think that could be a fair conclusion – yes. To the same level of exposure.

Chairman

363. Well I think that Professor Henderson wants to make a comment. I was about to conclude that the response from the experts had already been made, but please.

Dr. Henderson

364. I was a little bit surprised by the question as put because it didn’t distinguish between mesothelioma or lung cancer and amphibole versus chrysotile asbestos. But now that the translation has been given, clearly it refers to epidemiological investigations and I must admit I was a little bit surprised because animal experimental studies usually involve exposure to fibres of quite high levels - this is simply because the lifespan of an experimental animal is sufficiently short in comparison to the humans that you need to expose these animals to very high fibre concentrations or through a peculiar route whereby dust deposition in lung and translocation does not occur. That is you’d use either an implantation or a high-dose inhalation model. Again I’d would draw the same conclusion as Dr. de Klerk that the experimental investigations indicate that, if anything, the substitute fibres are likely to be safer than chrysotile and that even if one takes that question at face value, it indicates that none of them is more hazardous than chrysotile.

Chairman

365. Thank you. I think I would just ask the other experts if they want to indicate a view that differs in any way or adds in any way to the comment Professor Henderson has just made. Yes, Dr. Musk.
Dr. Musk

366. In practice, then, and I don’t know the answer to this, but one of the issues might be how easy it is to control exposure in the asbestos industry versus the substitute fibre industry.

Chairman

367. Thank you. Well it seems we have exhausted the comments. I would like to thank everybody very much for their participation in this meeting. I did say that I would offer the floor to the experts to give them the opportunity, if not the obligation, to make any concluding comments. We have about five minutes left so, and I would like a couple of minutes myself to deal with the issues that are raised concerning procedure. So I think I will quickly invite any of you who do wish to make any brief comment to do so, otherwise I will continue. Professor Henderson.

Dr. Henderson

368. One point that I would raise as a final issue which has not been covered in the proceedings today is the clearance half-life of chrysotile from lung tissue, because it is stated in a number of the submissions that chrysotile has an extraordinarily short half-life in lung tissue and I think a figure of 28 – 48 hours has been mentioned and a figure of less than ten days. When I read the figure of ten days I was reminded of a story I was told in my childhood of a wise man who performed a service for a Persian king, he was asked as his reward what he would like and he suggested a little thing, Sire, I’d like a grain of rice on the first square of a chess board and for you to double it on each successive square. Now if you go back to the paper by – I might add that he ended up owning all the rice in the kingdom – but if you go back to the study by Green and others on the Charleston textile workers at a median interval after sixteen years following cessation of exposure to asbestos, they still had a mean concentration of over 33 million fibres per gram dry lung tissue. If you then go back and say that the half-life is only ten days then you need to double that count for every ten days that you go back in time, 36.5 doublings per year for sixteen years

Mr. Hankey (Canada)

369. Excuse me, I would like to raise a point of order, Sir. I want to know, Sir, am I to have an opportunity to respond to the experts’ closing statements?

Chairman

370. These are closing statements and we cannot offer the opportunity to respond in this …

Mr. Hankey (Canada)

371. In that case, Chairman, will you please request that the experts not raise new issues in their closing statements. Dr. Henderson has just said that he is raising an issue which has not been discussed today. I think it is not really due process, if I might say so Sir, that the experts raise at the end of the day issues not discussed today to which I shall not have an opportunity to respond. So let’s either give the parties a fair opportunity to respond or else let’s keep the summaries to issues that have already been covered today.

Chairman

372. As I say, there is no room for any further debate on these issues today, but we are also under extreme time pressure which eliminates the possibility for raising any new issues. Can I just invite Professor Henderson to wrap up his remarks briefly?
Dr. Henderson

373. Well, this is not a new issue, it was covered in my Endnote to my original report and it was covered in the supplementary remarks I made. What I questioned is …

Mr. Hankey (Canada)

374. Point of order, Mr. Chairman. Shall I have an opportunity to respond to Professor Henderson’s Endnote?

Chairman

375. I think that … Excuse me. We can’t get into discussion at this point on whether something was or wasn’t a new issue and as the clock is also ticking, I think we noted the point that was made. I think I would like to ask Professor Henderson not to keep addressing this issue but to wrap up his conclusion in the next thirty seconds if he can.

Dr. Henderson

376. OK. I shan’t pursue this issue.

Mr. Christoforou (European Communities)

377. Sorry. I really object to this. The experts are free to express their views on what they have written in their reports. I don’t understand the objection of my colleague. There is no rule which allows the experts to express their views on what they have written in their report. If Canada didn’t feel necessary to raise this issue with … because the thing was clear.

Chairman

378. Mr. Christoforou, I have invited Professor Henderson to conclude his remarks in a manner that we can complete our work on time, and he is in the process of doing that. Please continue.

Dr. Henderson

379. Without pursuing the particular story further I would point out that I have already cited a paper by Finkelstein and Dufresne, published in 1999, which for the Quebec chrysotile miners and millers pointed to a half-life in human lung tissue of eight years for fibres greater than 10 micrometres in length, which suggests that chrysotile is far more biopersistent in tissues than many people realize. In fact we need to recognize that there is a short-term rapid clearance of chrysotile and then the remaining fibres that are deposited will remain persistent in lung tissue for a period of years thereafter – long enough for a carcinogenic effect.

Chairman

380. Thank you. Do any other of the experts wish to say anything briefly in conclusion? I say there is no obligation to do so and the shorter the better. Dr. Infante.

Dr. Infante

381. Yes. I would just like to sum up maybe where I began today. That is that, in my opinion, chrysotile asbestos is a very potent carcinogen, controlled use in my opinion is not realistic and the substitute fibres do not demonstrate – none of them demonstrate - carcinogenicity in humans and because of that I feel that in terms of public health it would be beneficial to substitute.
Chairman

382. Thank you, Dr. Musk.

Dr. Musk

383. I have nothing to add to that except to say that I agree with it.

Chairman

384. Dr. de Klerk.

Dr. de Klerk

385. I think I agree with that too.

Chairman

386. Well could I, on behalf of the Panel and the parties, thank our four expert very much, firstly for the very hard work that they put in before this meeting and their forbearance with us all during this meeting, being bombarded with questions and comments on these very complex issues. We are very confident that your work in the service of the Panel will be of great assistance, both to the parties and to the Panel members and to the Secretariat as we move towards the conclusion of our work. I did say that I would address at the end of the meeting the various procedural issues. Concerning the point raised by Canada earlier today following the couple of pages of comments which Dr. Infante circulated, the Panel discussed this in the lunchtime and I reiterate the distinction I made between the rules that were made for the parties and the arrangements which were made for the experts. The parties were given clear deadlines to submit material, comments on the experts’ reports. The experts themselves kept to their deadline for the submission of their own reports. We did not place any restriction on what the experts might do between all those submissions of reports and comments and what happened at this meeting. We certainly viewed the paper put together by Dr. Henderson, I think, it was dated 10 January, as he explained as a contribution to this meeting. I think the Panel would view Dr. Infante’s note in a similar light, as a contribution to this meeting, to the content of the discussion at this meeting. So it is not our intention to allow any further submission of evidence in relation to the papers submitted by, or the comments submitted by Dr. Infante, and we would note that Dr. Infante does not really depart at all or vary from the opinion that he expressed in his initial written reply. However, I would remind the parties that the purpose of having the gap of two days between this meeting concerning the scientific issues and the next formal meeting (the second formal meeting of the parties) was precisely so that the parties had time to comment, if you like, in the course of the second formal meeting on some of the discussion of the scientific issues that had taken place here. So, in the Panel’s view there was adequate opportunity for the parties to incorporate any comment that they wish in the course of the second formal meeting later this week.

387. Just also to set this proceeding a little bit in the broader context, I could just explain what will happen in the coming weeks. It is probably very familiar to the parties. As I said, following today’s meeting and the second substantive meeting which will take place on Thursday and Friday, the Panel will proceed to prepare its report. The first part of the report will be a summary of the facts of the case and the arguments of the parties, and will be provided in draft form to the parties for their comments. The responses of the experts to the Panel’s questions will also be included in the report. The experts will all receive a draft of the relevant section and will be given the opportunity to make any necessary corrections. Subsequently we need to provide a first and interim Panel report to the parties, including findings and conclusions. Then the parties have an opportunity to comment on that and we then submit a final report. As stated at the beginning of the meeting, there will be a verbatim transcript of today’s meeting which will be included as an annex to the final report, and both the parties and the experts will receive a draft of the
transcript of today’s proceedings for information and corrections as necessary, because the
draft is taken straight off the tape. So we would ask – this one final task – we would ask of
our experts, will be to check through the record of their remarks. So that, I hope, is a clear
explanation of how we intend to proceed. Mr. Hankey please.

Mr. Hankey (Canada)

388. Thank you Sir. I just wondered, Sir, if we might expect any further contributions
from the experts as part of their contributions to the meeting today or whether their
contribution is now definitively closed, apart from the checking of the record that you had
referred to.

Chairman

389. Thank you. Yes, as I said before the meeting started. We basically have a finite
resource, a finite time and it was as set out in our programme and schedule, that this is the
final point of the expert process with the exception of the checking of the record of the
transcript. So as of now the Panel’s expert consultation has been concluded. Yes, Mr.
Hankey.

Mr. Hankey (Canada)

390. When, Sir, might the transcript be available to the parties?

Chairman

391. I’ll ask the Secretariat to check that. It will be judged on how long it is going to
be.

Secretariat

392. Yes it’s technically rather long. I don’t expect the transcript to be ready before
mid-February, before the descriptive part is provided, if not slightly after. Not this week
anyway.

Chairman

393. Can I thank all the delegates, the parties and the experts, my own colleagues on
the Panel and the Secretariat staff very much once again for your cooperation and enabling
us to get through our work in the very short time. And I should also like to express a
special thank you to the interpreters who went on a little bit beyond the call of duty. Thank
you very much.
NOTES

1 See John A. Hoskins, *Chrysotile in the 21st Century*, (below “Hoskins”) and his bibliographic references 103, 105, 109 and 110. These bibliographic references are reproduced in full in other Canadian annexes as follows:

- Vanherle HE: In: Proceedings of the 8th biennial conference of the AIA, Paris, 11-12 May 1993, (reference 103 in Hoskins);
- Equitable Environmental Health Inc., *Dust Exposures during the Cutting and Machining of Asbestos/Cement Pipe - Additional Studies* (reference 105 in Hoskins);


7 Bernstein, *Correlation between Short Term Biopersistence and Chronic Toxicity Studies* (1997) Joint Research Centre, Environmental Institute, European Chemicals Bureau in Ispra (Italy).


9 That is why Dr. Julian Peto had to estimate the incidence of mesothelioma in Western Europe from data on mortality from cancer of the pleura, a very imperfect indicator of the incidence of mesothelioma.


11 Study by the Cancer Department of Health Canada (Schanzer, Semenciw and Ugnat, 1997) presented to the biennial conference of the Canadian Society for Epidemiology and Biostatistics.

12 Study on mesothelioma and cancers of the pleura and peritoneum for the British Columbia Cancer Registry (Coleman and Philips, 1996).

13 Study on mesothelioma and cancers of the pleura and peritoneum submitted to Health Canada’s Environmental Health Directorate (Camus, 1997).

14 Study of mesothelioma in Quebec from 1986 to 1993 and of pleural cancer and cancer of the peritoneum from 1986 to 1996 submitted to the Advisory Committee on Asbestos in the Ministry of Health and Social Services (Lebel, Gingras and Lévesque, Laval University Hospital Centre, 1998).

15 Our comparison with France reflects the fact that the statement of the European Communities incorrectly calculated the rate of growth for Canada as “a particularly high rate”. However, as shown on page 177 of the INSERM Report, the incidence of mesothelioma rose more rapidly in France than in Canada.


Source; EHC No. 203, Summary, 1.3.

Idem, Summary 1.4.

Idem, page 128.

Idem, Preamble.

Idem, Conclusions, b.


Idem., page 765.

Idem., page 760.


IPCS-EHC No. 151, 1993.


We recall that, where asbestos is concerned, the first findings of its harmful effects on health go back to 1906, as attested by the report of a French labour inspector who drew attention to the number of deaths among workers who had worked in factories making asbestos-based products.


At this point, we note that asbestos cement accounted for 90 per cent of the asbestos imported into France at the time of the ban. Thus, in accordance with INSERM’s recommendations, France has not generalized the use of man-made mineral fibres in place of asbestos.

Used much less frequently as a substitute for asbestos (less than 10 per cent of the asbestos used in France).

This hierarchy of risks showed that chrysotile asbestos (a proven human carcinogen) poses a much greater threat than ceramic fibres (a suspected animal carcinogen) and *a fortiori* mineral wool (no carcinogenic effect on animals, no effect on the human lung).

The data on which an international consensus was reached in 1991 came from studies and observations made at least four or five years earlier.

With the exception of the G2SAT study, the scientific studies appended to the first written submission of the European Communities are all post-1991.

In fact, because this disease has a very long latency period, the deaths that occurred up to the 1970’s correspond to exposures suffered prior to about the 1940’s, when asbestos was still little used.

Because of the long latency period, it took several decades for the asbestos marketed on a large scale after the Second World War to produce its effects on large numbers of “secondary” users.
It should again be pointed out that this estimate should be at least doubled to take into account the deaths by lung cancer and the cases among women, which are less numerous.

According to recent estimates, in the next 30 years about 500,000 men will die in Western Europe and 95 per cent of these will be workers in the “secondary” user category.

Far fewer workers (about 1,500 in France) are employed in de-flocking and demolition than in “secondary” user occupations (several hundred thousand people exposed daily).


54 Opinion on chrysotile asbestos and likely substitute products, Scientific Committee on Toxicity, Ecotoxicity and the Environment (CSTEE), 19 September 1998.

55 That is the product’s properties, nature and quality, tariff classification, the product’s end-uses given market and consumers’ tastes and habits.


61 Figure 1, page 198, Figure 2, page 199; Table 3, page 200.


65 A roofer using a grinder on corrugated asbestos-cement sheeting in the open air is subjected to a maximum exposure peak of 41 f/ml, i.e. 410 times greater than the French limit value.

66 The permissible limit values under ISO 7337 are very considerably exceeded.

67 See the reply to question 24 of the Panel.

68 Idem.

69 See IARC classification in the List of Products Recognized as Carcinogenic for Man, Global Evaluation of Carcinogenicity in Man, monographs of the International Agency for Research on Cancer, Volumes 1 to 63.

70 IARC Group 2-b.

71 See United States - Section 337 of the Tariff Act 1930, BISD 36S/386, paragraph 5.26 (adopted on 7.11.89); Thailand - Restrictions on Importation of and Internal Taxes on Cigarettes, BISD 37S/214, paragraph 75 (adopted on 7.11.90); United States - Standards for Reformulated and Conventional Gasoline, WT/D92/R of 29.01.96, paragraph 6.22 to 6.24.
See on this point, by analogy, the Appellate Body Report in *Japan - Measures Affecting Agricultural Products*, AB-1998-8, para. 126.


<sup>74</sup> Idem., p. 18.

Canada notes that, in using the term “coverage”, the Panel wondered whether Article III:4 governed the treatment of both the categories of products mentioned in III:2, namely both “like” products and “directly competitive or directly substitutable” products.


<sup>77</sup> *European Communities – Regime Applicable to the Importation, Sale and Distribution of Bananas (Bananas III)*, op. cit., paragraph 204.


<sup>82</sup> See pages 193 to 202 of the INSERM Report – see also our reply to question 3 by Canada.

<sup>83</sup> The European Communities note that the case law of the Appellate Body in the case *Japan – Measures Affecting Agricultural Products* (AB-1998-8, paragraph 126) is also relevant by analogy, where it was held that the burden lies on the Plaintiff Member to establish that the measure at issue is more trade-restrictive than necessary to fulfil the legitimate objective, taking account of the risks non-fulfilment would create.


119 See Section III.C.2.(b)(i).
123 See Annex 1, point 2 of the TBT Agreement.
124 The European Communities note that, as a matter of fact, a general and horizontal prohibition of a given product (as in this case) is by definition likely to increase (rather than restrict) trade in alternative/substitute products regardless of their origin.
125 The European Communities note that if such a dangerous product to human health is allowed to be imported by a Member, this is most likely because either the dangerous nature of the product is not known, not detectable or not visible, or the importing Member has a level of health protection that can cope with a risk of human health of this nature or degree.
126 Report of the Appellate Body, para. 221.
128 Replies by Canada to the questions posed by the Panel and the EC at the first substantive meeting of the Panel on 1 and 2 June 1999.
129 See Section III.C.2(b)(i).
130 Rep. by Canada to the queries posed by the Panel and the EC on the Panel on 1 and 2 June 1999.
131 See Canada’s observations on the replies by the experts to question 3 of the Panel.
132 See Canada’s observations on the replies by the experts to question 6 of the Panel.
133 Article 3.2 of the Understanding on Dispute Settlement and Article 31.1 of the Vienna Convention 1969.
135 See Annex 1, point 2 of the TBT Agreement.
136 The European Communities note that, as a matter of fact, a general and horizontal prohibition of a given product (as in this case) is by definition likely to increase (rather than restrict) trade in alternative/substitute products regardless of their origin.
137 See above Section III.C.2(b)(i) above.
138 The European Communities note that if such a dangerous product to human health is allowed to be imported by a Member, this is most likely because either the dangerous nature of the product is not known, not detectable or not visible, or the importing Member has a level of health protection that can cope with a risk of human health of this nature or degree.
141 See, for example, Article 1.5 of the TBT Agreement and Article 1:4 of the SPS Agreement.
142 See, by analogy, the findings of the Appellate Body in the Reformulated Gasoline Case, in
which the “substantial relationship” of two positions of the gasoline rule was decided in the light of the overall aim of the basic gasoline rule: AB-1996-1, pages 17 to 18.

This is, for example, the way the issue of the relevant legal basis and of the applicable EC treaty provisions are determined in European Community law: see, for example, judgment of 11 June 1991, Case C-300/89, Titanium Dioxide [1991] ECR I-2867. For the problem of characterization in general under international law see, for example, J.A. Salmon, Some Observations on Characterisation in Public International Law, in UN Law / Fundamental Rights, Two Topics in International Law (ed. A. Cassese, 1979), page 3 et seq.

The EC notes, that in fact, the number of exceptions has fallen rapidly since the adoption of the French Decree prohibiting all kinds of asbestos. Thus, in 1997, 87 enterprises used 1,200 tonnes of asbestos under all the permitted exceptions. In 1998, only 63 enterprises used 200 tonnes, 40 per cent of which was used by a single enterprise to produce “waterproof seals” (Latty-Soffa seal). In 1999, the number of enterprises fell to 25 with a volume of only 50 tonnes, 80 per cent of which was used by a single enterprise to produce chlorine (chloride).

According to the EC, this proposition finds implicit support in the decision of the Appellate Body in the Reformulated Gasoline case, where it discussed the term “measures” for the purpose of deciding whether the entire Gasoline Rule or only the baseline establishment rules fell within the scope of Article XX(g). It explained that “The Panel report did not purport to find the Gasoline Rule itself as a whole, or any part thereof other than the baseline establishment rules, to be inconsistent with Article III:4; accordingly, there was no need at all to examine whether the whole of the Gasoline Rule or any of its other rules, was saved or justified by Article XX(g)”. See AB-1996-1, p. 12.


Report of the Panel, paragraph 6.26 to 6.29.


Report of the Appellate Body, paragraph 628: “(...) the statutory baselines would only be applied if it was impossible to determine the source of the imported gasoline or to establish a baseline for lack of data” Report of the Appellate Body, page 25: “we agree with the finding made in the Panel report”. See, e.g, Section 337 Panel report, BISD 36S/345, para. 5.26; Thai Cigarettes Panel report, BISD 37S/200, paragraph 75; Reformulated Gasoline Panel report, WT/D52/R, paragraph 6.22-6.28.

Panel report on Article 337, supra, paragraph 6.1.

The findings of the Panel on these points were confirmed on appeal, pages 29-30.

See, for example, the Panel Report in Thai Cigarettes, paragraph 73.

The European Communities note that Canada has never claimed in its request for consultations or in its request for the establishment of the Panel that the scope of this dispute is about high density chrysotile-containing cement products. Also, the terms of reference of the Panel do not contain such a provision. Furthermore, Canada has been exporting only chrysotile asbestos as a raw material, not high density chrysotile-containing cement products.

Point 3 of the conclusions and recommendations, WHO Report Number 203.


Règlement sur la qualité en milieu de travail.

Code de sécurité pour les travaux de construction.


See also Article 3.23.3.1 of the Code of Safety for Construction Works.
159 Code of Safety for Construction Works, Article 3.23.16.
160 Idem., article 3.23.9 and 3.23.10.
161 Idem., article 3.23.15 and 3.23.16.
162 Idem., article 3.23.16.4.
163 See Decree 97-855 and Decree 97-1219.
164 INSERM, Report on the Effects on Health of the Main Types of Exposure to Asbestos, page 73.
165 See [OSHA, 3349 standards cited, 19101001 or 19261101 issued FY (October 1998 to September 1999)], Nationwide.
166 This Regulation was repealed by Decree No. 96-98 of 7 February 1996 on protection of workers against risks related to the inhalation of asbestos dust, cf page 62 (Article 33) of the collection of French regulatory texts on asbestos, in Recueil des textes français dans le domaine de l’amiante, Official Journals, Ministry of Employment and Solidarity, 1998.
168 The EC point out that the Decree of 28 April 1988 prohibits the sale of crocidolite, subject to limited exceptions, and certain products containing chrysotile, and makes it mandatory to label products containing asbestos accordingly. The Decree of 28 April 1988 was subsequently amended by Decrees No. 94-645 of 26 July 1994, No. 98-668 of 26 July 1996 and No. 96-113 of 24 December 1996. The Decree of 26 July 1994 prohibits the use of all varieties of asbestos, except for chrysotile asbestos. It also extends the list of products containing asbestos which are prohibited from sale.
170 Summary of the principal conclusions of the meeting of French asbestos experts on 20 December 1994.
171 Outline note of the Conseil supérieur de prévention des risques professionnels of 3 July 1995.
173 Idem., page 53.
174 Idem., page 79.
178 Articles R. 232 etc. of the Labour Code.
180 Articles R. 231-56 et seq. of the Labour Code.
181 See, e.g., Article 5.1-5.2 SPS and the Appellate Body report in the Hormones case, AB-1997-4, paragraph 187.
182 The European Communities note that a general overview of the principles applied by WHO and its specialized agencies in the evaluation of food additives and contaminants in food may be gained by looking at the WHO Environmental Health Criteria no. 70, Principles for the Safety Assessment of Food Additives and Contaminants in Food, Geneva, 1987. A study by the US General Accounting Office on pesticides has also demonstrated serious differences in the way a risk assessment is carried out by several industrialised nations: see US General Accounting Office, Pesticides – A Comparative Study of Industrialised Nations’ Regulatory Systems, Washington, 1993.
184 See Section IV.A of this Report.

177 Miriam Cruzê Barros de Oliveira et al., Technical Report Nr. 36 889, Instituto de Pesquisas Tecnológicas, São Paulo (Brazil), 1998.


179 Meldrum, M., Review of Fibre Toxicology, Health & Safety Executive (1996) at 1.

180 Los Angeles Times, Link to Lung Disease Traced to 1906, but Asbestos' Strengths Spurred Use, Part 1, page 2, column 1 (13 July 1986).

181 Executive Summary, Workshop on Health Risks Associated with Chrysotile Asbestos, St. Helier, Channel Islands (1994) at page 2. Brazil also notes that most of this data is from exposure to chrysotile mixed with amphibole.


185 INSERM Report at page 275; see also id. at pages 86-7; id. at page 155 (asbestos-related disease is linked to “massive use”). The effect of massive doses is well documented in the scientific literature on “lung overload.” Thus, one might question how the dose-effect relationship can be defined by studying the effects of “massive quantities of fibers at concentrations far higher than those seen clinically in humans.” At the least, one would not expect such data to define the safe threshold. The existence of a safe threshold for chrysotile is discussed at length in Brazil’s response to EC Question 10.


189 Review of Fibre Toxicology at 1 (1996).


191 Brazil notes that EC Questions 11 through 15 are irrelevant to this proceeding. Brazil nonetheless supplies following answers in order to be as responsive as possible to the EC.

192 Letter from ABRA (Associação Brasileira do Amianto) to SAMA, dated 16 March 1999.


196 Id. at 145.


199 Albin, M. et al., Retention Patterns of Asbestos Fibres in Lung Tissue among Asbestos Cement
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211 Workshop on Health Risks Associated with Chrysotile Asbestos, held at St. Helier, Chan-
nel Islands, 14-17 November 1993, Executive Summary, at page 2.
212 Churg, A. and Vedal, S., Fiber Burden and Patterns of Asbestos-Related Disease in Workers
with Heavy Mixed Amosite and Chrysotile Exposure, Am. J. Respir. Crit. Care Med., Vol. 150,
No. 3 at 667 (1994).
213 Stayner, L.T., Dankovic, D.A., and Lemen, R.A., Occupational Exposure to Chrysotile As-
bestos and Cancer Risk: A Review of the Amphibole Hypothesis, 86 American Journal of Public
214 The United States notes that these represent the most important types of asbestos fibres,
as recognized by the International Union Against Cancer (UICC ).
215 Wagner, J.C., Berry, G., Skidmore, J.W., and Timbrell, V., The Effects of the Inhalation of
Asbestos in Rats, 29 Br. J. Cancer 252-269 (1974); Airborne Asbestos Health Assessment Update,
(EPA, June 1986) at 128 and 130.
216 According to the United States, no data exist, from any studies, concerning risks to
workers exposed solely to crocidolite.
217 In the mid-1980s, OSHA performed a quantitative risk assessment to justify amend-
ments to its standards. This risk assessment was published in 1986 as “Occupational Ex-
posure to Asbestos, Tremolite, Anthophyllite and Actinolite: Final Rules”, in volume 51 of
the U.S. Federal Register at pp. 22615 to 22650 (51 FR 22615-22650) and has been submitted
as U.S. Exhibit 38.
218 Studies by Dement et al (1983) referred to at 51 FR 22645 and Berry et al. discussed at 51 FR 22645, in OSHA risk assessment submitted
as U.S. Exhibit 38.
219 At the Panel’s meeting with third parties, Brazil alleged that worker exposure standards
of the U.S. Department of Health and Human Services differ for chrysotile and other as-
bestos. EPA and OSHA are not aware of any such standards. A 1996 HHS Public Health
Service report referred to an exposure standard of the American Conference of Govern-
mental Industrial Hygienists (ACGIH), but it should be clear that the ACGIH is not a U.S.
government agency and its threshold values are not regulatory standards.
220 Statement by Richard Lemen before the Subcommittee on Toxic Substances, Environ-
mental Oversight, Research and Development, Committee on Environment and Public
Works, April 26, 1990, p. 104-105, Item E in Asbestos Bibliography (Revised), Sept. 1997, DHHS
(NIOSH) Publication No. 97-162, published at Lemen testimony is in document and is
submitted as U.S. Exhibit 44.
222 HEI, Asbestos in Public and Commercial Buildings: A Literature Review and Synthesis of
223 For example, Managing Asbestos in Place: A Building Owner’s Guide to Operations and

OSHA Final Rule on Occupational Exposure to Asbestos, 59 FR 40964, 41036-38, August 10, 1994 at 41036-37.


See Section III.B.7 of this Report.

This risk assessment has been submitted by the United States in Occupational Exposure to Asbestos, Tremolite, Anthophyllite and Actinolite: Final Rules, 51 CFR 22615-22650.


Baujon et Authier, Détermination des concentrations de fibres d’amiante dans l’atmosphère lors de la pose sur chantier de plaques ondulées et d’ardoises au amiante-ciment (Rapport 93.189), Laboratoire d’hygiène et de contrôle des fibres minérales, Paris, 1993, referred to by Canada (see Section III.B.7).

See arguments of the EC in Section III.C.1.(c)(ii).

Affections professionnelles consécutives à l’inhalation de poussières d’amiante, Tableau n° 30 (document submitted by the EC to the Panel).

See Section V. C. 1(b) of this Report.

See Part V.C.2 of this report, answer to Question 5(c).

See Section V. D. 1, Canada’s comments to Question 5(a).

See Section V.D.1 of this Report, Canada’s comments to Question 5(a).

See Section V.C.2, reply to Question 5(a).

See Section V. C. 2, reply to Question 5(a).

See Section V.C.2, reply to Question 5(a).


Thank you, Mr. Chairman. My name is Marcel Goldberg and I am one of the authors of this report and I am in charge of this part of the report indeed. We have drafted this sentence that has been quoted, but, once again, this quotation is out of context. It is true that we said that, but of course this is one sentence in the part dealing with epidemiological data and the whole report has more or less 450 pages, and we took into account all the data available, including experimental data, and the conclusion of this whole work has enabled us to conclude that, in all probability, the risk of cancer linked to this kind of fibre was largely under that of chrysotile. Thank you.]
THE APPELATE REPORT
I. Introduction

1. Canada appeals certain issues of law and legal interpretations developed in the Panel Report in European Communities – Measures Affecting Asbestos and Asbestos-Containing Products (the “Panel Report”). The Panel was established to consider claims made by Canada regarding French Decree No. 96-1133 concerning asbestos and products containing asbestos (décret no.96-1133 relatif à l’interdiction de l’amiante, pris en application du code de travail et du code de la consommation) (“the Decree”), which entered into force on 1 January 1997.

2. Articles 1 and 2 of the Decree set forth prohibitions on asbestos and on products containing asbestos fibres, followed by certain limited and temporary exceptions from those prohibitions:

Article 1

I. For the purpose of protecting workers, and pursuant to Article L. 231-7 of the Labour Code, the manufacture, processing, sale, import, placing on the domestic market and transfer under any title whatsoever of all varieties of asbestos fibres shall be prohibited, regardless of whether these substances have been incorporated into materials, products or devices.

II. For the purpose of protecting consumers, and pursuant to Article L. 221.3 of the Consumer Code, the manufacture, import, domestic marketing, exportation, possession for sale, offer, sale and transfer under any title whatsoever of all varieties of asbestos fibres or any product containing asbestos fibres shall be prohibited.

III. The bans instituted under Articles I and II shall not prevent fulfilment of the obligations arising from legislation on the elimination of wastes.

Article 2

I. On an exceptional and temporary basis, the bans instituted under Article 1 shall not apply to certain existing materials, products or devices containing chrysotile fibre when, to perform an
equivalent function, no substitute for that fibre is available which:

- On the one hand, in the present state of scientific knowledge, poses a lesser occupational health risk than chrysotile fibre to workers handling those materials, products or devices;

- on the other, provides all technical guarantees of safety corresponding to the ultimate purpose of the use thereof.

II. The scope of application of paragraph I of this Article shall cover only the materials, products or devices falling within the categories shown in an exhaustive list decreed by the Ministers for Labour, Consumption, the Environment, Industry, Agriculture and Transport. To ascertain the justification for maintaining these exceptions, the list shall be re-examined on an annual basis, after which the Senior Council for the Prevention of Occupational Hazards and the National Commission for Occupational Health and Safety in Agriculture shall be consulted.

The remaining operative provisions of the Decree contain additional rules governing the grant of an exception (Articles 3 and 4), the imposition of penalties for violation of the prohibition in Article 1 (Article 5), and the temporary exclusion of certain “vehicles” and “agricultural and forestry machinery” from aspects of the prohibition (Article 7). Further factual aspects of this dispute are set forth in paragraphs 2.1 – 2.7 of the Panel Report, and the Decree is reproduced in its entirety as Annex I in the Addendum to the Panel Report. 3

3. Canada claimed that the Decree is inconsistent with a number of obligations of the European Communities under Article 2 of the Agreement on Technical Barriers to Trade (the “TBT Agreement”), Articles III and XI of the General Agreement on Tariffs and Trade 1994 (the “GATT 1994”), and that, under Article XXIII:1(b) of the GATT 1994, the Decree nullified or impaired advantages accruing to Canada directly or indirectly under the Marrakesh Agreement Establishing the World Trade Organization (the “WTO Agreement”), or impeded the attainment of an objective of that Agreement. 4

4. In the Panel Report, circulated to WTO Members on 18 September 2000, the Panel concluded that:

(a) … the “prohibition” part of the Decree does not fall within the scope of the TBT Agreement. The part of the Decree relating to “exceptions” does fall within the scope of the TBT Agreement. However, as Canada has not made any claim concerning the compatibility with the TBT Agreement of the part of the Decree relating to exceptions, the Panel refrains from reaching any conclusion with regard to the latter.
...chrysotile asbestos fibres as such and fibres that can be substituted for them as such are like products within the meaning of Article III:4 of the GATT 1994. Similarly, the Panel concludes that the asbestos-cement products and the fibro-cement products for which sufficient information has been submitted to the Panel are like products within the meaning of Article III:4 of the GATT 1994.

With respect to the products found to be like, the Panel concludes that the Decree violates Article III:4 of the GATT 1994.

However, ... the Decree, insofar as it introduces a treatment of these products that is discriminatory under Article III:4, is justified as such and in its implementation by the provisions of paragraph (b) and the introductory clause of Article XX of the GATT 1994.

Finally, ... Canada has not established that it suffered non-violation nullification or impairment of a benefit within the meaning of Article XXIII:1(b) of the GATT 1994. 5

Having found that the Decree is subject to, and inconsistent with, the obligations set forth in Article III:4 of the GATT 1994, the Panel did not deem it necessary to examine the claims of Canada under Article XI of the GATT 1994. 6

On 23 October 2000, Canada notified the Dispute Settlement Body (the “DSB”) of its decision to appeal certain issues of law covered in the Panel Report and certain legal interpretations developed by the Panel, pursuant to Article 16.4 of the Understanding on Rules and Procedures Governing the Settlement of Disputes (the “DSU”), and filed a Notice of Appeal with the Appellate Body pursuant to Rule 20 of the Working Procedures for Appellate Review (the “Working Procedures”). 7 On 16 November 2000, Canada filed an appellant’s submission. 8 On 21 November 2000, the European Communities filed an other appellant’s submission. 9 On 1 December 2000, Canada and the European Communities each filed an appellee’s submission. 10 On the same day, Brazil and the United States each filed a third participant’s submission. 11

On 21 November 2000, the Appellate Body received a letter from Zimbabwe indicating its interest in attending the oral hearing in this appeal. Zimbabwe participated in the proceedings before the Panel as a third party which had notified its interest to the DSB under Article 10.2 of the DSU, but it did not file a third participant’s submission in the appeal. No participant or third participant objected to Zimbabwe’s request. On 15 December 2000, the Members of the Division hearing this appeal informed Zimbabwe, the participants and third participants, that Zimbabwe would be allowed to attend the oral hearing as a passive observer.

On 20 December 2000, the Appellate Body informed the DSB that, due to the exceptional workload of the Appellate Body, and in light of the agreement of the participants, Canada and the European Communities, the Appellate Body Report in this appeal would be circulated to WTO Members no later than Monday, 12 March 2001. 12
9. The oral hearing in the appeal was held on 17 and 18 January 2001. The participants and the third participants presented oral arguments and responded to questions put to them by Members of the Division hearing the appeal.

II. Arguments of the Participants and the Third Participants

A. Claims of Error by Canada – Appellant

1. **TBT Agreement**

10. Canada requests that the Appellate Body reverse the Panel’s findings and conclusions on the definition of the term “technical regulation”, hold that the Decree as a whole falls within the scope of the TBT Agreement, and find that the Decree is inconsistent with paragraphs 1, 2, 4 and 8 of Article 2 of the TBT Agreement.

11. Canada asserts that the Panel erred in law in failing to examine Canada’s allegations under the TBT Agreement. The Panel wrongly split the Decree into two and considered the prohibitions and exceptions in the Decree to be separate measures for the purposes of determining whether the Decree is a technical regulation within the meaning of the TBT Agreement. Canada believes that the Panel’s analysis is arbitrary, contrary to the internal coherence of the Decree, and allows the applicability of the TBT Agreement to be determined by the way in which a Member drafts its legislation.

12. Canada argues that the Panel also erred in its interpretation of the definition of “technical regulation” in Annex 1 to the TBT Agreement, in particular, in articulating two criteria that must be satisfied before a measure can be a “technical regulation”: (i) the measure must concern identifiable products; and (ii) the measure must identify the technical characteristics that products must have to be marketed in the territory of the Member taking the measure. This interpretation adds requirements to the definition of “technical regulation” that have no basis in the text of the TBT Agreement, and are inconsistent with the object and purpose of that Agreement, namely to restrain non-tariff barriers to trade that may be disguised as technical regulations. In addition, with respect to the first criterion, requiring a measure to relate to identifiable products to constitute a technical regulation could lead to arbitrary results in practice. As for the second criterion, Canada alleges that it is too narrow and would exclude from characterization as “technical regulations”, and thereby insulate from the disciplines of the TBT Agreement, measures regulating activities other than the marketing of products, such as measures relating to transportation of products, disposal of hazardous waste, and use of special equipment to repair certain products.

13. Canada challenges the Panel’s conclusion that the TBT Agreement does not apply to a general prohibition like the one in the Decree. The Panel relied on a false distinction between general prohibitions, which it considered fall exclusively under the GATT 1994, and technical regulations, which are subject to the disciplines of the TBT Agreement. In fact, a technical regulation can have the effect on trade of a general prohibition.

14. Canada maintains that, had the Panel viewed the Decree as a unified measure, and correctly interpreted the term “technical regulation”, the Panel would have concluded that the Decree is a technical regulation within the meaning of the TBT Agreement. However, even if the general prohibition contained in the Decree were not characterized as a technical regulation, the Panel nevertheless erred in failing to examine Canada’s claims under the TBT Agreement, given that the Panel also found that the TBT Agreement applies to the part of the Decree concerning exceptions, and that Canada’s claims related to the Decree as a whole. Canada therefore requests the Appellate Body to reverse the Panel’s conclusions on the applicability of the TBT Agreement to the Decree, and to assess the compatibility of the Decree with that Agreement. Canada argues that, as in United States – Import Prohibition of Certain Shrimp and Shrimp Products (“United States – Shrimp”), “the facts on the record of the panel proceedings” allow the Appellate Body “to undertake the completion of the analysis required to resolve this dispute.”
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15. Canada argues that the Decree is inconsistent with Article 2.1 of the TBT Agreement. Since the principle of national treatment in Article 2.1 is a specific, particular expression of Article III:4 of the GATT 1994, the interpretation of the words “like products” in Article 2.1 must be identical to the interpretation of the same words in Article III:4. The meaning of “like products” in Article III:4 is relevant context and, in the view of Canada, both Article III:4 of the GATT 1994 and Article 2.1 of the TBT Agreement have the same object and purpose, namely to avoid protectionism and to provide equality of competitive conditions for imported products in relation to domestic products. Thus, Canada maintains, the findings of “likeness”, and of less favourable treatment, made by the Panel pursuant to Article III:4 of the GATT 1994 must be extended to Article 2.1 of the TBT Agreement.

16. In Canada’s view, the Decree is inconsistent with Article 2.2 of the TBT Agreement. Canada insists, first, that there is no rational connection between the Decree and France’s objective of protecting human health since: (i) it is friable materials containing amphiboles which pose a risk to human health; (ii) the manipulation of chrysotile-cement products and other high-density products containing chrysotile asbestos fibres does not pose a danger to human health; and (iii) the Decree exposes the French public to substitute fibres, the health risks of which are still poorly understood. Canada adds, second, that the Decree has effects that are more trade-restrictive than necessary to achieve its objective, in particular, because: (i) the manipulation of chrysotile-cement products and other high-density products containing chrysotile asbestos fibres does not create a risk to human health; and (ii) there is a less trade-restrictive alternative that protects human health, namely the “controlled use” of chrysotile-cement products and other high-density products containing chrysotile asbestos fibres. What must be demonstrated under Article 2.2 of the TBT Agreement is the same as what must be demonstrated under Article XX(b) of the GATT 1994. In this regard, according to Canada, the reports of the panel and the Appellate Body in United States - Standards for Reformulated and Conventional Gasoline (“United States – Gasoline”) establish that a less trade-restrictive alternative can only be ruled out if it is shown to be impossible to implement. However, France did not demonstrate, and the Panel did not find, that it is impossible to implement “controlled use”. Furthermore, Canada contends, it would be less trade-restrictive to ban products containing chrysotile asbestos fibres on the basis of a product-by-product demonstration of the ineffectiveness and unfeasibility of “controlled use”, rather than on the basis of the existence of substitute products.

17. Canada also argues that the Decree is inconsistent with Article 2.4 of the TBT Agreement, because there are relevant international standards on the “controlled use” of chrysotile, which constitute an effective and appropriate means to achieve France’s objective of protecting human health. In any event, the French government acted inconsistently with Article 2.4 because it did not use international standards as a basis for the Decree. Lastly, Canada considers that the Decree is inconsistent with Article 2.8 of the TBT Agreement because it institutes a prohibition based on the descriptive characteristics of products, rather than on requirements in terms of performance.

2. Article XX(b) of the GATT 1994 and Article 11 of the DSU

18. Canada requests that the Appellate Body reverse the Panel’s findings and conclusions under Article XX(b) of the GATT 1994 and find that the Decree is not justified under that provision. Canada also asks the Appellate Body to find that the Panel did not make an “objective assessment of the matter”, as required under Article 11 of the DSU, because it failed to assess the scientific data in accordance with the principle of the balance of probabilities, and failed to assess the facts objectively.

19. Canada argues that the Panel erred in finding that there is a risk to human health associated with the manipulation of chrysotile-cement products. Canada identifies seven factors it claims the Panel mistakenly relied on in reaching this conclusion: (i) a statement by Dr. Henderson that “building workers now count among those most exposed to
chrysotile fibres and hence to the risk of mesothelioma" 16; (ii) an “anecdotal” statement by Dr. Henderson concerning “cases of mesothelioma in patients who had been only incidentally exposed, without any relation to their occupational activity” 17; (iii) the opinion of experts that it has not been established that there is a threshold below which exposure does not constitute a risk for mesothelioma or lung cancer; (iv) the “Charleston study” 18; (v) “statistical data” adduced by Dr. Henderson, which, according to the Panel, confirmed “the impact of chrysotile on mechanics exposed to that material in a car brake maintenance context” despite a contrary study on automobile brake maintenance relied on by Canada 19; (vi) the use of the no-threshold linear relationship model as a basis for concluding that there is a “real risk” and “an undeniable public health risk” associated with exposure to chrysotile asbestos fibres at low or intermittent levels 20; and (vii) data supplied by the European Communities concerning intermittent manipulation and a reference by Dr. Henderson to a Japanese study as a basis for concluding that the manipulation of chrysotile-cement using inappropriate tools could cause exposure levels above statutory limits. 21

Canada sets forth detailed explanations as to why none of these factors supports the Panel’s conclusion.

20. Canada also contends that the Panel erred in its application of the test of “necessity” under Article XX(b) of the GATT 1994. Canada accepts the Panel’s view that the extent of the risk to human health is relevant to the assessment of “necessity”. However, Canada disputes that there is any risk involved in the manipulation of such products, highlights that the evidence relied on by the Panel certainly could not form the basis for a finding that the health risk was so high that it could justify strict measures, and argues that the Panel failed to comply with its obligation to quantify this type of risk. In Canada’s view, these errors distorted the Panel’s analysis of the test of necessity and led it to take a much too restrictive approach to its consideration of reasonably available alternatives to the Decree.

21. Canada asserts that, in its examination of whether less restrictive international trade alternatives can achieve the level of protection inherent in the Decree, the Panel erred in accepting that such level of protection is a halt to the spread of the risk associated with chrysotile asbestos fibres. This premise does not take account of the risk associated with the use of substitute fibres, of the absence in France of any regulatory framework for “controlled use” of such fibres, or of the false sense of security created among the French public due to the absence of such a framework. The Panel also erred in law in finding that there was no reasonably available alternative to the Decree that is consistent or less inconsistent with the GATT 1994. In this regard, Canada makes the same arguments that it made above with respect to Article 2.2 of the TBT Agreement, and emphasizes that the Panel was overly strict in its examination of the alternatives, considering that France could have adopted a measure establishing bans on specific products containing chrysotile asbestos fibres, based on demonstrations of the ineffectiveness and unfeasibility of the “controlled use” of each product.

22. Canada submits that the Panel failed to discharge its responsibility to make an objective assessment of the matter when it declined to take a position on the opinions expressed by the scientific community. For Canada, the principle of the balance of probabilities, or the preponderance of evidence, requires the trier of fact to take a position as to the respective weight of the evidence. Had the Panel properly applied this principle, it would not have been able to conclude that the Decree was justified under Article XX(b) of the GATT 1994, in view of the multiple studies submitted by Canada showing, for example, that there is no increased risk among garage and brake mechanics, or among construction workers, resulting from the manipulation of chrysotile asbestos. Canada adds that the Panel also failed to make an objective assessment of the matter before it because, in its determinations on the “controlled use” of chrysotile, it relied extensively on the opinions of the experts consulted, who in fact did not possess expertise in the area of “controlled use”.

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B. Arguments of the European Communities – Appellee

1. TBT Agreement

23. The European Communities urges the Appellate Body to reject Canada’s appeal on the TBT Agreement. The Panel correctly concluded that the “prohibition part” of the Decree is not a technical regulation within the meaning of Annex 1.1 to the TBT Agreement. Canada’s arguments with respect to the “exceptions part” of the Decree are legally irrelevant, since it would be impossible for the Appellate Body to complete the legal analysis due to the lack of sufficient and undisputed facts. The European Communities adds that the claims made by Canada under the TBT Agreement should, in any event, be denied.

24. The European Communities sees no error in the Panel’s separation of the prohibitions part of the Decree from the exceptions part. The exceptions are ancillary to the prohibitions, and separating the two parts for the purpose of their legal characterization under the TBT Agreement in no way affects the internal coherence of the Decree. In this case, the issue before the Panel was whether the prohibitions laid down in the Decree constitute a technical regulation, not whether, in the abstract, a general ban may be a technical regulation. The European Communities also considers that the Panel correctly interpreted the term “technical regulation”, and that the interpretation suggested by Canada would deprive other GATT 1994 provisions, such as Article XI, of effect.

25. The European Communities agrees with the Panel’s treatment of the exceptions part of the Decree, and insists that, having made no specific violation claim regarding the narrowness of the exceptions throughout these proceedings, Canada cannot now argue that the exceptions violate the TBT Agreement. The European Communities argues that the Appellate Body is in any case prevented from addressing Canada’s claims under the TBT Agreement because to do so would require the Appellate Body to make findings of a factual and technical nature which, in the absence of undisputed facts and findings in the record, it cannot do on appeal. The Appellate Body could not simply use the findings of the Panel under Articles III:4 and XX of the GATT 1994 as a basis for an analysis under the TBT Agreement. While the two sets of rules are related, they are not “part of a logical continuum” and are not sufficiently closely related as to allow the Appellate Body to extrapolate the findings of the Panel under Article III:4 and Article XX(b) of the GATT 1994 into the sphere of the TBT Agreement. Should the Appellate Body examine Canada’s claims under the TBT Agreement, the European Communities argues that these claims should be dismissed and refers, in this regard, to its arguments with respect to Article XX(b) of the GATT 1994, and to the arguments it made before the Panel with regard to Articles 2.1, 2.2, 2.4 and 2.8 of the TBT Agreement.

2. Article XX(b) of the GATT 1994 and Article 11 of the DSU

26. The European Communities submits that the Panel’s finding that the violation of Article III:4 is justified under Article XX(b) of the GATT 1994 is legally sound and correct. Canada’s arguments on this issue amount to a request that the Appellate Body make new factual and scientific findings on appeal, contrary to the limits on the scope of appellate review set out in Article 17.6 of the DSU.

27. The European Communities believes that the Panel concluded that the ban on asbestos was “necessary” based on a series of objective and verifiable findings, made after a detailed and careful evaluation of the factual and scientific evidence presented. In assessing whether the ban was “necessary”, the Panel was not obliged to undertake a “quantitative” assessment of the identified risk. Neither the ordinary meaning of the terms “necessary to protect” in Article XX(b) nor the concept of risk assessment mandate such an approach. An assessment of risk may be made either in quantitative or qualitative terms. The European Communities adds that the Panel correctly found that, after the European Communities had established a prima facie case for the existence of a health risk in connection with the use of chrysotile, Canada bore the burden of refuting that case by showing the absence of such a health risk.
28. On the issue of whether another measure was reasonably available, the European Communities submits that Canada cannot, on appeal, make arguments based on the health risks associated with the substitute products for asbestos, or on the safety of the "controlled use" of asbestos, as both arguments seek to have the Appellate Body revisit factual findings made by the Panel on the basis of the evidence submitted and the opinions advanced by the experts consulted.

29. With respect to the alleged inconsistency with Article 11 of the DSU, the European Communities considers that Canada’s claim that the Panel committed a fundamental error in its appreciation of the facts seems to be based solely on the fact that the Panel based itself exclusively on the opinions of the experts consulted in this case. In this regard, the European Communities emphasizes that Canada did not object to the selection of the experts by the Panel, that Canada proposed one of those experts itself, and that the experts themselves answered a question on “controlled use” rather than professing a lack of expertise on the issue. As for Canada’s argument that the Panel erred in law in failing to evaluate the scientific evidence in accordance with the principle of preponderance of the evidence, the Panel’s approach does not seem inconsistent with such a principle and, in any case, the principle of preponderance of the evidence is inapposite in the context of risk assessment since such an approach would preclude Members from basing their regulatory decisions on diverging scientific opinions. The European Communities refuses to accept that the evidence relied on by the Panel – representing the unanimous views of the four experts consulted and of all international institutions that have evaluated asbestos – reflects, as Canada seems to suggest, a divergent, minority scientific point of view on asbestos.

C. Claims of Error by the European Communities – Appellant

1. “Like Products” in Article III:4 of the GATT 1994

30. The European Communities requests the Appellate Body to reverse the Panel’s findings that chrysotile asbestos fibres are “like” polyvinyl alcohol (“PVA”), cellulose and glass fibres, and that chrysotile-cement products are “like” fibro-cement products, as well as the Panel’s consequent finding that, with respect to the products found to be “like”, the Decree violates Article III:4 of the GATT 1994.

31. The Panel’s interpretation of the term “like products” in Article III:4, is of serious concern to the European Communities; is contrary to the ordinary meaning of Article III:4, read in context and in light of its object and purpose; and is inconsistent with established case law. As the Appellate Body has previously found, the first paragraph of Article III defines the object and purpose of the whole of Article III, namely, to provide equality of competitive conditions for imported products in relation to domestic products. In the view of the European Communities, the Panel, however, erroneously analyzed the term “like products” in light of the objective of ensuring market access for products, and, in so doing, adopted an exclusively commercial approach to the comparison of “like” products and erroneously expanded the scope of application of Article III:4.

32. The European Communities submits that this erroneous focus on market access led the Panel to exclude from its “like” product analysis the very reason why the Decree singles out asbestos fibres, namely, the fact that asbestos fibres are carcinogenic. While Article III:4 protects expectations concerning the competitive relationship between imported and domestic products, the impact of a measure on such expectations is not relevant in determining “likeness”, but only later in the Article III:4 analysis, for the purposes of establishing whether the measure discriminates between imported and domestic products. For the European Communities, the decisive criterion for determining the “likeness” of products must be whether the basis for the regulatory distinction between products denies to imported products the treatment accorded to domestic products that are the subject of the relevant measure.
33. The European Communities contends that, because the Panel ignored the basis for the regulatory treatment set forth in the Decree, it compared the wrong products in its analysis of “ likeness”. The Decree prohibits all carcinogenic asbestos fibres, and it denies competitive opportunities to all such fibres equally. Thus, the prohibited carcinogenic asbestos fibres are not “like” the three substitute fibres because the application of the French regulatory distinction between them does not alter or affect the competitive opportunities of those substitute fibres. The European Communities concludes that, instead of comparing the products claimed by Canada to be “like” products (PVA, cellulose and glass fibres) with the category of products prohibited by the French Decree at issue (all carcinogenic asbestos fibres), the Panel erroneously compared the allegedly “like” products with an arbitrary third category of products, namely “fibres with certain industrial applications”. 23

34. The European Communities challenges the Panel’s conclusion that, in view of the relationship between Articles III and XX(b) of the GATT 1994, it is not appropriate to take the “risk” criterion into account either when examining the properties, nature and quality of the product, or when examining other criteria of “ likeness”. 24 The Panel found that the health, safety or other concerns that lead regulators to apply different treatment to products may only be taken into account in the analysis under Article XX, not in the analysis under Article III:4 of the GATT 1994. The Panel’s approach misconstrues the relationship between Articles III:4 and XX of the GATT 1994, requires the “ likeness” of two products to be determined solely on the basis of commercial factors and, in the view of the European Communities, entails a serious curtailment of national regulatory autonomy. If non-commercial considerations may only be considered at the Article XX stage of the analysis, then the list of policy purposes for which regulators may distinguish between products is unduly limited to the categories listed in Article XX. The application of a “risk” criterion in the analysis of “ likeness” under Article III would not, as the Panel suggests, make the other criteria of “ likeness” “totally redundant” 25, since all relevant criteria, including the “risk” criterion, must be considered in the assessment of “ likeness”.

35. The European Communities contends that the Panel committed a number of errors in its application of the four criteria used to assess “ likeness”, and placed excessive importance on the criterion of end-use. The Panel failed to follow the approach used in previous case law, and ignored the fact that Article III:4 of the GATT 1994, unlike Article III:2 and its accompanying Interpretive Note, does not contain the phrase “directly competitive or substitutable” products. The Panel’s analysis of “end-use” is inadequately reasoned, in particular since the Panel failed to identify the small number of identical or similar end-uses for chrysotile asbestos, PVA, cellulose and glass fibres and ignored that, overall, the end-uses for asbestos and its substitutes are very different. The European Communities adds that the Panel relied on its conclusions on end-use in its analysis of the properties, nature and quality of the products, as well as their tariff classification, and, in effect disregarded these other criteria.

2. Article XXIII:1(b) of the GATT 1994

36. The European Communities appeals the Panel’s findings on Article XXIII:1(b) of the GATT 1994 in paragraphs 8.262, 8.264, 8.273 and 8.274 of the Panel Report, but not the Panel’s conclusion that Canada did not establish nullification or impairment of a benefit within the meaning of Article XXIII:1(b). The Panel’s reasoning is inconsistent with the proper interpretation of the GATT 1994, past practice, and relevant case law. Historically, the non-violation remedy was conceived as a legal instrument designed to prevent the circumvention of tariff concessions. Only three non-violation complaints have succeeded. All previous non-violation complaints have related to measures imposed for commercial purposes, and all such complaints would today most likely be resolved as violation complaints under the expanded WTO competence, reflected in the covered agreements. The European Communities urges the Appellate Body to accept that the concept of non-violation nullification and impairment is an exceptional one, as WTO Members have recognized, and should be applied with utmost circumspection.
37. The European Communities challenges, in particular, the Panel’s conclusion that "Article XXIII:1(b) applies to a measure whether it is consistent with the GATT because the GATT does not apply to it or is justified by Article XX." In so finding, the Panel wrongly implied that Article XXIII:1(b) of the GATT 1994 protects the expectation that, once a tariff concession has been made for a product, the regulatory framework applicable to that product will not be adapted in response to new scientific knowledge concerning health risks. In the view of the European Communities, the Panel’s interpretation wrongly expanded the coverage of Article XXIII:1(b) in a manner that has grave systemic implications.

38. The European Communities urges the Appellate Body to reject, as a matter of legal principle, the possibility of finding nullification or impairment under Article XXIII:1(b) with respect to health and safety regulations, or with respect to measures that fall under any of the other grounds listed in Article XX, or under provisions such as Articles XIX and XXI of the GATT 1994. Article XXIII:1(b) cannot apply in cases involving health measures, since the legitimacy of an exporting Member’s expectation that the health measure will not be taken cannot be assessed without examining the health measure itself and the balance of interests underlying that law. The participants in the Uruguay Round knew that the value of the concessions negotiated in that Round could be adversely affected by measures taken to protect, inter alia, human, animal or plant life or health, or a national security interest. Therefore, the European Communities concludes, if a Member takes a measure that is consistent with the GATT 1994, it does not disturb the balance of rights and obligations under the GATT 1994, and no redress is available under Article XXIII:1(b).

D. Arguments of Canada – Appellee

1. "Like Products" in Article III:4 of the GATT 1994

39. Canada requests the Appellate Body to dismiss the European Communities’ appeal relating to Article III:4 of the GATT 1994. Canada is of the view that the Panel correctly separated the analysis of "likeness" from the issue of whether the competitive opportunities afforded to imports on the domestic market have been upset. In its appeal, the European Communities confounds these two distinct questions and attaches undue significance to the Panel’s statement regarding the importance of "market access" under Article III:4 of the GATT 1994.

40. Canada considers that the Panel properly applied the criteria set out in the case law for determining whether products are "like". The European Communities appears to confuse the concept of "likeness" under Article III:4 of the GATT 1994 with "likeness" under Article III:2. "Likeness", however, under Article III:4 is different from, and broader than, "likeness" under the first sentence of Article III:2, and the Panel’s approach properly reflects this distinction. In assessing the "likeness" of the fibres, the Panel recognized that the criteria of "properties" and "end-use" are interdependent, and analyzed them accordingly. Canada does not accept that the Panel created a hierarchy among the traditional "likeness" criteria, but, even so, this would not be an error of law, since "likeness" must be approached on a case-by-case basis, and it is within a panel’s discretion to establish a hierarchy among the criteria in any given case. Finally, Canada notes, the appeal of the European Communities focuses on the Panel’s conclusion that chrysotile asbestos fibres are "like" PVA, cellulose and glass fibres, and the criticisms that the European Communities makes of this conclusion cannot be extended to the Panel’s separate conclusion that chrysotile-cement products are "like" fibro-cement products.

41. Canada submits that the Panel correctly decided that the “dangerousness” of a product is not a factor to be considered in determining “likeness” and that to introduce a criterion of this nature into the analysis of “likeness” would nullify the effect of Article XX(b) of the GATT 1994. The object and purpose of Article III of the GATT 1994 is to provide equality of competitive conditions for imported and domestic products, and the four traditional criteria of "likeness" all relate to the state of commercial competition between such products. The “dangerousness” of products is unrelated to such commercial competition. Furthermore, to introduce such factors into the analysis of “likeness” under Article III:4
would lead to unpredictability as to the scope of that provision, and imply that determining the “likeness” of products requires complex scientific analysis for which panels have no special expertise. Canada adds that even if the “dangerousness” of a product were relevant to the determination of “likeness”, it would not necessarily follow that chrysotile asbestos fibres are not “like” the substitute fibres. Since Article XX of the GATT 1994 was specially designed to balance the interest of promoting international trade with legitimate societal interests, it is a more appropriate framework than Article III for taking account of these types of considerations. Canada also stresses that, contrary to the argument of the European Communities, such an approach does not lead to a curtailment of national regulatory autonomy, because the list in Article XX covers a broad range of interests on the basis of which a Member may justify a measure.

42. Canada also submits that, in its appeal, the European Communities errs in asserting that the examination of “likeness” must be done on the basis of the regulatory distinction in question, and in claiming that the Panel should only have compared chrysotile asbestos fibres with carcinogenic fibres, rather than with other fibres that serve similar industrial uses. Such an approach is inconsistent with the proper interpretation of Article III:4. In seeking to focus the analysis on the reason for any given regulatory distinction, the European Communities would allow national regulatory authorities to predetermine the scope of Article III:4 through the distinctions they choose to make. Such an approach is also inconsistent with the object and purpose of Article III:4, which aims to discipline measures that have trade-restrictive effects, even when those measures are not aimed at restricting trade. Finally, in Canada’s view, the Panel correctly compared chrysotile asbestos fibres with the fibres with which they compete in certain industrial applications, since such a comparison is consistent with the aim of providing equality of competitive conditions, and since the Decree itself makes no reference to carcinogenic fibres.

2. Article XXIII:1(b) of the GATT 1994

43. Canada requests the Appellate Body to reject the European Communities’ appeal with respect to Article XXIII:1(b) of the GATT 1994. Canada suggests, first, that the Appellate Body should apply the principle of judicial economy and refrain from ruling on these grounds of appeal. Canada argues that a ruling by the Appellate Body in respect of Article XXIII:1(b) of the GATT 1994 would not further the objective of dispute settlement, as set forth in Article 3.7 of the DSU, namely to secure a positive solution to a dispute. There is no dispute concerning Article XXIII:1(b) because neither party has appealed the Panel’s conclusions on this issue. Canada also refers to Article 3.2 of the DSU and cautions the Appellate Body against “making law” by clarifying provisions of the WTO Agreement outside the context of resolving a particular dispute.

44. Should the Appellate Body address the interpretation of Article XXIII:1(b) of the GATT 1994, Canada invites it to affirm the Panel’s reasoning, in particular the Panel’s recognition that there may be particularly exceptional cases in which a measure justified under Article XX(b) would nonetheless nullify or impair benefits within the meaning of Article XXIII:1(b). Article XX(b) and XXIII:1(b) may be applied simultaneously, since Article 26.1 of the DSU does not require the withdrawal of a measure that nullifies or impairs benefits within the meaning of Article XXIII:1(b). As regards the concept of legitimate expectations, Canada rejects as artificial, and without any textual basis, the distinction that the European Communities seeks to draw between pure trade measures and measures linked to the protection of health.

E. Arguments of the Third Participants

1. Brazil

(a) TBT Agreement

45. Brazil believes that the Panel erred in its findings regarding the scope of the TBT Agreement. Brazil argues that the Panel erred in dividing the Decree into two separate
parts in determining whether the TBT Agreement applies to the Decree. This division was arbitrary and inconsistent with the logic and objectives of the Decree, which deals with the same products in both the prohibition and the exception parts. Furthermore, Brazil is particularly concerned by the findings of the Panel in paragraphs 8.38, 8.39, 8.43, 8.49, 8.57, 8.60, 8.61 and 8.71 of the Panel Report, and by the serious systemic implications of the finding that a general prohibition does not constitute a technical regulation within the meaning of Annex 1.1 of the TBT Agreement. Contrary to the Panel’s interpretation, nothing in the TBT Agreement specifies that a product must be “identifiable”, or that a measure must relate to one, or more than one product, in order to be a technical regulation. Such a narrow interpretation unduly excludes from the scope of the TBT Agreement a wide range of measures affecting products that could potentially represent barriers to trade. Brazil also contests the Panel’s finding that a technical regulation must include specifications to be met in order for a product to be authorized for marketing. Brazil adds that, in its view, both France and the European Communities conceded, when they notified the Decree under the TBT Agreement, that the measure is a technical regulation.

2. United States

(a) TBT Agreement

46. The United States argues that the Panel erred in its interpretation of the phrase “technical regulation” in Annex 1 to the TBT Agreement, and, in consequence, improperly excluded from the scope of the TBT Agreement technical regulations that apply generally to products. Specifically, the United States contends that the Panel erred in finding that the phrase “product characteristics” in the definition of “technical regulation” refers to characteristics of “one or more given products”, rather than characteristics of products generally.

47. Should the Appellate Body find that the TBT Agreement applies to the Decree and decide to complete the analysis of Canada’s claims under that Agreement, the United States submits that the Appellate Body should find that the Decree is consistent with the TBT Agreement. Asbestos and asbestos-containing products, on the one hand, and substitute fibres and asbestos-free products, on the other, are not “like products” for the purposes of Article III:4 of the GATT 1994. The test to be applied under Article 2.2 of the TBT Agreement is very similar to the test to be applied under Article XX(b) and the introductory clause to Article XX. However, unlike Article XX of the GATT 1994, where the burden was on the European Communities to present a prima facie case that the Decree was justified, under Article 2.2 of the TBT Agreement, it is for Canada to make a prima facie case that the Decree creates an unnecessary barrier to trade, and it has not done so. The Decree is also consistent with Article 2.4 of the TBT Agreement, since the international standards identified by Canada are neither relevant to, nor an effective or appropriate means of achieving, France’s public health objective. Lastly, the United States argues that the Decree is consistent with Article 2.8 of the TBT Agreement, since it would be inappropriate to express the technical regulation in any way other than as a prohibition on the use of asbestos.

(b) “Like Products” in Article III:4 of the GATT 1994

48. The United States submits that the Panel erred in concluding that asbestos fibres and substitute fibres are “like products” under Article III:4 of the GATT 1994. The Panel erred in law in concluding that, in examining the properties, nature and quality of asbestos, it could not take into account the fact that asbestos differs from other fibres because it splits longitudinally into narrow, or thin, fibres, and has a high potential to release particles that possess certain characteristics, and in concluding that, in examining consumer tastes and habits, it could not take account of the proven carcinogenic nature of asbestos. In so proceeding, the Panel ignored the single most important distinguishing feature between asbestos and its substitutes. The Panel also wrongly inflated the significance of another factor – the end uses of products concerned. In the view of the United States, the
application of a proper "like product" analysis should lead the Appellate Body to find that asbestos is not "like" its substitute fibres, and that asbestos-containing products are not "like" asbestos-free products and, therefore, conclude that the Decree does not violate Article III:4 of the GATT 1994.

(c) Article XX(b) of the GATT 1994 and Article 11 of the DSU

49. Should the Appellate Body resort to Article XX(b) of the GATT 1994, the United States urges the Appellate Body to find that the Decree is permissible under Article XX(b). Canada’s appeal on this issue is based on criticism of the Panel’s findings with respect to the scientific information before it, and that Canada erroneously asserts that Article 11 of the DSU requires the Panel to decide which scientific view is the correct one. However, the role of a panel, under Article 11 of the DSU, is to make an objective assessment of the facts before it, and to evaluate whether there is a rational or objective relationship between the measure at issue and the scientific basis asserted for the measure. The United States argues that the Panel acted consistently with this mandate in finding that the Decree is necessary to protect human health, and the Appellate Body should not disturb this finding.

III. Preliminary Procedural Matter

50. On 27 October 2000, we wrote to the parties and the third parties indicating that we were mindful that, in the proceedings before the Panel in this case, the Panel received five written submissions from non-governmental organizations, two of which the Panel decided to take into account. In our letter, we recognized the possibility that we might receive submissions in this appeal from persons other than the parties and the third parties to this dispute, and stated that we were of the view that the fair and orderly conduct of this appeal could be facilitated by the adoption of appropriate procedures, for the purposes of this appeal only, pursuant to Rule 16(1) of the Working Procedures, to deal with any possible submissions received from such persons. To this end, we invited the parties and the third parties in this appeal to submit their comments on a number of questions. These related to: whether we should adopt a “request for leave” procedure; what procedures would be needed to ensure that the parties and third parties would have a full and adequate opportunity to respond to submissions that might be received; and whether we should take any other points into consideration if we decided to adopt a “request for leave” procedure. On 3 November 2000, all of the parties and third parties responded in writing to our letter of 27 October. Canada, the European Communities and Brazil considered that issues pertaining to any such procedure should be dealt with by the WTO Members themselves. The United States welcomed adoption of a request for leave procedure, and Zimbabwe indicated that it had no specific reasons to oppose adoption of a request for leave procedure. Without prejudice to their positions, Canada, the European Communities and the United States each made a number of suggestions regarding any such procedure that might be adopted.

51. On 7 November 2000, and after consultations among all seven Members of the Appellate Body, we adopted, pursuant to Rule 16(1) of the Working Procedures, an additional procedure, for the purposes of this appeal only, to deal with written submissions received from persons other than the parties and third parties to this dispute (the “Additional Procedure”). The Additional Procedure was communicated to the parties and third parties in this appeal on 7 November 2000. On 8 November 2000, the Chairman of the Appellate Body informed the Chairman of the Dispute Settlement Body, in writing, of the Additional Procedure adopted, and this letter was circulated, for information, as a dispute settlement document to the Members of the WTO. In that communication, the Chairman of the Appellate Body stated that:

... This additional procedure has been adopted by the Division hearing this appeal for the purposes of this appeal only pursuant to Rule 16(1) of the Working Procedures for Appellate Review, and is not a new working procedure drawn up by the Appel-
late Body pursuant to paragraph 9 of Article 17 of the Understanding on Rules and Procedures Governing the Settlement of Disputes. (original emphasis)

The Additional Procedure was posted on the WTO website on 8 November 2000.

52. The Additional Procedure provided:

1. In the interests of fairness and orderly procedure in the conduct of this appeal, the Division hearing this appeal has decided to adopt, pursuant to Rule 16(1) of the Working Procedures for Appellate Review, and after consultations with the parties and third parties to this dispute, the following additional procedure for purposes of this appeal only.

2. Any person, whether natural or legal, other than a party or a third party to this dispute, wishing to file a written brief with the Appellate Body, must apply for leave to file such a brief from the Appellate Body by noon on Thursday, 16 November 2000.

3. An application for leave to file such a written brief shall:

(a) be made in writing, be dated and signed by the applicant, and include the address and other contact details of the applicant;

(b) be in no case longer than three typed pages;

(c) contain a description of the applicant, including a statement of the membership and legal status of the applicant, the general objectives pursued by the applicant, the nature of the activities of the applicant, and the sources of financing of the applicant;

(d) specify the nature of the interest the applicant has in this appeal;

(e) identify the specific issues of law covered in the Panel Report and legal interpretations developed by the Panel that are the subject of this appeal, as set forth in the Notice of Appeal (WT/DS135/8) dated 23 October 2000, which the
applicant intends to address
in its written brief;

(f) state why it would be desirable, in the interests of achieving, a satisfactory settlement of the matter at issue, in accordance with the rights and obligations of WTO Members under the DSU and the other covered agreements, for the Appellate Body to grant the applicant leave to file a written brief in this appeal; and

indicate, in particular, in what way the applicant will make a contribution to the resolution of this dispute that is not likely to be repetitive of what has been already submitted by a party or third party to this dispute; and

(g) contain a statement disclosing whether the applicant has any relationship, direct or indirect, with any party or any third party to this dispute, as well as whether it has, or will, receive any assistance, financial or otherwise, from a party or a third party to this dispute in the preparation of its application for leave or its written brief.

4. The Appellate Body will review and consider each application for leave to file a written brief and will, without delay, render a decision whether to grant or deny such leave.

5. The grant of leave to file a brief by the Appellate Body does not imply that the Appellate Body will address, in its Report, the legal arguments made in such a brief.

6. Any person, other than a party or a third party to this dispute, granted leave to file a written brief with the Appellate Body, must file its brief with the Appellate Body Secretariat by noon on Monday, 27 November 2000.

7. A written brief filed with the Appellate Body by an applicant granted leave to file such a brief shall:

   (a) be dated and signed by the person filing the brief;
(b) be concise and in no case longer than 20 typed pages, including any appendices; and

(c) set out a precise statement, strictly limited to legal arguments, supporting the applicant’s legal position on the issues of law or legal interpretations in the Panel Report with respect to which the applicant has been granted leave to file a written brief.

8. An applicant granted leave shall, in addition to filing its written brief with the Appellate Body Secretariat, also serve a copy of its brief on all the parties and third parties to the dispute by noon on Monday, 27 November 2000.

9. The parties and the third parties to this dispute will be given a full and adequate opportunity by the Appellate Body to comment on and respond to any written brief filed with the Appellate Body by an applicant granted leave under this procedure. (original emphasis)

53. The Appellate Body received 13 written submissions from non-governmental organizations relating to this appeal that were not submitted in accordance with the Additional Procedure. Several of these were received while we were considering the possible adoption of an additional procedure. After the adoption of the Additional Procedure, each of these 13 submissions was returned to its sender, along with a letter informing the sender of the procedure adopted by the Division hearing this appeal and a copy of the Additional Procedure. Only one of these associations, the Korea Asbestos Association, subsequently submitted a request for leave in accordance with the Additional Procedure.

54. By letter dated 15 November 2000, Canada and the European Communities jointly requested that they be provided with copies of all applications filed pursuant to the Additional Procedure, and of the decision taken by the Appellate Body in respect of each such application. All such documents were subsequently provided to the parties and third parties in this dispute.

55. Pursuant to the Additional Procedure, the Appellate Body received 17 applications requesting leave to file a written brief in this appeal. Six of these 17 applications were received after the deadline specified in paragraph 2 of the Additional Procedure and, for this reason, leave to file a written brief was denied to these six applicants. Each such applicant was sent a copy of our decision denying its application for leave because the application was not filed in a timely manner.

56. The Appellate Body received 11 applications for leave to file a written brief in this appeal within the time limits specified in paragraph 2 of the Additional Procedure. We carefully reviewed and considered each of these applications in accordance with the Additional Procedure and, in each case, decided to deny leave to file a written brief. Each applicant was sent a copy of our decision denying its application for leave for failure to comply sufficiently with all the requirements set forth in paragraph 3 of the Additional Procedure.
57. We received a written brief from the Foundation for International Environmental Law and Development, on its behalf and on behalf of Ban Asbestos (International and Virtual) Network, Greenpeace International, International Ban Asbestos Secretariat, and World Wide Fund for Nature, International, dated 6 February 2001. As we had already denied, in accordance with the Additional Procedure, an application from these organizations for leave to file a written brief in this appeal 33, we did not accept this brief.

IV. Issues Raised in this Appeal

58. This appeal raises the following issues:

(a) whether the Panel erred in its interpretation of the term “technical regulation” in Annex 1.1 of the TBT Agreement in finding, in paragraph 8.72(a) of the Panel Report, that “the part of the Decree relating to the ban on imports of asbestos and asbestos-containing products” does not constitute a “technical regulation”;

(b) whether the Panel erred in its interpretation and application of the term “like products” in Article III:4 of the GATT 1994 in finding, in paragraph 8.144 of the Panel Report, that chrysotile asbestos fibres are “like” PVA, cellulose and glass fibres, and in finding, in paragraph 8.150 of the Panel Report, that cement-based products containing chrysotile asbestos fibres are “like” cement-based products containing polyvinyl alcohol, cellulose and glass fibres;

(c) whether the Panel erred in finding that the measure at issue is “necessary to protect human … life or health” under Article XX(b) of the GATT 1994, and whether, in carrying out its examination under Article XX(b) of the GATT 1994, the Panel failed to make an objective assessment of the matter under Article 11 of the DSU; and

(d) whether the Panel erred in its interpretation of Article XXIII:1(b) of the GATT 1994 in finding that that provision applies to a measure which falls within the scope of application of other provisions of the GATT 1994, and in finding that Article XXIII:1(b) applies to measures which pursue health objectives.

V. TBT Agreement

59. Before the Panel, Canada claimed that the measure at issue is inconsistent with Articles 2.1, 2.2, 2.4 and 2.8 of the TBT Agreement. Each of these provisions applies solely to “technical regulations”. Thus, a threshold issue in the examination of Canada’s claims under the TBT Agreement is whether the measure at issue is a “technical regulation”.

60. In addressing this threshold issue, the Panel examined the nature and structure of the measure to assess how the TBT Agreement might apply to it. For this examination, the Panel decided that it would be appropriate to examine the measure in two stages. First, the Panel examined “the part of the Decree prohibiting the marketing of asbestos and asbestos-containing products”; next, the Panel analyzed the “exceptions” in the Decree. 44 The Panel concluded that the part of the Decree containing the prohibitions is not a “technical regulation”, and that, therefore, the TBT Agreement does not apply to this part of the Decree. 45 However, the Panel also concluded that the part of the Decree containing the exceptions does constitute a “technical regulation”, and that, therefore, the TBT Agreement applies to that part of the Decree. On this basis, the Panel decided not to examine Canada’s claims under the TBT Agreement because, it said, those claims relate solely to the part of the Decree containing the prohibitions, which, in the Panel’s view, does not constitute a “technical regulation”, and, therefore, the TBT Agreement does not apply. 46
61. In concluding that the part of the Decree containing the prohibitions is not a “technical regulation”, the Panel found that:

- The measure affects one or more given products;
- The measure specifies the technical characteristics of the product(s) which allow them to be marketed in the Member that took the measure;
- Compliance is mandatory.

62. Canada appeals the Panel’s finding that the TBT Agreement does not apply to the part of the Decree relating to the prohibitions on imports of asbestos and asbestos-containing products. According to Canada, the Panel erred in considering the part of the Decree relating to those prohibitions separately from the part of the Decree relating to the exceptions to those prohibitions, and, therefore, the Panel should have examined the Decree as a single, unified measure. Furthermore, Canada argues that the Panel erred in its interpretation of a “technical regulation”, as defined in Annex 1.1 to the TBT Agreement, because, in Canada’s view, a general prohibition can be a “technical regulation”.

63. We start with the measure at issue. It is clear from Canada’s request for the establishment of a panel that Canada’s complaint concerns Decree 96-1133 as a whole. The Decree, in essence, consists of prohibitions on asbestos fibres and on products containing asbestos (Article 1), coupled with limited and temporary exceptions from the prohibitions for certain “existing materials, products or devices containing chrysotile fibre” (Article 2). The remaining operative provisions of the Decree contain additional rules governing the grant of an exception (Articles 3 and 4) and the imposition of penalties for violation of the prohibitions in Article 1 (Article 5). Furthermore, certain used “vehicles” and “agricultural and forestry machinery” are entirely excluded, until 31 December 2001, from certain aspects of the prohibitions in Article 1, namely, from the prohibitions on “possession for sale, offering for sale and transfer under any title” (Article 7).

64. In our view, the proper legal character of the measure at issue cannot be determined unless the measure is examined as a whole. Article 1 of the Decree contains broad, general prohibitions on asbestos and products containing asbestos. However, the scope and generality of those prohibitions can only be understood in light of the exceptions to it which, albeit for a limited period, permit, inter alia, the use of certain products containing asbestos and, principally, products containing chrysotile asbestos fibres. The measure is, therefore, not a total prohibition on asbestos fibres, because it also includes provisions that permit, for a limited duration, the use of asbestos in certain situations. Thus, to characterize the measure simply as a general prohibition, and to examine it as such, overlooks the complexities of the measure, which include both prohibitive and permissive elements. In addition, we observe that the exceptions in the measure would have no autonomous legal significance in the absence of the prohibitions. We, therefore, conclude that the measure at issue is to be examined as an integrated whole, taking into account, as appropriate, the prohibitive and the permissive elements that are part of it.

65. Accordingly, we reverse the Panel’s two-stage interpretive approach of examining, first, the application of the TBT Agreement to the prohibitions contained in the measure and, second and separately, its application to the exceptions contained in the measure.

66. We turn now to the term “technical regulation” and to the considerations that must go into interpreting the term. Article 1.2 of the TBT Agreement provides that, for the
purposes of this Agreement, the meanings given in Annex 1 apply. Annex 1.1 of the TBT Agreement defines a “technical regulation” as a:

Document which lays down product characteristics or their related processes and production methods, including the applicable administrative provisions, with which compliance is mandatory. It may also include or deal exclusively with terminology, symbols, packaging, marking or labelling requirements as they apply to a product, process or production method. (emphasis added)

67. The heart of the definition of a “technical regulation” is that a “document” must “lay down” – that is, set forth, stipulate or provide – “product characteristics”. The word “characteristic” has a number of synonyms that are helpful in understanding the ordinary meaning of that word, in this context. Thus, the “characteristics” of a product include, in our view, any objectively definable “features”, “qualities”, “attributes”, or other “distinguishing mark” of a product. Such “characteristics” might relate, inter alia, to a product’s composition, size, shape, colour, texture, hardness, tensile strength, flammability, conductivity, density, or viscosity. In the definition of a “technical regulation” in Annex 1.1, the TBT Agreement itself gives certain examples of “product characteristics” – “terminology, symbols, packaging, marking or labelling requirements”. These examples indicate that “product characteristics” include, not only features and qualities intrinsic to the product itself, but also related “characteristics”, such as the means of identification, the presentation and the appearance of a product. In addition, according to the definition in Annex 1.1 of the TBT Agreement, a “technical regulation” may set forth the “applicable administrative provisions” for products which have certain “characteristics”. Further, we note that the definition of a “technical regulation” provides that such a regulation “may also include or deal exclusively with terminology, symbols, packaging, marking or labelling requirements”. (emphasis added) The use here of the word “exclusively” and the disjunctive word “or” indicates that a “technical regulation” may be confined to laying down only one or a few “product characteristics”.

68. The definition of a “technical regulation” in Annex 1.1 of the TBT Agreement also states that “compliance” with the “product characteristics” laid down in the “document” must be “mandatory”. A “technical regulation” must, in other words, regulate the “characteristics” of products in a binding or compulsory fashion. It follows that, with respect to products, a “technical regulation” has the effect of prescribing or imposing one or more “characteristics” – “features”, “qualities”, “attributes”, or other “distinguishing mark”.

69. “Product characteristics” may, in our view, be prescribed or imposed with respect to products in either a positive or a negative form. That is, the document may provide, positively, that products “must possess” certain “characteristics”, or the document may require, negatively, that products “must not possess” certain “characteristics”. In both cases, the legal result is the same: the document “lays down” certain binding “characteristics” for products, in one case affirmatively, and in the other by negative implication.

70. A “technical regulation” must, of course, be applicable to an identifiable product, or group of products. Otherwise, enforcement of the regulation will, in practical terms, be impossible. This consideration also underlies the formal obligation, in Article 2.9.2 of the TBT Agreement, for Members to notify other Members, through the WTO Secretariat, “of the products to be covered” by a proposed “technical regulation”. (emphasis added) Clearly, compliance with this obligation requires identification of the product coverage of a technical regulation. However, in contrast to what the Panel suggested, this does not mean that a “technical regulation” must apply to “given” products which are actually named, identified or specified in the regulation. (emphasis added) Although the TBT Agreement clearly applies to “products” generally, nothing in the text of that Agreement suggests that
those products need be named or otherwise expressly identified in a “technical regulation”. Moreover, there may be perfectly sound administrative reasons for formulating a “technical regulation” in a way that does not expressly identify products by name, but simply makes them identifiable – for instance, through the “characteristic" that is the subject of regulation.

71. With these considerations in mind, we examine whether the measure at issue is a “technical regulation”. Decree 96-1133 aims primarily at the regulation of a named product, asbestos. The first and second paragraphs of Article 1 of the Decree impose a prohibition on asbestos fibres, as such. This prohibition on these fibres does not, in itself, prescribe or impose any “characteristics” on asbestos fibres, but simply bans them in their natural state. Accordingly, if this measure consisted only of a prohibition on asbestos fibres, it might not constitute a “technical regulation”.

72. There is, however, more to the measure than this prohibition on asbestos fibres. It is not contested that asbestos fibres have no known use in their raw mineral form. Thus, the regulation of asbestos can only be achieved through the regulation of products that contain asbestos fibres. This, too, is addressed by the Decree before us. An integral and essential aspect of the measure is the regulation of "products containing asbestos fibres", which are also prohibited by Article 1, paragraphs I and II of the Decree. It is important to note here that, although formulated negatively – products containing asbestos are prohibited – the measure, in this respect, effectively prescribes or imposes certain objective features, qualities or “characteristics” on all products. That is, in effect, the measure provides that all products must not contain asbestos fibres. Although this prohibition against products containing asbestos applies to a large number of products, and although it is, indeed, true that the products to which this prohibition applies cannot be determined from the terms of the measure itself, it seems to us that the products covered by the measure are identifiable: all products must be asbestos free; any products containing asbestos are prohibited. We also observe that compliance with the prohibition against products containing asbestos is mandatory and is, indeed, enforceable through criminal sanctions.

73. Articles 2, 3 and 4 of the Decree also contain certain exceptions to the prohibitions found in Article 1 of the Decree. As we have already noted, these exceptions would have no meaning in the absence of the rest of the measure because they define the scope of the prohibitions in the measure. The nature of these exceptions is to permit the use of certain products containing chrysotile asbestos fibres, subject to compliance with strict administrative requirements. The scope of the exceptions is determined by an “exhaustive list” of products that are permitted to contain chrysotile asbestos fibres, which is promulgated and reviewed annually by a government Minister. The inclusion of a product in the list of exceptions depends on the absence of an acceptable alternative fibre for incorporation into a particular product, and the demonstrable provision of “all technical guarantees of safety”. Any person seeking to avail himself of these limited exceptions must provide a detailed justification to the authorities, complete with necessary supporting documentation concerning “the state of scientific and technological progress”. Compliance with these administrative requirements is mandatory.

74. Like the Panel, we consider that, through these exceptions, the measure sets out the “applicable administrative provisions, with which compliance is mandatory” for products with certain objective “characteristics”. The exceptions apply to a narrowly defined group of products with particular “characteristics”. Although these products are not named, the measure provides criteria which permit their identification, both by reference to the qualities the excepted products must possess and by reference to the list promulgated by the Minister.

75. Viewing the measure as an integrated whole, we see that it lays down “characteristics” for all products that might contain asbestos, and we see also that it lays down the “applicable administrative provisions” for certain products containing chrysotile asbestos fibres which are excluded from the prohibitions in the measure. Accordingly, we find that the measure is a “document” which “lays down product characteristics ... including the
applicable administrative provisions, with which compliance is mandatory.” For these reasons, we conclude that the measure constitutes a “technical regulation” under the TBT Agreement.

76. We, therefore, reverse the Panel’s finding, in paragraph 8.72(a) of the Panel Report, that the TBT Agreement “does not apply to the part of the Decree relating to the ban on imports of asbestos and asbestos-containing products because that part does not constitute a ‘technical regulation’ within the meaning of Annex 1.1 to the TBT Agreement.”

77. We note, however – and we emphasize – that this does not mean that all internal measures covered by Article III:4 of the GATT 1994 “affecting” the “sale, offering for sale, purchase, transportation, distribution or use” of a product are, necessarily, “technical regulations” under the TBT Agreement. Rather, we rule only that this particular measure, the Decree at stake, falls within the definition of a “technical regulation” given in Annex 1.1 of that Agreement.

78. As we have reached a different conclusion from the Panel’s regarding the applicability of the TBT Agreement to the measure, we now consider whether it is appropriate for us to rule on the claims made by Canada relating to the TBT Agreement. In previous appeals, we have, on occasion, completed the legal analysis with a view to facilitating the prompt settlement of the dispute, pursuant to Article 3.3 of the DSU. However, we have insisted that we can do so only if the factual findings of the panel and the undisputed facts in the panel record provide us with a sufficient basis for our own analysis. If that has not been the case, we have not completed the analysis.

79. The need for sufficient facts is not the only limit on our ability to complete the legal analysis in any given case. In Canada – Periodicals, we reversed the panel’s conclusion that the measure at issue was inconsistent with Article III:2, first sentence, of the GATT 1994, and we then proceeded to examine the United States’ claims under Article III:2, second sentence, which the panel had not examined at all. However, in embarking there on an analysis of a provision that the panel had not considered, we emphasized that “the first and second sentences of Article III:2 are closely related” and that those two sentences are “part of a logical continuum.” (emphasis added)

80. In this appeal, Canada’s outstanding claims were made under Articles 2.1, 2.2, 2.4 and 2.8 of the TBT Agreement. We observe that, although the TBT Agreement is intended to “further the objectives of GATT 1994”, it does so through a specialized legal regime that applies solely to a limited class of measures. For these measures, the TBT Agreement imposes obligations on Members that seem to be different from, and additional to, the obligations imposed on Members under the GATT 1994.

81. As the Panel decided not to examine Canada’s four claims under the TBT Agreement, it made no findings, at all, regarding any of these claims. Moreover, the meaning of the different obligations in the TBT Agreement has not previously been the subject of any interpretation or application by either panels or the Appellate Body. Similarly, the provisions of the Tokyo Round Agreement on Technical Barriers to Trade, which preceded the TBT Agreement and which contained obligations similar to those in the TBT Agreement, were also never the subject of even a single ruling by a panel.

82. In light of their novel character, we consider that Canada’s claims under the TBT Agreement have not been explored before us in depth. As the Panel did not address these claims, there are no “issues of law” or “legal interpretations” regarding them to be analyzed by the parties, and reviewed by us under Article 17.6 of the DSU. We also observe that the sufficiency of the facts on the record depends on the reach of the provisions of the TBT Agreement claimed to apply – a reach that has yet to be determined.

83. With this particular collection of circumstances in mind, we consider that we do not have an adequate basis properly to examine Canada’s claims under Article 2.1, 2.2, 2.4 and 2.8 of the TBT Agreement and, accordingly, we refrain from so doing.
VI. “Like Products” in Article III:4 of the GATT 1994

A. Background

84. In addressing Canada’s claims under Article III:4 of the GATT 1994, the Panel examined whether two different sets of products are “like”. First, the Panel examined whether chrysotile asbestos fibres are “like” certain other fibres, namely polyvinyl alcohol fibres (“PVA”), cellulose and glass fibres (PVA, cellulose and glass fibres are all collectively referred to, in the remainder of this Report, as “PCG fibres”). The Panel concluded that chrysotile asbestos and PCG fibres are all “like products” under Article III:4. The Panel next examined whether cement-based products containing chrysotile asbestos fibres are “like” cement-based products containing one of the PCG fibres. The Panel also concluded that all these cement-based products are “like”.

85. In examining the “likeness” of these two sets of products, the Panel adopted an approach based on the Report of the Working Party on Border Tax Adjustments. Under that approach, the Panel employed four general criteria in analyzing “likeness”: (i) the properties, nature and quality of the products; (ii) the end-uses of the products; (iii) consumers’ tastes and habits; and, (iv) the tariff classification of the products. The Panel declined to apply “a criterion on the risk of a product”, “neither in the criterion relating to the properties, nature and quality of the product, nor in the other likeness criteria ….”

86. On appeal, the European Communities requests that we reverse the Panel’s findings that the two sets of products examined by the Panel are “like products” under Article III:4 of the GATT 1994, and requests, in consequence, that we reverse the Panel’s finding that the measure is inconsistent with Article III:4 of the GATT 1994. The European Communities contends that the Panel erred in its interpretation and application of the concept of “like products”, in particular, in excluding from its analysis consideration of the health risks associated with chrysotile asbestos fibres. According to the European Communities, in this case, Article III:4 calls for an analysis of the health objective of the regulatory distinction made in the measure between asbestos fibres, and between products containing asbestos fibres, and all other products. The European Communities argues that, under Article III:4, products should not be regarded as “like” unless the regulatory distinction drawn between them “entails [a] shift in the competitive opportunities” in favour of domestic products.

B. Meaning of the Term “Like Products” in Article III:4 of the GATT 1994

87. Article III:4 of the GATT 1994 reads, in relevant part:

The products of the territory of any Member imported into the territory of any other Member shall be accorded treatment no less favourable than that accorded to like products of national origin in respect of all laws, regulations and requirements affecting their internal sale, offering for sale, purchase, transportation, distribution or use. (emphasis added)

88. The European Communities’ appeal on this point turns on the interpretation of the word “like” in the term “like products” in Article III:4 of the GATT 1994. Thus, this appeal provides us with our first occasion to examine the meaning of the word “like” in Article III:4 of the GATT 1994. Yet, this appeal is, of course, not the first time that the term “like products” has been addressed in GATT or WTO dispute settlement proceedings. Indeed, the term “like product” appears in many different provisions of the covered agreements, for example, in Articles I:1, II:2, III:2, III:4, VI:1, IX:1, XI:2(c), XIII:1, XVI:4 and XIX:1 of the GATT 1994. The term is also a key concept in the Agreement on Subsidies
and Countervailing Measures, the Agreement on Implementation of Article VI of the General Agreement on Tariffs and Trade 1994 (the “Anti-Dumping Agreement”), the Agreement on Safeguards and other covered agreements. In some cases, such as in Article 2.6 of the Anti-Dumping Agreement, the term is given a specific meaning to be used “[t]hroughout [the] Agreement”, while in others, it is not. In each of the provisions where the term “like products” is used, the term must be interpreted in light of the context, and of the object and purpose, of the provision at issue, and of the object and purpose of the covered agreement in which the provision appears. Accordingly, and as we observed in an earlier case concerning Article III:2 of the GATT 1994:

... there can be no one precise and absolute definition of what is “like”. The concept of “likeness” is a relative one that evokes the image of an accordion. The accordion of “likeness” stretches and squeezes in different places as different provisions of the WTO Agreement are applied. The width of the accordion in any one of those places must be determined by the particular provision in which the term “like” is encountered as well as by the context and the circumstances that prevail in any given case to which that provision may apply. ... 60 (emphasis added)

89. It follows that, while the meaning attributed to the term “like products” in other provisions of the GATT 1994, or in other covered agreements, may be relevant context in interpreting Article III:4 of the GATT 1994, the interpretation of “like products” in Article III:4 need not be identical, in all respects, to those other meanings.

90. Bearing these considerations in mind, we turn now to the ordinary meaning of the word “like” in the term “like products” in Article III:4. According to one dictionary, “like” means:

Having the same characteristics or qualities as some other ... thing; of approximately identical shape, size, etc., with something else; similar. 61

91. This meaning suggests that “like” products are products that share a number of identical or similar characteristics or qualities. The reference to “similar” as a synonym of “like” also echoes the language of the French version of Article III:4, “produits similaires”, and the Spanish version, “productos similares”, which, together with the English version, are equally authentic. 62

92. However, as we have previously observed, “dictionary meanings leave many interpretive questions open.” 63 In particular, this definition does not resolve three issues of interpretation. First, this dictionary definition of “like” does not indicate which characteristics or qualities are important in assessing the “likeness” of products under Article III:4. For instance, most products will have many qualities and characteristics, ranging from physical properties such as composition, size, shape, texture, and possibly taste and smell, to the end-uses and applications of the product. Second, this dictionary definition provides no guidance in determining the degree or extent to which products must share qualities or characteristics in order to be “like products” under Article III:4. Products may share only very few characteristics or qualities, or they may share many. Thus, in the abstract, the term “like” can encompass a spectrum of differing degrees of “likeness” or “similarity”. Third, this dictionary definition of “like” does not indicate from whose perspective “likeness” should be judged. For instance, ultimate consumers may have a view about the “likeness” of two products that is very different from that of the inventors or producers of those products.

93. To begin to resolve these issues, we turn to the relevant context of Article III:4 of the GATT 1994. In that respect, we observe that Article III:2 of the GATT 1994, which deals
with the internal tax treatment of imported and domestic products, prevents Members, through its first sentence, from imposing internal taxes on imported products “in excess of those applied ... to like domestic products.” (emphasis added) In previous Reports, we have held that the scope of “like” products in this sentence is to be construed “narrowly”. This reading of “like” in Article III:2 might be taken to suggest a similarly narrow reading of “like” in Article III:4, since both provisions form part of the same Article. However, both of these paragraphs of Article III constitute specific expressions of the overarching, “general principle”, set forth in Article III:1 of the GATT 1994. As we have previously said, the “general principle” set forth in Article III:1 “informs” the rest of Article III and acts “as a guide to understanding and interpreting the specific obligations contained” in the other paragraphs of Article III, including paragraph 4. Thus, in our view, Article III:1 has particular contextual significance in interpreting Article III:4, as it sets forth the “general principle” pursued by that provision. Accordingly, in interpreting the term “like products” in Article III:4, we must turn, first, to the “general principle” in Article III:1, rather than to the term “like products” in Article III:2.

94. In addition, we observe that, although the obligations in Articles III:2 and III:4 both apply to “like products”, the text of Article III:2 differs in one important respect from the text of Article III:4. Article III:2 contains two separate sentences, each imposing distinct obligations: the first lays down obligations in respect of “like products”, while the second lays down obligations in respect of “directly competitive or substitutable” products. By contrast, Article III:4 applies only to “like products” and does not include a provision equivalent to the second sentence of Article III:2. We note that, in this dispute, the Panel did not examine, at all, the significance of this textual difference between paragraphs 2 and 4 of Article III.

95. For us, this textual difference between paragraphs 2 and 4 of Article III has considerable implications for the meaning of the term “like products” in these two provisions. In Japan – Alcoholic Beverages, we concluded, in construing Article III:2, that the two separate obligations in the two sentences of Article III:2 must be interpreted in a harmonious manner that gives meaning to both sentences in that provision. We observed there that the interpretation of one of the sentences necessarily affects the interpretation of the other. Thus, the scope of the term “like products” in the first sentence of Article III:2 affects, and is affected by, the scope of the phrase “directly competitive or substitutable” products in the second sentence of that provision. We said in Japan – Alcoholic Beverages:

Because the second sentence of Article III:2 provides for a separate and distinctive consideration of the protective aspect of a measure in examining its application to a broader category of products that are not “like products” as contemplated by the first sentence, we agree with the Panel that the first sentence of Article III:2 must be construed narrowly so as not to condemn measures that its strict terms are not meant to condemn. Consequently, we agree with the Panel also that the definition of “like products” in Article III:2, first sentence, should be construed narrowly.

96. In construing Article III:4, the same interpretive considerations do not arise, because the “general principle” articulated in Article III:1 is expressed in Article III:4, not through two distinct obligations, as in the two sentences in Article III:2, but instead through a single obligation that applies solely to “like products”. Therefore, the harmony that we have attributed to the two sentences of Article III:2 need not and, indeed, cannot be replicated in interpreting Article III:4. Thus, we conclude that, given the textual difference between Articles III:2 and III:4, the “accordion” of likeness stretches in a different way in Article III:4.
97. We have previously described the “general principle” articulated in Article III:1 as follows:

The broad and fundamental purpose of Article III is to avoid protectionism in the application of internal tax and regulatory measures. More specifically, the purpose of Article III “is to ensure that internal measures not be applied to imported and domestic products so as to afford protection to domestic production”. Toward this end, Article III obliges Members of the WTO to provide equality of competitive conditions for imported products in relation to domestic products. Article III protects expectations not of any particular trade volume but rather of the equal competitive relationship between imported and domestic products. (emphasis added)

98. As we have said, although this “general principle” is not explicitly invoked in Article III:4, nevertheless, it “informs” that provision. Therefore, the term “like product” in Article III:4 must be interpreted to give proper scope and meaning to this principle. In short, there must be consonance between the objective pursued by Article III, as articulated in the “general principle” articulated in Article III:1, and the interpretation of the specific expression of this principle in the text of Article III:4. This interpretation must, therefore, reflect that, in endeavouring to ensure “equality of competitive conditions”, the “general principle” in Article III seeks to prevent Members from applying internal taxes and regulations in a manner which affects the competitive relationship, in the marketplace, between the domestic and imported products involved, “so as to afford protection to domestic production.”

99. As products that are in a competitive relationship in the marketplace could be affected through treatment of imports “less favourable” than the treatment accorded to domestic products, it follows that the word “like” in Article III:4 is to be interpreted to apply to products that are in such a competitive relationship. Thus, a determination of “likeness” under Article III:4 is, fundamentally, a determination about the nature and extent of a competitive relationship between and among products. In saying this, we are mindful that there is a spectrum of degrees of “competitiveness” or “substitutability” of products in the marketplace, and that it is difficult, if not impossible, in the abstract, to indicate precisely where on this spectrum the word “like” in Article III:4 of the GATT 1994 falls. We are not saying that all products which are in some competitive relationship are “like products” under Article III:4. In ruling on the measure at issue, we also do not attempt to define the precise scope of the word “like” in Article III:4. Nor do we wish to decide if the scope of “like products” in Article III:4 is co-extensive with the combined scope of “like” and “directly competitive or substitutable” products in Article III:2. However, we recognize that the relationship between these two provisions is important, because there is no sharp distinction between fiscal regulation, covered by Article III:2, and non-fiscal regulation, covered by Article III:4. Both forms of regulation can often be used to achieve the same ends. It would be incongruous if, due to a significant difference in the product scope of these two provisions, Members were prevented from using one form of regulation – for instance, fiscal – to protect domestic production of certain products, but were able to use another form of regulation – for instance, non-fiscal – to achieve those ends. This would frustrate a consistent application of the “general principle” in Article III:1. For these reasons, we conclude that the scope of “like” in Article III:4 is broader than the scope of “like” in Article III:2, first sentence. Nonetheless, we note, once more, that Article III:2 extends not only to “like products”, but also to products which are “directly competitive or substitutable”, and that Article III:4 extends only to “like products”. In view of this different language, and although we need not rule, and do not rule, on the precise product scope of Article III:4, we do conclude that the product scope of Article III:4, although broader than the first sentence of Article III:2, is certainly not broader than the combined product scope of the two sentences of Article III:2 of the GATT 1994.
100. We recognize that, by interpreting the term “like products” in Article III:4 in this way, we give that provision a relatively broad product scope – although no broader than the product scope of Article III:2. In so doing, we observe that there is a second element that must be established before a measure can be held to be inconsistent with Article III:4. Thus, even if two products are “like”, that does not mean that a measure is inconsistent with Article III:4. A complaining Member must still establish that the measure accords to the group of “like” imported products “less favourable treatment” than it accords to the group of “like” domestic products. The term “less favourable treatment” expresses the general principle, in Article III:1, that internal regulations “should not be applied … so as to afford protection to domestic production”. If there is “less favourable treatment” of the group of “like” imported products, there is, conversely, “protection” of the group of “like” domestic products. However, a Member may draw distinctions between products which have been found to be “like”, without, for this reason alone, according to the group of “like” imported products “less favourable treatment” than that accorded to the group of “like” domestic products. In this case, we do not examine further the interpretation of the term “treatment no less favourable” in Article III:4, as the Panel’s findings on this issue have not been appealed or, indeed, argued before us.

C. Assessing the “Likeness” of Products under Article III:4 of the GATT 1994

101. We turn to consideration of how a treaty interpreter should proceed in determining whether products are “like” under Article III:4. As in Article III:2, in this determination, “[n]o one approach … will be appropriate for all cases.” Rather, an assessment utilizing “an unavoidable element of individual, discretionary judgement” has to be made on a case-by-case basis. The Report of the Working Party on Border Tax Adjustments outlined an approach for analyzing “likeness” that has been followed and developed since by several panels and the Appellate Body. This approach has, in the main, consisted of employing four general criteria in analyzing “likeness”: (i) the properties, nature and quality of the products; (ii) the end-uses of the products; (iii) consumers’ tastes and habits – more comprehensively termed consumers’ perceptions and behaviour – in respect of the products; and (iv) the tariff classification of the products. We note that these four criteria comprise four categories of “characteristics” that the products involved might share: (i) the physical properties of the products; (ii) the extent to which the products are capable of serving the same or similar end-uses; (iii) the extent to which consumers perceive and treat the products as alternative means of performing particular functions in order to satisfy a particular want or demand; and (iv) the international classification of the products for tariff purposes.

102. These general criteria, or groupings of potentially shared characteristics, provide a framework for analyzing the “likeness” of particular products on a case-by-case basis. These criteria are, it is well to bear in mind, simply tools to assist in the task of sorting and examining the relevant evidence. They are neither a treaty-mandated nor a closed list of criteria that will determine the legal characterization of products. More important, the adoption of a particular framework to aid in the examination of evidence does not dissolve the duty or the need to examine, in each case, all of the pertinent evidence. In addition, although each criterion addresses, in principle, a different aspect of the products involved, which should be examined separately, the different criteria are interrelated. For instance, the physical properties of a product shape and limit the end-uses to which the products can be devoted. Consumer perceptions may similarly influence – modify or even render obsolete – traditional uses of the products. Tariff classification clearly reflects the physical properties of a product.

103. The kind of evidence to be examined in assessing the “likeness” of products will, necessarily, depend upon the particular products and the legal provision at issue. When all the relevant evidence has been examined, panels must determine whether that evidence, as a whole, indicates that the products in question are “like” in terms of the legal provision at issue. We have noted that, under Article III:4 of the GATT 1994, the term “like products” is concerned with competitive relationships between and among products. Accordingly, whether the Border Tax Adjustments framework is adopted or not, it is impor-
tant under Article III:4 to take account of evidence which indicates whether, and to what extent, the products involved are – or could be – in a competitive relationship in the marketplace.

D. The Panel’s Findings and Conclusions on “Likeness” under Article III:4 of the GATT 1994

1. Overview

104. In this case, the European Communities argues that the Panel erred in its consideration of “likeness”, in particular, because it adopted an exclusively “commercial or market access approach” to the comparison of allegedly “like products”; placed excessive reliance on a single criterion, namely, end-use; and failed to include consideration of the health “risk” factors relating to asbestos. 73

105. Before considering these arguments, we think it helpful to summarize the way in which the Panel assessed the “likeness” of chrysotile asbestos fibres, on the one hand, and the PCG fibres – PVA, cellulose and glass fibres – on the other. It will be recalled that the Panel adopted the approach in the Border Tax Adjustments report, using the four general criteria mentioned above. 74 After reviewing the first criterion, “properties, nature and quality of the products”, the Panel conclude[d] that … chrysotile fibres are like PVA, cellulose and glass fibres.” 77 (emphasis added) In reaching this “conclusion”, the Panel found that it was not decisive that the products “do not have the same structure or chemical composition”, nor that asbestos is “unique”. Instead, the Panel focused on “market access” and whether the products have the “same applications” and can “replace” each other for some industrial uses. 73 The Panel also declined to “[i]ntroduce a criterion on the risk of a product”. 73

106. Under the second criterion, “end-use”, the Panel stated that it had already found, under the first criterion, that the products have “certain identical or at least similar end-uses” and that it did not, therefore, consider it necessary to elaborate further on this criterion. 80 The Panel declined to “take a position” on “consumers’ tastes and habits”, the third criterion, “[b]ecause this criterion would not provide clear results”. 81 The Panel observed that consumers’ tastes and habits are “very varied”. 82 Finally, the Panel did not regard as “decisive” the different “tariff classification” of the fibres. 83

107. Based on this reasoning, the Panel concluded that chrysotile asbestos fibres and PCG fibres are “like products” under Article III:4 of the GATT 1994. 84

108. The Panel next examined whether cement-based products containing chrysotile asbestos fibres are “like” cement-based products containing PCG fibres. 85 Applying the reasoning from its findings on fibres, and noting that the individual cement-based products have the same tariff classification, irrespective of their fibre content, the Panel concluded that these cement-based products are also “like” under Article III:4. 86

2. Chrysotile and PCG fibres

109. In our analysis of this issue on appeal, we begin with the Panel’s findings on the “likeness” of chrysotile asbestos and PCG fibres and, in particular, with the Panel’s overall approach to examining the “likeness” of these fibres. It is our view that, having adopted an approach based on the four criteria set forth in Border Tax Adjustments, the Panel should have examined the evidence relating to each of those four criteria and, then, weighed all of that evidence, along with any other relevant evidence, in making an overall determination of whether the products at issue could be characterized as “like”. Yet, the Panel expressed a “conclusion” that the products were “like” after examining only the first of the four criteria. The Panel then repeated that conclusion under the second criterion – without further analysis – before dismissing altogether the relevance of the third criterion and also before rejecting the differing tariff classifications under the fourth criterion. In our view, it was inappropriate for the Panel to express a “conclusion” after examining only one of the
four criteria. By reaching a “conclusion” without examining all of the criteria it had decided to examine, the Panel, in reality, expressed a conclusion after examining only some of the evidence. Yet, a determination on the “likeness” of products cannot be made on the basis of a partial analysis of the evidence, after examination of just one of the criteria the Panel said it would examine. For this reason, we doubt whether the Panel’s overall approach has allowed the Panel to make a proper characterization of the “likeness” of the fibres at issue.

110. We must next examine more closely the Panel’s treatment of the four individual criteria. We see the first criterion, “properties, nature and quality”, as intended to cover the physical qualities and characteristics of the products. In analyzing the “properties” of the products, the Panel said that, “because of its physical and chemical characteristics, asbestos is a unique product.” (emphasis added) The Panel expressly acknowledged that, based on physical properties alone, “[i]t could … be concluded that [the fibres] are not like products.” (emphasis added) However, to overcome that fact, the Panel adopted a “market access” approach to this first criterion. Thus, in the course of its examination of “properties”, the Panel went on to rely on “end-uses” – the second criterion – and on the fact that, in a “small number” of cases, the products have the “same applications” and can “replace” each other. The Panel then stated:

We therefore conclude that, taking into account the properties criterion, chrysotile fibres are like PVA, cellulose and glass fibres.

111. We believe that physical properties deserve a separate examination that should not be confused with the examination of end-uses. Although not decisive, the extent to which products share common physical properties may be a useful indicator of “likeness”. Furthermore, the physical properties of a product may also influence how the product can be used, consumer attitudes about the product, and tariff classification. It is, therefore, important for a panel to examine fully the physical character of a product. We are also concerned that it will be difficult for a panel to draw the appropriate conclusions from the evidence examined under each criterion if a panel’s approach does not clearly address each criterion separately, but rather entwines different, and distinct, elements of the analysis along the way.

112. In addition, we do not share the Panel’s conviction that when two products can be used for the same end-use, their “properties are then equivalent, if not identical.” (emphasis added) Products with quite different physical properties may, in some situations, be capable of performing similar or identical end-uses. Although the end-uses are then “equivalent”, the physical properties of the products are not thereby altered; they remain different. Thus, the physical “uniqueness” of asbestos that the Panel noted does not change depending on the particular use that is made of asbestos.

113. The European Communities argues that the inquiry into the physical properties of products must include a consideration of the risks posed by the product to human health. In examining the physical properties of the product at issue in this dispute, the Panel found that “it was not appropriate to apply the ‘risk’ criterion proposed by the EC.” The Panel said that to do so “would largely nullify the effect of Article XX(b)” of the GATT 1994. In reviewing this finding by the Panel, we note that neither the text of Article III:4 nor the practice of panels and the Appellate Body suggest that any evidence should be excluded a priori from a panel’s examination of “likeness”. Moreover, as we have said, in examining the “likeness” of products, panels must evaluate all of the relevant evidence. We are very much of the view that evidence relating to the health risks associated with a product may be pertinent in an examination of “likeness” under Article III:4 of the GATT 1994. We do not, however, consider that the evidence relating to the health risks associated with chrysotile asbestos fibres need be examined under a separate criterion, because we believe that this evidence can be evaluated under the existing criteria of physical properties, and of consumers’ tastes and habits, to which we will come below.
114. Panels must examine fully the physical properties of products. In particular, panels must examine those physical properties of products that are likely to influence the competitive relationship between products in the marketplace. In the case of chrysotile asbestos fibres, their molecular structure, chemical composition, and fibrillation capacity are important because the microscopic particles and filaments of chrysotile asbestos fibres are carcinogenic in humans, following inhalation. In this respect, we observe that, at paragraph 8.188 of its Report, the Panel made the following statements regarding chrysotile asbestos fibres:

... we note that the carcinogenicity of chrysotile fibres has been acknowledged for some time by international bodies. This carcinogenicity was confirmed by the experts consulted by the Panel, with respect to both lung cancers and mesotheliomas, even though the experts appear to acknowledge that chrysotile is less likely to cause mesotheliomas than amphiboles. We also note that the experts confirmed that the types of cancer concerned had a mortality rate of close to 100 per cent. We therefore consider that we have sufficient evidence that there is in fact a serious carcinogenic risk associated with the inhalation of chrysotile fibres. Moreover, in the light of the comments made by one of the experts, the doubts expressed by Canada with respect to the direct effects of chrysotile on mesotheliomas and lung cancers are not sufficient to conclude that an official responsible for public health policy would find that there was not enough evidence of the existence of a public health risk.

This carcinogenicity, or toxicity, constitutes, as we see it, a defining aspect of the physical properties of chrysotile asbestos fibres. The evidence indicates that PCG fibres, in contrast, do not share these properties, at least to the same extent. We do not see how this highly significant physical difference cannot be a consideration in examining the physical properties of a product as part of a determination of “likeness” under Article III:4 of the GATT 1994.

115. We do not agree with the Panel that considering evidence relating to the health risks associated with a product, under Article III:4, nullifies the effect of Article XX(b) of the GATT 1994. Article XX(b) allows a Member to “adopt and enforce” a measure, inter alia, necessary to protect human life or health, even though that measure is inconsistent with another provision of the GATT 1994. Article III:4 and Article XX(b) are distinct and independent provisions of the GATT 1994 each to be interpreted on its own. The scope and meaning of Article III:4 should not be broadened or restricted beyond what is required by the normal customary international law rules of treaty interpretation, simply because Article XX(b) exists and may be available to justify measures inconsistent with Article III:4.
The fact that an interpretation of Article III:4, under those rules, implies a less frequent recourse to Article XX(b) does not deprive the exception in Article XX(b) of *effet utile*. Article XX(b) would only be deprived of *effet utile* if that provision could *not* serve to allow a Member to “adopt and enforce” measures “necessary to protect human ... life or health”. Evaluating evidence relating to the health risks arising from the physical properties of a product does not prevent a measure which is inconsistent with Article III:4 from being justified under Article XX(b). We note, in this regard, that, different inquiries occur under these two very different Articles. Under Article III:4, evidence relating to health risks may be relevant in assessing the *competitive relationship in the marketplace* between allegedly “like” products. The same, or similar, evidence serves a different purpose under Article XX(b), namely, that of assessing whether a Member has a sufficient basis for “adopting or enforcing” a WTO-inconsistent measure on the grounds of human health.

116. We, therefore, find that the Panel erred, in paragraph 8.132 of the Panel Report, in excluding the health risks associated with chrysotile asbestos fibres from its examination of the physical properties of that product.

117. Before examining the Panel’s findings under the second and third criteria, we note that these two criteria involve certain of the key elements relating to the competitive relationship between products: first, the extent to which products are capable of performing the same, or similar, functions (end-uses), and, second, the extent to which consumers are willing to use the products to perform these functions (consumers’ tastes and habits). Evidence of this type is of particular importance under Article III of the GATT 1994, precisely because that provision is concerned with competitive relationships in the marketplace. If there is – or could be – *no* competitive relationship between products, a Member cannot intervene, through internal taxation or regulation, to protect domestic production. Thus, evidence about the extent to which products can serve the same end-uses, and the extent to which consumers are – or would be – willing to choose one product instead of another to perform those end-uses, is highly relevant evidence in assessing the “likeness” of those products under Article III:4 of the GATT 1994.

118. We consider this to be especially so in cases where the evidence relating to properties establishes that the products at issue are physically quite different. In such cases, in order to overcome this indication that products are *not* “like”, a higher burden is placed on complaining Members to establish that, despite the pronounced physical differences, there is a competitive relationship between the products such that *all* of the evidence, taken together, demonstrates that the products are “like” under Article III:4 of the GATT 1994. In this case, where it is clear that the fibres have very different properties, in particular, because chrysotile is a known carcinogen, a very heavy burden is placed on Canada to show, under the second and third criteria, that the chrysotile asbestos and PCG fibres are in such a competitive relationship.

119. With this in mind, we turn to the Panel’s evaluation of the second criterion, end-uses. The Panel’s evaluation of this criterion is far from comprehensive. First, as we have said, the Panel entwined its analysis of “end-uses” with its analysis of “physical properties” and, in purporting to examine “end-uses” as a distinct criterion, essentially referred to its analysis of “properties”. This makes it difficult to assess precisely how the Panel evaluated the end-uses criterion. Second, the Panel’s analysis of end-uses is based on a “small number of applications” for which the products are substitutable. Indeed, the Panel stated that “[i]t suffices that, for a *given utilization*, the properties are the same to the extent that one product can replace the other.” (emphasis added) Although we agree that it is certainly relevant that products have similar end-uses for a “small number of ... applications”, or even for a “given utilization”, we think that a panel must also examine the other, *different* end-uses for products. It is only by forming a complete picture of the various end-uses of a product that a panel can assess the significance of the fact that products share a limited number of end-uses. In this case, the Panel did not provide such a complete picture of the various end-uses of the different fibres. The Panel did not explain, or elaborate in any way on, the “small number of ... applications” for which the various fibres have similar end-uses. Nor did the Panel examine the end-uses for these products.
which were not similar. In these circumstances, we believe that the Panel did not adequately examine the evidence relating to end-uses.

120. The Panel declined to examine or make any findings relating to the third criterion, consumers’ tastes and habits, “[b]ecause this criterion would not provide clear results”. 100 There will be few situations where the evidence on the “likeness” of products will lend itself to “clear results”. In many cases, the evidence will give conflicting indications, possibly within each of the four criteria. For instance, there may be some evidence of similar physical properties and some evidence of differing physical properties. Or the physical properties may differ completely, yet there may be strong evidence of similar end-uses and a high degree of substitutability of the products from the perspective of the consumer. A panel cannot decline to inquire into relevant evidence simply because it suspects that evidence may not be “clear” or, for that matter, because the parties agree that certain evidence is not relevant. 101 In any event, we have difficulty seeing how the Panel could conclude that an examination of consumers’ tastes and habits “would not provide clear results”, given that the Panel did not examine any evidence relating to this criterion.

121. Furthermore, in a case such as this, where the fibres are physically very different, a panel cannot conclude that they are “like products” if it does not examine evidence relating to consumers’ tastes and habits. In such a situation, if there is no inquiry into this aspect of the nature and extent of the competitive relationship between the products, there is no basis for overcoming the inference, drawn from the different physical properties of the products, that the products are not “like”.

122. In this case especially, we are also persuaded that evidence relating to consumers’ tastes and habits would establish that the health risks associated with chrysotile asbestos fibres influence consumers’ behaviour with respect to the different fibres at issue. 102 We observe that, as regards chrysotile asbestos and PCG fibres, the consumer of the fibres is a manufacturer who incorporates the fibres into another product, such as cement-based products or brake linings. We do not wish to speculate on what the evidence regarding these consumers would have indicated; rather, we wish to highlight that consumers’ tastes and habits regarding fibres, even in the case of commercial parties, such as manufacturers, are very likely to be shaped by the health risks associated with a product which is known to be highly carcinogenic. 103 A manufacturer cannot, for instance, ignore the preferences of the ultimate consumer of its products. If the risks posed by a particular product are sufficiently great, the ultimate consumer may simply cease to buy that product. This would, undoubtedly, affect a manufacturer’s decisions in the marketplace. Moreover, in the case of products posing risks to human health, we think it likely that manufacturers’ decisions will be influenced by other factors, such as the potential civil liability that might flow from marketing products posing a health risk to the ultimate consumer, or the additional costs associated with safety procedures required to use such products in the manufacturing process.

123. Finally, we note that, although we consider consumers’ tastes and habits significant in determining “likeness” in this dispute, at the oral hearing, Canada indicated that it considers this criterion to be irrelevant, in this dispute, because the existence of the measure has disturbed normal conditions of competition between the products. In our Report in Korea – Alcoholic Beverages, we observed that, “[p]articularly in a market where there are regulatory barriers to trade or to competition, there may well be latent demand” for a product. 104 We noted that, in such situations, “it may be highly relevant to examine latent demand” that is suppressed by regulatory barriers. 105 In addition, we said that “evidence from other markets may be pertinent to the examination of the market at issue, particularly when demand on that market has been influenced by regulatory barriers to trade or to competition.” 106 We, therefore, do not accept Canada’s contention that, in markets where normal conditions of competition have been disturbed by regulatory or fiscal barriers, consumers’ tastes and habits cease to be relevant. In such situations, a Member may submit evidence of latent, or suppressed, consumer demand in that market, or it may submit evidence of substitutability from some relevant third market. In making this point, we do not...
wish to be taken to suggest that there is latent demand for chrysotile asbestos fibres. Our
point is simply that the existence of the measure does not render consumers’ tastes and
habits irrelevant, as Canada contends.

124. We observe also that the Panel did not regard as decisive the different tariff clas-
sifications of the chrysotile asbestos, PVA, cellulose and glass fibres, each of which is clas-
sified under a different tariff heading. 107 In the absence of a full analysis, by the Panel, of
the other three criteria addressed, we cannot determine what importance should be at-
tached to the different tariff classifications of the fibres.

125. In sum, in our view, the Panel reached the conclusion that chrysotile asbestos and
PCG fibres are “like products” under Article III:4 of the GATT 1994 on the following
basis: the Panel disregarded the quite different “properties, nature and quality” of chrysotile
asbestos and PCG fibres, as well as the different tariff classification of these fibres; it con-
sidered no evidence on consumers’ tastes and habits; and it found that, for a “small number”
of the many applications of these fibres, they are substitutable, but it did not consider the
many other end-uses for the fibres that are different. Thus, the only evidence supporting
the Panel’s finding of “likeness” is the “small number” of shared end-uses of the fibres.

126. For the reasons we have given, we find this insufficient to justify the conclusion
that the chrysotile asbestos and PCG fibres are “like products” and we, therefore, reverse
the Panel’s conclusion, in paragraph 8.144 of the Panel Report, “that chrysotile fibres, on
the one hand, and PVA, cellulose and glass fibres, on the other, are ‘like products’ within
the meaning of Article III:4 of the GATT 1994.”

3. Cement-based products containing chrysotile and PCG fibres

127. Having reversed the Panel’s finding on the “likeness” of the fibres, we now ex-
amine the Panel’s findings regarding the “likeness” of cement-based products containing
chrysotile asbestos fibres and cement-based products containing PCG fibres. In examining the
“likeness” of these cement-based products, the Panel stated that, physically, the only dif-
ference between these products is the incorporation of a different fibre. 108 In this respect,
the Panel indicated that “many of the arguments put forward in relation to chrysotile as-
bestos, PVA, cellulose and glass fibres are applicable mutatis mutandis to products con-
taining those fibres.” 109 The Panel noted that, for any given cement-based product, the
tariff classification is the same, irrespective of the fibre incorporated into the product. 110
The Panel declined to examine the “risk” criterion advanced by the European Communi-
ties, and also considered it unnecessary to analyze consumers’ tastes and habits. 111 On
this basis, the Panel concluded that “chrysotile-fibre products and fibro-cement products
are like products within the meaning of Article III:4 of the GATT 1994.” 112

128. As the Panel said, the primary physical difference between cement-based prod-
ucts containing chrysotile asbestos fibres and cement-based products containing PCG fi-
bres lies in the particular fibre incorporated into the product. This difference is important
because, as we have said in our examination of fibres, we believe that the health risks
associated with a product may be relevant to the inquiry into the physical properties of a
product when making a determination of “likeness” under Article III:4 of the GATT 1994. 113
This is also true for cement-based products containing the different fibres. In examining the
physical properties of the two sets of cement-based products, it cannot be ignored that
one set of products contains a fibre known to be highly carcinogenic, while the other does
not. 114 In this respect, we recall that the Panel concluded that “there is an undeniable
public health risk in relation to chrysotile contained in high-density chrysotile-cement prod-
ucts.” 115 We, therefore, reverse the Panel’s finding, in paragraph 8.149 of the Panel Report,
that these health risks are not relevant in examining the “likeness” of the cement-based
products.

129. Furthermore, the Panel did not indicate whether or to what extent the incorpo-
ration of one type of fibre, instead of another, affects other physical properties of a particu-
lar cement-based product and, consequently, affects the suitability of that product for a
specific end-use. The Panel noted that the fibres give the products their specific function – "mechanical strength, resistance to heat, compression, etc." – but the Panel did not examine the extent to which the presence of a particular fibre affects the ability of a cement-based product to perform one or more of these functions efficiently. 116

130. In addition, even if the cement-based products were functionally interchangeable, we consider it likely that the presence of a known carcinogen in one of the products would have an influence on consumers' tastes and habits regarding that product. We believe this to be true irrespective of whether the consumer of the cement-based products is a commercial party, such as a construction company, or is an individual, for instance, a do-it-yourself ("DIY") enthusiast or someone who owns or lives or works in a building. This influence may well vary, but the possibility of such an influence should not be overlooked by a panel when considering the "likeness" of products containing chrysotile asbestos. In the absence of an examination of consumers' tastes and habits, we do not see how the Panel could reach a conclusion on the "likeness" of the cement-based products at issue. 117

131. For all of these reasons, we reverse the Panel's conclusion, in paragraph 8.150 of the Panel Report, "that chrysotile-fibre products and fibro-cement products are like products within the meaning of Article III:4 of the GATT 1994."

132. As we have reversed both of the Panel’s conclusions on “likeness” under Article III:4 of the GATT 1994, we think it appropriate to complete the analysis, on the basis of the factual findings of the Panel and of the undisputed facts in the Panel record. We have already examined the meaning of the term “like products”, and we have also approved the approach for inquiring into “likeness” that is based on the Report of the Working Party in Border Tax Adjustments and that was also approved, though not entirely followed, by the Panel in this case. Under that approach, the evidence is to be examined under four criteria: physical properties; end-uses; consumers' tastes and habits; and tariff classification.

E. Completing the “Like Product” Analysis under Article III:4 of the GATT 1994

133. As we have reversed both of the Panel’s conclusions on “likeness” under Article III:4 of the GATT 1994, we think it appropriate to complete the analysis, on the basis of the factual findings of the Panel and of the undisputed facts in the Panel record. We have already examined the meaning of the term “like products”, and we have also approved the approach for inquiring into “likeness” that is based on the Report of the Working Party in Border Tax Adjustments and that was also approved, though not entirely followed, by the Panel in this case. Under that approach, the evidence is to be examined under four criteria: physical properties; end-uses; consumers' tastes and habits; and tariff classification.

1. Chrysotile and PCG fibres

134. We address first the “likeness” of chrysotile asbestos fibres and PCG fibres. As regards the physical properties of these fibres, we recall that the Panel stated that:

The Panel notes that no party contests that the structure of chrysotile fibres is unique by nature and in comparison with artificial fibres that can replace chrysotile asbestos. The parties agree that none of the substitute fibres mentioned by Canada in connection with Article III:4 has the same structure, either in terms of its form, its diameter, its length or its potential to release particles that possess certain characteristics. Moreover, they do not have the same chemical composition, which means that, in purely physical terms, none of them has the same nature or quality. … 118
We also see it as important to take into account that, since 1977, chrysotile asbestos fibres have been recognized internationally as a known carcinogen because of the particular combination of their molecular structure, chemical composition, and fibrillation capacity. In that respect, the Panel noted that:

… the carcinogenicity of chrysotile fibres has been acknowledged for some time by international bodies. This carcinogenicity was confirmed by the experts consulted by the Panel, with respect to both lung cancers and mesotheliomas, even though the experts appear to acknowledge that chrysotile is less likely to cause mesotheliomas than amphiboles. We also note that the experts confirmed that the types of cancer concerned had a mortality rate of close to 100 per cent. We therefore consider that we have sufficient evidence that there is in fact a serious carcinogenic risk associated with the inhalation of chrysotile fibres. …

In contrast, the Panel found that the PCG fibres “are not classified by the WHO at the same level of risk as chrysotile.” The experts also confirmed, as the Panel reported, that current scientific evidence indicates that PCG fibres do “not present the same risk to health as chrysotile” asbestos fibres.

It follows that the evidence relating to properties indicates that, physically, chrysotile asbestos and PCG fibres are very different. As we said earlier, in such cases, in order to overcome this indication that products are not “like”, a high burden is imposed on a complaining Member to establish that, despite the pronounced physical differences, there is a competitive relationship between the products such that, all of the evidence, taken together, demonstrates that the products are “like” under Article III:4 of the GATT 1994.

The Panel observed that the end-uses of chrysotile asbestos and PCG fibres are the same “for a small number” of applications. The Panel simply adverted to these overlapping end-uses and offered no elaboration on their nature and character. We note that Canada argued before the Panel that there are some 3,000 commercial applications for asbestos fibres. Canada and the European Communities indicated that the most important end-uses for asbestos fibres include, in no particular order, incorporation into: cement-based products; insulation; and various forms of friction lining. Canada noted that 90 percent, by quantity, of French imports of chrysotile asbestos were used in the production of cement-based products. This evidence suggests that chrysotile asbestos and PCG fibres share a small number of similar end-uses and, that, as Canada asserted, for chrysotile asbestos, these overlapping end-uses represent an important proportion of the end-uses made of chrysotile asbestos, measured in terms of quantity.

There is, however, no evidence on the record regarding the nature and extent of the many end-uses for chrysotile asbestos and PCG fibres which are not overlapping. Thus, we do not know what proportion of all end-uses for chrysotile asbestos and PCG fibres overlap. Where products have a wide range of end-uses, only some of which overlap, we do not believe that it is sufficient to rely solely on evidence regarding the overlapping end-uses, without also examining evidence of the nature and importance of these end-uses in relation to all of the other possible end-uses for the products. In the absence of such evidence, we cannot determine the significance of the fact that chrysotile asbestos and PCG fibres share a small number of similar end-uses.

As we have already stated, Canada took the view, both before the Panel and before us, that consumers’ tastes and habits have no relevance to the inquiry into the “likeness” of the fibres. We have already addressed, and dismissed, the arguments advanced by Canada in support of this contention. We have also stated that, in a case such as this
one, where the physical properties of the fibres are very different, an examination of the evidence relating to consumers’ tastes and habits is an indispensable – although not, on its own, sufficient – aspect of any determination that products are “like” under Article III:4 of the GATT 1994. 129 If there is no evidence on this aspect of the nature and extent of the competitive relationship between the fibres, there is no basis for overcoming the inference, drawn from the different physical properties, that the products are not “like”. However, in keeping with its argument that this criterion is irrelevant, Canada presented no evidence on consumers’ tastes and habits regarding chrysotile asbestos and PCG fibres. 130

140. Finally, we note that chrysotile asbestos fibres and the various PCG fibres all have different tariff classifications. While this element is not, on its own, decisive, it does tend to indicate that chrysotile and PCG fibres are not “like products” under Article III:4 of the GATT 1994.

141. Taken together, in our view, all of this evidence is certainly far from sufficient to satisfy Canada’s burden of proving that chrysotile asbestos fibres are “like” PCG fibres under Article III:4 of the GATT 1994. Indeed, this evidence rather tends to suggest that these products are not “like products” for the purposes of Article III:4 of the GATT 1994.

2. Cement-based products containing chrysotile and PCG fibres

142. We turn next to consider whether cement-based products containing chrysotile asbestos fibres are “like” cement-based products containing PCG fibres under Article III:4 of the GATT 1994. We begin, once again, with physical properties. In terms of composition, the physical properties of the different cement-based products appear to be relatively similar. Yet, there is one principal and significant difference between these products: one set of cement-based products contains a known carcinogenic fibre, while the other does not. The Panel concluded that the presence of chrysotile asbestos fibres in cement-based products poses “an undeniable public health risk”. 131

143. The Panel stated that the fibres give the cement-based products their specific function – “mechanical strength, resistance to heat, compression, etc.” 132 These functions are clearly based on the physical properties of the products. There is no evidence of record to indicate whether the presence of chrysotile asbestos fibres, rather than PCG fibres, in a particular cement-based product, affects these particular physical properties of the products. For instance, a tile incorporating chrysotile asbestos fibres may be more heat resistant than a tile incorporating a PCG fibre.

144. In addition, there is no evidence to indicate to what extent the incorporation of one type of fibre, instead of another, affects the suitability of a particular cement-based product for a specific end-use. 133 Once again, it may be that tiles containing chrysotile asbestos fibres perform some end-uses, such as resistance to heat, more efficiently than tiles containing a PCG fibre. Thus, while we accept that the two different types of cement-based products may perform largely similar end-uses, in the absence of evidence, we cannot determine whether each type of cement-based product can perform, with equal efficiency, all of the functions performed by the other type of cement-based product.

145. As with the fibres, Canada contends that evidence on consumers’ tastes and habits concerning cement-based products is irrelevant. Accordingly, Canada submitted no such evidence to the Panel. We have dismissed Canada’s arguments in support of this contention. 134 We have also indicated that it is of particular importance, under Article III of the GATT 1994, to examine evidence relating to competitive relationships in the marketplace. 135 We consider it likely that the presence of a known carcinogen in one of the products will have an influence on consumers’ tastes and habits regarding that product. 136 It may be, for instance, that, although cement-based products containing chrysotile asbestos fibres are capable of performing the same functions as other cement-based products, consumers are, to a greater or lesser extent, not willing to use products containing chrysotile asbestos fibres because of the health risks associated with them. Yet, this is only specula-
tion; the point is, there is no evidence. We are of the view that a determination on the "likeness" of the cement-based products cannot be made, under Article III:4, in the absence of an examination of evidence on consumers' tastes and habits. And, in this case, no such evidence has been submitted.

146. As regards tariff classification, we observe that, for any given cement-based product, the tariff classification of the product is the same. However, this indication of "likeness" cannot, on its own, be decisive.

147. Thus, we find that, in particular, in the absence of any evidence concerning consumers' tastes and habits, Canada has not satisfied its burden of proving that cement-based products containing chrysotile asbestos fibres are "like" cement-based products containing PCG fibres, under Article III:4 of the GATT 1994.

148. As Canada has not demonstrated either that chrysotile asbestos fibres are "like" PCG fibres, or that cement-based products containing chrysotile asbestos fibres are "like" cement-based products containing PCG fibres, we conclude that Canada has not succeeded in establishing that the measure at issue is inconsistent with Article III:4 of the GATT 1994.

149. One Member of the Division hearing this appeal wishes to make a concurring statement. At the outset, I would like to make it abundantly clear that I agree with the findings and conclusions reached, and the reasoning set out in support thereof, by the Division, in: Section V (TBT Agreement); Section VII (Article XX(b) of the GATT 1994 and Article 11 of the DSU); Section VIII (Article XXIII(b) of the GATT 1994); and Section IX (Findings and Conclusions) of the Report. This concurring statement, in other words, relates only to Section VI ("Like Products" in Article III:4 of the GATT 1994) of the Report.

150. More particularly, in respect of Section VI of the Report, I join in the findings and conclusions set out in: paragraphs 116, 126, 128, 131, 132, 141, 147 and 148. I am bound to say that, in truth, I agree with a great deal more than just the bare findings and conclusions contained in these eight paragraphs of the Report. It is, however, as a practical matter, not feasible to sort out and identify which part of which paragraph, of the sixty-odd paragraphs comprising Section VI of our Report in which I join. Nor is it feasible to offer a detailed statement with respect to the portions that would then remain. Accordingly, I set out only two related matters below.

151. In paragraph 113 of the Report, we state that "[w]e are very much of the view that evidence relating to the health risks associated with a product may be pertinent in an examination of 'likeness' under Article III:4 of the GATT 1994." We also point out, in paragraph 114, that "[p]anels must examine fully the physical properties of products. In particular, … those physical properties of products that are likely to influence the competitive relationship between products in the market place. In the cases of chrysotile asbestos fibres, their molecular structure, chemical composition, and fibrillation capacity are important because the microscopic particles and filaments of chrysotile asbestos fibres are carcinogenic in humans, following inhalation.” This carcinogenicity we describe as “a defining aspect of the physical properties of chrysotile asbestos fibres,” which property is not shared by the PCG fibres, "at least to the same extent.” We express our inability to “see how this highly significant physical difference cannot be a consideration in examining the physical properties of a product as part of a determination of 'likeness' under Article III:4 of the GATT 1994.” (emphasis in the original) We observe also that the Panel, after noting that the carcinogenicity of chrysotile asbestos fibres has been acknowledged by international bodies and confirmed by the experts the Panel consulted, ruled that it "has" sufficient evidence that "there is in fact a serious carcinogenic risk associated with the inhalation of chrysotile fibres." (emphasis added) In fact, the scientific evidence of record for this finding of carcinogenicity of chrysotile asbestos fibres is so clear, voluminous, and is confirmed, a number of times, by a variety of international organizations, as to be practically overwhelming.

152. In the present appeal, considering the nature and quantum of the scientific evidence showing that the physical properties and qualities of chrysotile asbestos fibres in-
clude or result in carcinogenicity, my submission is that there is ample basis for a definitive characterization, on completion of the legal analysis, of such fibres as not “like” PCG fibres. PCG fibres, it may be recalled, have not been shown by Canada to have the same lethal properties as chrysotile asbestos fibres. That definitive characterization, it is further submitted, may and should be made even in the absence of evidence concerning the other two Border Tax Adjustments criteria (categories of “potentially shared characteristics”) of end-uses and consumers’ tastes and habits. It is difficult for me to imagine what evidence relating to economic competitive relationships as reflected in end-uses and consumers’ tastes and habits could outweigh and set at naught the undisputed deadly nature of chrysotile asbestos fibres, compared with PCG fibres, when inhaled by humans, and thereby compel a characterization of “likeness” of chrysotile asbestos and PCG fibres.

153. The suggestion I make is not that any kind or degree of health risk, associated with a particular product, would a priori negate a finding of the “likeness” of that product with another product, under Article III:4 of the GATT 1994. The suggestion is a very narrow one, limited only to the circumstances of this case, and confined to chrysotile asbestos fibres as compared with PCG fibres. To hold that these fibres are not “like” one another in view of the undisputed carcinogenic nature of chrysotile asbestos fibres appears to me to be but a small and modest step forward from mere reversal of the Panel’s ruling that chrysotile asbestos and PCG fibres are “like”, especially since our holding in completing the analysis is that Canada failed to satisfy a complainant’s burden of proving that PCG fibres are “like” chrysotile asbestos fibres under Article III:4. That small step, however, the other Members of the Division feel unable to take because of their conception of the “fundamental”, perhaps decisive, role of economic competitive relationships in the determination of the “likeness” of products under Article III:4.

154. My second point is that the necessity or appropriateness of adopting a “fundamentally” economic interpretation of the “likeness” of products under Article III:4 of the GATT 1994 does not appear to me to be free from substantial doubt. Moreover, in future concrete contexts, the line between a “fundamentally” and “exclusively” economic view of “like products” under Article III:4 may well prove very difficult, as a practical matter, to identify. It seems to me the better part of valour to reserve one’s opinion on such an important, indeed, philosophical matter, which may have unforeseeable implications, and to leave that matter for another appeal and another day, or perhaps other appeals and other days. I so reserve my opinion on this matter.

VII. Article XX(b) of the GATT 1994 and Article 11 of the DSU

155. Under Article XX(b) of the GATT 1994, the Panel examined, first, whether the use of chrysotile-cement products poses a risk to human health and, second, whether the measure at issue is “necessary to protect human … life or health”. Canada contends that the Panel erred in law in its findings on both these issues. We will examine these two issues in turn before addressing Canada’s appeal that the Panel failed to make an “objective assessment”, under Article 11 of the DSU, in reaching its conclusions under Article XX(b) of the GATT 1994.

156. We recall that Article XX(b) of the GATT 1994 reads:

Subject to the requirement that such measures are not applied in a manner which would constitute a means of arbitrary or unjustifiable discrimination between countries where the same conditions prevail, or a disguised restriction on international trade, nothing in this Agreement shall be construed to prevent the adoption or enforcement by any Member of measures:

...
necessary to protect human, animal or plant life or health;

(emphasis added)

...  

A. "To Protect Human Life or Health"

157. On the issue of whether the use of chrysotile-cement products poses a risk to human health sufficient to enable the measure to fall within the scope of application of the phrase "to protect human ... life or health" in Article XX(b), the Panel stated that it "considers that the evidence before it tends to show that handling chrysotile-cement products constitutes a risk to health rather than the opposite." (emphasis added) On the basis of this assessment of the evidence, the Panel concluded that:

... the EC has made a prima facie case for the existence of a health risk in connection with the use of chrysotile, in particular as regards lung cancer and mesothelioma in the occupational sectors downstream of production and processing and for the public in general in relation to chrysotile-cement products. This prima facie case has not been rebutted by Canada. Moreover, the Panel considers that the comments by the experts confirm the health risk associated with exposure to chrysotile in its various uses. The Panel therefore considers that the EC have shown that the policy of prohibiting chrysotile asbestos implemented by the Decree falls within the range of policies designed to protect human life or health. ... (emphasis added)

Thus, the Panel found that the measure falls within the category of measures embraced by Article XX(b) of the GATT 1994.

158. According to Canada, the Panel deduced that there was a risk to human life or health associated with manipulation of chrysotile-cement products from seven factors. These seven factors all relate to the scientific evidence which was before the Panel, including the opinion of the scientific experts. Canada argues that the Panel erred in law by deducing from these seven factors that chrysotile-cement products pose a risk to human life or health.

159. Although Canada does not base its arguments about these seven factors on Article 11 of the DSU, we bear in mind the discretion that is enjoyed by panels as the trier of facts. In United States – Wheat Gluten, we said:

... in view of the distinction between the respective roles of the Appellate Body and panels, we have taken care to emphasize that a panel’s appreciation of the evidence falls, in principle, "within the scope of the panel’s discretion as the trier of facts". (emphasis added) In assessing the panel’s appreciation of the evidence, we cannot base a finding of inconsistency under Article 11 simply on the conclusion that we might have reached a different factual finding from the one the panel reached. Rather, we must be satisfied that the panel has exceeded the bounds of its discretion, as the trier of facts, in its appreciation of the evidence. As is clear from previous appeals, we will not interfere lightly with the panel’s exercise of its discretion.
160. In Korea – Alcoholic Beverages, we were faced with arguments that sought to cast doubt on certain studies relied on by the panel in that case. We stated:

The Panel’s examination and weighing of the evidence submitted fall, in principle, within the scope of the Panel’s discretion as the trier of facts and, accordingly, outside the scope of appellate review. This is true, for instance, with respect to the Panel’s treatment of the Dodwell Study, the Sofres Report and the Nielsen Study. We cannot second-guess the Panel in appreciating either the evidentiary value of such studies or the consequences, if any, of alleged defects in those studies. Similarly, it is not for us to review the relative weight ascribed to evidence on such matters as marketing studies … (emphasis added)

161. The same holds true in this case. The Panel enjoyed a margin of discretion in assessing the value of the evidence, and the weight to be ascribed to that evidence. The Panel was entitled, in the exercise of its discretion, to determine that certain elements of evidence should be accorded more weight than other elements – that is the essence of the task of appreciating the evidence.

162. With this in mind, we have examined the seven factors on which Canada relies in asserting that the Panel erred in concluding that there exists a human health risk associated with the manipulation of chrysotile-cement products. We see Canada’s appeal on this point as, in reality, a challenge to the Panel’s assessment of the credibility and weight to be ascribed to the scientific evidence before it. Canada contests the conclusions that the Panel drew both from the evidence of the scientific experts and from scientific reports before it. As we have noted, we will interfere with the Panel’s appreciation of the evidence only when we are “satisfied that the panel has exceeded the bounds of its discretion, as the trier of facts, in its appreciation of the evidence.” (emphasis added) In this case, nothing suggests that the Panel exceeded the bounds of its lawful discretion. To the contrary, all four of the scientific experts consulted by the Panel concurred that chrysotile asbestos fibres, and chrysotile-cement products, constitute a risk to human health, and the Panel’s conclusions on this point are faithful to the views expressed by the four scientists. In addition, the Panel noted that the carcinogenic nature of chrysotile asbestos fibres has been acknowledged since 1977 by international bodies, such as the International Agency for Research on Cancer and the World Health Organization. In these circumstances, we find that the Panel remained well within the bounds of its discretion in finding that chrysotile-cement products pose a risk to human life or health.

163. Accordingly, we uphold the Panel’s finding, in paragraph 8.194 of the Panel Report, that the measure “protect[s] human … life or health”, within the meaning of Article XX(b) of the GATT 1994.

B. “Necessary”

164. On the issue of whether the measure at issue is “necessary” to protect public health within the meaning of Article XX(b), the Panel stated:

In the light of France’s public health objectives as presented by the European Communities, the Panel concludes that the EC has made a prima facie case for the non-existence of a reasonably available alternative to the banning of chrysotile and chrysotile-cement products and recourse to substitute products. Canada has not rebutted the presumption established by the EC. We also consider
165. Canada argues that the Panel erred in applying the “necessity” test under Article XX(b) of the GATT 1994 “by stating that there is a high enough risk associated with the manipulation of chrysotile-cement products that it could in principle justify strict measures such as the Decree.” Canada advances four arguments in support of this part of its appeal. First, Canada argues that the Panel erred in finding, on the basis of the scientific evidence before it, that chrysotile-cement products pose a risk to human health. Second, Canada contends that the Panel had an obligation to “quantify” itself the risk associated with chrysotile-cement products and that it could not simply “rely” on the “hypotheses” of the French authorities. Third, Canada asserts that the Panel erred by postulating that the level of protection of health inherent in the Decree is a halt to the spread of asbestos-related health risks. According to Canada, this “premise is false because it does not take into account the risk associated with the use of substitute products without a framework for controlled use.” Fourth, and finally, Canada claims that the Panel erred in finding that “controlled use” is not a reasonably available alternative to the Decree.

166. With respect to Canada’s first argument, we note simply that we have already dismissed Canada’s contention that the evidence before the Panel did not support the Panel’s findings. We are satisfied that the Panel had a more than sufficient basis to conclude that chrysotile-cement products do pose a significant risk to human life or health.

167. As for Canada’s second argument, relating to “quantification” of the risk, we consider that, as with the SPS Agreement, there is no requirement under Article XX(b) of the GATT 1994 to quantify, as such, the risk to human life or health. A risk may be evaluated either in quantitative or qualitative terms. In this case, contrary to what is suggested by Canada, the Panel assessed the nature and the character of the risk posed by chrysotile-cement products. The Panel found, on the basis of the scientific evidence, that “no minimum threshold of level of exposure or duration of exposure has been identified with regard to the risk of pathologies associated with chrysotile, except for asbestosis.” The pathologies which the Panel identified as being associated with chrysotile are of a very serious nature, namely lung cancer and mesothelioma, which is also a form of cancer. Therefore, we do not agree with Canada that the Panel merely relied on the French authorities’ “hypotheses” of the risk.

168. As to Canada’s third argument, relating to the level of protection, we note that it is undisputed that WTO Members have the right to determine the level of protection of health that they consider appropriate in a given situation. France has determined, and the Panel accepted, that the chosen level of health protection by France is a “halt” to the spread of asbestos-related health risks. By prohibiting all forms of amphibole asbestos, and by severely restricting the use of chrysotile asbestos, the measure at issue is clearly designed and apt to achieve that level of health protection. Our conclusion is not altered by the fact that PCG fibres might pose a risk to health. The scientific evidence before the Panel indicated that the risk posed by the PCG fibres is, in any case, less than the risk posed by chrysotile asbestos fibres, although that evidence did not indicate that the risk posed by PCG fibres is non-existent. Accordingly, it seems to us perfectly legitimate for a Member to seek to halt the spread of a highly risky product while allowing the use of a less risky product in its place. In short, we do not agree with Canada’s third argument.

169. In its fourth argument, Canada asserts that the Panel erred in finding that “controlled use” is not a reasonably available alternative to the Decree. This last argument is based on Canada’s assertion that, in United States – Gasoline, both we and the panel held that an alternative measure “can only be ruled out if it is shown to be impossible to implement.” We understand Canada to mean by this that an alternative measure is only excluded as a “reasonably available” alternative if implementation of that measure is “impossible”. We certainly agree with Canada that an alternative measure which is impossible to implement is not “reasonably available”. But we do not agree with Canada’s read-
ing of either the panel report or our report in *United States – Gasoline*. In *United States – Gasoline*, the panel held, in essence, that an alternative measure did not cease to be “reasonably” available simply because the alternative measure involved administrative difficulties for a Member. The panel’s findings on this point were not appealed, and, thus, we did not address this issue in that case.

Looking at this issue now, we believe that, in determining whether a suggested alternative measure is “reasonably available”, several factors must be taken into account, besides the difficulty of implementation. In *Thailand – Restrictions on Importation of and Internal Taxes on Cigarettes*, the panel made the following observations on the applicable standard for evaluating whether a measure is “necessary” under Article XX(b):

> The import restrictions imposed by Thailand could be considered to be “necessary” in terms of Article XX(b) only if there were no alternative measure consistent with the General Agreement, or less inconsistent with it, which Thailand could reasonably be expected to employ to achieve its health policy objectives. (emphasis added)

In our Report in *Korea – Beef*, we addressed the issue of “necessity” under Article XX(d) of the GATT 1994. In that appeal, we found that the panel was correct in following the standard set forth by the panel in *United States – Section 337 of the Tariff Act of 1930*:

> It was clear to the Panel that a contracting party cannot justify a measure inconsistent with another GATT provision as “necessary” in terms of Article XX(d) if an alternative measure which it could reasonably be expected to employ and which is not inconsistent with other GATT provisions is available to it. By the same token, in cases where a measure consistent with other GATT provisions is not reasonably available, a contracting party is bound to use, among the measures reasonably available to it, that which entails the least degree of inconsistency with other GATT provisions.

We indicated in *Korea – Beef* that one aspect of the “weighing and balancing process … comprehended in the determination of whether a WTO-consistent alternative measure” is reasonably available is the extent to which the alternative measure “contributes to the realization of the end pursued”. In addition, we observed, in that case, that “[t]he more vital or important [the] common interests or values” pursued, the easier it would be to accept as “necessary” measures designed to achieve those ends. In this case, the objective pursued by the measure is the preservation of human life and health through the elimination, or reduction, of the well-known, and life-threatening, health risks posed by asbestos fibres. The value pursued is both vital and important in the highest degree. The remaining question, then, is whether there is an alternative measure that would achieve the same end and that is less restrictive of trade than a prohibition.

Canada asserts that “controlled use” represents a “reasonably available” measure that would serve the same end. The issue is, thus, whether France could reasonably be expected to employ “controlled use” practices to achieve its chosen level of health protection – a halt in the spread of asbestos-related health risks.

In our view, France could not reasonably be expected to employ any alternative measure if that measure would involve a continuation of the very risk that the Decree seeks to “halt”. Such an alternative measure would, in effect, prevent France from achieving its chosen level of health protection. On the basis of the scientific evidence before it, the
Panel found that, in general, the efficacy of “controlled use” remains to be demonstrated. Moreover, even in cases where “controlled use” practices are applied “with greater certainty”, the scientific evidence suggests that the level of exposure can, in some circumstances, still be high enough for there to be a “significant residual risk of developing asbestos-related diseases.” The Panel found too that the efficacy of “controlled use” is particularly doubtful for the building industry and for DIY enthusiasts, which are the most important users of cement-based products containing chrysotile asbestos. Given these factual findings by the Panel, we believe that “controlled use” would not allow France to achieve its chosen level of health protection by halting the spread of asbestos-related health risks. “Controlled use” would, thus, not be an alternative measure that would achieve the end sought by France.

For these reasons, we uphold the Panel’s finding, in paragraph 8.222 of the Panel Report, that the European Communities has demonstrated a prima facie case that there was no “reasonably available alternative” to the prohibition inherent in the Decree. As a result, we also uphold the Panel’s conclusion, in paragraph 8.223 of the Panel Report, that the Decree is “necessary to protect human … life or health” within the meaning of Article XX(b) of the GATT 1994.

C. Article 11 of the DSU

As part of its argument that the Panel erred in finding that the measure is justified under Article XX(b) of the GATT 1994, Canada also asserts that the Panel failed to make an objective assessment of the matter, as required by Article 11 of the DSU. According to Canada, the requirement imposed on panels by Article 11 to make an objective assessment of the matter implies “that scientific data must be assessed in accordance with the principle of the balance of probabilities.” In particular, Canada asserts that, where the evidence is divergent or contradictory, the “principle of the preponderance of evidence” implies that a panel must take a position as to the respective weight of the evidence. Canada also contends that the Panel failed to assess the facts objectively because the Panel accepted “the opinions of experts on the controlled use of chrysotile, when those experts had no controlled-use expertise.”

These arguments by Canada on the “balance of probabilities” and the “preponderance of evidence” concern the credibility and weight that the Panel ascribed to different elements of evidence. In essence, Canada argues that the Panel has not taken sufficient account of certain evidence and that the Panel has placed too much weight on certain other evidence. Thus, Canada is challenging the Panel’s exercise of discretion in assessing and weighing the evidence. As we have already noted, “[w]e cannot second-guess the Panel in appreciating either the evidentiary value of … studies or the consequences, if any, of alleged defects in [the evidence]”. And, as we have already said, in this case, the Panel’s appreciation of the evidence remained well within the bounds of its discretion as the trier of facts.

In addition, in the context of the SPS Agreement, we have said previously, in European Communities – Hormones, that “responsible and representative governments may act in good faith on the basis of what, at a given time, may be a divergent opinion coming from qualified and respected sources.” (emphasis added) In justifying a measure under Article XX(b) of the GATT 1994, a Member may also rely, in good faith, on scientific sources which, at that time, may represent a divergent, but qualified and respected, opinion. A Member is not obliged, in setting health policy, automatically to follow what, at a given time, may constitute a majority scientific opinion. Therefore, a panel need not, necessarily, reach a decision under Article XX(b) of the GATT 1994 on the basis of the “preponderant” weight of the evidence.

With regard to Canada’s argument that certain of the experts lacked expertise in “controlled use”, we note that, from the beginning of the process for the selection of experts, the Panel made clear that it wished to consult experts on the “effectiveness of the controlled use of chrysotile.” The selection of the experts was the subject of a rigorous procedure which involved the consultation of five institutions with experience in this field.
and also of the parties. At no stage did Canada object to the selection of any of the experts, nor indicate that any of them was unqualified to deal with issues relating to “controlled use”. We also note that the experts were instructed by the Panel to answer only those questions that fell within their area of expertise. As Canada indicates, several experts indicated that particular questions, or parts of questions, posed to them went beyond their area of expertise.

In these circumstances, we have serious difficulty accepting that the Panel failed to make an objective assessment by relying on experts who had no expertise. The Panel was entitled to assume that the experts possessed the necessary expertise to answer the questions, or parts of questions, they chose to answer. In other words, it was not incumbent on the Panel expressly to confirm, with respect to every opinion expressed by each expert, that the expert possessed the necessary expertise to give that particular opinion. If Canada thought that one of the experts did not possess the expertise necessary to answer certain questions posed to him, Canada should have raised those concerns, either with the expert, at the meeting the Panel held with the parties and the experts on 17 January 2000, or with the Panel at some other time. We observe, finally, that, where an expert declined to answer a specific question, or part of a question, because of a professed lack of expertise, the Panel had no opinion from that expert on which to rely.

For these reasons, we decline Canada’s appeal on Article 11 of the DSU.

VIII. Article XXIII:1(b) of the GATT 1994

Before the Panel, Canada claimed, under Article XXIII:1(b) of the GATT 1994, that the application of the measure at issue nullified or impaired benefits accruing to Canada. The European Communities raised preliminary objections, arguing on two grounds that the measure falls outside the scope of application of Article XXIII:1(b). First, the European Communities contended that Article XXIII:1(b) only applies to measures which do not otherwise fall under other provisions of the GATT 1994. Second, the European Communities argued that, while it may be possible to have “legitimate expectations” in connection with a purely “commercial” measure, it is not possible to claim “legitimate expectations” with respect to a measure taken to protect human life or health, which can be justified under Article XX(b) of the GATT 1994. Such measures are, the European Communities asserted, excluded from the scope of Article XXIII:1(b).

Before examining the substance of Canada’s claim under Article XXIII:1(b) of the GATT 1994, the Panel first considered, and rejected, both of these preliminary objections raised by the European Communities, and found, as a consequence, that Canada could invoke Article XXIII:1(b) in respect of the measure. The European Communities appeals the Panel’s findings and conclusions relating to the two preliminary objections.

Before considering this aspect of the appeal, we note that the Panel went on to examine the substance of Canada’s claim under Article XXIII:1(b) and concluded that Canada had not established “the existence of nullification or impairment of a benefit within the meaning of Article XXIII:1(b) of the GATT 1994 as a result of the application of the measure”. We note also that this ultimate conclusion by the Panel has not been appealed by either party. Accordingly, we address only the two narrow issues that have been appealed by the European Communities, and we will not address any other aspects of the Panel’s findings under Article XXIII:1(b) of the GATT 1994.

This appeal is our first occasion to examine Article XXIII:1(b) of the GATT 1994. For this reason, before turning to the appeal by the European Communities, it seems to us useful to make certain preliminary observations about the relationship between Articles XXIII:1(a) and XXIII:1(b) of the GATT 1994. Article XXIII:1(a) sets forth a cause of action for a claim that a Member has failed to carry out one or more of its obligations under the GATT 1994. A claim under Article XXIII:1(a), therefore, lies when a Member is alleged to have acted inconsistently with a provision of the GATT 1994. Article XXIII:1(b) sets forth a separate cause of action for a claim that, through the application of a measure, a
Member has “nullified or impaired” “benefits” accruing to another Member, “whether or not that measure conflicts with the provisions” of the GATT 1994. Thus, it is not necessary, under Article XXIII:1(b), to establish that the measure involved is inconsistent with, or violates, a provision of the GATT 1994. Cases under Article XXIII:1(b) are, for this reason, sometimes described as “non-violation” cases; we note, though, that the word “non-violation” does not appear in this provision. The purpose of this rather unusual remedy was described by the panel in *European Economic Community - Payments and Subsidies Paid to Processors and Producers of Oils and Related Animal Feed Proteins ("EEC – Oils")* in the following terms:

The idea underlying [the provisions of Article XXIII:1(b)] is that the improved competitive opportunities that can legitimately be expected from a tariff concession can be frustrated not only by measures prescribed by the General Agreement but also by measures consistent with that Agreement. In order to encourage contracting parties to make tariff concessions they must therefore be given a right of redress when a reciprocal concession is impaired by another contracting party as a result of the application of any measure, whether or not it conflicts with the General Agreement.  

Like the panel in *Japan – Measures Affecting Consumer Photographic Film and Paper ("Japan – Film")*, we consider that the remedy in Article XXIII:1(b) “should be approached with caution and should remain an exceptional remedy”. That panel stated: Although the non-violation remedy is an important and accepted tool of WTO/GATT dispute settlement and has been “on the books” for almost 50 years, we note that there have only been eight cases in which panels or working parties have substantively considered Article XXIII:1(b) claims. This suggests that both the GATT contracting parties and WTO Members have approached this remedy with caution and, indeed, have treated it as an exceptional instrument of dispute settlement. We note in this regard that both the European Communities and the United States in the *EEC – Oils* case, and the two parties in this case, have confirmed that the non-violation nullification or impairment remedy should be approached with caution and treated as an exceptional concept. The reason for this caution is straightforward. Members negotiate the rules that they agree to follow and only exceptionally would expect to be challenged for actions not in contravention of those rules.  

Against this background, we turn now to the European Communities’ argument that Article XXIII:1(b) does not apply to measures that fall within the scope of application of other provisions of the GATT 1994. The text of Article XXIII:1(b) stipulates that a claim under that provision arises when a “benefit” is being “nullified or impaired” through the “application … of any measure, whether or not it conflicts with the provisions of this Agreement”. The wording of the provision, therefore, clearly states that a claim may succeed, under Article XXIII:1(b), even if the measure “conflicts” with some substantive provisions of the GATT 1994. It follows that a measure must, at one and the same time, be inconsistent with, or in breach of, a provision of the GATT 1994 and, nonetheless, give rise to a cause of action under Article XXIII:1(b). Of course, if a measure “conflicts” with a provision of the GATT 1994, that measure must actually fall within the scope of application of that provision of the GATT 1994. We agree with the Panel that this reading of Article XXIII:1(b) is consistent with the panel reports in *Japan – Film* and *EEC – Oils*, which both support the view that Article XXIII:1(b) applies to measures which simultaneously fall within the scope of application of other provisions of the GATT 1994. Accordingly, we decline the European Communities’ first ground of appeal under Article XXIII:1(b) of the GATT 1994.

The European Communities also contends that the Panel erred in finding that Article XXIII:1(b) applies to measures which pursue health, rather than commercial, objec-
atives and which can, therefore, be justified under Article XX(b) of the GATT 1994. Once again, we look to the text of Article XXIII:1(b), which provides that “the application by another Member of any measure” may give rise to a cause of action under that provision. The use of the word “any” suggests that measures of all types may give rise to such a cause of action. The text does not distinguish between, or exclude, certain types of measure. Clearly, therefore, the text of Article XXIII:1(b) contradicts the European Communities’ argument that certain types of measure, namely, those with health objectives, are excluded from the scope of application of Article XXIII:1(b).

189. In any event, an attempt to draw the distinction suggested by the European Communities between so-called health and commercial measures would be very difficult in practice. By definition, measures which affect trade in goods, and which are subject to the disciplines of the GATT 1994, have a commercial impact. At the same time, the health objectives of many measures may be attainable only by means of commercial regulation. Thus, in practice, clear distinctions between health and commercial measures may be very difficult to establish. Nor do we see merit in the argument that, previously, only “commercial” measures have been the subject of Article XXIII:1(b) claims, as that does not establish that a claim cannot be made under Article XXIII:1(b) regarding a “non-commercial” measure.

190. An important aspect of the European Communities’ argument is that a Member cannot have reasonable expectations of continued market access for products which are shown to pose a serious risk to human life or health. However, the paragraphs of the Panel Report appealed by the European Communities involve exclusively the Panel’s findings on the threshold issues of the scope of application of Article XXIII:1(b). This particular argument of the European Communities, important as it is, simply does not relate to those threshold issues. Rather, the European Communities’ argument relates to the substance of a claim that has been determined to fall within the scope of application of Article XXIII:1(b) and, in particular, concerns the issue whether a “benefit” has been “nullified or impaired” by a measure restricting market access for products posing a health risk. Here, we emphasize that the European Communities does not appeal the Panel’s findings relating to the “nullification or impairment” of a “benefit” through the frustration of reasonable expectations by application of the measure at issue. We do not, therefore, find it necessary to examine the European Communities’ argument relating to reasonable expectations.

191. For these reasons, we dismiss the European Communities’ appeal under Article XXIII:1(b) of the GATT 1994 and uphold the Panel’s finding that Article XXIII:1(b) applies to measures which fall within the scope of application of other provisions of the GATT 1994 and which pursue health objectives.

IX. Findings and Conclusions

192. For the reasons set out in this Report, the Appellate Body:

(a) reverses the Panel’s finding, in paragraph 8.72(a) of the Panel Report, that the TBT Agreement “does not apply to the part of the Decree relating to the ban on imports of asbestos and asbestos-containing products because that part does not constitute a ‘technical regulation’ within the meaning of Annex 1.1 to the TBT Agreement”, and finds that the measure, viewed as an integrated whole, does constitute a “technical regulation” under the TBT Agreement;

(b) reverses the Panel’s findings, in paragraphs 8.132 and 8.149 of the Panel Report, that “it is not appropriate” to take into consideration the health risks associated with chrysotile asbestos fibres in examining the “likeness”, under Article III:4 of the GATT 1994, of those fibres and PCG fibres, and, also, in examining the “likeness”, under that provision, of cement-based products containing chrysotile asbestos fibres or PCG fibres;
reverses the Panel’s finding, in paragraph 8.144 of the Panel Report, that chrysotile asbestos fibres and PCG fibres are “like products” under Article III:4 of the GATT 1994; and finds that Canada has not satisfied its burden of proving that these fibres are “like products” under that provision;

(d) reverses the Panel’s finding, in paragraph 8.150 of the Panel Report, that cement-based products containing chrysotile asbestos fibres and cement-based products containing PCG fibres are “like products” under Article III:4 of the GATT 1994; and finds that Canada has not satisfied its burden of proving that these cement-based products are “like products” under Article III:4 of the GATT 1994;

(e) reverses, in consequence, the Panel’s finding, in paragraph 8.158 of the Panel Report, that the measure is inconsistent with Article III:4 of the GATT 1994;

(f) upholds the Panel’s finding, in paragraphs 8.194, 8.222 and 8.223 of the Panel Report, that the measure at issue is “necessary to protect human ... life or health”, within the meaning of Article XX(b) of the GATT 1994; and, finds that the Panel acted consistently with Article 11 of the DSU in reaching this conclusion;

(g) upholds the Panel’s finding, in paragraphs 8.265 and 8.274 of the Panel Report, that the measure may give rise to a cause of action under Article XXIII:1(b) of the GATT 1994.

It follows from our findings that Canada has not succeeded in establishing that the measure at issue is inconsistent with the obligations of the European Communities under the covered agreements and, accordingly, we do not make any recommendations to the DSB under Article 19.1 of the DSU.

Signed in the original at Geneva this 16th day of February 2001 by:

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Florentino P. Feliciano
Presiding Member

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James Bacchus Claus-Dieter Ehlermann
Member Member
NOTES

2Journal officiel, 26 December 1996.
3WT/DS135/R/Add.1, pp. 3-6.
4Panel Report, paras. 1.1 and 1.2. In its request for the establishment of a panel (WT/DS/135/3, 9 October 1998), Canada also claimed that the Decree is inconsistent with the obligations of the European Communities under Articles 2 and 5 of the Agreement on the Application of Sanitary and Phytosanitary Measures (the “SPS Agreement”). However, Canada did not pursue this claim in its written or oral arguments before the Panel.
5Panel Report, para. 9.1.
6Ibid., para. 8.159.
7WT/DS135/8, 23 October 2000.
8Pursuant to Rule 21(1) of the Working Procedures.
9Pursuant to Rule 23(1) of the Working Procedures.
10Pursuant to Rule 27 of the Working Procedures.
13WT/DS135/9, 8 November 2000.
14Such submissions were received from: Asbestos Information Association (United States); FVL Asbestos (Swaziland) Limited (Bulembu Mine); South African Asbestos Producers Advisory Committee (South Africa); J & S Briddle Associates (United Kingdom); Associação das Indústrias de Produtos de Amianio Crisótilo (Portugal); Asbestos Cement Industries Limited (Sri Lanka); The Federation of Thai Industries, Roofing and Accessories Club (Thailand); Korea Asbestos Association (Korea); Senac (Senegal); Syndicat des Métallos (Canada); Duralita de Centroamerica, S.A. de C.V. (El Salvador); Asociación Colombiana de Fibras (Colombia); and Japan Asbestos Association (Japan).
15Applications from the following persons were received by the Division after the deadline specified in the Additional Procedure for receipt of such applications: Association of Personal Injury Lawyers (United Kingdom); All India A.C. Pressure Pipe Manufacturer’s Association (India); International Confederation of Free Trade Unions/European Trade Union Confederation (Belgium); Maharashtra Asbestos Cement Pipe Manufacturers’ Association (India); Roofit Industries Ltd. (India); and Society for Occupational and Environmental Health (United States).
16Applications from the following persons were received by the Division within the deadline specified in the Additional Procedure for receipt of such applications: Professor Robert Lloyd Howse (United States); Occupational & Environmental Diseases Association (United Kingdom); American Public Health Association (United States); Centro de Estudios Comunitarios de la Universidad Nacional de Rosario (Argentina); Only Nature Endures (India); Korea Asbestos Association (Korea); International Council on Metals and the En-
vironment and American Chemistry Council (United States); European Chemical Industry Council (Belgium); Australian Centre for Environmental Law at the Australian National University (Australia); Associate Professor Jan McDonald and Mr. Don Anton (Australia); and a joint application from Foundation for Environmental Law and Development (United Kingdom), Center for International Environmental Law (Switzerland), International Ban Asbestos Secretariat (United Kingdom), Ban Asbestos International and Virtual Network (France), Greenpeace International (The Netherlands), World Wide Fund for Nature, International (Switzerland), and Lutheran World Federation (Switzerland).

These organizations, together with the Center for International Environmental Law and the Lutheran World Federation, filed a joint application for leave to file a written brief. We decided to deny leave to these applicants to file a written brief. See supra, para. 56 and footnote 32.

Panel Report, heading (a) on p. 404 and heading (b) on p. 411.

Ibid., para. 8.72(a).

Ibid., para. 8.72.

Ibid., para. 8.57.

WT/DS135/3. In its request for the establishment of a panel, Canada stated:

…the Government of Canada requested consultations with the European Communities concerning certain measures taken by France prohibiting asbestos and products containing asbestos, and concerning the general asbestos regulations in force in France. These measures and regulations include, but are not limited to, Decree No. 96-1133. (emphasis added)

Canada requested that the Panel “find that Decree No. 96-1133” is inconsistent with the European Communities’ WTO obligations. (emphasis added) See, further, Canada’s request for consultations, WT/DS135/1, G/SPS/GEN/72, G/TBT/D/15, which also identifies the measure at issue as Decree No. 96-1133.

The full text of the Decree is reproduced in Annex I in the Addendum to the Panel Report. Articles 1 and 2 of the Decree are reproduced in paragraph 2 of this Report.

Panel Report, para. 8.57. We note that the Panel stated that a “technical regulation” must apply to “identifiable” products (Panel Report, para. 8.38; emphasis added). However, the Panel went on to state that a “technical regulation” must apply to “given” products (Panel Report, para. 8.57; emphasis added). The Panel also noted that the measure does not “identify by name nor even by function or category” the products covered by the measure (Panel Report, para. 8.40; emphasis added). Thus, in parts of the Panel Report, the Panel appears to require that a “technical regulation” apply to given products rather than identifiable products.

Canada asserted that “chrysotile fibre has no use in its raw form; it serves as an input in the production of chrysotile materials” (Panel Report, paras. 3.418 and 3.439). This assertion is not contested by the European Communities.

Article 5 of the Decree characterizes a contravention of any aspect of Articles 1.I or 1.II as a “5th class offence”.

Article 2.II of the Decree.

Article 2.I of the Decree.

Article 3.I of the Decree.

Article 3.II of the Decree limits the benefit of the exception to activities that have been the subject of the necessary formalities.

Panel Report, para. 8.69.


The Panel’s approach is set forth in para. 8.111 of the Panel Report.

Panell Report, para. 8.144.

Ibid., para. 8.150.


Panel Report, paras. 8.130 and 8.132.

European Communities’ other appellant’s submission, para. 45.

We have already had occasion to interpret other aspects of Article III:4 of the GATT 1994 in two other appeals, but in neither appeal were we asked to address the meaning of the term “like products” (see Appellate Body Report, European Communities – Regime for the Importation, Sale and Distribution of Bananas, WT/DS27/AB/R, adopted 25 September 1997, and Appellate Body Report, Korea – Beef, supra, footnote 49).


In addition, the term “like commodity” appears in Article VI:7 and the term “like merchandise” is used in Article VII:2 of the GATT 1944.

The fourth criterion, tariff classification, was not mentioned by the Working Party on Border Tax Adjustments, but was included by subsequent panels (see, for instance, EEC – Animal Feed, supra, footnote 58, para. 4.2, and 1987 Japan – Alcoholic Beverages, supra, footnote 58, para. 5.6).
1175 European Communities’ other appellant’s submission, para. 33.
1177 Ibid., para. 8.126.
1178 Ibid., paras. 8.123, 8.124 and 8.126.
1179 Ibid., para. 8.130.
1180 Ibid., para. 8.136.
1181 Panel Report, para. 8.139.
1182 Ibid.
1183 Ibid., para. 8.143.
1184 Ibid., para. 8.144.
1185 Ibid.
1186 Ibid., para. 8.150. The Panel devoted six paragraphs to the “likeness” of the cement-based products, whereas it devoted 27 paragraphs to the “likeness” of chrysotile asbestos and PCG fibres.
1187 Ibid., para. 8.126.
1188 Panel Report, para. 8.123.
1189 Ibid., para. 8.121.
1190 Ibid., paras. 8.122 and 8.124.
1191 Ibid., paras. 8.123 and 8.125.
1192 Ibid., para. 8.126.
1193 Ibid., para. 8.125.
1194 Panel Report, para. 8.132.
1195 Ibid., para. 8.130.
1198 Ibid., para. 8.124.
1199 Ibid., paras. 8.124 and 8.125.
1200 Ibid., para. 8.139.
1201 In that respect, we note that, at the oral hearing before us, Canada stated that it believed that the parties were in agreement that consideration of consumers’ tastes and habits “would add nothing” to the determination of “likeness”.
1202 We have already noted the health risks associated with chrysotile asbestos fibres in our consideration of properties (supra, para. 114).
1203 We recognize that consumers’ reactions to products posing a risk to human health vary considerably depending on the product, and on the consumer. Some dangerous products, such as tobacco, are widely used, despite the known health risks. The influence known dangers have on consumers’ tastes and habits is, therefore, unlikely to be uniform or entirely predictable.
1204 Supra, footnote 58, para. 115.
1205 Ibid., para. 120. We added that “studies of cross-price elasticity … involve an assessment of latent demand” (para. 121).
1206 Supra, footnote 58, para. 137.
1207 Panel Report, para. 8.143.
1208 Panel Report, para. 8.145.
1209 Ibid.
1210 Ibid., para. 8.148.
1211 Ibid., para. 8.149.
1212 Ibid., para. 8.150.
1213 Supra, para. 113.
1214 Supra, para. 114.
1215 Panel Report, para. 8.203.
1216 Panel Report, para. 8.145.
1217 See, further, supra, paras. 117 and 118. See, also, supra, paras. 121 and 122.
1218 Panel Report, para. 8.121.
1219 Supra, para. 114.
1220 Panel Report, para. 8.188.
1221 Ibid., para. 8.220.
Our reasons for reaching this conclusion are set forth, supra, in paras. 117, 118, 121 and 122. Canada did present evidence that the impact of the Decree was to reduce demand for chrysotile (Panel Report, paras. 3.20 and 3.422). However, as Canada recognized, this is a necessary consequence of the prohibition on chrysotile and is not evidence of consumers’ attitudes and choices regarding the products at issue. As we have said, regulatory measures may suppress latent consumer demand for a product (supra, para. 123).


Ibid., paras. 8.213 and 8.214.

Canada’s appellant’s submission, para. 204.

Ibid.

Ibid., para. 209.

Ibid., para. 204.


Supra, footnote 48, para. 194.

Panel Report, para. 5.1.

Ibid., para. 2.20.

Ibid.


Panel Report, para. 8.255.

Ibid., para. 8.257.

Ibid., paras. 8.265 and 8.274.

Ibid., para. 8.304.


Including EEC – Oilseeds II, and the present dispute, there have, therefore, been 14 cases in which a claim under Article XXIII:1(b) has been considered by working parties, panels, and, now, the Appellate Body. In six of these cases, the claim under Article XXIII:1(b) was successful and, on three of these occasions, the report was adopted. The successful claims were made in: Australia – Ammonium Sulphate (adopted 3 April 1950), Germany – Sardines (adopted 31 October 1952), EC – Citrus Products (unadopted), EEC – Canned Fruit (unadopted), EEC – Oilseeds (adopted 25 January 1990) and EEC – Oilseeds II (unadopted).

See Panel Report, para. 8.263, which refers to the Panel Report in Japan – Film, supra, footnote 187, para. 10.50, and footnote 1214; and EEC – Oilseeds, supra, footnote 186, para. 144.