This document is meant purely as a documentation tool and the institutions do not assume any liability for its contents

COMMISSION REGULATION (EC) No 829/2007

of 28 June 2007

amending Annexes I, II, VII, VIII, X and XI to Regulation (EC) No 1774/2002 of the European Parliament and of the Council as regards the placing on the market of certain animal by-products

(Text with EEA relevance)

(OJ L 191, 21.7.2007, p. 1)

Amended by:

<u>B</u>

Official Journal

No page date

►<u>M1</u> Commission Regulation (EC) No 1256/2007 of 25 October 2007 L 282 30 26.10.2007

COMMISSION REGULATION (EC) No 829/2007

of 28 June 2007

amending Annexes I, II, VII, VIII, X and XI to Regulation (EC) No 1774/2002 of the European Parliament and of the Council as regards the placing on the market of certain animal by-products

(Text with EEA relevance)

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

Having regard to the Treaty establishing the European Community,

Having regard to Regulation (EC) No 1774/2002 of the European Parliament and of the Council of 3 October 2002 laying down health rules concerning animal by-products not intended for human consumption (1) and in particular the second paragraph of Article 28, the first subparagraph of Article 29(3) and Article 32(1),

Whereas:

- (1) Regulation (EC) No 1774/2002 lays down animal and public health requirements for the importation into and transit through the Community of certain animal by-products and products derived there from. That Regulation provides for general hygiene requirements for the processing of Category 1, 2 and 3 materials and also sets out requirements for the placing on the market of those animal by-products and products derived therefrom including model health certificates for their importation into the Community.
- (2) Due to the threat of introduction of the small hive beetle into the Community, which is currently free of that parasite, it is necessary to lay down requirements for the importation of beeswax, intended for technical purposes, and also to amend the definition of apiculture products in Annex I to Regulation (EC) No 1774/2002. Therefore, the relevant definition in Annex I to that Regulation, the import requirements laid down in Chapter IX of Annex VIII and the model health certificate set out in Chapter 13 of Annex X to that Regulation should be amended accordingly.
- (3) Chapter X of Annex II to Regulation (EC) No 1774/2002 lays down a model commercial document to accompany animal byproducts and processed products during transportation. In order to improve the traceability of hides and skins during transportation, it is necessary to amend that model document. Annex II to that Regulation should therefore be amended accordingly.
- (4) Regulation (EC) No 999/2001 of the European Parliament and of the Council of 22 May 2001 laying down rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies (²) applies to the production and placing on the market of live animals and products of animal origin. It prohibits the feeding of processed animal protein to farmed animals which are kept, fattened or bred for the production of food
- (5) While that prohibition in Regulation (EC) No 999/2001 remains in force, less stringent processing requirements as laid down in Chapter II of Annex VII to Regulation (EC) No 1774/2002 than

⁽¹⁾ OJ L 273, 10.10.2002, p. 1. Regulation as last amended by Commission Regulation (EC) No 2007/2006 (OJ L 379, 28.12.2006, p. 98).

⁽²⁾ OJ L 147, 31.5.2001, p. 1. Regulation as last amended by Commission Regulation (EC) No 1923/2006 (OJ L 404, 30.12.2006, p. 1).

- the current processing Method 1 should apply to processed animal protein derived from porcine blood, as there is no scientific indication of a risk of transmitting TSEs from pigs. It is necessary however for public and animal health reasons to introduce a minimum temperature requirement for the processing of porcine blood. Chapter II of Annex VII to Regulation (EC) No 1774/2002 should therefore be amended accordingly.
- (6) Chapter II A(1) of Annex VIII to Regulation (EC) No 1774/2002 permits the use of parts of slaughtered animals which are fit for human consumption but are not intended for human consumption for commercial reasons in the manufacture of raw petfood within the Community. It appears that parts of such animals which are rejected as unfit for human consumption but which are not affected by any signs of disease communicable to humans or animals may also be safely used in petfood. Chapter II A (1) and Chapter XI of Annex VIII to that Regulation, the latter providing for the respective provisions for the importation into the Community, should therefore be amended accordingly.
- (7) It is necessary to amend the existing model health certificates for the importation of hides and skins into the Community, in order to introduce some technical amendments. In the interest of clarity, the specific requirements for those products set out in Chapter VI of Annex VIII, and the model health certificates set out in Chapters 5(A), 5(B) and 5(C) of Annex X to Regulation (EC) No 1774/2002 should therefore be amended accordingly.
- (8) The European Food Safety Authority (EFSA) in its Scientific Opinion on Animal health and welfare aspects of avian influenza, adopted on 13 and 14 September 2005, concluded that feathers should be treated before trade, in order to reduce the risk of the spread of avian influenza. Chapter VIII of Annex VIII to Regulation (EC) No 1774/2002 lays down the permanent Community measures concerning the placing of the market and importation of feathers. In the light of that opinion and the current worldwide epidemiological situation regarding avian influenza, it is appropriate to amend the relevant Community measures for the importation and treatment of feathers prior to importation. Annex VIII to that Regulation should therefore be amended accordingly.
- (9) The relevant health certificate set out in Chapter 1 of Annex X to Regulation (EC) No 1774/2002 for imports into the Community of processed animal protein, not intended for human consumption, and products other than petfood containing such protein should be amended accordingly in order to provide for equivalent processing possibilities for third countries as for the Community. Chapter I of Annex X to Regulation (EC) No 1774/2002 should therefore be amended accordingly.
- (10) Chapter 3(B) of Annex X to Regulation (EC) No 1774/2002 sets out the model health certificate for the importation into the Community of processed pet food, other than canned pet food. It appears that processed pet food is increasingly produced in third countries by mixing already processed ingredients which have individually been treated in compliance with the requirements of that Regulation. Heat treatment of such ingredients could have adverse effects on their nutritional qualities. Therefore, the importation into the Community of processed petfood consisting of such safe ingredients should be allowed and the model health certificate set out in Chapter 3(B) of Annex X to Regulation (EC) No 1774/2002 should be amended accordingly.
- (11) Regulation (EC) No 1774/2002 provides that certain animal byproducts for the manufacture of petfood may be imported into the Community although they contain material which has been

derived from animals treated with certain prohibited substances. However, the importation into the Community of processed petfood, dogchews and flavouring innards containing such material is currently not allowed. Since it appears that the health risk is not higher when importing such material for the production of petfood within the Community than in case such material is an ingredient of imported processed pet food, dogchews and flavouring innards, the importation into the Community of these processed products containing such material should also be allowed. Therefore, the model health certificates set out in Chapters 3(A), 3(B), 3(C) and 3(E) of Annex X to Regulation (EC) No 1774/2002 should be amended accordingly.

- (12) Chapter 3(B) of Annex X to Regulation (EC) No 1774/2002 provides that the health certificate for processed pet food, other than canned petfood, must contain a certificate stating that the pet food was packed in new packaging, which bears labels indicating 'not intended for human consumption'. Council Directive 79/373/ EEC of 2 April 1979 on the marketing of compound feeding-stuffs (¹) already provides for equivalent marking requirements which prevent unintentional or negligent misuse of such products for human consumption. The requirements for the model health certificate in Chapter 3(B) of Annex X to Regulation (EC) No 1774/2002 should take account of that legislation and should, therefore, be amended accordingly.
- (13) Annex VIII to Regulation (EC) No 1774/2002 lays down requirements for the placing on the market and importation into the Community of pet food, dogchews and technical products. Chapter II B(4) of that Annex provides that dogchews must have undergone a heat treatment during processing sufficient to destroy pathogenic organisms. Several Member States have requested to allow other treatments, during processing of dogchews, which provide equivalent safety guarantees. Therefore, Chapter II B(4) of Annex VIII to Regulation (EC) No 1774/2002 should be amended accordingly.
- (14) Chapter 3(C) of Annex X to Regulation (EC) No 1774/2002 sets out the model health certificate for the importation into the Community of dogchews. Since other treatments than heat treatment shall be allowed for the processing of dogchews within the Community, those other treatments should also be allowed for imported dogchews. Some dogchews are produced according to traditional methods from fish. Animal health considerations do not require a heat treatment in such case. Therefore, the model health certificate in Chapter 3(C) of Annex X to that Regulation should be amended accordingly.
- (15) Chapter II B(6) of Annex VIII to Regulation (EC) No 1774/2002 lays down microbiological standards applicable to raw petfood produced in the Community. Those standards should also be imposed on imports into the Community of raw petfood for direct sale and on animal by-products to be fed to farmed fur animals intended for dispatch to the Community. Chapter 3(D) of Annex X of that Regulation should therefore be amended accordingly.
- (16) Annex XI to Regulation (EC) No 1774/2002 sets out lists of third countries from which Member States may authorise imports of animal by-products not intended for human consumption. Part XIII of that Annex provides for a list of third countries from which Member States may authorise the imports of serum of equidae. According to that part XIII, this list includes third countries referred to in Annex I to Commission Decision

OJ L 86, 6.4.1979, p. 30. Directive as last amended by Regulation (EC) No 807/2003 (OJ L 122, 16.5.2003, p. 36).

- 2004/211/EC of 6 January 2004 establishing the list of third countries and parts of territory thereof from which Member States authorise imports of live equidae and semen, ova and embryos of the equine species, and amending Decisions 93/195/EECand 94/63/EC (¹), from which the importation of equidae for slaughter is allowed.
- (17) Serum of equidae may, however, be sourced from living animals. Therefore, the animal health status allowing for the inclusion of a third country in that list of third countries from which the importation of equidae for breeding and production is allowed should be sufficient to determine whether serum of equidae may be imported from this third country. That would, in particular, allow for the importation of serum of equidae from Mexico. Part XIII of Annex XI to Regulation (EC) No 1774/2002 should therefore be amended accordingly.
- (18) Veterinary certificates shall be provided for in the format laid down in Commission Decision 2007/240/EC (²) regarding the adoption of harmonized templates for veterinary certificates related to the import into the European Community of live animals and products of animal origin. The commercial document laid down in Annex II, Chapter X to Regulation (EC) No 1774/2002 and the model health certificates laid down in Annex X to that Regulation should be amended accordingly.
- (19) A transitional period should be provided for after the date of entry into force of this Regulation, in order to allow for the continued importation into the Community of the animal byproducts and products derived there from covered by Regulation (EC) No 1774/2002 and accompanied by health certificates in compliance with that Regulation
- (20) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on the Food Chain and Animal Health,

HAS ADOPTED THIS REGULATION:

Article 1

Annexes I, II, VII, VIII, X and XI to Regulation (EC) No 1774/2002 are amended in accordance with the Annex to this Regulation.

▼<u>M1</u>

Article 2

For a transitional period until 30 April 2008, Member States shall accept consignments accompanied by commercial documents and health certificates completed and signed in conformity with the provisions of Regulation (EC) No 1774/2002 applying until 23 July 2007.

Until 30 June 2008, Member States shall accept such consignments if the accompanying commercial documents and health certificates were completed and signed before 1 May 2008.

▼B

Article 3

This Regulation shall enter into force and apply on the third day following its publication in the Official Journal of the European Union.

OJ L 73, 11.3.2004, p. 1. Decision as amended by Regulation (EC) No 1792/2006 (OJ L 362, 20.12.2006, p. 1).

⁽²⁾ OJ L 104, 21.4.2007, p. 37.

This Regulation shall be binding in its entirety and directly applicable in all Member States.

ANNEX

The Annexes to Regulation (EC) No 1774/2002 are amended as follows:

- (1) Annex I is amended as follows:
 - (a) point 1 is replaced by the following:
 - '1. "apiculture by-products" means honey, beeswax, royal jelly, propolis or pollen not intended for human consumption;';
 - (b) point 42 is replaced by the following:
 - '42. "processed animal protein" means animal protein derived entirely from Category 3 material, which have been treated in accordance with Chapter II of Annex VII so as to render them suitable for direct use as feed material or for any other use in feedingstuffs, including petfood, or for use in organic fertilisers or soil improvers; however, it does not include blood products, milk, milk-based products, colostrum, gelatine, hydrolysed proteins and dicalcium phosphate, eggs and egg-products, tricalcium phosphate and collagen;'.
- (2) In Annex II, Chapter X is replaced by the following:

'CHAPTER X

Commercial document

- The following commercial document shall accompany animal by-products and processed products during transportation. However, Member States may decide to use a different commercial document, in paper or in electronic form, for animal by-products and processed products transported within the same Member State provided that such commercial document complies with the requirements laid down in Chapter III(2).
- 2. Where more than one transporter is involved, each transporter shall fill in a declaration as referred to in point 7 of the commercial document, which shall be part of the document.

MODEL COMMERCIAL DOCUMENT FOR THE TRANSPORTATION WITHIN THE EUROPEAN COMMUNITY OF ANIMAL BY-PRODUCTS AND PROCESSED PRODUCTS

Notes

- (a) Commercial documents shall be produced, according to the layout of the model appearing in this Annex. It shall contain, in the numbered order that appears in the model, the attestations that are required for the transportation of animal by-products and processed products derived there from.
- (b) It shall be drawn up in one of the official languages of the MemberState of origin or the MemberState of destination, as appropriate. However, it may also be drawn up in other official Community languages, if accompanied by an official translation or if previously agreed by the competent authority of the MemberState of destination.
- (c) The commercial document must be produced at least in triplicate (one original document and two copies). The original document must accompany the consignment to its final destination. The receiver must retain it. The producer must retain one of the copies and the carrier the other.
- (d) The original of each commercial document shall consist of a single page, both sides, or, where more text is required it shall be in such a form that all pages needed are part of an integrated whole and indivisible.
- (e) If for reasons of identification of the items of the consignment, additional pages are attached to the commercial document, these pages shall also be considered as forming part of the original document by the application of the signature of the person responsible for the consignment, on each of the pages.
- (f) When the commercial document, including additional pages referred to in (e), comprises more than one page, each page shall be numbered (page number) of (total number of pages) at the bottom and shall bear the code number of the document that has been designated by the responsible person at the top.
- (g) The original of the commercial document must be completed and signed by the responsible person. In doing so, the responsible person shall ensure that the principles of documentation as laid down in Chapter III of Annex II to Regulation (EC) No 1774/2002 are followed. The commercial document must specify:
 - (i) the date on which the material was taken from the premises;
 - (ii) the description of the material, including the identification of the material, the animal species for Category 3 material and processed products derived therefrom destined for use as feed material and, if applicable, the ear-tag number of the animal;
 - (iii) the quantity of the material;
 - (iv) the place of origin of the material;
 - (v) the name and the address of the carrier of the material;
 - (vi) the name and the address of the receiver and, if applicable, its approval number; and
 - (vii) if appropriate, the approval number of the plant of origin, and the nature and the methods of the treatment
- (h) The colour of the signature of the responsible person shall be different to that of the printing.
- The commercial document must be kept for a period of at least two years for presentation to the competent authority to verify the records referred to in Article 9 of Regulation (EC) No 1774/2002.
- (j) Where Member States decide to use a commercial document in electronic form, the requirements listed in points (a) to (i) shall be complied with as appropriate for such electronic form.

Commercial document

For the transportation within the European Community of animal by-products and processed products not intended for human consumption in accordance with Regulation (EC) No 1774/2002 (²)

EURO	UROPEAN COMMUNITY Commercial document							
	I.1. Consignor Name	I.2. Document reference number: l.2.a. Local reference number:						
	Address	I.3. Central competent authority						
pe	Postal code	I.4. Local competent authority						
sent	I.5. Consignee	1.6.						
pre	Name Address							
nent	Address							
signr	Postal code	1.7.						
Part I : Details of consignment presented	I.8. Country of ISO code I.9. Region of origin Code origin	I.10. Country of ISO code I.11. Region of Code destination destination						
tails	I.12. Place of origin	I.13. Place of destination						
å.	Establishment	Establishment Other						
art	Name Approval number	Name Approval number						
٠ ا	Address	Address						
	Postal code	Postal code						
	I.14. Place of loading Postal code	I.15. Date and time of departure						
	I.16. Means of transport	I.17. Transporter						
	Aeroplane Ship	Name Approval number						
	Road vehicle Other Railway wagon Identification:	Address Postal code Member State						
	I.18. Description of commodity	I.19. Commodity code (CN code)						
	,							
		I.20. Number/quantity						
	I.21. Temperature of products Ambient Chilled Chilled	I.22. Number of packages						
	I.23. Identification of container/Seal number	I.24. Type of packaging						
	I.25. Commodities certified for Animal feedingstuff Technical use	Other 🔲						
	I.26. Transit through third country	I.27. Transit through Member States						
	Third country ISO code Exit point Code	Member State ISO code Member State ISO code						
	Exit point Code Entry point BIP unit No:	Member State ISO code						
	I.28. Export SC code	1.29.						
	Exit point Code							
	1.50.							
	I.31. Identification of the commodities							
	Species Nature of commodity Category Treats (Scientific name)	Approval number of establishments ment type Manufacturing plant Batch number						

EUROPEAN COMMUNITY

Animal by-products/processed products not intended for human consumption

II.a.	Document reference number	II.b.	Local reference number

II.1. Declaration by the consignor

I, the undersigned, declare that:

- II.1.1. A label attached to the container/carton/other packaging material carries the following indication (1):
 - (a) the Category of the animal by-products (see box reference I.31: Category);
 - (b) in the case of processed products, the Category of animal by-products from which the processed products were derived (see box reference I.31: Category);
 - (c) (i) in the case of Category 3 material, the words 'not for human consumption';
 - (ii) in the case of Category 2 material, other than manure and digestive tract content and processed products derived therefrom, the words 'not for animal consumption';
 - (iii) in the case of Category 2 material intended for feeding of animals referred to in point (c) of Article 23(2) under the conditions provided for in that Article of Regulation (EC) No 1774/2002 (2), the words "for feeding to ..." completed with the name of the specific species of those animal(s) for the feeding of which the material is intended;
 - (iv) in the case of manure and digestive tract content, the word "manure"; or
 - (v) in the case of Category 1 material and processed products derived therefrom, the words "for disposal only";
- II.1.2. in the case where the packaging is done by the consignor the animal by-products and/or processed products are:
- (1) either [in sealed new packaging;]
- (¹) or [transported in bulk in covered leak-proof containers or vehicles or other means of transport that were thoroughly cleaned and dry before use;]
- II.1.3. in the case of treatment
 - (a) hides and skins have been treated in accordance with "note Part I, box reference I.31: Treatment type" to this document;
 - (b) the consignment has not been in contact with other animal products or live animals presenting a risk of spreading a serious transmissible disease:
- II.1.4. the animal by-products and/or processed products were stored properly prior to loading and dispatch;
- II.1.5. all precautions have been taken to avoid contamination of the animal by-products or processed products with pathogenic agents and cross-contamination between various Categories.

Notes

Part I:

- Box reference I.9 and I.11: if appropriate.
- -- Box reference I.14: complete if different from "I.1. Consignor".
- Box reference I.31:

Animal species: For Category 3 material and processed products derived therefrom destined for use as feed material.

Nature of commodity: Enter unprocessed animal by-product or processed product chosen among the following list: "apiculture products", "blood products", "blood", "bloodmeal", "canned petfood", "digestion residues", "digestive tract content", "dogchews", "fishmeal", "gelatin", "greaves", "hides and skins", "hydrolysed proteins", "organic fertilizers", "petfood", "processed animal protein", "processed petfood", "processed products", "raw petfood", "rendered fats".

Category: Categories 1, 2 or 3. In case of Category 3, specify which letter from a to k (as under Article 6, paragraph 1 of Regulation (EC) No 1774/2002):

In the case of animal by-product for use in raw petfood indicate 3a or 3b whether the animal by-products derive from:

Category 3a, Article 6(1)(a) i.e. parts of slaughtered animals, which are fit for human consumption in accordance with Community legislation, but are not intended for human consumption for commercial reasons; or

Category 3b, Article 6(1)(b) i.e. parts of slaughtered animals, which are rejected as unfit for human consumption but are not affected by any signs of diseases communicable to humans or animals and derive from carcasses that are fit for human consumption in accordance with Community legislation;

In the case of hides and skins and processed products derived there from, indicate 3c or 3k whether the animal by-products derive from:

Category 3c, Article 6(1)(c) i.e. hides and skins originating from animals that are slaughtered in a slaughterhouse, after undergoing antemortem inspection, and were fit, as a result of such inspection, for slaughter for human consumption in accordance with Community legislation; or

Category 3k, Article 6(1)(k) i.e. hides and skins originating from animals that did not show clinical signs of any disease communicable through that product to humans or animals.

Part II: Declaration

Where the consignment is made of more than one Category, indicate the quantity and if applicable the number of containers per Category of materials.

Treatment type: For treated hides and skins, which (a) are not fulfilling the requirements of Regulation (EC) No 853/2004 of 29 April 2004 laying down specific hygiene rules for food of animal origin (OJ No L226, 25.6.2004, p. 22) or (b) have not undergone the complete process of tanning or (c) are not 'wet blue'; or (d) are not 'pickled pelts' or (e) are not limed (treated with lime and in brine at a pH of 12 to 13 for at least eight hours): enter treatment among the following: (a) dried; (b) dry-salted or wet-salted for at least 14 days prior to dispatch; (c) salted for seven days in sea salt with the addition of 2 % sodium carbonate; or (d) preserved by a process other than tanning specified in accordance with the procedure referred to in Article 33(2) of Regulation (EC) No 1774/2002.

For Category 3 materials and processed products derived therefrom destined for use as feed: if appropriate describe the nature and the methods of the treatment.

Batch number: enter batch number or ear tag number if applicable.

Part II:

(1)	Delete as appropriate.						
(²) OJ L 273, 10.10.2002, p. 1.							
The sig	nature must be in a different colour to that of the printing						
Signati	ıre						
Done a	ıt	on					
	(date)	(place)					
		(signature of the responsible person/consignor)					
		(name, in capital letters)					
		(name, in deptar letters)					
Declar	ation by the transporter						
	I, the undersigned, declare that:						
II.2.1.	in the case where the packaging is done by the transporte	r, the animal by-products and/or processed products are:					
(1) eithe	er [in sealed new packaging;]						
(1) or	[transported in bulk in covered leak-proof containers or ve cleaned, washed and disinfected after each use;]	hicles or other means of transport that were clean and dry before use and					
II.2.2.	all precautions have been taken:						
	 to avoid contamination of the animal by-products between various Categories during transportation, a 	or processed products with pathogenic agents and cross-contamination and					
	 to ensure transportation under appropriate tempera 	ture to avoid risk to animal or public health.					
Notes							
Part II:							
(1)	Delete as appropriate.						
_	The signature must be in a different colour to that of the pr	ínting					
		y the consignment (*) from the place of loading for dispatch until it reaches					
	the point of destination.	y the consignment () from the place of loading for dispatch than it reaches					
(*)	from the same consignor and covered by the same com-	type, which may contain different Categories of animal by-products, coming mercial document conveyed by the same means of transport to the same					
	recipient."						
Signatu	ure						
Done at							
	(place)	(date)					
		(signature of the responsible person/transporter)					

(name, in capital letters)'

- (3) In Annex VII, Chapter II, paragraph A(1), the first subparagraph is replaced by the following:
 - 'Mammalian processed animal protein must have been submitted to processing method 1. However, porcine blood may be submitted to any of processing methods 1 to 5 or to processing method 7 provided that in the case of processing method 7, a heat treatment throughout its substance at a minimum temperature of 80 °C has been applied.'.
- (4) Annex VIII is amended as follows:
 - (a) Chapter II is amended as follows:
 - (i) paragraph A(1) is replaced by the following:
 - '1. The only animal by-products that may be used to produce petfood and dogchews are those referred to in Article 6(1)(a) to (j). However, raw petfood may only be manufactured from animal by-products referred to in Article 6(1)(a) or Article 6(1) (b).':
 - (ii) paragraph B(4) is replaced by the following:
 - Dogchews must be subjected to a treatment during processing sufficient to destroy pathogenic organisms, including salmonella.

After that treatment, every precaution must be taken to ensure that such dogchews are not exposed to contamination. The dogchews must be packed in new packaging.';

- (b) Chapter III is amended as follows:
 - (i) paragraph I(A)(3) is replaced by the following:
 - '3. Unprocessed manure of equidae which is traded must not originate from a holding subject to animal health restrictions pertaining to glanders, vesicular stomatitis, anthrax or rabies in accordance with Article 4 (5) of Directive 90/426/EEC.';
 - (ii) paragraph I(B)(4) is replaced by the following:
 - '4. The importation of unprocessed manure is prohibited.';
 - (iii) paragraph II(B)(6)(d) is replaced by the following:
 - '(d) are accompanied by a health certificate that conforms to the model laid down in Chapter 17 of Annex X.';
- (c) Chapter VI is amended as follows:
 - (i) in A, paragraph (1)(a) is replaced by the following:
 - '(a) to hides and skins of ungulates complying with the requirements of Regulation (EC) No 853/2004 of 29 April 2004 of the European Parliament and of the Council laying down specific hygiene rules for food of animal origin (*)

- (ii) in B, paragraph 3 is replaced by the following:
 - '3. Trade in fresh or chilled hides and skins is subject to the same health conditions as those applicable to fresh meat pursuant to Council Directive 2002/99/EC of 16 December 2002 laying down the animal health rules governing the production, processing, distribution and introduction of products of animal origin for human consumption (*).

- (iii) in C, paragraph 5(b), the introductory phrase is replaced by the following:
 - '(b) they come from a third country or, in the case of regionalisation in accordance with Community legislation, from a part of a

^(*) OJ L 139, 30.4.2004, p. 55; corrected version (OJ L 226, 25.6.2004, p. 22).';

^(*) OJ L 18, 23.1.2003, p. 11.';

third country, appearing on the list set out in Part XIV(A) of Annex XI and which, as appropriate to the species concerned.';

- (iv) in C, paragraphs 6(b) to (e) are replaced by the following:
 - '(b) they come either from:
 - a third country or, in the case of regionalisation in accordance with Community legislation, from a part of a third country, appearing on the list set out in Part XIV(B) of Annex XI from which imports of fresh meat of the corresponding species are authorised and they have been treated in accordance with paragraph 2(a), (b) and (c) of A; or
 - (ii) a third country appearing on the list set out in Part XIV(B) of Annex XI and they have been treated in accordance with paragraph 2(c) or (d) of A; or
 - (iii) equidae or ruminant animals from a third country appearing on the list set out in Part XIV(C) of Annex XI, which have been treated in accordance with paragraph 2(a), (b) and (c) of A and after treatment have been kept separate for at least 21 days;
 - (c) in the case of salted hides and skins transported by ship, they have been treated in accordance with paragraphs 2(b) or (c) of A and have been kept separated after treatment during transportation for at least 14 days in the case of paragraph (b) or seven days in the case of paragraph (c) before importation and the health certificate accompanying the consignment attests such treatment and the duration of the transportation; and
 - (d) a health certificate conforming to the model health certificate laid down in Chapter 5(B) of Annex X, or, in the case of hides and skins referred to in paragraph 6(b)(iii) of C of this Annex, an official declaration conforming to the model laid down in Chapter 5 (C) of Annex X, accompanies them.';
- (d) Chapter VIII is amended as follows:
 - (i) in B, paragraph 4, the introductory phrase is replaced by the following:
 - 'Member States must authorise the importation of unprocessed wool and hair, if they are:';
 - (ii) in B, the following paragraph 5 is added:
 - The importation of unprocessed feathers and parts of feathers is prohibited.

Member States must authorise the importation of processed feathers and parts of feathers if:

- (a) they are treated decorative feathers, treated feathers carried by travellers for their private use or consignments of treated feathers sent to private individuals for non-industrial purposes; or
- (b) they are accompanied by a commercial document stating that the feathers or parts of feathers have been treated with a steam current or by another method ensuring the inactivation of pathogens and are securely enclosed in packaging and dry.';
- (e) Chapter IX is amended as follows:
 - (i) in A, paragraph 1, the introductory phrase is replaced by the following:
 - '1. Apiculture by-products intended exclusively for use in apiculture must:';
 - (ii) in B, paragraph 3 is replaced by the following:
 - '3. Member States must authorise the importation of apiculture byproducts, other than beeswax in the form of honeycomb, intended for use in apiculture if they:

- (a) come from third countries that appear on the list in Part XII of Annex XI;
- (b) either:
 - (i) have been subjected to a temperature of −12 °C or lower for at least 24 hours; or
 - (ii) in the case of wax, the material has been refined or rendered before importation; and
- (c) are accompanied by a health certificate that conforms to the model set out in Chapter 13 of Annex X.';
- (iii) in B, the following paragraphs 4 and 5 are added:
 - '4. Member States must authorise the importation of beeswax for technical purposes, other than beeswax in the form of honeycomb, if it:
 - (a) has been refined or rendered before importation; and
 - (b) is accompanied by a commercial document attesting that refinement or rendering.
 - The importation of beeswax in the form of honeycomb shall be prohibited.';
- (f) Chapter XI is amended as follows:
 - (i) in point 2, the second subparagraph is replaced by the following: 'however, animal by-products for use in feed for farmed fur animals

or for use in raw petfood must consist of animal by-products referred to in Article 6(1)(a) and (b) only;';

- (ii) paragraph 6 is replaced by the following:
 - '6. are accompanied by a certificate that conforms to one of the models set out in Chapter 3(D), 3(F) or 8 of Annex X.'.
- (5) Annex X is replaced as follows:

'ANNEX X

MODEL HEALTH CERTIFICATES FOR THE IMPORTATION FROM THIRD COUNTRIES AND FOR THE TRANSIT THROUGH THE EUROPEAN COMMUNITY OF CERTAIN ANIMAL BY-PRODUCTS AND PRODUCTS DERIVED THEREFROM

Notes

- (a) Veterinary certificates shall be produced by the exporting country, based on the models appearing in this Annex X, according to the layout of the model that corresponds to the animal by-products concerned. They shall contain, in the numbered order that appears in the model, the attestations that are required for any third country and, as the case may be, those supplementary guarantees that are required for the exporting third country or part thereof.
- (b) The original of each certificate shall consist of a single page, both sides, or, where more text is required, it shall be in such a form that all pages needed are part of an integrated whole and indivisible.
- (c) It shall be drawn up in at least one of the official languages of the EU Member State in which the inspection at the border post shall be carried out and of the EU Member State of destination. However, these Member States may allow other languages, accompanied, if necessary, by an official translation.
- (d) If for reasons of identification of the items of the consignment, additional pages are attached to the certificate, these pages shall also be considered as forming part of the original of the certificate by the application of the signature and stamp of the certifying official veterinarian, in each of the pages.
- (e) When the certificate, including additional schedules referred to in d), comprises more than one page, each page shall be numbered (page number) of (total number of pages) on its bottom and shall bear the code number of the certificate that has been designated by the competent authority on its top.

- (f) The original of the certificate must be completed and signed by an official veterinarian. In doing so, the competent authorities of the exporting country shall ensure that the principles of certification equivalent to those laid down in Council Directive 96/93/EC are followed.
- (g) The colour of the signature shall be different to that of the printing. The same rule applies to stamps other than those embossed or watermark.
- (h) The original of the certificate must accompany the consignment at the EU border inspection post.
- (i) If health certificates are used for consignments in transit, box No I.5 (Consignee) of the relevant health certificate shall be completed with the name and address of the border inspection post through which the consignment is intended to leave the European Community.

CHAPTER 1

Health certificate

For processed animal protein not intended for human consumption, including mixtures and products other than petfood containing such protein, for dispatch to or for transit through (²) the European Community

COUN	TRY	Veterinary certificate to EU
	I.1. Consignor Name	I.2. Certificate reference I.2.a.
	Address	I.3. Central Competent Authority
ent	Tel. No	I.4. Local Competent Authority
guu	I.5. Consignee	I.6. Person responsible for the consignment in EU
onsi	Name	Name
ρ	Address	Address
ıtch	Postal code	Postal code
gdsi	Tel. No	Tel. No
of d	I.7. Country of ISO code I.8. Region of origin Cod	de I.9. Country of ISO code I.10. Region of Code
Part I: Details of dispatched consignment	origin	destination destination
ğ	I.11. Place of origin	I.12. Place of destination
art		Custom warehouse
-	Name Approval number	Name Approval number
	Address	Address
		Postal code
	I.13. Place of loading	I.14. Date of departure
	I.15. Means of transport	I.16. Entry BIP in EU
	Aeroplane Ship Railway wagon]
	Road vehicle Other III	147
	Documentary references:	1.17.
	I.18. Description of commodity	I.19. Commodity code (HS code)
		I.20. Quantity
	I.21. Temperature of product	I.22. Number of packages
	Ambient Chilled	Frozen 🗌
	I.23. Identification of container/Seal number	1.24. Type of packaging
	I.25. Commodities certified for:	L
	Animal feedingstuff	Technical use Other
	I.26. For transit to third country vis-à-vis EU	I.27. For import or admission into EU
	3rd country ISO code	
	I.28. Identification of the commodities	
		Approval number of
	Species Nature of commodity (Scientific name)	establishments Manufacturing plant Net weight Batch number

COUNTRY

Processed animal protein not intended for human consumption including mixtures and products other than petfood containing such protein

			II.a. Certificate reference number	II.b.							
	II.	Health attestation	ptorinarian dealars that I have read and understood I	Pagulation (EC) No 1774/2002 (1) and in particular							
		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (*) and in part Article 6 and Annex VII Chapter II thereof and certify that:									
tion	11.1.	the processed animal prote consumption that:	n or product described above contains exclusively	processed animal protein not intended for human							
Part II: Certification			d stored in a plant approved, validated and supervis ppropriate Article 11 of Regulation (EC) No 1774/200								
: Ce		(b) has been prepared ex	lusively with the following animal by-products:								
Part II			ughtered animals, which were fit for human consump nded for human consumption for commercial reason								
		of diseases	ughtered animals, which were rejected as unfit for hur communicable to humans or animals and derive from with Community legislation,]								
		slaughterh	skins, hooves and horns, pig bristles and feathers or buse, underwent antemortem inspection and were f with Community legislation,]								
			ined from animals other than ruminants that were in inspection and were fit, as a result of such inspec								
		(²) and/or [animal by-p bones and	roducts derived from the production of products inter greaves,]	nded for human consumption, including degreased							
		waste, whi	Istuffs of animal origin, or former foodstuffs contain ch are no longer intended for human consumptior ing or packaging defects or other defects which do n	n for commercial reasons or due to problems of							
		(2) and/or [— fish or other	er sea animals, except sea mammals, caught in the open sea for the purposes of fishmeal production,]								
		(2) and/or [fresh by-pr	roducts from fish from plants manufacturing fish products for human consumption,]								
			Ichery by-products and cracked egg by-products originating from animals which did not show clinical ny disease communicable through that product to humans or animals;]								
		and									
			the following processing standard: e temperature of more than 133 °C for at least 20 min	urtee without interruption at a procesure (absolute) of							
			produced by saturated steam, with a particle size prior								
			non-mammalian protein other than fishmeal, the p degulation (EC) 1774/2002;]	rocessing method as set out in Annex V,							
		(2) or [in the case of f (EC) 1774/2002	shmeal the processing method ;]	as set out in Annex V, Chapter III, of Regulation							
			porcine blood, the processing methodas se where in case of method 7 a heat treatment of								
	II.2.	the competent authority ex standards (3):	amined a random sample immediately prior to dis	spatch and found it to comply with the following							
		Salmonella: Absence in 25	n = 5, c = 0, m = 0, M = 0;								
		Enterobacteriaceae:	n = 5, $c = 2$, $m = 10$, $M = 300$ in 1	g;							
	II.3.	the end product:									
	(²) either [was packed in		new or sterilised bags;]								
			d in bulk in containers or other means of transport the proved by the competent authority before use,]	nat were thoroughly cleaned and disinfected with a							
		which bear labels indicating "NOT FOR HUMAN CONSUMPTION"									
	II. 4 .	the end product was stored	in enclosed storage;								
	II.5.	the product has undergone	all precautions to avoid recontamination with pathogo-	enic agents after treatment.							

wotes								
Part I:								
	Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity.							
_		Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses.						
_	Box reference I.15: Registration number (railway wagons is to be provided in the event of unloading and reloading	or container and lorries), flight number (aircraft) or name (ship); information						
_	Box reference I.19: use the appropriate HS code: 05.05	05.06; 05.07 or 23.01						
_	Box reference I.26 and I.27: fill in according to whether i	t is a transit or an import certificate.						
Part II:								
(¹)	OJ L 273, 10.10.2002, p. 1.							
(²)	Delete as appropriate.							
(3)	Where:							
	n = number of samples to be tested;							
	m = threshold value for the number of bacteria; the res not exceed m;	ult is considered satisfactory if the number of bacteria in all samples does						
	M = maximum value for the number of bacteria; the re samples is M or more; and	sult is considered unsatisfactory if the number of bacteria in one or more						
	c = number of samples the bacterial count of which ma bacterial count of the other samples is m or less.	by be between m and M, the sample still being considered acceptable if the						
_	Note for the person responsible for the consignment in the consignment until it reaches the border inspection per	EU: This certificate is only for veterinary purposes and has to accompany ost.						
_	The signature and the stamp must be in a different color	ur to that of the printing.						
Official v	eterinarian							
	Name (in capitals):	Qualification and title:						
	Date:	Signature:						
	Stamp:							

CHAPTER 2(A)

Health certificate

For milk and milk-based products, which have undergone a single heat treatment and are not intended for human consumption for dispatch to or for transit through (3) the European Community

COUN	TRY			Veterinary certificate to El		
	l.1.	Consignor Name	1.2.	Certificate reference number		
		Address	I.3. Central Competent Authority			
ment		Tel. No	1.4.	I.4. Local Competent Authority		
signi	1.5.	Consignee	1.6.	Person responsible for the consignment in EU		
šios		Name		Name		
ped (Address		Address		
atch		Postal code		Postal code		
disp		Tel. No		Tel. No		
Part I: Details of dispatched consignment	1.7.	Country of ISO code I.8. Region of origin Code origin	1.9.	Country of ISO code I.10. destination		
ä	1.11.	Place of origin	l.12.	2. Place of destination		
art				Custom warehouse		
		Name Approval number		Name Approval number		
		Address		Address		
				Postal code		
	1.13.	Place of loading	1.14.	4. Date of departure		
	I.15.	Means of transport	1.16.	6. Entry BIP in EU		
		Aeroplane Ship Railway wagon Road vehicle Other				
	Identification:		1.17.	7		
	Documentary references:					
	I.18.	Description of commodity		I.19. Commodity code (HS code)		
				I.20. Quantity		
	1.21.	Temperature of product		I.22. Number of packages		
		Ambient Chilled Chilled		Frozen 🔲		
	1.23.	Identification of container/Seal number		I.24. Type of packaging		
	1.25.	Commodities certified for:				
		Animal feedingstuff		Technical use Other		
	1.26.	For transit to third country vis-à-vis EU	1.27.	7. For import or admission into EU		
		3rd country ISO code				
	1.28.	Identification of the commodities				
		Approval number of	of estab	ablishments		
		Species Nature of Manufactur (Scientific name) commodity	ring pla	lant Number of Net weight Batch number packages		

COUNTRY

Milk and milk-based products, which have undergone a single heat treatment

			II.a.	Certificate reference nur	mber	II.b.				
	II.	Health attestation								
Part II: Certification		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (1) and certify that:								
	II.1.	(exporting country), (region) (²), has been free from foot-and-mouth disease and rinderpest for 12 months immediately prior to export and has not practiced vaccination against foot-and-mouth disease or rinderpest in the 12 months immediately prior to export;								
	II.2.	the milk and milk-based product referred to in this certificate:								
ertif		(a) has been prepared from raw milk that comes from animals:								
ن ≅		 not showing clinical signs of a diseases that can be transmitted through the milk to humans or animals, and 								
art		 belonging to holdings that are not under official restriction due to footandmouth disease or rinderpest; and 								
Δ.		(b) has undergone a process involving heating to (temperature) for (time), which ensured a negative reaction to the phosphatase test, followed by, in the case of dried milk or dried milk-based product, a drying process;								
	II.3.	every precaution was taken to avoid contamination of the milk/milkbased product after processing;								
	II.4.	the milk/milk-based product was packed:								
	(³) either	[in new containers;]								
	(3) or	(3) or [in vehicles or bulk containers disinfected prior to loading using a product approved by the competent authority;]								
		and the containers are ma HUMAN CONSUMPTION		indicate the nature of the	milk/milk-ba	sed product and bear labels indicating "NOT FOR				
	Notes									
	Part I:									
	_			for the consignment in EU ificate is for import commod		s to be filled in only if it is a certificate for transit				
	 Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The transit can only be stored in free zones, free warehouses and custom warehouses. 									
	_	Box reference I.15: Regist information is to be provided			ntainer and	lorries), flight number (aircraft) or name (ship);				
		Box reference I.19: use the	appropriate	HS code: 23.09.10; 23.09.9	90; 35.01; 3	5.02 or 35.04.				
	_	Box reference I.23: for bull	containers,	the container number and t	he seal num	ber (if applicable) should be included.				
	_	Box reference I.26 and I.2	7: fill in accor	ding to whether it is a transi	t or an impo	ort certificate.				
		Box reference I.28: Manufa	acturing plan	t: provide the registration nu	mber of trea	atment or processing establishment.				
	Part II:									
	(¹)	OJ L 273, 10.10.2002, p. 1.								
	(2)	For completion if the author	risation to in	port into the Community is	restricted to	certain regions of the third country concerned.				
	(3)	Delete as appropriate.								
	_	The signature and the star	np must be i	n a different colour to that of	the printing	i.				
Note for the person responsible for the consignment until it reaches the					rtificate is o	nly for veterinary purposes and has to accompany				
	Official vete	rinarian								
		Name (in capitals):			Qua	alification and title:				
		Date:			Sig	nature:				
		Stamp:								

CHAPTER 2(B)

Health certificate

For heat-treated milk-based products with a pH reduced to less than 6 not intended for human consumption and for dispatch to or for transit through (²) the European Community

COUN	COUNTRY Veterinary certificate to EU								
	I.1. Consignor Name	I.2. Certificate reference I.2.a. number							
	Address	I.3. Central Competent Authority							
nent	Tel. No	I.4. Local Competent Authority							
ignr	I.5. Consignee	I.6. Person responsible for the consignment in EU							
Suo	Name	Name							
pe	Address	Address							
atch	Postal code	Postal code							
disp	Tel. No	Tel. No							
Part I: Details of dispatched consignment	I.7. Country of ISO code I.8. Region of origin Code origin	I.9. Country of ISO code I.10. destination							
. De	I.11. Place of origin	I.12. Place of destination							
art		Custom warehouse							
"	Name Approval number	Name Approval number							
	Address	Address							
		Postal code							
	I.13. Place of loading	I.14. Date of departure							
	I.15. Means of transport	I.16. Entry BIP in EU							
	Aeroplane Ship Railway 🔲								
	Road vehicle Other wagon								
	Identification: Documentary references:	1.17.							
	I.18. Description of commodity	I.19. Commodity code (HS code)							
		I.20. Quantity							
	I.21. Temperature of product	I.22. Number of packages							
	Ambient C Chilled	Frozen							
	I.23. Identification of container/Seal number	I.24. Type of packaging							
	I.25. Commodities certified for: Animal feedingstuff Further process Further process	Technical use Other Other							
	I.26. For transit to third country vis-à-vis EU	I.27. For import or admission into EU							
	3rd country ISO code								
	I.28. Identification of the commodities								
	Approval number of establis	shments							
	Species Nature of commodity Manufacturing plant	Number of packages Net weight Batch number							
	(Scientific name)								

COUNTRY

Heat-treated milk-based products with a pH reduced to less than 6

			II.a.	Certificate reference number	II.b.				
	H.	Health attestation	L						
	I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (¹) and certify that:								
	II.1.	the milk-based product referred to in this certificate:							
tion		(a) has been prepared from raw milk that comes from animals:							
fica		(i) not showing clinical signs of a diseases that can be transmitted through the milk to humans or animals; and							
Serti		(ii) belonging to holdings that are not under official restriction due to foot-and-mouth disease or rinderpest;							
Part II: Certification			process involving heating to (temperature) for (time), which ensured a negative reaction to the followed by, in the case of dried milk or dried milk-based product, a drying process; and						
ď		(c) has undergone an ac) has undergone an acidification process whereby its pH has been maintained at less than 6 for at least one hour;						
	II.2.	every precaution was take	n to avoid o	ontamination of the milk-based produ	ct after processing;				
	II.3.	the milk-based product wa	s packed:						
	(²) either	[in new containers;]							
	(²) or	[in vehicles or bulk containers disinfected prior to loading using a product approved by the competent authority;]							
		and the containers are man CONSUMPTION".	ked so as to	o indicate the nature of the milk-based	product and bear labels indicating "NOT FOR HUMAN				
	Notes								
				e for the consignment in EU: this be rtificate is for import commodity.	ox is to be filled in only if it is a certificate for transit				
	_			on: this box is to be filled in only if it s, free warehouses and custom ware	is a certificate for transit commodity. The products in houses.				
	_	Box reference I.15: Reginformation is to be provide			and lorries), flight number (aircraft) or name (ship);				
		Box reference I.19: use the	e appropria	te HS code: 23.09.10; 23.09.90; 35.0	1; 35.02 or 35.04.				
		Box reference I.23: for bul	k containers	s, the container number and the seal	number (if applicable) should be included.				
		Box reference I.26 and I.2	7: fill in acc	ording to whether it is a transit or an in	mport certificate.				
	_	Box reference I.28: Manuf	acturing pla	nt: provide the registration number of	treatment or processing establishment.				
	Part II:								
	(¹)	OJ L 273, 10.10.2002, p. 1	l.						
	(2)	Delete as appropriate.							
		_		in a different colour to that of the prin	-				
		Note for the person respo the consignment until it re			is only for veterinary purposes and has to accompany				
	Official vete	erinarian							
		Name (in capitals):			Qualification and title:				
		Date:			Signature:				
		Stamp:							

CHAPTER 2(C)

Health certificate

For milk and milk-based products, which have undergone a sterilisation or a double heat treatment and are not intended for human consumption, for dispatch to or for transit through (2) the European Community

COUN	COUNTRY Veterinary certificate to EU								
	I.1. Consignor Name	I.2. Certificate reference number I.2.a.							
	Address	I.3. Central Competent Authority							
nent	Tel. No	I.4. Local Competent Authority							
ignr	I.5. Consignee	I.6. Person responsible for the consignment in EU							
Suo:	Name	Name							
o pa	Address	Address							
atch	Postal code	Postal code							
lisp	Tel. No	Tel. No							
Part I: Details of dispatched consignment	I.7. Country of ISO code I.8. Region of origin Code origin	I.9. Country of ISO code I.10. destination							
. De	I.11. Place of origin	I.12. Place of destination							
art		Custom warehouse							
-	Name Approval number	Name Approval number							
	Address	Address							
		Postal code							
	I.13. Place of loading	I.14. Date of departure							
	I.15. Means of transport	I.16. Entry BIP in EU							
	Aeroplane Ship Railway	1.10. EIRLY BIF III EO							
	Road vehicle Other wagon								
	Identification:	I.17.							
	Documentary references:								
	I.18. Description of commodity	I.19. Commodity code (HS code)							
		I.20. Quantity							
	I.21. Temperature of product	I.22. Number of packages							
	Ambient Chilled Chilled	Frozen 🔲							
	I.23. Identification of container/Seal number	I.24. Type of packaging							
	I.25. Commodities certified for:								
	Animal feedingstuff Further process	Technical use Other Other							
	I.26. For transit to third country vis-à-vis EU	I.27. For import or admission into EU							
	3rd country ISO code								
	I.28. Identification of the commodities								
	Approval number of establis	hments							
	Species Nature of commodity Manufacturing plant	Number of packages Net weight Batch number							
	(Scientific name)								

COUNTRY

Milk and milk-based products, which have undergone a sterilisation or a double heat treatment

			II.a.	Certificate reference number	II.b.				
	11.	Health attestation	L						
	I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (1) that:								
	II.1. the milk/milk-based product referred to in this certificate: (a) has been prepared from raw milk that comes from animals:								
ion									
Part II: Certification		hrough the milk to humans or animals; and							
ertif		(ii) belonging to holdings that are not under official restriction due to footandmouth disease or rinderpest; and							
II: C		(b) has undergone:							
art	(²) either	[(i) a sterilisation	n process wh	nereby an Fc value equal to or greater t	nat 3 is achieved;]				
ь	(²) or	the phosph which ensu	ocess involving heating to (temperature) for						
	II.2.	every precaution was tal	en to avoid c	ontamination of the milk/milk-based pro	duct after processing;				
	II.3.	the milk/milk-based prod	uct was pack	ed:					
	(²) either	[in new containers;]							
	(2) or	[in vehicles or bulk conta	iners disinfec	ted prior to loading using a product app	roved by the competent authority;]				
and the containers are marked so as to indicate the nature of the milk/milk-based product and bear labels indicating HUMAN CONSUMPTION".									
	Notes								
	Part I:								
				e for the consignment in EU: this box rtificate is for import commodity.	is to be filled in only if it is a certificate for transit				
	_			on: this box is to be filled in only if it is s, free warehouses and custom wareho	a certificate for transit commodity. The products in uses.				
	_			nber (railway wagons or container an f unloading and reloading.	d lorries), flight number (aircraft) or name (ship);				
	_	Box reference I.19; use t	he appropriat	e HS code: 23.09.10; 23.09.90; 35.01;	35.02 or 35.04.				
	-	Box reference 1.23: for b	ulk containers	k containers, the container number and the seal number (if applicable) should be included.					
	_	Box reference I.26 and I.	27: fill in acco	7: fill in according to whether it is a transit or an import certificate.					
	_	Box reference I.28: Man	ıfacturing pla	acturing plant: provide the registration number of treatment or processing establishment.					
	Part II:								
	(¹)	OJ L 273, 10.10.2002, p	1.						
	(2)	Delete as appropriate.							
		The signature and the st	amp must be	in a different colour to that of the printing	g.				
 Note for the person responsible for the consignment in EU: this certificate is only for vertice the consignment until it reaches the border inspection post. 				only for veterinary purposes and has to accompany					
	Official vete	fficial veterinarian							
		Name (in capitals):		Q	ualification and title:				
		Date:		Si	gnature:				
		Stamp:							

CHAPTER 3(A)

Health certificate

For canned petfood intended for dispatch to or for transit through $(^2)$ the European Community

COUN	TRY	Veterinary certificate to EU
	I.1. Consignor Name	I.2. Certificate reference number I.2.a.
	Address	I.3. Central Competent Authority
nent	Tel. No	I.4. Local Competent Authority
sign	I.5. Consignee	I.6. Person responsible for the consignment in EU
con	Name Address	Name Address
ched		
ispat	Postal code Tel. No	Postal code Tel. No
Part I: Details of dispatched consignment	I.7. Country of ISO code I.8. Region of origin Code origin	I.9. Country of ISO code I.10. Region of Code destination
: Det	I.11. Place of origin	I.12. Place of destination
Part		Custom warehouse
	Name Approval number Address	Name Approval number Address
	Addiese	, Address
		Postal code
	I.13. Place of loading	I.14. Date of departure
	I.15. Means of transport Aeroplane Ship Raitway wagon ☐	I.16. Entry BIP in EU
	Road vehicle Other	
	Identification: Documentary references:	1.17.
	I.18. Description of commodity	I.19. Commodity code (HS code)
	Social plants of commonly	23.09.10
		I.20. Quantity
	I.21. Temperature of product	I.22. Number of packages
	Ambient Chilled	Frozen 🔲
	I.23. Identification of container/Seal number	I.24. Type of packaging
	I.25. Commodities certified for: Animal feedingstuff	Technical use Other
	I.26. For transit to third country vis-à-vis EU	1.27. For import or admission into EU
	3rd country ISO code	
	I.28. Identification of the commodities	
	Approval number of establishr Species Manufacturing plant (Scientific name)	nents Net weight Batch number

COUNTRY Canned Petfood

		II.a. Certificate reference number II.b.			
	II.	Health attestation			
		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (1) and in particular			
		Article 6 and Annex VIII Chapter II thereof and certify that the petfood described above:			
ll.1. has been prepared and stored in a plant approved and supervised by the competent authority in accordance wi where appropriate Article 11 of Regulation (EC) No 1774/2002;					
ertifi	11.2.	has been prepared exclusively with the following animal by-products:			
Part II: Certification	(²) either	 parts of slaughtered animals, which were fit for human consumption in accordance with Community legislation, but are not intended for human consumption for commercial reasons,] 			
Pa	(²) and/or	[— parts of slaughtered animals, which were rejected as unfit for human consumption but are not affected by any signs of diseases communicable to humans or animals and derive from carcases that were fit for human consumption in accordance with Community legislation,]			
	(²) and/or	 hides and skins, hooves and horns, pig bristles and feathers originating from animals that were slaughtered in a slaughterhouse, underwent ante-mortem inspection and were fit, as a result of such inspection, for slaughter in accordance with Community legislation,] 			
	(²) and/or	[— blood obtained from animals other than ruminants that were slaughtered in a slaughterhouse, underwent antemortem inspection and were fit, as a result of such inspection, for slaughter in accordance with Community legislation,]			
	(²) and/or	[— animal by-products derived from the production of products intended for human consumption, including degreased bones and greaves,]			
	(²) and/or	[— former foodstuffs of animal origin, or former foodstuffs containing products of animal origin, other than catering waste, which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects which do not present any risk to humans or animals,]			
	(²) and/or	[— raw milk originating from animals that do not show clinical signs of any disease communicable through that product to humans or animals.]			
	(2) and/or	[— fish or other sea animals, except sea mammals, caught in the open sea for the purposes of fishmeal production,]			
	(2) and/or	[— fresh by-products from fish from plants manufacturing fish products for human consumption,]			
	(²) and/or	[— shells, hatchery by-products and cracked egg by-products originating from animals which did not show clinical signs of any disease communicable through that product to humans or animals,]			
	(²) andlor	[— material from animals which have been treated with certain substances which are prohibited pursuant to Directive 96/22/EC, the import of the material being permitted in accordance with Article 28 of Regulation (EC) No 1774/2002];			
	II.3.	has been subjected to heat treatment to a minimum Fc value of 3 in hermetically sealed containers;			
	II.4.	was analysed by a random sampling of at least five containers from each processed batch by laboratory diagnostic methods to ensure adequate heat treatment of the whole consignment as foreseen under point II.3;			
	II.5.	has undergone all precautions to avoid contamination with pathogenic agents after treatment.			
	Notes				
	Part I:				
	_	Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity.			
	_	Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses.			
	_	Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in the event of unloading and reloading.			
	_	Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be given.			
	_	Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.			
l					

Part II:		
(¹)	OJ L 273, 10.10.2002, p. 1.	
(2)	Delete as appropriate.	
_	The signature and the stamp must be in a different colour to that of the pri	nting.
_	Note for the person responsible for the consignment in EU: This certificat the consignment until it reaches the border inspection post.	e is only for veterinary purposes and has to accompany
Official vet	erinarian	
	Name (in capitals):	Qualification and title:
	Date:	Signature:
	Stamp:	

CHAPTER 3(B)

Health certificate

For processed petfood other than canned petfood, intended for dispatch to or for transit through (²) the European Community

COUN	TRY			Veterinary certificate to El
	l.1.	Consignor Name	1.2.	Certificate reference I.2.a. number
		Address	1.3.	Central Competent Authority
ent		Tel. No	1.4.	Local Competent Authority
Part I: Details of dispatched consignment	1.5.	Consignee	1.6.	Person responsible for the consignment in EU
ons		Name		Name
pa		Address		Address
tch		Postal code		Postal code
lispa		Tel. No	ļ	Tel. No
ofo	1.7.	Country of ISO code I.8. Region of origin Code	1.9.	Country of ISO code I.10. Region of Code
tails		origin		destination destination
. De	l.11.	Place of origin	1.12.	Place of destination
arti				Custom warehouse
۵		Name Approval number		Name Approval number
		Address		Address
				Postal code
	1.40		<u> </u>	
	1.13.	Place of loading	1.14.	Date of departure
	I.15.	Means of transport	1.16.	6. Entry BIP in EU
		Aeroplane Ship Railway wagon		
		Road vehicle Other I	<u> </u>	
		Documentary references:	1.17.	
	I.18.	Description of commodity		I.19. Commodity code (HS code)
				23.09.10
				I.20. Quantity
	1.21.	Temperature of product		I.22. Number of packages
		Ambient Chilled Chilled		Frozen 🔲
	1.23.	Identification of container/Seal number		I.24. Type of packaging
	1.25.	Commodities certified for: Animal feedingstuff		Technical use Other
	126	For transit to third country vis-à-vis EU	127	. For import or admission into EU
	1.20.		1.21.	. For import of duminosoff into 25
		3rd country ISO code		
	1.28.	Identification of the commodities		
		Species		
		Approval number of establishments Manufacturin	a plant	t Net weight Batch number
		(Scientific name)		5

COUNTRY

Processed petfood other than canned petfood

		II.a. Certificate reference number II.b.						
	II.	Health attestation						
		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (¹) and in particular Article 6 and Annex VIII Chapter II thereof and certify that the petfood described above:						
ication	II.1.	has been prepared and stored in a plant approved and supervised by the competent authority in accordance with Article 18 and where appropriate Article 11 of Regulation (EC) No 1774/2002;						
ertif	II.2.	has been prepared exclusively with the following animal by-products:						
Part II: Certification	(²) either	[— parts of slaughtered animals, which were fit for human consumption in accordance with Community legislation, but are not intended for human consumption for commercial reasons,]						
-	(²) and/or	[— parts of slaughtered animals, which were rejected as unfit for human consumption but are not affected by any signs of diseases communicable to humans or animals and derive from carcases that were fit for human consumption in accordance with Community legislation.]						
	(²) and/or	[— hides and skins, hooves and horns, pig bristles and feathers originating from animals that were slaughtered in a slaughterhouse, underwent ante-mortem inspection and were fit, as a result of such inspection, for slaughter in accordance with Community legislation,]						
	(²) and/or	[— blood obtained from animals other than ruminants that were slaughtered in a slaughterhouse, underwent ante-mortem inspection and were fit, as a result of such inspection, for slaughter in accordance with Community legislation,]						
	(²) and/or	[— animal by-products derived from the production of products intended for human consumption, including degreased bones and greaves,]						
	(²) and/or	[— former foodstuffs of animal origin, or former foodstuffs containing products of animal origin, other than catering waste, which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects which do not present any risk to humans or animals,]						
	(²) and/or	[— raw milk originating from animals that do not show clinical signs of any disease communicable through that product to humans or animals,]						
	(2) and/or	[— fish or other sea animals, except sea mammals, caught in the open sea for the purposes of fishmeal production,]						
	(²) and/or	[— fresh by-products from fish from plants manufacturing fish products for human consumption,]						
	(²) and/or	[— shells, hatchery by-products and cracked egg by-products originating from animals which did not show clinical signs of any disease communicable through that product to humans or animals,]						
	(²) and/or	[— material from animals which have been treated with certain substances which are prohibited pursuant to Directive 96/22/EC, the import of the material being permitted in accordance with Article 28 of Regulation (EC) No 1774/2002;]						
	II.3.							
	(²) either [was subjected to a heat treatment of at least 90 °C throughout its substance;]							
	(²) or [was produced as regards ingredients of animal origin using exclusively products which had been							
	(a) in the case of meat or meat products subjected to a heat treatment of at least 90 °C throughout its substance;							
	(b) in the case of milk and milk based products,							
	 (i) if they are from third countries or parts of third countries listed in column B of Annex I to Decision 200 submitted to a pasteurisation treatment sufficient to produce a negative phosphatase test; 							
		(ii) with a pH reduced to less than 6 from third countries or parts of third countries listed in column C of Annex I to Decision 2004/438/EC, first submitted to a pasteurisation treatment sufficient to produce a negative phosphatase test;						
	(iii) if they are from third countries or parts of third countries listed in column C of Annex I to Decision 2004/43 submitted to a sterilisation process or a double heat treatment where each treatment was sufficient to produce a neg phosphatase test on its own;							

(iv) if they are from third countries or parts of third countries listed in column C of Annex I to Decision 2004/438/EC where there has been an outbreak of foot-and-mouth disease in the last 12 months or where vaccination against foot-andmouth disease has been carried out in the last 12 months submitted to:

either

a sterilisation process whereby an Fc value equal or greater than 3 is achieved,

or

an initial heat treatment with a heating effect at least equal to that achieved by a pasteurisation process of at least
 72 °C for at least 15 seconds and sufficient to produce a negative reaction to a phosphatase test, followed by

either

a second heat treatment with a heating effect at least equal to that achieved by the initial heat treatment, and which
would be sufficient to produce a negative reaction to a phosphatase test, followed, in the case of dried milk, or dried
milk-based products by a drying process,

Of

- an acidification process such that the pH has been maintained at less than 6 for at least one hour;
- (c) in the case of gelatine, produced using a process that ensures that unprocessed Category 3 material is subjected to a treatment with acid or alkali, followed by one or more rinses with subsequent adjustment of the pH and subsequent, if necessary repeated, extraction by heat, followed by purification by means of filtration and sterilisation;
- (d) in the case of hydrolysed protein produced using a production process involving appropriate measures to minimise contamination of raw Category 3 material, using only material with a molecular weight below 10 000 Dalton and, in the case of hydrolysed protein entirely or partly derived from ruminant hides and skins produced in a processing plant dedicated only to hydrolysed protein production, using a process involving the preparation of raw Category 3 material by brining, liming and intensive washing followed by
 - (i) exposure of the material to a pH of more than 11 for more than three hours at a temperature of more than 80 °C and subsequently by heat treatment at more than 140 °C for 30 minutes at more than 3,6 bar; or
 - (ii) exposure of the material to a pH of 1 to 2, followed by a pH of more than 11, followed by heat treatment at 140° C for 30 minutes at 3 bar:
- (e) in the case of egg products submitted to any of the processing methods 1 to 5 or 7, as referred to in Annex V Chapter III to Regulation (EC) No 1774/2002; or treated in accordance with Chapter II of Section X of Annex III to Regulation (EC) No 853/2004 (4);
- (f) in the case of collagen submitted to a process ensuring that unprocessed Category 3 material is subjected to a treatment involving washing, pH adjustment using acid or alkali followed by one or more rinses, filtration and extrusion, the use of preservatives other than those permitted by Community legislation being prohibited;
- (g) in the case of blood products, produced using any of the processing methods 1 to 5 or 7, as referred to in Annex V Chapter III to Regulation (EC) No 1774/2002;
- (h) in the case of mammalian processed animal protein submitted to any of the processing methods 1 to 5 or 7 and, in the case of porcine blood, submitted to any of the processing methods 1 to 5 or 7 provided that in the case of method 7 a heat treatment throughout its substance at a minimum temperature of 80 °C has been applied;
- in the case of non-mammalian processed protein with the exclusion of fishmeal submitted to any of the processing methods 1 to 5 or 7 as referred to in Annex V Chapter III to Regulation (EC) No 1774/2002;
- (k) in the case of fishmeal submitted to any of the processing methods or to a method and parameters which ensure that the products complies with the microbiological standards set in Annex VII Chapter I paragraph 10 to Regulation (EC) No 1774/2002;
- (I) in the case of rendered fat, including fish oils, submitted to processing methods 1 to 5 or 7 (and method 6 in the case of fish oil) as referred to in Annex V Chapter III to Regulation (EC) No 1774/2002 or produced in accordance with Chapter II of Section XII of Annex III to Regulation (EC) No 853/2004 (*); rendered fats from ruminant animals must be purified in such a way that the maximum level of remaining total insoluble impurities does not excess 0,15 % in weight;
- (m) in the case of dicalcium phosphate produced by a process that
 - ensures that all Category 3 bone-material is finely crushed and degreased with hot water and treated with dilute hydrochloric acid (at a minimum concentration of 4 % and a pH of less than 1,5) over a period of at least two days;
 - (ii) following the procedure under (i), applies a treatment of the obtained phosphoric liquor with lime, resulting in a precipitate of dicalcium phosphate at pH 4 to 7; and
 - (iii) finally, air dries the precipitate of dicalcium phosphate with inlet temperature of 65 °C to 325 °C and end temperature between 30 °C and 65 °C;

	(11) 111 ti	ie case of incarcium phospitate produce	by a process that ensures
	(i)	that all Category 3 bone-material is fin 14 mm);	ely crushed and degreased in counter-flow with hot water (bone chips less than
	(ii)	continuous cooking with steam at 145	°C during 30 minutes at 4 bar;
	(iii)	separation of the protein broth from the	e hydroxyapatite (tricalciuum phosphate) by centrifugation; and
	(iv)	granulation of the tricalcium phosphate	e after drying in a fluid bed with air at 200 °C];
II.4.		ysed by a random sampling of at least ng plant and complies with the following s	five samples from each processed batch taken during or after storage at the standards $(^5)$:
	Salmone	lla: absence in 25 g	g: n = 5, c = 0, m = 0, M = 0;
	Enteroba	cteriaceae: n = 5, c = 2, m	= 10, M = 300 in 1 gram;
II.5.	has unde	rgone all precautions to avoid contamina	ation with pathogenic agents after treatment;
II.6.			od is not dispatched in ready-to-sale packages on which it is clearly indicated that are labels indicating "NOT FOR HUMAN CONSUMPTION".
Notes			
Part I:			
_		rence I.6: Person responsible for the co ty; it may be filled in if the certificate is fo	onsignment in EU: this box is to be filled in only if it is a certificate for transit r import commodity.
_		ence I.12: Place of destination: this box is be stored in free zones, free warehouses	s to be filled in only if it is a certificate for transit commodity. The products in transit s and custom warehouses.
_		ence I.15: Registration number (railway w rovided in the event of unloading and rel	ragons or container and lorries), flight number (aircraft) or name (ship); information oading.
_	Box refer	ence I.23: for bulk containers, the contain	ner number and the seal number (if applicable) should be given.
_	Box refer	ence I.26 and I.27: fill in according to wh	ether it is a transit or an import certificate.
Part II:	O I I 273	, 10.10.2002, p. 1.	
(2)		s appropriate.	
(3)		, 30.4.2004, p. 55. Corrected by OJ L 22	6, 25.6.2004, p. 22.
(4)		, 25.6.2004, p. 22.	
(5)	Where:	·	
	n = nur	mber of samples to be tested;	
		eshold value for the number of bacteria; exceed m;	the result is considered satisfactory if the number of bacteria in all samples does
		ximum value for the number of bacteria; nples is M or more; and	the result is considered unsatisfactory if the number of bacteria in one or more
		mber of samples the bacterial count of whoterial count of the other samples is m or	lich may be between m and M, the sample still being considered acceptable if the less.
_	The signa	ature and the stamp must be in a differer	at colour to that of the printing.
_		the person responsible for the consignm gnment until it reaches the border inspec	ent in EU: This certificate is only for veterinary purposes and has to accompany tion post.
Official ve	torinarion		
Onicial ve	rei ii ialiali		
	Name (in	capitals):	Qualification and title:
	Date:		Signature:
	Stamp:		

CHAPTER 3(C)

Health certificate

For dogchews intended for dispatch to or for transit through (2) the European Community

COUN	TRY				Veterinary certificate to EU
	l.1.	Consignor Name	1.2.	Certificate reference number	1.2.a.
		Address	1.3.	Central Competent Authori	ty
Part I: Details of dispatched consignment		Tel. No	1.4.	Local Competent Authority	
signr	1.5.	Consignee	1.6.	Person responsible for the	consignment in EU
Ö		Name		Name	
p _e		Address		Address	
tch		Postal code		Postal code	
ispa		Tel. No		Tel. No	
of d	1.7.	Country of ISO code I.8. Region of origin Code	1.9.	Country of ISO code	I.10. Region of Code
ails		origin		destination	destination
Det	I 11	Place of origin	112	Place of destination	<u> </u>
Ë		Thad of origin		Custom wareho	iise 🗖
Pa		Name Approval number		Name	Approval number
		Address		Address	Approvariamber
		1.000		, taarooo	
				Postal code	
	I.13.	Place of loading	1.14.	Date of departure	
	l.15.	Means of transport	1.16.	Entry BIP in EU	
		Aeroplane Ship Railway wagon			
		Road vehicle Other	L		
		Identification: Documentary references:	1.17.		
	1.10	Description of commodity		L10 Common	dity code (HS code)
	1, 10.	Description of commodity		1.19. Commod	42.05.00
					I.20. Quantity
	1.21.	Temperature of product			I.22. Number of packages
		Ambient Chilled		Frozen	
				1102011 🔲	
	1.23.	Identification of container/Seal number			I.24. Type of packaging
	1.25.	Commodities certified for:			
		Animal feedingstuff		Technical use	Other
	1.26.	For transit to third country vis-à-vis EU	1.27.	For import or admission into	o EU
		3rd country ISO code			
	1.28.	Identification of the commodities			
		Approval number of			577
		Species Manufactu (Scientific name)	ring pia	int Net	weight Batch number
		(Selection results)			
	1				

		II.a. Certificate reference number II.b.							
		II.a. Certificate reference number II.b.							
	II.	Health attestation							
		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (') and in particular							
١		Article 6 and Annex VIII Chapter II thereof and certify that the dogchews described above:							
	II.1.	have been prepared and stored in a plant approved and supervised by the competent authority in accordance with Article 18 and where appropriate Article 11 of Regulation (EC) No 1774/2002;							
١	II.2.	have been prepared exclusively with the following animal by-products:							
	(²) either	 parts of staughtered animals, which were fit for human consumption in accordance with Community legislation, but are n intended for human consumption for commercial reasons,] 							
	(°) and/or	[— parts of slaughtered animals, which were rejected as unfit for human consumption but are not affected by any signs diseases communicable to humans or animals and derive from carcasses that were fit for human consumption in accordance with Community legislation,]							
	(°) and/or	i hides and skins originating from animals that were slaughtered in a slaughterhouse, underwent ante-mortem inspection ar were fit, as a result of such inspection, for slaughter in accordance with Community legislation,]							
	(²) and/or	[— animal by-products derived from the production of products intended for human consumption, including degreased bones are greaves.]							
١	(²) and/or	[— fresh by-products from fish from plants manufacturing fish products for human consumption,]							
_	(²) and/or	material from animals which have been treated with certain substances which are prohibited pursuant to Directive 96/22/E0 the import of the material being permitted in accordance with Article 28 of Regulation (EC) No 1774/2002];							
	II.3.	have been subjected:							
	(²) either	[in the case of dogchews made from hides and skins of ungulates or from fish, to a treatment sufficient to destroy pathogenic organisms (including salmonella); and the dogchews are dry];							
	(²) or	[in the case of dogchews made from animal by-products other than hides and skins of ungulates or from fish, to a heat treatment of at least 90 °C throughout their substance;]							
	11.4.	were examined by random sampling of at least five samples from each processed batch taken during or after storage at the processing plant and complies with the following standards (*):							
		Salmonella: absence in 25 g: n = 5, c = 0, m = 0, M = 0;							
		Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram;							
	II.5.	have undergone all precautions to avoid contamination with pathogenic agents after treatment;							
	11.6.	were packed in new packaging.							
	Notes								
	Part I:								
	_	Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for transcommodity; it may be filled in if the certificate is for import commodity.							
	_	Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transcan only be stored in free zones, free warehouses and custom warehouses.							
	_	Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in the event of unloading and reloading.							
		Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be given.							

(') (')	OJ L 273, 10.10.2002, p. 1.				
(2)					
O	Delete as appropriate.				
(°)	Where:				
	n = number of samples to be tested;				
	m = threshold value for the number of bacteria; the not exceed m;	e result is considered satisfactory if the number of bacteria in all samples does			
	M = maximum value for the number of bacteria; the samples is M or more; and	ne result is considered unsatisfactory if the number of bacteria in one or more			
	c = number of samples the bacterial count of whic bacterial count of the other samples is m or le	h may be between m and M, the sample still being considered acceptable if the ss.			
	The signature and the stamp must be in a different	colour to that of the printing.			
_	Note for the person responsible for the consignme the consignment until it reaches the border inspection	nt in EU: This certificate is only for veterinary purposes and has to accompany on post.			
Official ve	eterinarian				
	Name (in capitals):	Qualification and title:			
	Date: Signature:				
	Stamp:				

CHAPTER 3(D)

Health certificate

For raw petfood for direct sale or animal by-products to be fed to farmed fur animals, intended for dispatch to or for transit through (*) the European Community

COUN	TRY			Veterinary certificate	to EU
	l.1.	Consignor Name	1.2.	Certificate reference I.2.a.	
		Address	1.3.	Central Competent Authority	
ment		Tel. No	1.4.	Local Competent Authority	
igu	1.5.	Consignee	1.6.	Person responsible for the consignment in EU	
ő		Name		Name	
ped		Address		Address	
atch		Postal code		Postal code	
disp		Tel. No		Tel. No	
Part I: Details of dispatched consignment	1.7.	Country of ISO code I.8. Region of origin Code origin	1.9.	Country of ISO code I.10. Region of Codestination destination	Code
. De	l.11.	Place of origin	1.12.	2. Place of destination	
art				Custom warehouse	
"		Name Approval number		Name Approval number	
		Address		Address	
				Postal code	
	1.13.	Place of loading	1.14.	Date of departure	
	1.15.	Means of transport	1.16.	6. Entry BIP in EU	
		Aeroplane Ship Railway		•	
		Road vehicle Other wagon Identification:		<u>,</u>	
		Documentary references:	1.17.		
	l.18.	Description of commodity		I.19. Commodity code (HS code)	
				I.20. Quantity	
	ļ				
	1.21.	Temperature of product		I.22. Number of packages	
		Ambient Chilled Chilled		Frozen	
	1.23.	Identification of container/Seal number		I.24. Type of packaging	
	1.25.	Commodities certified for:			
		Animal feedingstuff		Technical use Other Other	
	1.26.	For transit to third country vis-à-vis EU	1.27.	. For import or admission into EU	
		3rd country ISO code			
	1.28.	Identification of the commodities			
		Approval	numbe	er of establishments	
		Species Nature of commodity	Manufa	acturing plant Net weight Batch number	
		(Scientific name)			

COUNTRY

Raw petfood for direct sale or animal by products to be fed to farmed fur animals

II. Health attestation I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (1) and in particul Article 6 and Annex VIII Chapter II thereof and certify that the raw perfood or animal by-product described above: II.1. consist of animal by-products that satisfy the health requirements below. II.2. consist of animal by-products: (a) derived from meat which satisfies the relevant animal and public health requirements laid down in: — Council Decision 79543/EEC (?) and provided the animals from which the meat is derived come from of a territory (So code) as listed in that Decision which has been free of foot and mouth disease, rinderper classical swine fever, African swine fever and swine vesscular disease for the last 12 months and where no vaccination has taken place during that time (only as relevant for the susceptible species). — and/or Commission Decision 2006/95/EC (?), and provided the animals from which the meat is derived come from territory or part of a territory (ISO code) as listed with that Decision which has been free from Newcastle disease and Avian Influenza for the last 12 months and provided the animals from which the meat is derived come from territory or part of a territory (ISO code) as listed with has been free from foot and mouth disease, inderper classical swine fever, African swine fever, swine vesicular disease. Newcastle disease and Avian Influenza for the last 12 months and where no vaccination has been free from foot and mouth disease, inderper classical swine fever, African swine fever, swine vesicular diseases. Newcastle disease and Avian influenza for the susceptible species). (b) derived from animals that, at the slaughterhouse, have passed the anter-morem health inspection during the 24 hour before the slaughter and have shown no evidence of the diseases referred in the Decisions above for which the animals as ausceptible; and (c) derived from animals that have been	1			II.a.	Certificate reference number	II.b.				
I. the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (*) and in particul Article 6 and Annex VIII Chapter II thereof and certify that the raw petfood or animal by-product described above: II.1. consist of animal by-products that satisfy the health requirements below, III.2. consist of animal by-products: (a) derived from meat which satisfies the relevant animal and public health requirements laid down in: — Council Decision 79/542/EEC (*) and provided the animals from which the meat is derived come from a territory or profit a territory (SO code) as listed in that Decision which has been free of foot and mouth disease, inderpost classical swine fever. African swine fever and swine vesicular disease for the last 12 months and where no vaccination has taken place during that time (only as relevant for the susceptible spoces). — and/or Commission Decision 2006/66/EC (*), and provided the animals from which the meat is derived come from territory or part of a territory (SO code) as listed which has been free from Newcastle disease and Avian Influenza for the last 12 months, — and/or Commission Decision 2006/65/EC (*), and provided the animals from which the meat is derived come from territory or part of a territory (SO code) as listed which has been free from foot and mouth disease, inderpost classical swine fever, African swine fever, swine vesicular disease, Newcastle disease and value influenza for the last 12 months and where no vaccination has taken place during that time (only as relevant for the susceptible species). (b) derived from animals that, at the slaughterhouse, have passed the anim-mortem health inspection during the 24 hou before the slaughter and have shown no evidence of the diseases referred in the Decisions above for which the animals assuceptible, and (c) derived from animals that have been treated in the slaughterhouse before and at the time of slaughter or killing in accordanc										
I. the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (*) and in particul Article 6 and Annex VIII Chapter II thereof and certify that the raw petfood or animal by-product described above: II.1. consist of animal by-products that satisfy the health requirements below, III.2. consist of animal by-products: (a) derived from meat which satisfies the relevant animal and public health requirements laid down in: — Council Decision 79/542/EEC (*) and provided the animals from which the meat is derived come from a territory or profit a territory (SO code) as listed in that Decision which has been free of foot and mouth disease, inderpost classical swine fever. African swine fever and swine vesicular disease for the last 12 months and where no vaccination has taken place during that time (only as relevant for the susceptible spoces). — and/or Commission Decision 2006/66/EC (*), and provided the animals from which the meat is derived come from territory or part of a territory (SO code) as listed which has been free from Newcastle disease and Avian Influenza for the last 12 months, — and/or Commission Decision 2006/65/EC (*), and provided the animals from which the meat is derived come from territory or part of a territory (SO code) as listed which has been free from foot and mouth disease, inderpost classical swine fever, African swine fever, swine vesicular disease, Newcastle disease and value influenza for the last 12 months and where no vaccination has taken place during that time (only as relevant for the susceptible species). (b) derived from animals that, at the slaughterhouse, have passed the anim-mortem health inspection during the 24 hou before the slaughter and have shown no evidence of the diseases referred in the Decisions above for which the animals assuceptible, and (c) derived from animals that have been treated in the slaughterhouse before and at the time of slaughter or killing in accordanc										
Article 6 and Ānnex VIII Chapter II thereof and certify that the raw perfood or animal by-product described above: II.1. consist of animal by-products: (a) derived from meat which satisfies the relevant animal and public health requirements laid down in: — Council Decision 79/54/2/EEC (?) and provided the animals from which the meat is derived come from a territory or per of a territory		II.	Health attestation							
Countries and services of the service of the products of the service of the servi										
Countries and services of the service of the products of the service of the servi		II.1.	consist of animal by-products	that sati	sfy the health requirements below;					
Countries and services of the service of the products of the service of the servi		II.2.	consist of animal by-products	:						
Countries and services of the service of the products of the service of the servi			(a) derived from meat which satisfies the relevant animal and public health requirements laid down in:							
territory or part of a territory			of a territory classical swine fev	(ISC er, Afric	code) as listed in that Decision which an swine fever and swine vesicular di	h has been free of foot and mouth disease, rinderpest, sease for the last 12 months and where no vaccination				
territory or part of a territory (ISO code) as listed which has been free from foot and mouth disease, indergee classical swine fever, African swine fever, as wine vesticular disease, Newcode ideaseas en davia influenza for the la 12 months and where no vaccination has taken place during that time (only as relevant for the susceptible species). (b) derived from animals that, at the slaughterhouse, have passed the ante-mortem health inspection during the 24 hou before the slaughter and have shown no evidence of the diseases referred in the Decisions above for which the animals as usceptible; and (c) derived from animals that have been treated in the slaughterhouse before and at the time of slaughter or killing in accordance with the relevant provisions of Council Directive 93/119/EC (?) on animal welfare; II.3. consist only of the following animal by-products: (a) parts of slaughtered animals, which were fit for human consumption in accordance with Community legislation, but are n intended for human consumption for commercial reasons, and (b) parts of slaughtered animals, which are rejected as unfit for human consumption but are not affected by any signs of disease communicable to humans or animals and derive from carcasses that are fit for human consumption in accordance with Community legislation; II.4. have been obtained and prepared without contact with other material not complying with the conditions required in the Decision above, and it has been handled so as to avoid contamination with pathogenic agents; II.5. have been packed in final packaging which bear labels indicating "RAW PETFOOD — NOT FOR HUMAN CONSUMPTION" and then in leak-proof are officially sealed boxes/containers or in new packaging preventing any leakage and officially sealed boxes/containers which be labels indicating "RAW PETFOOD — NOT FOR HUMAN CONSUMPTION" in he name and the address of the establishment of destination; II.6. in the case of raw petfood: (a) have been prepared and stored in a plant ap			territory or part of	a territor	/ (ISO code) as listed in that					
before the slaughter and have shown no evidence of the diseases referred in the Decisions above for which the animals as usceptible; and (c) derived from animals that have been treated in the slaughterhouse before and at the time of slaughter or killing in accordance with the relevant provisions of Council Directive 93/119/EC (*) on animal welfare; (a) parts of slaughtered animals, which were fit for human consumption in accordance with Community legislation, but are n intended for human consumption for commercial reasons, and (b) parts of slaughtered animals, which are rejected as unfit for human consumption but are not affected by any signs of disease communicable to humans or animals and derive from carcasses that are fit for human consumption in accordance with Community legislation; II.4. have been obtained and prepared without contact with other material not complying with the conditions required in the Decision above, and it has been handled so as to avoid contamination with pathogenic agents; II.5. have been packed in final packaging which bear labels indicating "RAW PETFOOD — NOT FOR HUMAN CONSUMPTION" "ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS — NOT FOR HUMAN CONSUMPTION" and then in leak-proof an officially sealed boxes/containers or in new packaging preventing any leakage and officially sealed boxes/containers which be labels indicating "RAW PETFOOD — NOT FOR HUMAN CONSUMPTION" for FEED FOR FEU FOR FEU FOR FUR ANIMALS — NOT FOR HUMAN CONSUMPTION". II.6. in the case of raw petfood: (a) have been prepared and stored in a plant approved and supervised by the competent authority in accordance with Article and where appropriate Article 11 of Regulation (EC) No 1774/2002; and (b) were examined by random sampling of at least five samples from each batch taken during storage (before dispatch) are complies with the following standards (*): Salmonella: absence in 25 g: n = 5, c = 0, m = 0, M = 0;			territory or part of a classical swine fe	a territory er, Afric	/ (ISO code) as listed which an swine fever, swine vesicular disea	has been free from foot and mouth disease, rinderpest, se, Newcastle disease and Avian Influenza for the last				
 with the relevant provisions of Council Directive 93/119/EC (*) on animal welfare; (a) parts of slaughtered animals, which were fit for human consumption in accordance with Community legislation, but are not intended for human consumption for commercial reasons, and (b) parts of slaughtered animals, which are rejected as unfit for human consumption but are not affected by any signs of disease communicable to humans or animals and derive from carcasses that are fit for human consumption in accordance with Community legislation; II.4. have been obtained and prepared without contact with other material not complying with the conditions required in the Decision above, and it has been handled so as to avoid contamination with pathogenic agents; II.5. have been packed in final packaging which bear labels indicating "RAW PETFOOD — NOT FOR HUMAN CONSUMPTION" "ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS — NOT FOR HUMAN CONSUMPTION" and then in leak-proof an officially sealed boxes/containers or in new packaging preventing any leakage and officially sealed boxes/containers which be labels indicating "RAW PETFOOD — NOT FOR HUMAN CONSUMPTION" or "ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS — NOT FOR HUMAN CONSUMPTION", the name and the address of the establishment of destination; II.6. in the case of raw petfood: (a) have been prepared and stored in a plant approved and supervised by the competent authority in accordance with Article and where appropriate Article 11 of Regulation (EC) No 1774/2002; and (b) were examined by random sampling of at least five samples from each batch taken during storage (before dispatch) are complies with the following standards (*9): Salmonella: absence in 25 g: n = 5, c = 0, m = 0, M = 0; 	before the slaughter and have shown no evidence of the diseases referred in the D									
 (a) parts of slaughtered animals, which were fit for human consumption in accordance with Community legislation, but are n intended for human consumption for commercial reasons, and (b) parts of slaughtered animals, which are rejected as unfit for human consumption but are not affected by any signs of disease communicable to humans or animals and derive from carcasses that are fit for human consumption in accordance with Community legislation; II.4. have been obtained and prepared without contact with other material not complying with the conditions required in the Decision above, and it has been handled so as to avoid contamination with pathogenic agents; III.5. have been packed in final packaging which bear labels indicating "RAW PETFOOD — NOT FOR HUMAN CONSUMPTION" "ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS — NOT FOR HUMAN CONSUMPTION" and then in leak-proof are officially sealed boxes/containers or in new packaging preventing any leakage and officially sealed boxes/containers which bear labels indicating "RAW PETFOOD — NOT FOR HUMAN CONSUMPTION" or "ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS — NOT FOR HUMAN CONSUMPTION", the name and the address of the establishment of destination; II.6. in the case of raw petfood: (a) have been prepared and stored in a plant approved and supervised by the competent authority in accordance with Article and where appropriate Article 11 of Regulation (EC) No 1774/2002; and (b) were examined by random sampling of at least five samples from each batch taken during storage (before dispatch) are complies with the following standards (⁶): Salmonella: absence in 25 g: n = 5, c = 0, m = 0, M = 0; 										
intended for human consumption for commercial reasons, and (b) parts of slaughtered animals, which are rejected as unfit for human consumption but are not affected by any signs of disease communicable to humans or animals and derive from carcasses that are fit for human consumption in accordance with Community legislation; II.4. have been obtained and prepared without contact with other material not complying with the conditions required in the Decision above, and it has been handled so as to avoid contamination with pathogenic agents; II.5. have been packed in final packaging which bear labels indicating "RAW PETFOOD — NOT FOR HUMAN CONSUMPTION" and then in leak-proof are officially sealed boxes/containers or in new packaging preventing any leakage and officially sealed boxes/containers which be labels indicating "RAW PETFOOD — NOT FOR HUMAN CONSUMPTION" or "ANIMAL BY-PRODUCTS FOR FEED FOR FU ANIMALS — NOT FOR HUMAN CONSUMPTION", the name and the address of the establishment of destination; II.6. in the case of raw petfood: (a) have been prepared and stored in a plant approved and supervised by the competent authority in accordance with Article and where appropriate Article 11 of Regulation (EC) No 1774/2002; and (b) were examined by random sampling of at least five samples from each batch taken during storage (before dispatch) are complies with the following standards (*): **Salmonella:** **absence in 25 g: n = 5, c = 0, m = 0, M = 0;		II.3.	consist only of the following a	nimal by	-products:					
communicable to humans or animals and derive from carcasses that are fit for human consumption in accordance with Community legislation; II.4. have been obtained and prepared without contact with other material not complying with the conditions required in the Decision above, and it has been handled so as to avoid contamination with pathogenic agents; II.5. have been packed in final packaging which bear labels indicating "RAW PETFOOD — NOT FOR HUMAN CONSUMPTION" "ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS — NOT FOR HUMAN CONSUMPTION" and then in leak-proof are officially sealed boxes/containers or in new packaging preventing any leakage and officially sealed boxes/containers which bear labels indicating "RAW PETFOOD — NOT FOR HUMAN CONSUMPTION" or "ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS — NOT FOR HUMAN CONSUMPTION", the name and the address of the establishment of destination; II.6. in the case of raw petfood: (a) have been prepared and stored in a plant approved and supervised by the competent authority in accordance with Article and where appropriate Article 11 of Regulation (EC) No 1774/2002; and (b) were examined by random sampling of at least five samples from each batch taken during storage (before dispatch) are complies with the following standards (8): **Salmonella:** **absence in 25 g: n = 5, c = 0, m = 0, M = 0;						in accordance with Community legislation, but are not				
above, and it has been handled so as to avoid contamination with pathogenic agents; II.5. have been packed in final packaging which bear labels indicating "RAW PETFOOD — NOT FOR HUMAN CONSUMPTION" "ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS — NOT FOR HUMAN CONSUMPTION" and then in leak-proof ar officially sealed boxes/containers or in new packaging preventing any leakage and officially sealed boxes/containers which be labels indicating "RAW PETFOOD — NOT FOR HUMAN CONSUMPTION" or "ANIMAL BY-PRODUCTS FOR FEED FOR FU ANIMALS — NOT FOR HUMAN CONSUMPTION", the name and the address of the establishment of destination; II.6. in the case of raw petfood: (a) have been prepared and stored in a plant approved and supervised by the competent authority in accordance with Article and where appropriate Article 11 of Regulation (EC) No 1774/2002; and (b) were examined by random sampling of at least five samples from each batch taken during storage (before dispatch) are complies with the following standards (6): Salmonella: absence in 25 g: n = 5, c = 0, m = 0, M = 0;			communicable to huma							
"ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS — NOT FOR HUMAN CONSUMPTION" and then in leak-proof ar officially sealed boxes/containers or in new packaging preventing any leakage and officially sealed boxes/containers which be labels indicating "RAW PETFOOD — NOT FOR HUMAN CONSUMPTION" or "ANIMAL BY-PRODUCTS FOR FEED FOR FU ANIMALS — NOT FOR HUMAN CONSUMPTION", the name and the address of the establishment of destination; II.6. in the case of raw petfood: (a) have been prepared and stored in a plant approved and supervised by the competent authority in accordance with Article and where appropriate Article 11 of Regulation (EC) No 1774/2002; and (b) were examined by random sampling of at least five samples from each batch taken during storage (before dispatch) are complies with the following standards (8): Salmonella: absence in 25 g: n = 5, c = 0, m = 0, M = 0;		II.4.								
(a) have been prepared and stored in a plant approved and supervised by the competent authority in accordance with Article and where appropriate Article 11 of Regulation (EC) No 1774/2002; and (b) were examined by random sampling of at least five samples from each batch taken during storage (before dispatch) are complies with the following standards (6): Salmonella: absence in 25 g: n = 5, c = 0, m = 0, M = 0;	"ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS — NOT FOR HUMAN CONSUMPTION" and then in lea officially sealed boxes/containers or in new packaging preventing any leakage and officially sealed boxes/containers labels indicating "RAW PETFOOD — NOT FOR HUMAN CONSUMPTION" or "ANIMAL BY-PRODUCTS FOR FEEL					HUMAN CONSUMPTION" and then in leak-proof and tage and officially sealed boxes/containers which bear 4" or "ANIMAL BY-PRODUCTS FOR FEED FOR FUR				
and where appropriate Article 11 of Regulation (EC) No 1774/2002; and (b) were examined by random sampling of at least five samples from each batch taken during storage (before dispatch) are complies with the following standards (6): Salmonella: absence in 25 g: n = 5, c = 0, m = 0, M = 0;		II.6.	in the case of raw petfood:							
complies with the following standards (6): Salmonella: absence in 25 g: n = 5, c = 0, m = 0, M = 0;										
-						ach batch taken during storage (before dispatch) and				
Enterobacteriacae: n = 5, c = 2, m = 10, M = 300 in 1 gram.			Salmonella:		absence in 25 g: n = 5, c = 0, m = 0	, M = 0;				
			Enterobacteriacae:		n = 5, c = 2, m = 10, M = 300 in 1 g	ram.				

Notes		
Part I:		
_	Box reference I.6: Person responsible for the consignment in EU: this commodity; it may be filled in if the certificate is for import commodity.	box is to be filled in only if it is a certificate for transit
_	Box reference I.12: Place of destination: this box is to be filled in only if it is can only be stored in free zones, free warehouses and custom warehous	
	Box reference I.15: Registration number (railway wagons or container and is to be provided in the event of unloading and reloading.	lorries), flight number (aircraft) or name (ship); information
_	Box reference I.19: use the appropriate HS code: 05.11.91; 05.11.99 or 2	23.09.90.
	Box reference I.23: for bulk containers, the container number and the sea	al number (if applicable) should be given.
	Box reference I.26 and I.27: fill in according to whether it is a transit or an	import certificate.
****	Box reference I.28: Nature of commodity: select raw petfood or animal by	<i>r-</i> product.
Part II:		
(*)	Delete as appropriate.	
(¹)	OJ L 273, 10.10.2002, p.1.	
(²)	Council Decision 79/542/EEC of 21 December 1976 drawing up a list of animal and public health and veterinary certification conditions, for import fresh meat.	
(3)	OJ L 295, 25.10.2006, p. 1.	
(4)	Commission Decision 2000/585/EC of 7 September 2000 laying down certifications for import of wild and farmed game meat and rabbit meat f 97/217/EC, 97/218/EC, 97/219/EC and 97/220/EC. OJ L 251, 6.10.2000,	rom third countries and repealing Commission Decisions
(5)	Council Directive 93/119/EC of 22 December 1993 on the protection of 31.12.1993, p. 21.	of animals at the time of slaughter or killing. OJ L 340,
(⁶)	Where:	
	n = number of samples to be tested;	
	m = threshold value for the number of bacteria; the result is considered not exceed m;	satisfactory if the number of bacteria in all samples does
	M = maximum value for the number of bacteria; the result is considered samples is M or more; and	d unsatisfactory if the number of bacteria in one or more
	c = number of samples the bacterial count of which may be between m bacterial count of the other samples is m or less.	and M, the sample still being considered acceptable if the
	The signature and the stamp must be in a different colour to that of the p	rinting.
_	Note for the person responsible for the consignment in EU: This certificate the consignment until it reaches the border inspection post.	ate is only for veterinary purposes and has to accompany
Official ve	eterinarian	
	Name (in capitals):	Qualification and title:
	Date:	Signature:
	Stamp:	

CHAPTER 3(E)

Health certificate

For flavouring innards for use in the manufacture of petfood, intended for dispatch to or for transit through (²) the European Community

COUN	ITRY	Veterinary certificate to EU
	I.1. Consignor	I.2. Certificate reference number
	Name	number
	Address	I.3. Central Competent Authority
ment	Tel. No	I.4. Local Competent Authority
sign	I.5. Consignee	I.6. Person responsible for the consignment in EU
Co	Name Address	Name Address
hed	Address	Address
patc	Postal code	Postal code
dis	Tel. No	Tel. No
Part I: Details of dispatched consignment	I.7. Country of ISO code I.8. Region of origin Cod origin	e I.9. Country of ISO code I.10. Region of Code destination
e De	I.11. Place of origin	I.12. Place of destination
art		Custom warehouse
"	Name Approval number	Name Approval number
	Address	Address
		Postal code
	I.13. Place of loading	I.14. Date of departure
	I.15. Means of transport Aeroplane Ship Railway	I.16. Entry BIP in EU
	Aeroplane Ship Railway Road vehicle Other wagon	J
	Identification:	L17.
	Documentary references:	
	I.18. Description of commodity	I.19. Commodity code (HS code)
		I.20. Quantity
	I.21. Temperature of product	I.22. Number of packages
	Ambient Chilled Chilled	Frozen 🔲
	I.23. Identification of container/Seal number	I.24. Type of packaging
	I.25. Commodities certified for:	
	Animal feedingstuff	Technical use Other
	I.26. For transit to third country vis-à-vis EU	I.27. For import or admission into EU
	3rd country ISO code	
	I.28. Identification of the commodities	
	Ap	proval number of establishments
	Species Nature of commodity (Scientific name)	Manufacturing plant Net weight Batch number

COUNTRY

Flavouring innards for use in the manufacture of petfood

Article 18 and where appropriate Article 11 of Regulation (EC) No 1774/2002; (i) either (ii) either (iii) either (iiii) either (iii) either (iiii) either (ii			II.a. Certificate reference number II.b.				
I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (1) and in partic Article 8 and Annex VIII (Tangter XIV thereof and certify that the flavouring innards products described above: III.1. consist of animal by-products that satisfy the animal health requirement below: III.2. have been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance v. Article 18 and where appropriate Article 11 of Regulation (EC) No 1774/2002; III.3. have been prepared including the following animal by-products which are exclusively: (?) either [— parts of slaughtered animals, which were fit for human consumption in accordance with Community legislation, but are interfect of humans or animals and derive from carcases that were lift for human consumption in accordance with Community legislation.] (?) and/or [— parts of slaughtered animals, which were rejected as unfit for human consumption but are not affected by any sign diseases communicable to humans or animals and derive from carcases that were lift for human consumption in accordance with Community legislation.] (?) and/or [— hides and skins, hooves and horns, plp bristles and feathers originating from animals that were slaughtered in a slaughterion underwent ante-morten inspection and were fit, as a result of such inspection, for slaughter in a slaughterion and underwent ante-morten inspection and were fit, as a result of such inspection, for slaughter in a scordance with Community legislation.] (?) and/or [— animal by-products derived from the production of products intended for human consumption, including degreased bones or greaves.] (?) and/or [— fromer foodstuffs of animal origin, or former foodstuffs containing products of numan lorigin, other than catering waste, with a not or animals.] (?) and/or [— fish or other sea animals, except sea mammals, caught in the open sea for the purposes of fishmeal product to human or animals.] (?) and/or [— fish or other							
I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (1) and in partic Article 8 and Annex VIII (Tangter XIV thereof and certify that the flavouring innards products described above: III.1. consist of animal by-products that satisfy the animal health requirement below: III.2. have been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance v. Article 18 and where appropriate Article 11 of Regulation (EC) No 1774/2002; III.3. have been prepared including the following animal by-products which are exclusively: (?) either [— parts of slaughtered animals, which were fit for human consumption in accordance with Community legislation, but are interfect of humans or animals and derive from carcases that were lift for human consumption in accordance with Community legislation.] (?) and/or [— parts of slaughtered animals, which were rejected as unfit for human consumption but are not affected by any sign diseases communicable to humans or animals and derive from carcases that were lift for human consumption in accordance with Community legislation.] (?) and/or [— hides and skins, hooves and horns, plp bristles and feathers originating from animals that were slaughtered in a slaughterion underwent ante-morten inspection and were fit, as a result of such inspection, for slaughter in a slaughterion and underwent ante-morten inspection and were fit, as a result of such inspection, for slaughter in a scordance with Community legislation.] (?) and/or [— animal by-products derived from the production of products intended for human consumption, including degreased bones or greaves.] (?) and/or [— fromer foodstuffs of animal origin, or former foodstuffs containing products of numan lorigin, other than catering waste, with a not or animals.] (?) and/or [— fish or other sea animals, except sea mammals, caught in the open sea for the purposes of fishmeal product to human or animals.] (?) and/or [— fish or other		_{II} .	Health attestation				
12. have been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance v. Article 18 and where appropriate Article 11 of Regulation (EC) No 1774/2002. 13. have been prepared including the following animal by-products which are exclusively: (2) and/or		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (1) ar					
Article 18 and where appropriate Article 11 of Regulation (EC) No 1774/2002; have been prepared including the following animal by-products which are exclusively; (*) either							
(?) and/or [ion	II.2.	have been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance with Article 18 and where appropriate Article 11 of Regulation (EC) No 1774/2002;				
(?) and/or (] and/or (ficat	11.3.	have been prepared including the following animal by-products which are exclusively:				
(?) and/or (] and/or (Certif	(²) either	[— parts of slaughtered animals, which were fit for human consumption in accordance with Community legislation, but are not intended for human consumption for commercial reasons,]				
underwent anter-mortem inspection and were fit, as a result of such inspection, for slaughter in accordance with Commu legislation.] (2) and/or (3) and/or (4) and/or (5) and/or (6) and/or (7) and/or (8) and/or (9) and/or (10) and	Part II:	(²) and/or	diseases communicable to humans or animals and derive from carcases that were fit for human consumption in accordance				
inspection and were fit, as a result of such inspection, for slaughter in accordance with Community legislation.] (?) and/or		(²) and/or	[— hides and skins, hooves and horns, pig bristles and feathers originating from animals that were slaughtered in a slaughterhouse, underwent ante-mortem inspection and were fit, as a result of such inspection, for slaughter in accordance with Community legislation,]				
greaves,] (?) and/or [(²) and/or	[— blood obtained from animals other than ruminants that were slaughtered in a slaughterhouse, underwent ante-mortem inspection and were fit, as a result of such inspection, for slaughter in accordance with Community legislation,]				
are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or package defects or other defects which do not present any risk to humans or animals.] (2) and/or [— raw milk originating from animals that do not show clinical signs of any disease communicable through that product to hum or animals.] (3) and/or [— fish or other sea animals, except sea mammals, caught in the open sea for the purposes of fishmeal production.] (3) and/or [— fish products from fish from plants manufacturing fish products for human consumption.] (3) and/or [— fresh by-products and cracked egg by-products for human consumption.] (3) and/or [— material from animals which have been treated with certain substances which are prohibited pursuant to Directive 96/22// the import of the material being permitted in accordance with Article 28 of Regulation (EC) No 1774/2002; in order to pathogenic agents; II.4. have been subjected to processing in accordance with Annex VIII, Chapter XIV of Regulation (EC) No 1774/2002, in order to pathogenic agents; II.5. have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply withe following standards (3): Salmonella: absence in 25 g: n = 5, c = 0, m = 0, M = 0; Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram; II.6. the end product was: (3) either [packed in new or sterilised bags:] (4) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfected approved by the competent authority before use:] and which bear labels indicating "NOT FOR HUMAN CONSUMPTION"; II.7. the end product was stored in enclosed storage; III.8. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment.		(2) and/or	[— animal by-products derived from the production of products intended for human consumption, including degreased bones and greaves,]				
or animals,] (²) and/or [— fish or other sea animals, except sea mammals, caught in the open sea for the purposes of fishmeal production,] (²) and/or [— fresh by-products from fish from plants manufacturing fish products for human consumption,] (²) and/or [— shells, hatchery by-products and cracked egg by-products originating from animals which did not show clinical signs of disease communicable through that product to humans or animals.] (²) and/or [— material from animals which have been treated with certain substances which are prohibited pursuant to Directive 96/22/ the import of the material being permitted in accordance with Article 28 of Regulation (EC) No 1774/2002;] II.4. have been subjected to processing in accordance with Annex VIII, Chapter XIV of Regulation (EC) No 1774/2002, in order to pathogenic agents; II.5. have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply the following standards (²): Salmonella: absence in 25 g: n = 5, c = 0, m = 0, M = 0; Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram; II.6. the end product was: (²) either [packed in new or sterilised bags;] (²) or [product in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfect approved by the competent authority before use:] and which bear labels indicating "NOT FOR HUMAN CONSUMPTION"; II.7. the end product was stored in enclosed storage; II.8. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment.		(²) and/or	[— former foodstuffs of animal origin, or former foodstuffs containing products of animal origin, other than catering waste, which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects which do not present any risk to humans or animals,]				
 (²) and/or [— fresh by-products from fish from plants manufacturing fish products for human consumption.] (²) and/or [— shells, hatchery by-products and cracked egg by-products originating from animals which did not show clinical signs of disease communicable through that product to humans or animals.] (²) and/or [— material from animals which have been treated with certain substances which are prohibited pursuant to Directive 96/22/f the import of the material being permitted in accordance with Article 28 of Regulation (EC) No 1774/2002.] II.4. have been subjected to processing in accordance with Annex VIII, Chapter XIV of Regulation (EC) No 1774/2002, in order to pathogenic agents; II.5. have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply with following standards (²): Salmonella: absence in 25 g: n = 5, c = 0, m = 0, M = 0; Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram; II.6. the end product was: (²) either [packed in new or sterilised bags:] (²) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfect approved by the competent authority before use:] and which bear labels indicating "NOT FOR HUMAN CONSUMPTION"; II.7. the end product was stored in enclosed storage; II.8. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment. Notes Part I: Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for transport. 		(2) and/or	[— raw milk originating from animals that do not show clinical signs of any disease communicable through that product to humans or animals.]				
(²) and/or [— shells, hatchery by-products and cracked egg by-products originating from animals which did not show clinical signs of disease communicable through that product to humans or animals.] (²) and/or [— material from animals which have been treated with certain substances which are prohibited pursuant to Directive 96/22// the import of the material being permitted in accordance with Article 28 of Regulation (EC) No 1774/2002;] II.4. have been subjected to processing in accordance with Annex VIII, Chapter XIV of Regulation (EC) No 1774/2002, in order to pathogenic agents; II.5. have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply the following standards (³): Salmonella: absence in 25 g: n = 5, c = 0, m = 0, M = 0; Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram; II.6. the end product was: (²) either [packed in new or sterilised bags;] (²) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfect approved by the competent authority before use;] and which bear labels indicating "NOT FOR HUMAN CONSUMPTION"; II.7. the end product was stored in enclosed storage; II.8. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment. Notes Part I: Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for tra		(2) and/or	[— fish or other sea animals, except sea mammals, caught in the open sea for the purposes of fishmeal production,]				
disease communicable through that product to humans or animals.] (2) and/or [(2) and/or	[— fresh by-products from fish from plants manufacturing fish products for human consumption,]				
the import of the material being permitted in accordance with Article 28 of Regulation (EC) No 1774/2002;] II.4. have been subjected to processing in accordance with Annex VIII, Chapter XIV of Regulation (EC) No 1774/2002, in order to pathogenic agents; II.5. have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply of the following standards (3): Salmonella: absence in 25 g: n = 5, c = 0, m = 0, M = 0; Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram; II.6. the end product was: (2) either [packed in new or sterilised bags;] (3) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfect approved by the competent authority before use;] and which bear labels indicating "NOT FOR HUMAN CONSUMPTION"; II.7. the end product was stored in enclosed storage; II.8. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment. Notes Part I: — Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for tra		(²) and/or	[— shells, hatchery by-products and cracked egg by-products originating from animals which did not show clinical signs of any disease communicable through that product to humans or animals,]				
pathogenic agents; II.5. have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply the following standards (²): Salmonella: absence in 25 g: n = 5, c = 0, m = 0, M = 0; Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram; II.6. the end product was: (²) either [packed in new or sterilised bags;] (²) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfect approved by the competent authority before use:] and which bear labels indicating "NOT FOR HUMAN CONSUMPTION"; II.7. the end product was stored in enclosed storage; II.8. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment. Notes Part I: Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for tra		(2) and/or	[— material from animals which have been treated with certain substances which are prohibited pursuant to Directive 96/22/EC, the import of the material being permitted in accordance with Article 28 of Regulation (EC) No 1774/2002;]				
the following standards (³): Salmonella: absence in 25 g; n = 5, c = 0, m = 0, M = 0; Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram; II.6. the end product was: (²) either [packed in new or sterilised bags;] (²) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfect approved by the competent authority before use;] and which bear labels indicating "NOT FOR HUMAN CONSUMPTION"; II.7. the end product was stored in enclosed storage; II.8. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment. Notes Part I: — Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for tra		II.4.	have been subjected to processing in accordance with Annex VIII, Chapter XIV of Regulation (EC) No 1774/2002, in order to kill pathogenic agents;				
Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram; II.6. the end product was: (²) either [packed in new or sterilised bags;] (²) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfect approved by the competent authority before use;] and which bear labels indicating "NOT FOR HUMAN CONSUMPTION"; II.7. the end product was stored in enclosed storage; II.8. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment. Notes Part I: — Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for tra		II.5.	have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply with the following standards (3):				
II.6. the end product was: (²) either [packed in new or sterilised bags;] (²) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfect approved by the competent authority before use;] and which bear labels indicating "NOT FOR HUMAN CONSUMPTION"; II.7. the end product was stored in enclosed storage; II.8. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment. Notes Part I: Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for tra			Salmonella: absence in 25 g: $n = 5$, $c = 0$, $m = 0$, $M = 0$;				
(2) either [packed in new or sterilised bags;] (2) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfect approved by the competent authority before use;] and which bear labels indicating "NOT FOR HUMAN CONSUMPTION"; II.7. the end product was stored in enclosed storage; II.8. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment. Notes Part I: Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for tra			Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram;				
(²) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfect approved by the competent authority before use;] and which bear labels indicating "NOT FOR HUMAN CONSUMPTION"; II.7. the end product was stored in enclosed storage; II.8. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment. Notes Part I: Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for tra		II.6.	the end product was:				
approved by the competent authority before use;] and which bear labels indicating "NOT FOR HUMAN CONSUMPTION"; II.7. the end product was stored in enclosed storage; II.8. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment. **Notes** Part I: Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for tra		(2) either	[packed in new or sterilised bags;]				
II.7. the end product was stored in enclosed storage; II.8. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment. Notes Part I: Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for tra		(²) or	[transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectant approved by the competent authority before use;]				
II.8. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment. Notes Part I: Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for tra		and which bear labels indicating "NOT FOR HUMAN CONSUMPTION";					
Notes Part I: Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for tra		II.7. the end product was stored in enclosed storage;					
Part I: Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for tra		II.8.	the product has undergone all precautions to avoid contamination with pathogenic agents after treatment.				
Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for tra		Notes	Notes				
		Part I:					
commodity, it may be filled in it the certificate is for import commodity.			Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity.				
 Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in tra can only be stored in free zones, free warehouses and custom warehouses. 		_	Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses.				
			Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information				
Box reference I.19: use the appropriate HS code: 05.04 or 05.11.91.			Box reference I.19: use the appropriate HS code: 05.04 or 05.11.91.				

▼B

	Box reference I.23: for bulk containers, the container	number and the seal number (if applicable) should be given.
_	Box reference I.26 and I.27: fill in according to whether	er it is a transit or an import certificate.
_	Box reference I.28: define the innard product.	
Part II:		
(1)	OJ L 273, 10.10.2002, p. 1.	
(2)	Delete as appropriate.	
(3)	Where:	
	n = number of samples to be tested;	
	m = threshold value for the number of bacteria; the not exceed m;	result is considered satisfactory if the number of bacteria in all samples does
	M = maximum value for the number of bacteria; the samples is M or more; and	result is considered unsatisfactory if the number of bacteria in one or more
	c = number of samples the bacterial count of which bacterial count of the other samples is m or less	may be between m and M, the sample still being considered acceptable if the
_	The signature and the stamp must be in a different co	lour to that of the printing.
-	Note for the person responsible for the consignment the consignment until it reaches the border inspection	in EU: This certificate is only for veterinary purposes and has to accompany post.
Official	veterinarian	
	Name (in capitals):	Qualification and title:
	Date:	Signature:
	Stamp:	
1		

CHAPTER 3(F)

Health certificate

For animal by-products (*) for the manufacture of petfood, intended for dispatch to or for transit through (3) the European Community

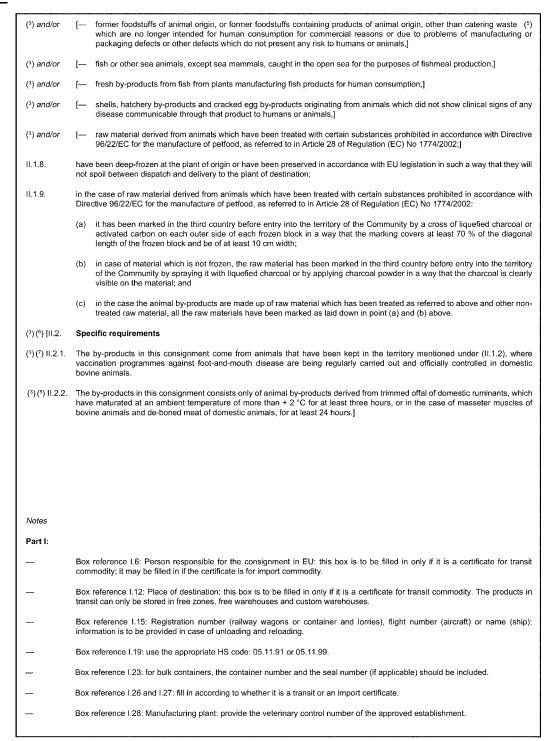
COUN	TRY				Veterinary certificate to EU
	l.1.	Consignor Name	1.2.	Certificate reference number	I.2.a.
		Address	1.3.	Central Competent Authori	ty
nent		Tel. No	1.4.	Local Competent Authority	,
ign	1.5.	Consignee	1.6.	Person responsible for the	consignment in EU
ons		Name		Name	
g g		Address		Address	
che					
spat		Postal code		Postal code	
đ.		Tel. No	ļ	Tel. No	
Part I: Details of dispatched consignment	1.7.	Country of ISO code I.8. Region of origin Code origin	1.9.	Country of ISO code destination	1.10.
o l	l.11.	Place of origin	I.12.	Place of destination	
Ĭ,				Custom warehou	use 🗍
ے ا		Name Approval number		Name	Approval number
1		Address		Address	
				Postal code	
	I.13.	Place of loading	1.14.	Date of departure	
	I.15.	Means of transport	I.16.	Entry BIP in EU	
		Aeroplane Ship Railway Ship Wagon			
		Trodd Vollido			· · · · · · · · · · · · · · · · · · ·
		Identification: Documentary references:	1.17.		
	140			140.0	Bt. and (IIO and a)
	1.18.	Description of commodity		1.19. Commod	lity code (HS code)
					I.20. Quantity
	1.21.	Temperature of product			I.22. Number of packages
		Ambient Chilled Chilled		Frozen 🔲	
	123	Identification of container/Seal number			I.24. Type of packaging
					1.24. Type of packaging
	1.25.	Commodities certified for: Animal feedingstuff Further process Further process		Technical use	Other
	1.26.	For transit to third country vis-à-vis EU	1.27.	For import or admission into	EU
		3rd country ISO code			
	1.28.	Identification of the commodities		· · · · · · · · · · · · · · · · · · ·	
		Approval number of establis	shment	S	
		Species Nature of commodity Manufacturing plan (Scientific name)	nt	Number of packages	Net weight Batch number

COUNTRY

Animal by-products for the manufacture of petfood

		II.a. Certificate reference number II.b.					
	II.1.	Health attestation					
		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (¹) and certify that the animal by-products described above:					
_	II.1.1.	consist of animal by-products that satisfy the animal health requirements below;					
tion	II.1.2.	have been obtained in the territory of:					
tifica	(³) either	[(a) that have remained in this territory since birth or for at least the last three months before slaughter;]					
Cert	(3) or	[(b) killed in the wild in this territory (4);]					
Part II: Certification	II.1.3.	have been obtained from animals:					
Ра	(3) either [(a) coming from holdings:						
		(i) where, for the following diseases for which the animals are susceptible, there has been neither case/outbreak of rinderpest, swine vesicular disease, Newcastle disease or highly pathogenic avian influenza during the prior 30 days, nor of classical or African swine fever during the prior 40 days; nor in the holdings situated in their vicinity within 10 km, during the prior 30 days; and					
		(ii) where there has been neither case/outbreak of foot and mouth disease during the prior 60 days, nor in the holdings situated in their vicinity within 25 km, during the prior 30 days; and					
		(b) which:					
		(i) were not killed to eradicate any epizootic disease;					
		 (ii) have remained in their holdings of origin for at least forty days before departure and which have been transported directly to the slaughterhouse without contact with other animals which did not comply with the same health conditions; 					
		(iii) at the slaughterhouse, have passed the ante-mortem health inspection during the 24 hours before the slaughter and have shown no evidence of the diseases referred to above for which the animals are susceptible; and					
		(iv) have been treated in the slaughterhouse before and at the time of slaughter or killing in accordance with the relevant provisions of Council Directive 93/119/EC on animal welfare;					
	(3) or	[(a) captured and killed in the wild in an area:					
		(i) in which within 25 km there has been no case/outbreak of any of the following diseases for which the animals are susceptible: foot and mouth disease, rinderpest, Newcastle disease or highly pathogenic avian influenza during the prior 30 days, nor of classical or African swine fever during the prior 40 days; and					
		(ii) that is situated at a distance that exceeds 20 km from the borders separating another territory of a country or part thereof, which is not authorised at these dates for exporting this material to the European Community; and					
		(b) which after killing were transported within 12 hours for chilling either to a collection centre and immediately afterwards to a game establishment, or directly to a game establishment;]					
	II.1.4.	have been obtained in an establishment around which, within a radius of 10 km, there has been no case/outbreak of diseases referred to in point II.1.3 for which the animals are susceptible during the prior 30 days or, in the event of a case of disease, the preparation of raw material for exportation to the European Community has been authorised only after removal of all meat, and the total cleaning and disinfection of the establishment under the control of an official veterinarian;					
	II.1.5.	have been obtained and prepared without contact with other material not complying with the conditions required above, and it has been handled so as to avoid contamination with pathogenic agents;					
	II.1.6.	have been packed in new packaging preventing any leakage and in officially sealed containers bearing the label indicating "RAW MATERIAL ONLY FOR THE MANUFACTURE OF PETFOOD" and the name and address of the EU establishment of destination;					
	II.1.7.	consist only of the following animal by-products:					
	(³) either	[— parts of slaughtered animals, which are fit for human consumption in accordance with Community legislation, but are not intended for human consumption for commercial reasons,]					
	(³) and/or	 parts of slaughtered animals, which are rejected as unfit for human consumption but are not affected by any signs of diseases communicable to humans or animals and derive from carcasses that are fit for human consumption in accordance with Community legislation,] 					
	(³) and/or) and/or [— animal by-products derived from the production of products intended for human consumption, including degreased bones and greaves.]					

▼B



▼B

(*)		
()	Excluding raw blood, raw milk, hides and the import of these products).	skins, hooves and horn, pig bristles and feathers (see relevant specific certificates for
(1)	OJ L 273, 10.10.2002, p. 1.	
(2)	The name and ISO code number of the ex	country as laid down in:
	 part 1 of Annex II of Council Decision 	79/542/EEC,
	 the Annex to Commission Decision 9 	4/984/EC, and
	 the Annex to Commission Decision 2 	000/585/EC.
	In addition the ISO code of regionalisation included.	on in this Annex (where applicable for the susceptible species concerned) should be
(3)	Delete as appropriate.	
(4)	Only for countries from where game meat i into the European Community.	ntended for human consumption of the same animal species is authorised for importation
(5)	Catering waste means all waste food, ir including central kitchens and household	icluding used cooking oils, originating in restaurants, catering facilities and kitchens, kitchens.
(6)	or South African country or part thereof fi consumption is permitted for exportation to which must be exclusively offal from which tissue, fat and mucus have been comple	when the material of domestic ruminants originated in the territory of a South American rom where only maturated and de-boned fresh meat of domestic ruminants for human to the European Community. In the case of offal only trimmed offal of domestic ruminants in the bones, cartilage, trachea and main bronchi, lymphatic glands, adhering connective tely removed is permitted. The whole masseter muscles of bovine animals, incised in ster VIII of Annex I to Council Directive 64/433/EEC, are also permitted.
(7)	Only for certain South American countries	i.
(8)	Only for certain South American and Sout	h African countries.
_	The signature and the stamp must be in a	different colour to that of the printing.
	Note for the person responsible for the co	onsignment in EU: this certificate is only for veterinary purposes and has to accompany
_	the consignment until it reaches the borde	
— Official ve		
Official ve	the consignment until it reaches the borde	
Official ve	the consignment until it reaches the borde	er inspection post.

CHAPTER 4(A)

Health certificate

For the import of serum from equidae to be used for technical purposes including pharmaceuticals, in vitro diagnosis and laboratory reagents, intended for dispatch to or for transit through (²) the European Community

COUN	TRY	Veterinary certificate to EU
	I.1. Consignor Name	I.2. Certificate reference number I.2.a.
	Address	I.3. Central Competent Authority
ment	Tel. No	I.4. Local Competent Authority
ısign	I.5. Consignee	I.6. Person responsible for the consignment in EU
d cor	Name Address	Name Address
tche		
lispa	Postal code Tel. No	Postal code Tel. No
Part I: Details of dispatched consignment	I.7. Country of ISO code I.8. Region of origin Code origin	I.9. Country of ISO code I.10. destination
1: De	I.11. Place of origin	I.12. Place of destination
Part	Name Approval number	Custom warehouse
	Name Approval number Address	Name Approval number Address
		Doublands
	I.13. Place of loading	Postal code I.14. Date of departure
		·
	I.15. Means of transport Aeroplane Ship Railway wagon	I.16. Entry BIP in EU
	Road vehicle Other	
	Identification: Documentary references:	1.17.
	I.18. Description of commodity	I.19. Commodity code (HS code) 30.02
		I.20. Quantity
	I.21. Temperature of product	I.22. Number of packages
		· · · · · · · · · · · · · · · · · · ·
	I.23. Identification of container/Seal number	I.24. Type of packaging
	I.25. Commodities certified for: Technical use]
	I.26. For transit to third country vis-à-vis EU	I.27. For import or admission into EU
	3rd country ISO code	
	I.28. Identification of the commodities	
	Species (Scientific name)	Approval number of establishments Manufacturing plant

COUNTRY

Serum from equidae for technical purposes including pharmaceuticals, in vitro diagnosis and laboratory reagents

		II.a. Certificate reference number II.b.				
	II.	Health attestation I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (1) and certify that the serum of equidae described above:				
	II.1.	consist of serum from equidae that satisfy the health requirements below;				
ication	II.2.	consist exclusively of serum of equidae not intended for human nor animal consumption;				
Part II: Certification	II.3.	comes from a country where the following diseases are compulsorily notifiable: African horse sickness, dourine, glanders, equine encephalomyelitis (all types including VEE), equine infectious anemia, vesicular stomatitis, rabies, anthrax;				
Part	II.4.	was obtained, under the supervision of a veterinarian, from equidae which, at the time of collection, were free from clinical signs of infectious disease or were obtained from equidae that passed ante-mortem inspection at the time of slaughter;				
	II.5.	was obtained from equidae that have remained since birth in the territory or, in case of official regionalisation according to Community legislation, in parts of the territory of a third country in which:				
		(a) Venezuelan equine encephalomyelitis has not occurred during the last two years;				
		(b) dourine has not occurred during the last six months; and				
		(c) glanders has not occurred during the last six months;				
L	II.6.	was obtained from equidae that had never been present on a holding that had been subject to prohibition for animal health reasons or where:				
	(²) eíther	[(a) in the case of equine encephalomyelitis, the date on which all the equidae suffering from the disease were slaughtered was at least six months before the date of collection;				
	(b) in the case of infectious anaemia, all the infected animals had been slaughtered and the remaining animals showed a negative reaction to two Coggins tests carried out three months apart;					
	(c) in the case of vesicular stomatitis, the prohibition was lifted at least six months before the date of collection;					
	(d) in the case of rabies, the last recorded case was at least a month before the date of collection; and					
		(e) in the case of anthrax, the last recorded case was at least 15 days before the date of collection;]				
	(²) or	[all the animals of species susceptible to the disease located on the holding were slaughtered and the premises disinfected, at least 30 days before the date of collection (or, in the case of anthrax, at least 15 days before);]				
	II.7.	has undergone all precautions to avoid contamination with pathogenic agents during production, handling and packaging;				
	II.8.	was packed in sealed impermeable containers clearly labelled 'serum from equidae' and bearing the registration number of the establishment of collection.				
	Notes					
	Part I:					
	_	Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity.				
	_	Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses.				
	_	Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading.				
	_	Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.				
	_	Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.				
	_	Box reference I.28: Manufacturing plant: provide the veterinary control number of the registered establishment of collection.				

Part II:		
(¹)	OJ L 273, 10.10.2002, p. 1.	
(²)	Delete as appropriate.	
	The signature and the stamp must be in a	different colour to that of the printing.
_	Note for the person responsible for the cor consignment until it reaches the border ins	signment in EU: this certificate is only for veterinary purposes and has to accompany the spection post.
Official	veterinarian	
	Name (in capitals):	Qualification and title:
	Date:	Signature:
	Stamp:	

CHAPTER 4(B)

Health certificate

For blood products not intended for human consumption that could be used as feed material, intended for dispatch to or for transit through (²) the European Community

COUN	TRY			Veterinary certificate to E
	l.1.	Consignor Name	1.2.	Certificate reference 1.2.a. number
	Address		1.3.	Central Competent Authority
ment		Tel. No	1.4.	Local Competent Authority
igi	1.5.	Consignee	1.6.	Person responsible for the consignment in EU
io:		Name		Name
ped		Address		Address
atch		Postal code		Postal code
disp		Tel. No		Tel. No
Part I: Details of dispatched consignment	1.7.	Country of ISO code I.8. Region of origin Code origin	1.9.	Country of ISO code I.10. destination
e l	l.11.	Place of origin	1.12.	. Place of destination
art				Custom warehouse
		Name Approval number		Name Approval number
		Address	ŀ	Address
				Postal code
	1.13.	Place of loading	1.14.	. Date of departure
	I.15.	Means of transport	1.16.	. Entry BIP in EU
		Aeroplane Ship Railway wagon Road vehicle Other		
		Identification:	1.17.	
	Documentary references:			
	l.18.	Description of commodity		I.19. Commodity code (HS code)
				I.20. Quantity
	1.21.	Temperature of product		I.22. Number of packages
	Ambient Chille		ı 🔲	Frozen
	1.23.	Identification of container/Seal number		I.24. Type of packaging
	1.25.	Commodities certified for:	1	
	1.00	Animal feedingstuff	107	Facility and a solution in the FU
	1.26.	For transit to third country vis-à-vis EU	1.27.	. For import or admission into EU
		3rd country ISO code		
	1.28.	Identification of the commodities		
		Species Nature of commodity (Scientific name)		Approval number of establishments Manufacturing plant Batch number

COUNTRY

Blood products that could be used as feed material

		II.a. Certificate reference number II.b.				
	II.	Health attestation				
	и.	I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (¹) and certify the blood products described above:				
_	II.1.	consist of blood products that satisfy the health requirements below;				
icatio	II.2.	consist exclusively of blood products not intended for human consumption;				
Part II: Certification	II.3.	have been prepared and stored in a plant, approved, validated and supervised by the competent authority in accordance with Article 17 and where appropriate Article 11 of Regulation (EC) No 1774/2002;				
<u> </u>	II.4.	have been prepared (derived) exclusively with the following animal by-products:				
Pa	(²) either	[blood of slaughtered animals, which is fit for human consumption in accordance with Community legislation, but is not intended for human consumption for commercial reasons;]				
	(²) and/or	[blood of slaughtered animals, which is rejected as unfit for human consumption but is not affected by any signs of diseases communicable to humans or animals, derived from carcasses that are fit for human consumption in accordance with Community legislation;]				
	II.5.	have been submitted				
	(²) either	[to processing in accordance with processing method(3) as set out in Annex V, Chapter III of Regulation (EC) No 1774/2002;]				
	(²) or	[to a method and parameters which ensure that the product complies with the microbiological standards set in Chapter I, paragraph 10 of Annex VII to Regulation (EC) No 1774/2002;]				
		in order to kill pathogenic agents;				
	II.6.	have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply with the following standards (4):				
		Salmonella: absence in 25 g: $n = 5$, $c = 0$, $m = 0$, $M = 0$,				
		Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram;				
	II.7.	the end product was:				
	(²) either [packed in new or sterilised bags;]					
	(²) or	[transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectant approved by the competent authority before use,]				
		and which bear labels indicating "NOT FOR HUMAN CONSUMPTION";				
	II.8.	the end product was stored in enclosed storage;				
	11.9.	the product has undergone all precautions to avoid contamination with pathogenic agents after treatment.				
	Notes	otes				
	Part I:	Part I:				
	_	 Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity. 				
	 Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products it transit can only be stored in free zones, free warehouses and custom warehouses. 					
	 Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship information is to be provided in case of unloading and reloading. 					
	_	Box reference I.19: use the appropriate HS code: 05.11.91 or 05.11.99.				
	_	Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.				
	_	Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.				

▼B

Part II:						
(1)	OJ L 273, 10.10.2002, p. 1.					
(2)	Delete as appropriate.	Delete as appropriate.				
(3)	Insert method 1 to 5 or 7 as applicable.					
(4)	Where:					
	n = number of samples to be tested;					
	m = threshold value for the number of bacterianot exceed m;	a; the result is considered satisfactory if the number of bacteria in all samples does				
	M = maximum value for the number of bacter samples is M or more; and	ria; the result is considered unsatisfactory if the number of bacteria in one or more				
	c = number of samples the bacterial count of which may be between m and M, the sample still being considered acceptable if bacterial count of the other samples is m or less.					
_	The signature and the stamp must be in a diffe	erent colour to that of the printing.				
-	Note for the person responsible for the consignment in EU: this certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.					
Official v	veterinarian					
	Name (in capitals):	Qualification and title:				
	Date:	Signature:				
	Stamp:					

CHAPTER 4(C)

Health certificate

For blood products, excluding serum of equidae and intermediate products as referred to in Article 1 of Commission Regulation (EC) No 2007/2006, to be used for technical purposes, intended for dispatch to or for transit through (²) the European Community

COUN	TRY				Veterinary certificate to EU
	l.1.	Consignor Name	1.2.	Certificate reference number	1.2.a.
		Address	1.3.	Central Competent Authori	y
nent		Tel. No	1.4.	Local Competent Authority	
sign	1.5.	Consignee	1.6.	Person responsible for the	consignment in EU
con		Name Address		Name Address	
hed		Audress		Address	
patc		Postal code		Postal code	
dis		Tel. No		Tel. No	
Part I: Details of dispatched consignment	1.7.	Country of ISO code I.8. Region of origin Code origin	1.9.	Country of ISO code destination	1.10.
ı. D	1.11.	Place of origin	l.12.	Place of destination	
Part				Custom warehou	_
-		Name Approval number		Name	Approval number
		Address		Address	
				Postal code	
	1.13.	Place of loading	1.14.	Date of departure	
	1.15.	Means of transport	I.16.	Entry BIP in EU	
		Aeroplane Ship Railway wagon Road vehicle Other			
		Identification:	1.17.		
		Documentary references:	1. (7.		
	l.18.	Description of commodity		I.19. Commod	ity code (HS code) 30.02
					I.20. Quantity
	121	Temperature of product			I.22. Number of packages
	1.21.	Ambient Chilled		Frozen	1.22. Humber of packages
	1.23.	Identification of container/Seal number			I.24. Type of packaging
	1.25.	Commodities certified for: Technical use			
	1.26.	For transit to third country vis-à-vis EU	1.27.	For import or admission into	EU
		3rd country ISO code		,	
	100	Manufication of the account of the			
	1.20.	Identification of the commodities			
				Approval number of establi	shments
		Species Nature of commodity (Scientific name)		Manufacturing plan	t Batch number

COUNTRY

Blood products for technical purposes, excluding serum of equidae and intermediate products as referred to in Article 1 of Commission Regulation (EC) No 2007/2006

			II.a. Certificate reference number	II.b.					
	11.	Health attestation							
		I, the undersigned of that the blood produc		l veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (¹) and certify sscribed above:					
	II.1.	consist of blood products that satisfy the health requirements below;							
tion	II.2.	consist exclusively of	mption;						
ifica	II.3.	have been prepared exclusively with the following animal by-products:							
Part II: Certification	(²) either		d of slaughtered animals, which is fit for human consumption in accordance with Community legislation, but is not need for human consumption for commercial reasons,]						
Part	(²) and/or	communicable to	 blood of slaughtered animals, which is rejected as unfit for human consumption but is not affected by any signs of diseases communicable to humans or animals, derived from carcasses that are fit for human consumption in accordance with Community legislation.] 						
	(²) and/or		om animals other than ruminants that are slaughtered in ere fit, as a result of such inspection, for slaughter for hi						
	(2) and/or	[blood and blood	products derived from the production of products intende	ed for human consumption,]					
	(²) and/or	[— blood and blood p	products originating from animals that did not show clinic is or animals;]	al signs of any disease communicable through that					
	(²) either [II.	4. in the case of blood p	roducts derived from ruminant animals they originate in	a third country or regions, where:					
		rinderpest which vac of ruminar	als and products come from a region where no case peste des petits ruminants, Rift Valley fever and blueto cination has not been carried out against those disease at animals are authorized pursuant to Community legited must have been collected:	ongue (3) has been recorded for 12 months and in es for at least 12 months and from which imports					
		(2) either [in slaught	erhouses approved in accordance with Community legis	approved in accordance with Community legislation;]					
		(2) or [from live a	[from live animals in facilities approved in accordance with Community legislation;]						
		Commissi	aughterhouses approved and supervised by the competent authority of the third country. In this case mission and Member States must be notified of the address and approval number of such slaughterhouse ertificate shall indicate this information;]]						
			cts have undergone one of the following treatments, guaranteeing the absence of pathogens of the rumir oot-and-mouth disease, vesicular stomatitis, rinderpest, peste des petits ruminants, Rift Valley fever e (3):						
		(2) either [heat treat	eat treatment at a temperature of 65 °C for at least three hours, followed by an effectiveness check;]						
		(2) or [irradiation	at 2,5 megarads or by gamma rays, followed by an effect	ectiveness check;]					
		(2) or [change in pH to pH 5 for two hours, followed by an effectiveness check;]							
		(2) or [heat treat	ment of at least 90°C throughout their substance, follow	ed by an effectiveness check;]]					
		pharmace		etongue animals are present, and the blood products are intended for technical purposes including in vitro diagnosis and laboratory reagents, to be processed in the approved plants [approval number] if (4):					
(2) or [II.4. in the case of blood products derived from animals excluding rumin		roducts derived from animals excluding ruminants they	originate in a third country or regions where:						
		(2) either [the animals and the products come from a region where no case of foot-and-mouth disease, swine vesicular disease, classical swine fever, African swine fever, Newcastle disease or highly pathogenic avian influenza has been recorded for 12 months in the susceptible species and in which vaccination has not been carried out against those diseases for at least 12 months. The health certificate shall follow the model according to the species of animal from which the blood products are derived:]							
		effectivene swine ves	cts have undergone a heat treatment at a temperature ess check, guaranteeing the absence of pathogens of cular disease, classical swine fever, African swine feven the susceptible species;]]	the following diseases: foot-and-mouth disease,					

11.5.	the end product was:				
(2) either	[packed in new or sterilised bags;]				
(2) or	[transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectan approved by the competent authority before use;]				
	and which bear labels indicating 'NOT FOR HUMAN OR ANIMAL CONSUMPTION';				
11.6.	the end product was stored in enclosed storage;				
II.7.	the product has undergone all precautions to avoid contamination with pathogenic agents after treatment.				
Notes					
Part I:					
_	Box reference I.6: Person responsible for the consignment in EU: this commodity; it may be filled in if the certificate is for import commodity.	s box is to be filled in only if it is a certificate for transit			
	Box reference I.12: Place of destination: this box is to be filled in only transit can only be stored in free zones, free warehouses and custom v				
AMERICA	Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading.				
_	Box reference I.23: for bulk containers, the container number and the s	seal number (if applicable) should be included.			
	Box reference I.26 and I.27: fill in according to whether it is a transit or	an import certificate.			
Part II:					
(1)	OJ L 273, 10.10.2002, p. 1.				
(2)	Delete as appropriate.				
(3)	In the case of countries in which bluetongue sero-positive ruminant anin animals have been tested sero-negative.	nals are present, blood products have been treated or the			
(4)	This must be the same Member State of first entry of the products into	the Community.			
	The signature and the stamp must be in a different colour to that of the	printing.			
_	Note for the person responsible for the consignment in EU: this certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.				
Official veter	erinarian				
	Name (in capitals):	Qualification and title:			
	Date:	Signature:			
	Stamp:				

CHAPTER 5(A)

Health certificate

For fresh or chilled hides and skins of ungulates, intended for dispatch to or for transit through (²) the European Community

COUN	TRY	Veterinary certificate to E
	I.1. Consignor Name	I.2. Certificate reference number I.2.a.
	Address	I.3. Central Competent Authority
nent	Tel. No	I.4. Local Competent Authority
ignn	I.5. Consignee	I.6. Person responsible for the consignment in EU
ons	Name	Name
D D	Address	Address
tche		5
sba	Postal code Tel. No	Postal code Tel. No
of di	I.7. Country of ISO code I.8. Region of origin Code	
Part I: Details of dispatched consignment	origin	destination destination
Det	I.11. Place of origin	I.12. Place of destination
벁	·	Custom warehouse
a l	Name Approval number	Name Approval number
	Address	Address
		Postal code
	I.13. Place of loading	I.14. Date of departure
	I.15. Means of transport	I.16. Entry BIP in EU
	Aeroplane Ship Railway wagon Road vehicle Other	
	Identification:	I.17. No(s) of CITES
	Documentary references:	1.17. NO(5) OF CITES
	I.18. Description of commodity	I.19. Commodity code (HS code)
		I.20. Quantity
	I.21. Temperature of product	I.22. Number of packages
	Ambient Chilled Chilled	Frozen 🗌
	I.23. Identification of container/Seal number	I.24. Type of packaging
	I.25. Commodities certified for:	L
	Animal feedingstuff	Technical use Other
	I.26. For transit to third country vis-à-vis EU	I.27. For import or admission into EU
	3rd country ISO code	
	I.28. Identification of the commodities	L
	Approval numb	per of establishments
	**	facturing plant Net weight
	(Scientific name)	5,

▼B

COUNTRY Fresh or chilled hides and skins of Ungulates Шh II.a. Certificate reference number II. I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (') and in particular Article 6 and Annex VIII Chapter VI thereof and certify that the hides and skins described above: II.1. have been obtained from animals that (2): Certification (a) were slaughtered and their carcasses are fit for human consumption in accordance with Community legislation; or were slaughtered in a slaughterhouse, after undergoing ante-mortem inspection, and were fit, as a result of such inspection, for slaughter for human consumption in accordance with Community legislation; Part II: 11.2. originate from a country or, in the case of regionalisation in accordance with Community legislation, from a part of a country from which imports of all categories of fresh meat of the corresponding species are authorised and which: (a) for at least 12 months before dispatch, has been free from the following diseases (*): [-- classical swine fever, and African swine fever,] [--- rinderpest,] and has been free for at least 12 months before dispatch from foot-and-mouth disease and where, for 12 months before dispatch, no vaccination has been carried out against foot-and-mouth disease (1); 11.3. have been obtained from: [animals that have remained in the territory of the country of origin for at least three months before being slaughtered or since birth in the case of animals less that three months old:] [in the case of hides and skins from bi-ungulates, animals that come from holdings in which there has been no outbreak of foot-and-mouth disease in the previous 30 days, and around which within a radius of 10 km there has been no case of foot-and-mouth disease for 30 days;] [in the case of hides and skins from swine, animals that come from holdings in which there has been no outbreak of swine vesicular disease in the previous 30 days, or of classical or African swine fever in the previous 40 days, and around which within a radius of 10 km there has been no case of these diseases for 30 days;] [animals that have shown no evidence of [foot-and-mouth disease], [rinderpest], [classical swine fever], [African swine fever] or [swine vesicular disease] (*) during ante-mortem health inspection at the slaughterhouse during the 24 hours before slaughter;] 11.4. have undergone all precautions to avoid recontamination with pathogenic agents. Notes Part I: Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity. Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in the event of unloading and reloading. Box reference I.19: use the appropriate HS code: 41.01; 41.02 or 41.03. Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be given Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.

Part II:	·			
(')	OJ L 273, 10.10.2002, p. 1.			
(²)	Delete as appropriate.			
(°)	Delete diseases not applicable to the species	s concerned.		
_	The signature and the stamp must be in a di	fferent colour to that of the printing.		
	Note for the person responsible for the consignment in EU: This certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.			
Official	veterinarian			
	Name (in capitals):	Qualification and title:		
	Date:	Signature:		
	Stamp:			

CHAPTER 5(B)

Health certificate

For treated hides and skins of ungulates, intended for dispatch to or for transit through (²) the European Community

COUN	TRY				Veterinary certificate to EU
	1.1.	Consignor Name	1.2.	Certificate reference number	1.2.a.
		Address	1.3.	Central Competent Author	ity
Part I: Details of dispatched consignment		Tel. No	1.4.	Local Competent Authority	,
ign	1.5.	Consignee	1.6.	Person responsible for the	consignment in EU
Suo		Name		Name	
Pe l		Address		Address	
l th		Postal code		Postal code	
ispē		Tel. No		Tel. No	
o do	1.7.	Country of ISO code I.8. Region of origin Code	1.9.	Country of ISO code	e I.10. Region of Code
is is		origin		destination	destination
Det	I 11	Place of origin	112	Place of destination	
=		Trade of origin		Custom wareho	use \square
Pa		Name Approval number		Name	Approval number
1 1		Address		Address	Approvarnamber
				Postal code	
	I.13.	Place of loading	l.14.	Date of departure	
	I.15.	Means of transport	1.16.	Entry BIP in EU	
		Aeroplane Ship Railway wagon			
		Road vehicle Other I	I.17. No(s) of CITES		
		Documentary references:	1.17.	No(s) of CITES	
	I 18	Description of commodity		I 19 Commo	dity code (HS code)
	1.10.	- Secondary		1.10. Golimio	
					I.20. Quantity
	1.21.	Temperature of product		_	I.22. Number of packages
		Ambient Chilled		Frozen	
	1.23.	Identification of container/Seal number			I.24. Type of packaging
	125	Commodities certified for:			
	1.20.	Animal feedingstuff		Techr	nical use Other
	1.26.	For transit to third country vis-à-vis EU	1.27.	For import or admission int	
				•	
		3rd country ISO code			
	1.28.	Identification of the commodities			
		Approval number	er of es	tablishments	
		Species Manufa	acturing	plant	Net weight
		(Scientific name)			

▼B

COUNTRY Treated hides and skins of Ungulates II.a. Certificate reference number Пb H. Health attestation I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (1) and in particular Article 6 and Annex VIII Chapter VI thereof and certify that the hides and skins described above: 11.1. have been obtained from animals that (2): Certification (a) were slaughtered and their carcasses are fit for human consumption in accordance with Community legislation; or were slaughtered in a slaughterhouse, after undergoing ante-mortem inspection, and were fit, as a result of such inspection, for slaughter for human consumption in accordance with Community legislation; or (b) Part II:

(2) either [dried:1

(2) or [dry-salted or wet-salted for at least 14 days prior to dispatch;]

corresponding species are authorised and have been:

epizootic disease:

...... and according to the declaration of the transporter, the hides and (2) or [dry-salted or wet-salted on the following date skins will be transported by ship and the duration of transport will be such that they will have undergone a minimum of 14 days of salting before they reach the EC border inspection post;]

(2) either [II.2 come from animals originate from a third country or, in the case of regionalisation in accordance with Community legislation,

did not show any clinical signs of any disease communicable to humans or animals, and were not killed to eradicate any

from a part of a third country listed in part 1 of Annex II to Decision 79/542/EEC (3) from which imports of fresh meat of the

- (2) or [salted for seven days in sea salt with the addition of 2 % of sodium carbonate;]
- (2) or will have undergone a minimum of 7 days of salting before they reach the EC border inspection post;]
- come from animals originate from a third country or, in the case of regionalisation in accordance with Community legislation, from (2) or [II.2. a part of a third country listed in part 1 of Annex II to Decision 79/542/EEC from which imports of fresh meat of the corresponding species are NOT authorised and have been:
- (2) either [salted for seven days in sea salt with the addition of 2 % of sodium carbonate;]
- (2) or will have undergone a minimum of 7 days of salting before they reach the EC border inspection post;]
- (2) or [dried for 42 days at a temperature of at least 20 °C;]
- 11.3. the consignment has not been in contact with other animal products or with live animals presenting a risk of spreading a serious transmissible disease

Notes

Part I:

- Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) and information is to be provided in the event of unloading and reloading.
- Box reference I.19: use the appropriate HS code: 41.01; 41.02 or 41.03.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be given.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.

Part II:		
(¹)	OJ L 273, 10.10.2002, p. 1.	
(²)	Delete as appropriate.	
(3)	OJ L 146, 14.6.1979, p. 15.	
_	The signature and the stamp must be in a diffe	rent colour to that of the printing.
	Note for the person responsible for the consigned the consignment until it reaches the border inst	ment in EU: This certificate is only for veterinary purposes and has to accompany pection post.
Official v	eterinarian	
	Name (in capitals):	Qualification and title:
	Date:	Signature:
	Stamp:	

CHAPTER 5(C)

Official declaration

For treated hides and skins of ruminants and of equidae that are intended for dispatch to or for transit through (') the European Community and have been kept separate for 21 daysor will undergo transport for 21 uninterrupted days before importation

COUN	TRY		Veterinary certificate to EU
	I.1. Consignor Name	I.2. Certificate reference number	I.2.a.
	Address	I.3. Central Competent Author	ity
ment	Tel. No	I.4. Local Competent Authority	,
sign	I.5. Consignee	I.6. Person responsible for the	consignment in EU
Sons	Name	Name	
peq (Address	Address	
atch	Postal code	Postal code	
disp	Tel. No	Tel. No	
Part I: Details of dispatched consignment	I.7. Country of ISO code I.8. Region of origin Code origin	I.9. Country of ISO code destination	e I.10. Region of Code destination
: Dei	I.11. Place of origin	I.12. Place of destination	1
art		Custom wareho	use
۵	Name Approval number	Name	Approval number
	Address	Address	
		Postal code	
	I.13. Place of loading	I.14. Date of departure	
	I.15. Means of transport	I.16. Entry BIP in EU	
	Aeroplane Ship Railway wagon		
	Road vehicle Other I	I.17. No(s) of CITES	
	Documentary references:	1.17. NO(S) OF CITES	
	I.18. Description of commodity	I.19. Commo	dity code (HS code)
			I.20. Quantity
	I.21. Temperature of product		I.22. Number of packages
	Ambient Chilled	Frozen	i.zz. Number of packages
	I.23. Identification of container/Seal number		I.24. Type of packaging
	I.25. Commodities certified for:		
	Animal feedingstuff	Techi	nical use Other
	I.26. For transit to third country vis-à-vis EU	I.27. For import or admission int	to EU
	3rd country ISO code		
	I.28. Identification of the commodities		
	Approval number	er of establishments	
	Species Manuf.	acturing plant	Net weight
	(Scientific name)		

COUNTRY

Treated hides and skins of Ruminants and of Equidae that have been kept separate for 21 days or will undergo transport for 21 uninterrupted days before importation

			II.a. Certi	icate reference number	II.b.				
	II. Declaration I the undersigned declare that the hides and skins described above:								
	I, the undersigned declare that the hides and skins described above:								
	II.1.	have been obtained from a		fit for burning an accommission in a					
ا ـ		(a) were slaughtered and	their carcasses are	iit for numan consumption in a	ccordance with Community legislation; or				
Part II: Certification			n a slaughterhouse, after undergoing ante-mortem inspection, and were fit, as a result of such inspection, man consumption in accordance with Community legislation; or						
II: Certi		(c) did not show any clinical signs of any disease communicable to humans or animals, and were not killed to eradicate any epizootic disease;							
r E	II.2.	have been:							
انت	(1) either	[dried;]							
	(1) or	[dry-salted or wet-salted for	at least 14 days pri	or to dispatch;]					
	(1) or	[salted for seven days in se	a salt with the additi	on of 2 % of sodium carbonate]				
	II.3.	II.3. have not been in contact with other animal products or with live animals presenting a risk or spreading a serious transmissible disease;							
	(¹) either	(1) either [II.4. have been kept separate immediately before dispatch for 21 days under official supervision after the treatment described under point (II.2);]							
	(1) or	[II.4. following the declara	ation of the transport	er, the duration of the transpor	period is foreseen to be at least 21 days.]				
	Notes								
	Part I:								
	******	Box reference I.6: Person commodity; it may be filled			is to be filled in only if it is a certificate for transit				
	_	Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses.							
	_	Box reference I.15: Registratis to be provided in the ever			s), flight number (aircraft) or name (ship); information				
	_	Box reference I.19: use the							
	_	Box reference I.23: for bulk	ulk containers, the container number and the seal number (if applicable) should be given.						
	_	Box reference I.26 and I.27	: fill in according to v	whether it is a transit or an impo	ort certificate.				
	Part II:								
	(1)	Delete as appropriate.							
		The signature and the stam	p must be in a differ	ent colour to that of the printing	ı.				
		Note for the person respon- the consignment until it rea			only for veterinary purposes and has to accompany				
	Official vet	erinarian							
		Name (in capitals):		Qu	alification and title:				
		Date:		Sig	nature:				
		Stamp:							

CHAPTER 6(A)

Health certificate

For treated game trophies of birds and ungulates, being solely bones, horns, hooves, claws, antlers, teeth, hides or skins, for dispatch to or for transit through (²) the European Community

COUN	TRY	Veterinary certificate to EU
	I.1. Consignor Name	I.2. Certificate reference I.2.a. number
	Address	I.3. Central Competent Authority
ment	Tel. No	I.4. Local Competent Authority
sign	I.5. Consignee	I.6. Person responsible for the consignment in EU
con	Name Address	Name Address
hed	Address	Address
patc	Postal code	Postal code
fdis	Tel. No	Tel. No
Part I: Details of dispatched consignment	I.7. Country of ISO code I.8. Region of origin Cod origin	de I.9. Country of ISO code I.10. destination
ĕ	I.11. Place of origin	I.12. Place of destination
Part		Custom warehouse
	Name Approval number	Name Approval number
	Address	Address
		Postal code
	I.13. Place of loading	I.14. Date of departure
	I.15. Means of transport	I.16. Entry BIP in EU
	Aeroplane Ship Railway wagon Road vehicle Other	
	Identification:	I.17. No.(s) of CITES
	Documentary references:	1.77 163(6) 61 61126
	I.18. Description of commodity	I.19. Commodity code (HS code)
		I.20. Quantity
	I.21.	I.22. Number of packages
	I.23. Identification of container/Seal number	I.24. Type of packaging
	I.25. Commodities certified for:	Other 🗍
	I.26. For transit to third country vis-à-vis EU	1.27. For import or admission into EU
	3rd country ISO code	
	I.28. Identification of the commodities	
	Species Nature of (Scientific name)	commodity Number of packages

COUNTRY

Treated game trophies of birds and ungulates, being solely bones, horns, hooves, claws, antlers, teeth, hides or skins

			II.a.	Certificate reference number	II.b.				
	II.	Health attestation							
		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (¹) and certify that the game trophies described above:							
ion	II.1. have been packaged, immediately after treatment, without being in contact with other products of animal origin likely to contaminate them, in individual, transparent and closed packages so as to avoid any subsequent contamination;								
licat	(²) either [II.2. in the case of game trophies consisting solely of hides or skin:								
erti	(2) either [have been dried;]								
<u>:</u>	(2) or [have been dry-salted or wet-salted for a minimum of 14 days before dispatch;]								
Part II: Certification	(²) or	(2) or [were dry-salted or wet-salted on							
	(2) or [II.2.	in the case of game tro	ophies consi	sting solely of bone, horns, hooves	, claws, antiers or teeth:				
				ng water for an appropriate time s th is removed; and	o as to ensure that any matter other than bone, horns,				
	 (b) have been disinfected with a product authorised by the competent authority, in particular with hydrogen parts consisting of bone are concerned.] 								
	Notes								
	Part I:								
	 Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for trans commodity; it may be filled in if the certificate is for import commodity. 								
	 Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses. 								
	Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship), information is to be provided in case of unloading and reloading.								
	alatenia.	Box reference I.23: fo	or bulk containers, the container number and the seal number (if applicable) should be						
		Box reference I.19: us	se the appropriate HS code: 05.05; 05.06; 05.07 or						
	_	Box reference I.26 ar	nd I.27: fill in according to whether it is a transit or an import						
	<u></u>		or nature of commodity, specify choosing one or more possibilities among the following: [bones], [horns], tters], [teeth], [hides] or [skins].						
	Part II:								
	(¹)	OJ L 273, 10.10.2002, p. 1.							
	(2)	Delete as appropriate.							
	_	•	•	be in a different colour to that of the	e printing. ate is only for veterinary purposes and has to accompany				
				e border inspection post.	ate is only for veterinary purposes and has to accompany				
	Official veteri	narian							
	1	Name (in capitals):			Qualification and title:				
	ı	Date:			Signature:				
	;	Stamp:							

CHAPTER 6(B)

Health certificate

For game trophies of birds and ungulates consisting of entire parts not having been treated, intended for dispatch to or for transit through (²) the European Community

COUN	TRY				Veterinary certificate to EU	
	l.1.	Consignor Name	1.2.	Certificate reference number	1.2.a.	
		Address	1.3.	I.3. Central Competent Authority		
ment		Tel. No	1.4.	Local Competent Authority		
sign	1.5.	Consignee	1.6.	Person responsible for the	consignment in EU	
Suos		Name		Name		
pec		Address		Address		
atcl		Postal code		Postal code		
disp		Tel. No		Tel. No		
Part I: Details of dispatched consignment	1.7.	Country of ISO code origin I.8. Region of origin Code	1.9.	Country of destination ISO code	1.10.	
ë :	l.11.	Place of origin	I.12.	Place of destination		
art				Custom warehou	use 🔲	
"		Name Approval number		Name	Approval number	
		Address		Address		
				Postal code		
	I.13.	Place of loading	1.14.	Date of departure		
	1.15.	Means of transport	1.16.	Entry BIP in EU		
		Aeroplane Ship Railway wagon		•		
		Road vehicle Other				
		Identification: Documentary references:	1.17.	No(s) of CITES		
	I.18.	Description of commodity		I.19. Commod	lity code (HS code)	
					I.20. Quantity	
	1.21.				I.22. Number of packages	
	1.27.				i.e.z. realisor of poortages	
	100		-		104 7 6 4 1	
	1.23.	Identification of container/Seal number			I.24. Type of packaging	
	1.25.	Commodities certified for:			Other	
	1.26.	For transit to third country vis-à-vis EU	1.27.	For import or admission into	EU	
		3rd country ISO code				
	1.28.	Identification of the commodities				
		Species	٨	lumber of packages		
		(Scientific name)				

COUNTRY

Game trophies of birds and ungulates consisting of entire parts not having been treated

		II.a. Certificate reference number II.b.						
	II. Health attestation							
		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (¹) and certify that the game trophies described above:						
(²) either [II.1. with respect to game trophies of cloven-hoofed animals, excluding swine:								
ficatio		 (a)(region) has been free from foot-and-mouth disease and rinderpest for the previous 12 months, and during the same period, no vaccination against any of those diseases has taken place; and 						
Serti		(b) the game trophies described above:						
Part II: Certification		(i) were obtained from animals which were killed in the territory of that region, which is authorised for export of fresh meat of the corresponding susceptible domestic species and where, during the last 60 days, there have been no animal health restrictions because of outbreaks of diseases to which the game animals are susceptible; and						
		(ii) originated from animals that were killed at a distance of at least 20 km from the borders of another third country or part of a third country not authorised to export untreated game trophies of cloven-hoofed animals other than swine to the Community.]						
	(2) or [II.1.	with respect to game trophies of wild swine:						
		(a)(region) during the last 12 months was free from classical swine fever, African swine fever, swine vescicular disease, foot-and-mouth disease and porcine enteroviral encephalmiyelitis (Teschen disease) and no vaccinations have been carried out against any of those diseases during the last 12 months; and						
		(b) the game trophies described above:						
		(i) were obtained from animals which were killed in that territory, which is authorised for export of fresh meat of the corresponding susceptible domestic species and where, during the last 60 days, there have been no animal health restrictions because of outbreaks of diseases to which the swine are susceptible; and						
		(ii) originated from animals that were killed at a distance of at least 20 km from the borders of another third country or part of a third country not authorised to export untreated game trophies of wild swine to the Community;]						
	(²) or [II.1.	2) or [II.1. with respect to game trophies of solipeds, the game trophies described above were obtained from wild solipeds that were killed in the territory of the exporting country mentioned above;]						
	(2) or [II.1.	with respect to game trophies of game birds:						
		(a) (region) is free from highly pathogenic avian influenza and Newcastle disease; and						
		(b) the game trophies described above were obtained from wild game birds that were killed in that region and where during the last 30 days there have been no animal health restrictions because of outbreaks of disease to which the wild birds are susceptible;]						
	II.2.	The game trophies described above have been packaged without being in contact with other products of animal origin likely to contaminate them, in individual, transparent and closed packages so as to avoid any subsequent contamination.						
	Notes							
	Part I:							
	*****	Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity.						
	tura.	Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses.						
		Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading.						
		Box reference I.19: use the appropriate HS code: 05.05; 05.06 or 05.07.						
		Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.						
	and the same of th	Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate. Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.						

Part II:	·	
(1)	OJ L 273, 10.10.2002, p. 1.	
(²)	Delete as appropriate.	
_	The signature and the stamp must be in a different	ent colour to that of the printing.
_	Note for the person responsible for the consignment until it reaches the border inspe	nent in EU: this certificate is only for veterinary purposes and has to accompany action post.
Official v	reterinarian	
	Name (in capitals):	Qualification and title:
	Date:	Signature:
	Stamp:	

CHAPTER 7(A)

Health certificate

For pig bristles from third countries or regions thereof that are free from African swine fever, intended for dispatch to or for transit through $(^2)$ the European Community

COUN	TRY	Veterinary certificate to EU
	I.1. Consignor Name	I.2. Certificate reference number I.2.a.
	Address	I.3. Central Competent Authority
nent	Tel. No	I.4. Local Competent Authority
ign	I.5. Consignee	I.6. Person responsible for the consignment in EU
cons	Name	Name
peu	Address	Address
atc	Postal code	Postal code
disp	Tel. No	Tel. No
Part I: Details of dispatched consignment	I.7. Country of ISO code I.8. Region of origin Code origin	I.9. Country of ISO code I.10. destination
: De	I.11. Place of origin	I.12. Place of destination
art		Custom warehouse
"	Name Approval number	Name Approval number
	Address	Address
		Postal code
	I.13. Place of loading	I.14. Date of departure
	I.15. Means of transport	I.16. Entry BIP in EU
	Aeroplane Ship Railway wagon	,,
	Road vehicle Other	
	Identification: Documentary references:	1.17.
	I.18. Description of commodity	I.19. Commodity code (HS code) 05.02
		I.20. Quantity
	I.21. Temperature of product	I.22. Number of packages
	Ambient Chilled	Frozen 🗌
	I.23. Identification of container/Seal number	I.24. Type of packaging
	I.25. Commodities certified for: Animal feedingstuff	Technical use \(\bigcap \) Other \(\bigcap \)
	I.26. For transit to third country vis-à-vis EU	I.27. For import or admission into EU
	3rd country ISO code	
	I.28. Identification of the commodities	
	Approval number of establishments	
	Manufacturing plant Numb	per of packages Net weight

COUNTRY

Pig bristles from third countries or regions thereof that are free from African swine fever

			II.a.	Certificate reference numb	er	II.b.			
	II.	Health attestation							
		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (1) and certify that:							
l	II.1.	the pig bristles described above have been obtained from pigs originating, and slaughtered in a slaughterhouse, in the country of origin;							
	11.2.	the pigs from which the pig bristles have been obtained did not show during inspection, carried out at the time of slaughtering, signs of diseases communicable to humans or animals and were not killed to eradicate any epizootic disease;							
	II.3.	the country of origin or, in case of regionalisation according to Community legislation, the region of origin, has been free from African swine fever for at least 12 months;							
l	11.4.	the pig bristles are dry and sec	curely en	closed in packaging.					
l									
l									
l									
l									
1									
l	Notes								
	Part I: Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for transit comit may be filled in if the certificate is for import commodity.			led in only if it is a certificate for transit commodity;					
	_	Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses.							
		Box reference I.15: Registration is to be provided in case of unline			r and lorries),	flight number (aircraft) or name (ship); information			
	_	Box reference I.23: for bulk co	ntainers,	the container number and the	seal number	(if applicable) should be included.			
l		Box reference I.26 and I.27: fil	I in accor	ding to whether it is a transit o	or an import ce	rtificate.			
		Box reference I.28: Manufactu	ring plan	t: provide the veterinary contro	ol number of th	ne registered establishment.			
	Part II:								
l	(¹)	OJ L 273, 10.10.2002, p. 1.							
١	(²)	Delete as appropriate. The signature and the stamp in	nust ha ir	a different colour to that of th	e printing				
		The signature and the stamp must be in a different colour to that of the printing. Note for the person responsible for the consignment in EU: this certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.							
İ	Official	veterinarian							
		Name (in capitals):			Qualific	ation and title:			
		Date:			Signatu	re:			
		Stamp:							
Ĺ									

CHAPTER 7(B)

Health certificate

For pig bristles from third countries or regions thereof that are not free from African swine fever, intended for dispatch to or for transit through (²) the European Community

COUN	TRY	Veterinary certificate to EU
	I.1. Consignor Name	I.2. Certificate reference number I.2.a.
	Address	I.3. Central Competent Authority
nent	Tel. No	I.4. Local Competent Authority
ign	I.5. Consignee	I.6. Person responsible for the consignment in EU
cons	Name	Name
ped	Address	Address
atcl	Postal code	Postal code
disp	Tel. No	Tel. No
Part I: Details of dispatched consignment	I.7. Country of ISO code I.8. Region of origin Code origin	I.9. Country of ISO code I.10. destination
e l	I.11. Place of origin	I.12. Place of destination
art		Custom warehouse
"	Name Approval number	Name Approval number
	Address	Address
		Postal code
	I.13. Place of loading	I.14. Date of departure
	I.15. Means of transport	I.16. Entry BIP in EU
	Aeroplane Ship Railway wagon	
	Road vehicle Other I	
	Documentary references:	1.17.
	I.18. Description of commodity	I.19. Commodity code (HS code) 05.02
		I.20. Quantity
	I.21. Temperature of product Ambient ☐ Chilled ☐	I.22. Number of packages Frozen
	I.23. Identification of container/Seal number	I.24. Type of packaging
	I.25. Commodities certified for: Animal feedingstuff	Technical use Other Other
	I.26. For transit to third country vis-à-vis EU	I.27. For import or admission into EU
	3rd country ISO code	 -
	I.28. Identification of the commodities	
	Approval number of establishments	
	Manufacturing plant Nu	imber of packages Net weight

COUNTRY

Pig bristles from third countries or regions thereof that are not free from African swine fever

			II.a.	Certificate reference number	II.b.			
	II. Health attestation I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (¹) and certify that:							
cation	II.1. the pig bristles described above have been obtained from pigs originating, and slaughtered in a slaughterhouse, in the country of origin;							
Part II: Certification	II.2. the pigs from which the pig bristles have been obtained did not show during inspection, carried out at the time of slaughtering, signs of diseases communicable to humans or animals and were not killed to eradicate any epizootic disease;							
Part l	II.3. (2) either (2) or (2) or	the pig bristles mentioned [boiled;] [dyed;] [bleached;]	d above have	been:				
	II.4.	the pig bristles are dry ar	id securely e	nclosed in packaging.				
	Notes Part I:							
	Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity.							
	Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses.							
	_			nber (railway wagons or containe f unloading and reloading.	r and lorries), flight number (aircraft) or name (ship);			
	Box reference i.23: for bulk containers, the container number and the seal number (if applicable) should be included.				al number (if applicable) should be included.			
		Box reference I.26 and I.	27: fill in acco	ording to whether it is a transit or a	n import certificate.			
	Box reference I.28: Manufacturing plant: provide the veterinary control number of the registered establishment.				umber of the registered establishment.			
	Part II:							
	(1)	OJ L 273, 10.10.2002, p.	1.					
	(2)	Delete as appropriate.						
	 The signature and the stamp must be in a different colour to that of the printing. Note for the person responsible for the consignment in EU: this certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post. 							
	Official veterinarian							
		Name (in capitals):			Qualification and title:			
		Date:			Signature:			
	Stamp:							

CHAPTER 8

Health certificate

For animal by-products (*) to be used for technical purposes, intended for dispatch to or for transit through (3) the European Community

COUN	TRY				Veterinary certificate to EU
		onsignor ame	1.2.	Certificate reference number	1.2.a.
	Ac	ddress	1.3.	Central Competent Author	ity
ment	Te	eł. No	1.4.	Local Competent Authority	
sign	1.5. Co	onsignee	1.6.	Person responsible for the	consignment in EU
Con		ame		Name	
per	Ac	ddress		Address	
atch	Po	ostal code		Postal code	
disp		el. No		Tel. No	
Part I: Details of dispatched consignment		ountry of ISO code I.8. Region of origin Code rigin	1.9.	Country of ISO code destination	1.10.
: De	I.11. Pla	lace of origin	I.12.	Place of destination	
art				Custom wareho	use
"	Na	ame Approval number		Name	Address
	Ac	ddress		Address	
				Postal code	
	1.13. Pla	lace of loading	1.14.	Date of departure	
	1.15 M	leans of transport	I 16	Entry BIP in EU	
	1, 13. 101	Aeroplane Ship Railway wagon	1.10.	Entry Dir III EO	
		Road vehicle Other			
		lentification: ocumentary references:	1.17.		
				1 40 0	
	1.18. De	escription of commodity		1.19. Commod	dity code (HS code)
					I.20. Quantity
	I.21. Te	emperature of product			I.22. Number of packages
		Ambient Chilled C		Frozen	
	1.23 Ide	lentification of container/Seal number			I.24. Type of packaging
					1.24. Type of puologing
	1.25. Co	ommodities certified for: Technical use	1		
	126 Fc	or transit to third country vis-à-vis EU	1 27	For import or admission int	o FU
	1120. 10	or various to using obtainery the a violate		To import of daminosion in	
	3r	rd country ISO code			
	1.28. ld	dentification of the commodities			
		Approval number of esta pecies Nature of commodity Manufacturing pl utific name)		ents Number of package	es Net weight Batch number

COUNTRY

Animal by-products for the manufacture of technical products

		II.a. Certificate reference number II.b.								
	II.1.	Health attestation								
		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (1) and certify that the animal by-products described above:								
	II.1.1.	consist of animal by-products that satisfy the animal health requirements below;								
ou	II.1.2.	have been obtained in the territory of:								
Part II: Certification	(³) either	[(a) that have remained in this territory since birth or for at least the last three months before slaughter;]								
ırtif	(3) or	[(b) killed in the wild in this territory (4);]								
ŭ	II.1.3.	have been obtained from animals:								
art	(3) either	[(a) coming from holdings:								
ď		(i) where, for the following diseases for which the animals are susceptible, there has been neither case/outbreak of rinderpest, swine vesicular disease, Newcastle disease or highly pathogenic avian influenza during the prior 30 days, nor of classical or African swine fever during the prior 40 days; nor in the holdings situated in their vicinity within 10 km, during the prior 30 days; and								
		(ii) where there has been neither case/outbreak of foot and mouth disease during the prior 60 days, nor in the holdings situated in their vicinity within 25 km, during the prior 30 days; and								
		(b) which:								
		(i) were not killed to eradicate any epizootic disease;								
		 (ii) have remained in their holdings of origin for at least forty days before departure and which have been transported directly to the slaughterhouse without contact with other animals which did not comply with the same health conditions; 								
		(iii) at the slaughterhouse, have passed the ante-mortem health inspection during the 24 hours before the slaughter and have shown no evidence of the diseases referred to above for which the animals are susceptible; and								
		(iv) have been treated in the slaughterhouse before and at the time of slaughter or killing in accordance with the relevant provisions of Council Directive 93/119/EC on animal welfare;]								
	(3) or	[(a) captured and killed in the wild in an area:								
		(i) in which within 25 km there has been no case/outbreak of any of the following diseases for which the animals are susceptible: foot and mouth disease, rinderpest, Newcastle disease or highly pathogenic avian influenza during the prior 30 days nor of classical or African swine fever during the prior 40 days; and								
		(ii) that is situated at a distance that exceeds 20 km from the borders separating another territory of a country or part thereof, which is not authorised at these dates for exporting this material to the European Community; and								
		(b) which after killing were transported within 12 hours for chilling either to a collection centre and immediately afterwards to a game establishment, or directly to a game establishment;]								
	II.1.4.	have been obtained in an establishment around which, within a radius of 10 km, there has been no case/outbreak of diseases referred to in point II.1.3 for which the animals are susceptible during the prior 30 days or, in the event of a case of disease, the preparation of raw material for exportation to the European Community has been authorised only after removal of all meat, and the total cleaning and disinfection of the establishment under the control of an official veterinarian;								
	have been obtained and prepared without contact with other material not complying with the conditions required above, and it has been handled so as to avoid contamination with pathogenic agents;									
	II.1.6.	have been packed in new packaging preventing any leakage and in officially sealed containers bearing the label indicating "RAW MATERIAL ONLY FOR THE MANUFACTURE OF TECHNICAL PRODUCTS" and the name and address of the EU establishment of destination;								
	II.1.7.	consist only of the following animal by-products:								
	(³) either	 parts of slaughtered animals, which are fit for human consumption in accordance with Community legislation, but are not intended for human consumption for commercial reasons,] 								
	(³) and/or	 parts of slaughtered animals, which are rejected as unfit for human consumption but are not affected by any signs of diseases communicable to humans or animals and derive from carcasses that are fit for human consumption in accordance with Community legislation.] 								
	(3) and/or	[— animal by-products derived from the production of products intended for human consumption, including degreased bones and greaves,]								
(3) and/or [— former foodstuffs of animal origin, or former foodstuffs containing products of animal origin, other than catering which are no longer intended for human consumption for commercial reasons or due to problems of manufa packaging defects or other defects which do not present any risk to humans or animals.]										
	(3) and/or	[— fish or other sea animals, except sea mammals, caught in the open sea for the purposes of fishmeal production,								

(3) and/or	[— fresh by-products from fish from plants manufact	uring fish products for human consumption,]			
(3) and/or	 [— shells, hatchery by-products and cracked egg by-products originating from animals which did not show clinical signs of any disease communicable through that product to humans or animals,] 				
(3) and/or	 fur originating from animals that did not show clinical signs of any disease communicable through that product to humans of animals.] 				
1.1.8.	have been deep-frozen at the plant of origin or have been preserved in accordance with EU legislation in such a way that the will not spoil between dispatch and delivery to the plant of destination.				
(³)(6) [II.2.	Specific requirements				
(³) (⁷) II.2.1.		ils that have been obtained in the territory mentioned under (II.1.2), wherease are being regularly carried out and officially controlled in domest			
(³) (8) II.2.2.	The by-products in this consignment consists of anima	al by-products derived from offal or de-boned meat.]			
Notes					
Part I:					
_	Box reference I.6: Person responsible for the consign commodity; it may be filled in if the certificate is for improvement of the certificate is for improvement.	nment in EU: this box is to be filled in only if it is a certificate for transport commodity.			
~~	Box reference I.12: Place of destination: this box is to transit can only be stored in free zones, free warehous	be filled in only if it is a certificate for transit commodity. The products ses and custom warehouses.			
	Box reference I.15: Registration number (railway wa information is to be provided in case of unloading and	agons or container and lorries), flight number (aircraft) or name (shipreloading.			
	Box reference I.19: use the appropriate HS code: 05.1	1.91; 05.11.99 or 30.01.			
	Box reference I.23: for bulk containers, the container r	number and the seal number (if applicable) should be included.			
_	Box reference I.26 and I.27: fill in according to whethe	r it is a transit or an import certificate.			
-	Box reference I.28: Manufacturing plant: provide the v	eterinary control number of the approved establishment.			
Part II:					
(*)	Excluding raw blood, raw milk, hides and skins of ungu import of these products) as well as wool, hair, feather	ılates or ruminants and pig bristles (see relevant specific certificates for tl rs or parts of feathers.			
(1)	OJ L 273, 10.10.2002, p. 1.				
(2)	The name and ISO code number of the exporting cour	•			
	part 1 of Annex II of Council Decision 79/542/EEC				
	 the Annex to Commission Decision 94/984/EC, ar 				
	— the Annex to Commission Decision 2000/585/EC.				
	included.	nex (where applicable for the susceptible species concerned) should			
(3)	Delete as appropriate.				
4)	Only for countries from where game meat intended importation into the European Community.	for human consumption of the same animal species is authorised f			
(5)	Catering waste means all waste food, including used including central kitchens and household kitchens.	d cooking oils, originating in restaurants, catering facilities and kitcher			
(6)	Supplementary guarantees to be provided when the material of domestic ruminants originated in the territory of a South America or South African country or part thereof from where only maturated and de-boned fresh meat of domestic ruminants for huma consumption is permitted for exportation to the European Community. In the case of offal only trimmed offal of domestic ruminant which must be exclusively offal from which the bones, cartilage, trachea and main bronchi, lymphatic glands adhering connectivities. If at and mucus have been completely removed is permitted. The whole masseter muscles of bovine animals, incised if accordance with paragraph 41(a) of Chapter VIII of Annex I to Council Directive 64/433/EEC, are also permitted.				
(7)	Only for certain South American countries.				
	Only for certain South American and South African con	untries.			
7)	The signature and the stamp must be in a different co	lour to that of the printing.			
-	Note for the person responsible for the consignment in EU: this certificate is only for veterinary purposes and has to accompar the consignment until it reaches the border inspection post.				
(8) — —					
C) Control the consignment until it reaches the border inspection					
_	the consignment until it reaches the border inspection				
_	the consignment until it reaches the border inspection inarian	post.			

Health certificate

For fish oil not intended for human consumption to be used as feed material or for technical purposes, intended for dispatch to or for transit through (²) the European Community

COUN	TRY	Veterinary certificate to EU
	I.1. Consignor Name	I.2. Certificate reference I.2.a. number
	Address	I.3. Central Competent Authority
nent	Tel. No	I.4. Local Competent Authority
ign	I.5. Consignee	I.6. Person responsible for the consignment in EU
Suo:	Name	Name
pg	Address	Address
atch	Postal code	Postal code
lisp	Tel. No	Tel. No
Part I: Details of dispatched consignment	I.7. Country of ISO code I.8. origin	I.9. Country of ISO code I.10. destination
Det	I.11. Place of origin	I.12. Place of destination
Έ	Ü	Custom warehouse
ا ۾	Name Approval number	Name Approval number
	Address	Address
		Postal code
	I.13. Place of loading	I.14. Date of departure
	I.15. Means of transport Aeroplane Ship Railway wagon	I.16. Entry BIP in EU
	Road vehicle Other	
	Identification:	1.17.
	Documentary references:	
	I.18. Description of commodity	I.19. Commodity code (HS code)
		I.20. Quantity
	I.21. Temperature of product	I.22. Number of packages
	Ambient Chilled	Frozen 🗌
	I.23. Identification of container/Seal number	1.24. Type of packaging
	I.25. Commodities certified for: Animal feedingstuff	Technical use ☐
	I.26. For transit to third country vis-à-vis EU	I.27. For import or admission into EU
	1.26. For transit to tilid country vis-a-vis E0	1.27. For import of admission into EO
	3rd country ISO code	
	I.28. Identification of the commodities	
	Approval number of establishments Nature of commodity Manufacturing plant	Number of packages Net weight Batch number

COUNTRY

Fish oil to be used as feed material or for technical purposes

		11.3	a. Certificate referer	nce number	II.b.				
11.	l.	Health attestation							
		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (¹) and certify that the fish oil described above:							
H.	.1.	consists of fish oil that satisfy the health requirements below;							
H.	.2.	contains exclusively fish oil not intended for human consumption;							
H.	.3.	has been prepared and stored in a dedicated fish plant approved, validated and supervised by the competent authority in accordance with Article 17 and where appropriate Article 11 of Regulation (EC) No 1774/2002;							
H.	.4.	has been prepared exclusively with the following animal by-products:							
(2	²) either	[— former foodstuffs of fish origin, other than catering waste (3), which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects which do not present any risk to humans or animals,]							
(2	2) and/or	[— fish or other sea animals, e	xcept sea mammals, caugh	nt in the open sea for	the purposes of fishmeal production,]				
(2) and/or [— fresh by-products from fish from plants manufacturing fish products for human consumption;]									
II.	.5.	the fish oil:							
		(a) has been subjected to propathogenic agents;	cessing in accordance with	n Annex VII, Chapte	er IV of Regulation 1774/2002/EC, in order to				
		(b) has not been in contact wit	h other types of oils includin	ng rendered fats fron	n other animal species; and				
- (²	²) either	[(c) is packaged in new cont contamination;]	ainers or in containers th	at have been clea	ned and all precautions taken to prevent the				
(2) or [(c) where bulk transport transportation of the					other bulk container or bulk road tanker used in t n to the ship or into shore tanks or direct to plar				
1		have been inspected and fe							
		have been inspected and for and which bear labels indicating	ound to be clean before use						
	lotes Part I:	and which bear labels indicating	ound to be clean before use	:] UMPTION".					
		and which bear labels indicating Box reference I.6: Person respondity; it may be filled in if the	ound to be clean before use "NOT FOR HUMAN CONS" onsible for the consignment certificate is for import co	it in EU: this box is ommodity.	to be filled in only if it is a certificate for trans				
		and which bear labels indicating Box reference I.6: Person resp. commodity; it may be filled in if the Box reference I.12: Place of dest can only be stored in free zones.	ound to be clean before use "NOT FOR HUMAN CONS onsible for the consignmen he certificate is for import or ination: this box is to be filler, free warehouses and custor	et in EU: this box is bornodity. It in only if it is a certion warehouses.	to be filled in only if it is a certificate for tran ficate for transit commodity. The products in tran				
		Box reference I.6: Person responsively; it may be filled in if the Box reference I.12: Place of dest can only be stored in free zones. Box reference I.15: Registration ris to be provided in case of unlost.	ound to be clean before use "NOT FOR HUMAN CONS onsible for the consignmen the certificate is for import or ination: this box is to be fille free warehouses and custo thumber (railway wagons or cading and reloading.	et in EU: this box is ommodity. If in only if it is a certion warehouses. Container and lorries)	to be filled in only if it is a certificate for trans				
		Box reference I.6: Person responding to the solution of the so	ound to be clean before uses "NOT FOR HUMAN CONS onsible for the consignment the certificate is for import or ination: this box is to be filler, free warehouses and custon the custon that the control of the custon that the custom that the custon that the custom that th	bit in EU: this box is ommodity. d in only if it is a certion warehouses. container and lorries)	to be filled in only if it is a certificate for tran ficate for transit commodity. The products in tran , flight number (aircraft) or name (ship); informati				
		Box reference I.6: Person responsive to the commodity; it may be filled in if the box reference I.12: Place of dest can only be stored in free zones. Box reference I.15: Registration is to be provided in case of unload Box reference I.19: use the approx reference I.23: for bulk contributions.	ound to be clean before uses "NOT FOR HUMAN CONS onsible for the consignment the certificate is for import or ination: this box is to be filler, free warehouses and custon the custon that the control of the control	b:] UMPTION". If in EU: this box is ommodity. If in only if it is a certion warehouses. Container and lorries) 5.18. If and the seal numbers.	to be filled in only if it is a certificate for transficate for transficate for transit commodity. The products in transfight number (aircraft) or name (ship); informativer (if applicable) should be included.				
		Box reference I.6: Person respondity; it may be filled in if the Box reference I.12: Place of dest can only be stored in free zones. Box reference I.15: Registration is to be provided in case of unload Box reference I.19: use the appr Box reference I.23: for bulk contributions and I.27: fill in Box reference I.26 and I.27: fill in	ound to be clean before uses "NOT FOR HUMAN CONS" onsible for the consignment the certificate is for import or ination: this box is to be filler, free warehouses and custon the cumber (railway wagons or cading and reloading, opriate HS code: 15.04 or 1 ainers, the container number in according to whether it is a	bit in EU: this box is commodity. If in only if it is a certion warehouses. International container and lorries. In and the seal number a transit or an importance.	to be filled in only if it is a certificate for transficate for transit commodity. The products in transfight number (aircraft) or name (ship); informativer (if applicable) should be included.				
P:		Box reference I.6: Person respondity; it may be filled in if the Box reference I.12: Place of dest can only be stored in free zones. Box reference I.15: Registration is to be provided in case of unload Box reference I.19: use the appr Box reference I.23: for bulk contributions and I.27: fill in Box reference I.26 and I.27: fill in	ound to be clean before uses "NOT FOR HUMAN CONS" onsible for the consignment the certificate is for import or ination: this box is to be filler, free warehouses and custon the cumber (railway wagons or cading and reloading, opriate HS code: 15.04 or 1 ainers, the container number in according to whether it is a	bit in EU: this box is commodity. If in only if it is a certion warehouses. International container and lorries. In and the seal number a transit or an importance.	to be filled in only if it is a certificate for transficate for transit commodity. The products in transfight number (aircraft) or name (ship); informativer (if applicable) should be included.				
P:		Box reference I.6: Person respondity; it may be filled in if the Box reference I.12: Place of dest can only be stored in free zones. Box reference I.15: Registration is to be provided in case of unload Box reference I.19: use the appr Box reference I.23: for bulk contributions and I.27: fill in Box reference I.26 and I.27: fill in	ound to be clean before uses "NOT FOR HUMAN CONS" onsible for the consignment the certificate is for import or ination: this box is to be filler, free warehouses and custon the cumber (railway wagons or cading and reloading, opriate HS code: 15.04 or 1 ainers, the container number in according to whether it is a	bit in EU: this box is commodity. If in only if it is a certion warehouses. International container and lorries. In and the seal number a transit or an importance.	to be filled in only if it is a certificate for transficate for transit commodity. The products in transfight number (aircraft) or name (ship); informativer (if applicable) should be included.				
P:	eart I:	Box reference I.6: Person responsively: it may be filled in if the Box reference I.12: Place of dest can only be stored in free zones. Box reference I.15: Registration is to be provided in case of unload Box reference I.19: use the appr Box reference I.23: for bulk control Box reference I.26 and I.27: fill it Box reference I.28: Manufacturing	ound to be clean before uses "NOT FOR HUMAN CONS" onsible for the consignment the certificate is for import or ination: this box is to be filler, free warehouses and custon the cumber (railway wagons or cading and reloading, opriate HS code: 15.04 or 1 ainers, the container number in according to whether it is a	bit in EU: this box is commodity. If in only if it is a certion warehouses. International container and lorries. In and the seal number a transit or an importance.	to be filled in only if it is a certificate for transiticate for transit commodity. The products in transit flight number (aircraft) or name (ship); informative (if applicable) should be included.				
P:	eart I:	Box reference I.6: Person responsive to model to the commodity; it may be filled in if the box reference I.12: Place of dest can only be stored in free zones. Box reference I.15: Registration is to be provided in case of unlocation of the control	ound to be clean before uses "NOT FOR HUMAN CONS "NOT FOR HUMAN CONS "INOT FOR HUMAN CONS "IN	at in EU: this box is ommodity. d in only if it is a certion warehouses. container and lorries) 5.18. er and the seal numb a transit or an importation number of the t	to be filled in only if it is a certificate for transficate for transit commodity. The products in transfight number (aircraft) or name (ship); informations (if applicable) should be included. It certificate.				
P: (2)	eart I:	Box reference I.6: Person responsive to the commodity; it may be filled in if the Box reference I.12: Place of dest can only be stored in free zones. Box reference I.15: Registration is to be provided in case of unload Box reference I.19: use the appr Box reference I.23: for bulk control Box reference I.26 and I.27: fill in Box reference I.28: Manufacturing OJ L 273, 10.10.2002, p. 1. Delete as appropriate. Catering waste means all waste from the signature and the stamp mutation.	ound to be clean before uses "NOT FOR HUMAN CONS "NOT FOR HUMAN CONS "NOT FOR HUMAN CONS "In this box is to be filled, free warehouses and custs number (railway wagons or cading and reloading, opriate HS code: 15.04 or 1 ainers, the container number in according to whether it is a griphant: provide the registration, including used cooking citchens.	ti in EU: this box is ommodity. d in only if it is a certion warehouses. container and lorries) 5.18. er and the seal numb a transit or an importation number of the tools, originating in retails.	to be filled in only if it is a certificate for transficate for transit commodity. The products in transfight number (aircraft) or name (ship); informations (if applicable) should be included. The certificate. The reatment processing establishment.				
P: (2)	eart I:	Box reference I.6: Person responsive to the commodity; it may be filled in if the Box reference I.12: Place of dest can only be stored in free zones. Box reference I.15: Registration is to be provided in case of unload Box reference I.19: use the appr Box reference I.23: for bulk control Box reference I.26 and I.27: fill in Box reference I.28: Manufacturing OJ L 273, 10.10.2002, p. 1. Delete as appropriate. Catering waste means all waste from the signature and the stamp mutation.	ound to be clean before uses "NOT FOR HUMAN CONS" "NOT FOR HUMAN CONS" onsible for the consignment he certificate is for import or ination: this box is to be filler, free warehouses and custo number (railway wagons or or ading and reloading, opriate HS code: 15.04 or 1 ainers, the container numbe in according to whether it is a g plant: provide the registra cood, including used cooking citchens. set be in a different colour to for the consignment in EU: the	ti in EU: this box is ommodity. d in only if it is a certion warehouses. container and lorries) 5.18. er and the seal numb a transit or an importation number of the tools, originating in retails.	to be filled in only if it is a certificate for transficate for transit commodity. The products in transfight number (aircraft) or name (ship); informativer (if applicable) should be included.				
P;	eart I:	Box reference I.6: Person respondity; it may be filled in if the Box reference I.12: Place of dest can only be stored in free zones. Box reference I.15: Registration is to be provided in case of unload Box reference I.19: use the appr Box reference I.23: for bulk cont. Box reference I.26 and I.27: fill in Box reference I.28: Manufacturin OJ L 273, 10.10.2002, p. 1. Delete as appropriate. Catering waste means all waste frontier in the signature and the stamp mu. Note for the person responsible tonsignment until it reaches the	ound to be clean before uses "NOT FOR HUMAN CONS" "NOT FOR HUMAN CONS" onsible for the consignment he certificate is for import or ination: this box is to be filler, free warehouses and custo number (railway wagons or or ading and reloading, opriate HS code: 15.04 or 1 ainers, the container numbe in according to whether it is a g plant: provide the registra cood, including used cooking citchens. set be in a different colour to for the consignment in EU: the	ti in EU: this box is ommodity. d in only if it is a certion warehouses. container and lorries) 5.18. er and the seal numb a transit or an importation number of the tools, originating in retails.	to be filled in only if it is a certificate for transit commodity. The products in transit flight number (aircraft) or name (ship); informativer (if applicable) should be included. The certificate are atment/processing establishment.				
P;		Box reference I.6: Person respondity; it may be filled in if the Box reference I.12: Place of dest can only be stored in free zones. Box reference I.15: Registration is to be provided in case of unload Box reference I.19: use the appr Box reference I.23: for bulk cont. Box reference I.26 and I.27: fill in Box reference I.28: Manufacturin OJ L 273, 10.10.2002, p. 1. Delete as appropriate. Catering waste means all waste frontier in the signature and the stamp mu. Note for the person responsible tonsignment until it reaches the	ound to be clean before uses "NOT FOR HUMAN CONS" "NOT FOR HUMAN CONS" onsible for the consignment he certificate is for import or ination: this box is to be filler, free warehouses and custo number (railway wagons or or ading and reloading, opriate HS code: 15.04 or 1 ainers, the container numbe in according to whether it is a g plant: provide the registra cood, including used cooking citchens. set be in a different colour to for the consignment in EU: the	bit in EU: this box is ommodity. d in only if it is a certion warehouses. container and lorries) 5.18. er and the seal number at ransit or an importation number of the toolis, originating in retat of the printing.	to be filled in only if it is a certificate for transit commodity. The products in transit flight number (aircraft) or name (ship); informativer (if applicable) should be included. The certificate are atment/processing establishment.				
P;		Box reference I.6: Person respondity; it may be filled in if the Box reference I.12: Place of dest can only be stored in free zones. Box reference I.15: Registration ris to be provided in case of unload Box reference I.19: use the appr Box reference I.23: for bulk cont. Box reference I.26 and I.27: fill in Box reference I.28: Manufacturin OJ L 273, 10.10.2002, p. 1. Delete as appropriate. Catering waste means all waste freentral kitchens and household in the signature and the stamp mu. Note for the person responsible tonsignment until it reaches the	ound to be clean before uses "NOT FOR HUMAN CONS" "NOT FOR HUMAN CONS" onsible for the consignment he certificate is for import or ination: this box is to be filler, free warehouses and custo number (railway wagons or or ading and reloading, opriate HS code: 15.04 or 1 ainers, the container numbe in according to whether it is a g plant: provide the registra cood, including used cooking citchens. set be in a different colour to for the consignment in EU: the	bit in EU: this box is commodity. d in only if it is a certion warehouses. container and lorries) 5.18. or and the seal number a transit or an importation number of the toolis, originating in rethat of the printing.	to be filled in only if it is a certificate for transit commodity. The products in transit ficate for transit commodity. The products in transit flight number (aircraft) or name (ship); information of the certificate of the certificate. The certificate of the				

CHAPTER 10(A)

Health certificate

For rendered fats not intended for human consumption to be used as feed material or for technical purposes, intended for dispatch to or for transit through (*) the European Community

COUN	TRY				Veterinary certificate to EU
	l.1.	Consignor Name	1.2.	Certificate reference number	1.2.a.
		Address	I.3. Central Competent Authority		
ment		Tel. No	1.4.	Local Competent Author	rity
ign	1.5.	Consignee	1.6.	Person responsible for the	ne consignment in EU
Ü		Name		Name	
ped		Address		Address	
atch		Postal code		Postal code	
disp		Tel. No		Tel. No	
Part I: Details of dispatched consignment	1.7.	Country of ISO code I.8. origin	1.9.	Country of ISO co	ode I.10.
. De	I.11.	Place of origin	l.12.	Place of destination	
art				Custom warel	house
		Name Approval number		Name	Approval number
		Address	ŀ	Address	
				Postal code	
	I.13.	Place of loading	1.14.	Date of departure	
	1.15.	Means of transport	I.16.	Entry BIP in EU	
		Aeroplane Ship Railway wagon		,	
		Road vehicle Other	L		
		Identification: Documentary references:	1.17.		
	I.18.	Description of commodity		I.19. Comn	nodity code (HS code)
				<u> </u>	I.20. Quantity
	1.21.	Temperature of product			I.22. Number of packages
		Ambient Chilled Chilled		Frozen 🗌	
	1.23.	Identification of container/Seal number			I.24. Type of packaging
	1.25.	Commodities certified for:		<u> </u>	
		Animal feedingstuff		Tec	chnical use
	1.26.	For transit to third country vis-à-vis EU	1.27.	For import or admission	into EU
		3rd country ISO code			
	1.28.	Identification of the commodities			
		Approval number of e	establis	hments	
		Species Nature of commodity Manufacturin (Scientific name)		Number of pac	kages Net weight Batch number

COUNTRY

Rendered fats not intended for human consumption to be used as feed material or for technical purposes

		II.a. Certificate reference number II.b.								
	H.	Health attestation								
		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (¹) and certify that the rendered fats described above:								
ation	II.1.	consist of rendered fats that satisfy the health requirements below;								
Certific	II.2.	consist of rendered fats not intended for human consumption;								
Part II: Certification	H.3.	have been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance with Article 17 and where appropriate Article 11 of Regulation (EC) No 1774/2002 or in accordance with Chapter II of Annex C to Council Directive 77/99/EEC (2) or Chapter 9 of Annex I to Council Directive 92/118/EEC (3), in order to kill pathogenic agents;								
	II.4.	have been prepared exclusively with the following animal by-products:								
	(4) either	 parts of slaughtered animals, which are fit for human consumption in accordance with Community legislation, but are not intended for human consumption for commercial reasons,] 								
	(4) and/or	 parts of slaughtered animals, which are rejected as unfit for human consumption but are not affected by any signs of diseases communicable to humans or animals and derive from carcasses that are fit for human consumption in accordance with Community legislation,] 								
	(4) and/or	[— hides and skins, hooves and horns, pig bristles and feathers originating from animals that were slaughtered in a slaughterhouse, after undergoing antemortem inspection, and were fit, as a result of such inspection, for slaughter for human consumption in accordance with Community legislation.]								
	(4) and/or	[— blood obtained from animals other than ruminants that are slaughtered in a slaughterhouse, after undergoing ante-mortem inspection, and were fit, as a result of such inspection, for slaughter for human consumption in accordance with Community legislation,]								
	(4) and/or	[— animal by-products derived from the production of products intended for human consumption, including degreased bones and greaves,]								
	(4) and/or	[— former foodstuffs of animal origin, or former foodstuffs containing products of animal origin, other than catering waste (5), which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects which do not present any risk to humans or animals,]								
	(4) and/or	[— milk originating from animals which do not show any clinical signs of any disease communicable through that product to humans or animals,]								
	(4) and/or	[— fish or other sea animals, except sea mammals, caught in the open sea for the purposes of fishmeal production,]								
	(4) and/or	[by-products from fish from plants manufacturing fish products for human consumption,]								
	(4) and/or	 shells, hatchery by-products and cracked egg by-products originating from animals which did not show clinical signs of any disease communicable through that product to humans or animals;] 								
	II.5.	if derived from ruminant animals were purified in such way that the maximum levels of remaining total insoluble impurities does not exceed 0,15 % in weight;								
	II.6.	the rendered fats:								
		(a) have been subjected to processing in accordance with Annex VII, Chapter IV of Regulation 1774/2002/EC, or treatment in accordance with Council Directives 77/99/EEC or 92/118/EEC, in order to kill pathogenic agents; and								
	(4) either	[(b) are packaged in new containers or in containers that have been cleaned and all precautions taken to prevent their contamination;]								
	(4) or	(b) where bulk transport is intended, the pipe, pumps and bulk tanks and any other bulk container or bulk road tanker used in the transportation of the product from the manufacturing plant either directly on to the ship or into shore tanks or direct to plants have been inspected and found to be clean before use;]								
		and which bear labels indicating "NOT FOR HUMAN CONSUMPTION".								

Notes			
Part I:			
	Box reference I.6: Person responsible for the consignment in commodity; it may be filled in if the certificate is for import comm	EU: this box is to be filled in only if it is a certificate for transit odity.	
_	Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transact on only be stored in free zones, free warehouses and custom warehouses.		
_	Box reference I.15: Registration number (railway wagons or contains to be provided in case of unloading and reloading.	iner and lorries), flight number (aircraft) or name (ship); information	
_	Box reference I.19: use the appropriate HS code: 15.02; 15.03;	15.04; 15.05; 15.06; 15.16.10; 15.17 or 15.18.	
	Box reference I.23: for bulk containers, the container number an	d the seal number (if applicable) should be included.	
_	Box reference I.26 and I.27: fill in according to whether it is a tra-	nsit or an import certificate.	
_	Box reference I.28: Manufacturing plant: provide the registration	number of the treatment/processing establishment.	
Part II:			
(¹)	OJ L 273, 10.10.2002, p. 1.		
(2)	OJ L 26, 31.1.1977, p. 85.		
(3)	OJ L 62, 15.3.1993, p. 49.		
(4)	Delete as appropriate.		
(5)	Catering waste means all waste food, including used cooking oils, central kitchens and household kitchens.	originating in restaurants, catering facilities and kitchens, including	
	The signature and the stamp must be in a different colour to that	of the printing.	
-	Note for the person responsible for the consignment in EU: this consignment until it reaches the border inspection post.	ertificate is only for veterinary purposes and has to accompany the	
Official	veterinarian		
	Name (in capitals):	Qualification and title:	
	Date:	Signature:	
	Stamp:		

CHAPTER 10(B)

Health certificate

For rendered fats not intended for human consumption to be used for technical purposes, intended for dispatch to or for transit through (²) the European Community

COUN	ΓRY	Veterinary certificate to EU
	I.1. Consignor Name	I.2. Certificate reference number I.2.a.
	Address	I.3. Central Competent Authority
Part I: Details of dispatched consignment	Tel. No	I.4. Local Competent Authority
ign	I.5. Consignee	I.6. Person responsible for the consignment in EU
Sons	Name	Name
ped	Address	Address
atch	Postal code	Postal code
disp	Tel. No	Tel. No
ot	I.7. Country of ISO code I.8.	I.9. Country of ISO code I.10.
tails	origin	destination
e :	I.11. Place of origin	I.12. Place of destination
art		Custom warehouse
"	Name Approval number	Name Approval number
	Address	Address
		Postal code
•	I.13. Place of loading	I.14. Date of departure
	I.15. Means of transport	I.16. Entry BIP in EU
	Aeroplane Ship Railway wagon	
	Road vehicle Other	
	Identification: Documentary references:	1.17.
	I.18. Description of commodity	I.19. Commodity code (HS code)
		I.20. Quantity
	I.21. Temperature of product	I.22. Number of packages
	Ambient Chilled	Frozen 🗍
	L 22 Identification of contained Contained	
	I.23. Identification of container/Seal number	I.24. Type of packaging
	I.25. Commodities certified for: Technical use	
	I.26. For transit to third country vis-à-vis EU	I.27. For import or admission into EU
	1.20. To dansic to dilid obality viola viole	1.27. For import of damission into 20
	3rd country ISO code	
	I.28. Identification of the commodities	
	Approval number of e	establishments
	Species Nature Manufactu	
	(Scientific name) of commodity plant	packages weight number

UN	TRY				Rendered fats to be used for technical purp					
			II.a.	Certificate reference number	II.b.					
	II.	Health attestation		**						
		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (¹) a								
1		the rendered fats describe		,,						
1	II.1.	consist of rendered fats that satisfy the health requirements below;								
1	II.2.	consist of rendered fats not intended for human or animal consumption;								
	II.3.	have been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance with Article 13 and where appropriate Article 11 of Regulation (EC) No 1774/2002, in order to kill pathogenic agents;								
1	11.4.	have been prepared exclusively with the following animal by-products:								
1	(²) either	[Category 2 materials (3);]								
1	(2) or	[a mixture of Category 2	naterials wit	h Category 3 materials (4);]						
	II.5.	if derived from ruminant a exceed 0,15 % in weight;	nimals were	purified in such way that the maximu	m levels of remaining total insoluble impurities does					
1	II.6.	the rendered fats:								
		(a) have been subjecte to kill pathogenic ag		sing in accordance with Annex VII, Cha	apter XII of Regulation (EC) No 1774/2002/EC, in o					
_	(²) either	[(b) are packaged in n contamination;]	ew containe	ers or in containers that have been	cleaned and all precautions taken to prevent t					
	(²) or	transportation of the	product fro		any other bulk container or bulk road tanker used in otty on to the ship or into shore tanks or direct to pla					
		and which bear labels indicating 'NOT FOR HUMAN OR ANIMAL CONSUMPTION'.								
	Notes									
1	Part I:									
	_			le for the consignment in EU: this b rtificate is for import commodity.	ox is to be filled in only if it is a certificate for tra					
	ADDRAGE.			on: this box is to be filled in only if it is a warehouses and custom warehouses	certificate for transit commodity. The products in tra					
	_	Box reference I.15: Registist to be provided in case			rries), flight number (aircraft) or name (ship); ínforma					
1	_	Box reference I.19: use the	ie appropria	te HS code: 15.01; 15.02; 15.03; 15.0	4; 15.05; 15.06; 15.16.10; 15.17 or 15.18.					
	_	Box reference I.23: for bu	lk container	s, the container number and the seal r	number (if applicable) should be included.					
		Box reference I.26 and I.2	7: fill in acc	ording to whether it is a transit or an ir	mport certificate.					
1	_	Box reference I.28: Manu	facturing pla	ant: provide the registration number of	the treatment/processing establishment.					
	Part II:									
	(¹)	OJ L 273, 10.10.2002, p.	1.							
	(2)	Delete as appropriate.								
	(3)	List of Category 2 materia	ıls:							
		Article 4(1)(d) or from	n Category		ughterhouses other than slaughterhouses covered ngs, materials from de-sanding, grease and oil mixtu					
				ning residues of veterinary drugs and dues exceed the permitted level laid do	contaminants listed in Group B(1) and (2) of Annex own by Community legislation;					
		inspections provide	d for in Con	nmunity legislation, fail to comply with	mported from third countries and, in the course of h the veterinary requirements for their importation cepted under restrictions laid down under Commu					

- (d) animals and parts of animals, other than those referred to in Article 4, that die other than by being slaughtered for human consumption, including animals killed to eradicate an epizootic disease;
- (e) mixtures of Category 2 material with Category 3 material, including any material destined for processing in a Category 2 processing plant; and
- (f) animal by-products other than Category 1 material or Category 3 material.
- (4) List of Category 3 materials:
 - (a) parts of slaughtered animals, which are fit for human consumption in accordance with Community legislation, but are not
 intended for human consumption for commercial reasons;
 - (b) parts of slaughtered animals, which are rejected as unfit for human consumption but are not affected by any signs of diseases communicable to humans or animals and derive from carcasses that are fit for human consumption in accordance with Community legislation;
 - (c) hides and skins, hooves and horns, pig bristles and feathers originating from animals that were slaughtered in a slaughterhouse, after undergoing antemortem inspection, and were fit, as a result of such inspection, for slaughter for human consumption in accordance with Community legislation;
 - (d) blood obtained from animals other than ruminants that are slaughtered in a slaughterhouse, after undergoing antemortem inspection, and were fit, as a result of such inspection, for slaughter for human consumption in accordance with Community legislation;
 - (e) animal by-products derived from the production of products intended for human consumption, including degreased bones and greaves;
 - (f) former foodstuffs of animal origin, or former foodstuffs containing products of animal origin, other than catering waste, which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects which do not present any risk to humans or animals;
 - (g) milk originating from animals which do not show any clinical signs of any disease communicable through that product to humans or animals:
 - (h) fish or other sea animals, except sea mammals, caught in the open sea for the purposes of fishmeal production;
 - (i) by-products from fish from plants manufacturing fish products for human consumption;
 - (j) shells, hatchery by-products and cracked egg by-products originating from animals which did not show clinical signs of any disease communicable through that product to humans or animals.
- The signature and the stamp must be in a different colour to that of the printing.
- Note for the person responsible for the consignment in EU: this certificate is only for veterinary purposes and has to accompany the
 consignment until it reaches the border inspection post.

ı		 						
ı	Official veterinarian							
l	Name (in capitals):	Qualification and title:						
	Date:	Signature:						
	Stamp:							

Health certificate

For gelatine and collagen not intended for human consumption to be used as feed material or for technical purposes, intended for dispatch to or for transit through (²) the European Community

COUN	TRY				Veterinary certificate to EU
	l.1.	Consignor Name	1.2.	Certificate reference number	1.2.a.
		Address	I.3. Central Competent Authority		
ment		Tel. No	1.4.	Local Competent Authority	,
igi	1.5.	Consignee	1.6.	Person responsible for the	consignment in EU
š		Name		Name	
pa (Address		Address	
atch		Postal code		Postal code	
disp		Tel. No	ŀ	Tel. No	
Part I: Details of dispatched consignment	1.7.	Country of ISO code I.8.	1.9.	Country of ISO code destination	e I.10.
: De	l.11.	Place of origin	l.12.	Place of destination	
art				Custom wareho	use 🗌
		Name Approval number		Name	Approval number
		Address	}	Address	
				Postal code	
	l.13.	Place of loading	1.14.	Date of departure	
	1.15.	Means of transport	1.16.	Entry BIP in EU	
		Aeroplane Ship Railway wagon		•	
		Road vehicle Other	L		
		Identification: Documentary references:	1.17.		
	l.18.	Description of commodity		I.19. Commod	dity code (HS code)
				1.2	20. Quantity
	1.21.	Temperature of product		1.2	22. Number of packages
		Ambient Chilled Chilled		Frozen 🔲	
	1.23.	Identification of container/Seal number		1.2	24. Type of packaging
	1.25.	Commodities certified for:			
		Animal feedingstuff		Techr	nical use
	1.26.	For transit to third country vis-à-vis EU	1.27.	For import or admission int	o EU
		3rd country ISO code			
	1.28.	Identification of the commodities			
		Approval number of Species Nature of commodity Manufacturin		nments Number of	Net weight Batch number
	(Sci	entific name)	ig plant	packages	Not weight Dator number

COUNTRY

Gelatine and collagen not intended for human consumption to be used as feed material or for technical purposes

			II.a.	Certificate reference number	II.b.					
	II.	Health attestation	<u></u>							
		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (1) and certif the gelatine/collagen (2) described above:								
	N.1.	consists of gelatine/collagen (²) that satisfy the health requirements below;								
ation	II.2.	II.2. consist exclusively of gelatine/collagen (²) not intended for human consumption;								
ertifica	II.2. consist exclusively of gelatine/collagen (²) not intended for human consumption; II.3. has been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance in Article 17 and where appropriate Article 11 of Regulation (EC) No 1774/2002, in order to kill pathogenic agents; II.4. has been prepared exclusively with the following animal by-products: [— parts of slaughtered animals, which are fit for human consumption in accordance with Community legislation, intended for human consumption for commercial reasons.]									
: C	II.4.	has been prepared	exclusively with	the following animal by-products:						
Part	(²) either			 which are fit for human consumption in a ption for commercial reasons,] 	accordance with Community legislation, but are not					
	(²) and/or	of diseases		to humans or animals and derive from	n consumption but are not affected by any signs carcasses that are fit for human consumption in					
	(²) and/or	slaughterh	ouse, after under		inating from animals that were slaughtered in a fit, as a result of such inspection, for slaughter for					
	(²) and/or	[— animal by-ş	products derived	from the production of products intended	for human consumption,]					
	(²) and/or [— former foodstuffs of animal origin, or former foodstuffs containing products of animal origin, other than catering which are no longer intended for human consumption for commercial reasons or due to problems of manufact packaging defects or other defects which do not present any risk to humans or animals.]									
	(²) and/or	[fish or othe	r sea animals, ex	cept sea mammals, caught in the open se	ea for the purposes of fishmeal production,]					
	(²) and/or	[fresh by-pr	oducts from fish	rom plants manufacturing fish products fo	or human consumption;]					
	II.5.	the gelatine/collage	en (²):							
					nygiene conditions, and in particular wrapping and rmitted under Community legislation were used.					
			and packages c ONSUMPTION",		ords "GELATINE/COLLAGEN (2) SUITABLE FOR					
	(²) either [(b) in the case of gelatine, has been produced by a process that ensuring that unprocessed Category 3 material is to a treatment with acid or alkali, followed by one or more rinses, involving pH adjustment, extraction by heat several times in succession, followed by purification by means of filtration and sterilisation, in order to kill pagents:]				olving pH adjustment, extraction by heating one or					
				ning, pH adjustment using acid or alkali folk	that unprocessed Category 3 material is subjected owed by one or more rinses, filtration and extrusion,					
	Notes									
	Part I:									
	_			ble for the consignment in EU: this box is sertificate is for import commodity.	is to be filled in only if it is a certificate for transit					
	Halas			tion: this box is to be filled in only if it is a les, free warehouses and custom wareho	a certificate for transit commodity. The products in uses.					
	amenda			umber (railway wagons or container and of unloading and reloading.	d lorries), flight number (aircraft) or name (ship);					
	_	Box reference I.19:	use the appropri	ate HS code: 35.03 or 35.04.						
	_	Box reference I.23:	for bulk contained	ers, the container number and the seal number	mber (if applicable) should be included.					
		Box reference I.26	and I.27: fill in ac	cording to whether it is a transit or an imp	ort certificate.					
	-	Box reference I.28:		odity: select gelatine or collagen. lant: provide the registration number of tre	eatment/processing establishment.					
	Manufacturing plant: provide the registration number of treatment/processing establishment.									

Part II:							
(¹)	OJ L 273, 10.10.2002, p. 1.	OJ L 273, 10.10.2002, p. 1.					
(2)	Delete as appropriate.						
(3)		Catering waste means all waste food, including used cooking oils, originating in restaurants, catering facilities and kitchens, including central kitchens and household kitchens.					
	The signature and the stamp must be in a diffe	erent colour to that of the printing.					
_	Note for the person responsible for the consig the consignment until it reaches the border ins	nment in EU: this certificate is only for veterinary purposes and has to accompany spection post.					
Official	veterinarian						
	Name (in capitals):	Qualification and title:					
	Date:	Signature:					
	Stamp:						

Health certificate

For hydrolysed protein, dicalcium phosphate and tricalcium phosphate not intended for human consumption to be used as feed material or for technical purposes, intended for dispatch to or for transit through (²) the European Community

COUN	TRY	Veterinary certificate to EU
	I.1. Consignor Name	I.2. Certificate reference I.2.a. number
	Address	I.3. Central Competent Authority
nent	Tel. No	I.4. Local Competent Authority
sign	I.5. Consignee	I.6. Person responsible for the consignment in EU
CO	Name Address	Name Address
hed	Address	Address
patc	Postal code	Postal code
fdis	Tel. No	Tel. No
Part I: Details of dispatched consignment	I.7. Country of ISO code I.8. origin	I.9. Country of ISO code I.10. destination
ιο	I.11. Place of origin	I.12. Place of destination
Part		Custom warehouse
1	Name Approval number Address	Name Approval number Address
	Address	Address
		Postal code
	I.13. Place of loading	I.14. Date of departure
	I.15. Means of transport	I.16. Entry BIP in EU
	Aeroplane Ship Railway wagon Road vehicle Other	4
	Identification:	1.17.
	Documentary references:	
	I.18. Description of commodity	I.19. Commodity code (HS code)
		I.20. Quantity
	I.21. Temperature of product	I.22. Number of packages
	Ambient Chilled	Frozen
	I.23. Identification of container/Seal number	I.24. Type of packaging
	I.25. Commodities certified for: Animal feedingstuff	Technical use
	1.26. For transit to third country vis-à-vis EU	1.27. For import or admission into EU
	1.26. For transit to tilito country vis-a-vis EO	1.27. For import of admission into EU
	3rd country ISO code	
	I.28. Identification of the commodities	
	Approval number of es	stablishments
	Species Nature of commodity Manufacturing (Scientific name)	plant Number of packages Net weight Batch number

COUNTRY

Hydrolysed protein, dicalcium phosphate and tricalcium phosphate to be used as feed material or for technical purposes

							or for technical purposes		
				II.a.	Certificate refe	erence number	II.b.		
	II.	Health att	estation						
		I, the unde	ersigned official	veterinarian.	. declare that I have	e read and understoo	d Regulation (EC) No 1774/2002 (1) and certify that		
						nosphate (2) describe			
ication	II.1.	consists of	f hydrolysed pr	otein/dicalciu	um phosphate/trical	cium phosphate (2) th	hat satisfy the health requirements below;		
Part II: Certification	II.2.	consists ex	consists exclusively of hydrolysed protein/dicalcium phosphate/tricalcium phosphate (²) not intended for human consumption;						
Part II:	II.3.						ed by the competent authority in accordance with , in order to kill pathogenic agents;		
	II.4.	has been p	prepared exclu	sively with th	e following animal	by-products:			
	(²) either				which are fit for huit tion for commercial		accordance with Community legislation, but are not		
	(²) and/or	of		municable to	o humans or anim		n consumption but are not affected by any signs carcasses that are fit for human consumption in		
	(²) and/or	sla	aughterhouse,	after underg		spection, and were	inating from animals that were slaughtered in a fit, as a result of such inspection, for slaughter for		
	(²) and/or	mo		on, and were			ered in a slaughterhouse, after undergoing ante- aughter for human consumption in accordance with		
	(²) and/or	[an	nimal by-produc	cts derived fr	om the production	of products intended	for human consumption;]		
	(²) and/or	wh	nich are no lon	ger intended	for human consur		ducts of animal origin, other than catering waste (3), al reasons or due to problems of manufacturing or umans or animals;]		
	(²) and/or		w milk originat ımans or anima		mals that do not sh	ow clinical signs of a	any disease communicable through that product to		
	(²) and/or	[— fis	h or other sea	animals, exc	ept sea mammals,	caught in the open s	ea for the purposes of fishmeal production;]		
	(2) and/or	[— fre	sh by-product	s from fish fro	om plants manufac	turing fish products fo	or human consumption;]		
	(²) and/or					y-products originating to humans or anima	g from animals which did not show clinical signs of ls;]		
	II.5.	5. the hydrolysed protein/dicalcium phosphate/tricalcium phosphate (²):							
		sto	ored and trans	ported under	satisfactory hygier	ne conditions, and in	ating "NOT FOR HUMAN CONSUMPTION" and particular wrapping and packaging took place in a legislation were used; and		
	(²) either		the case of I			produced by a proc	ess involving appropriate measures to minimise		
		pro	ocessing plant	dedicated or	nly to hydrolysed p		minants hides and skins, has been produced in a sing a process involving the preparation of the raw I by:		
		(i)					an 3 hours at temperature of more than 80 °C and ninutes at more than 3,6 bar; and		
		(ii)		e of the mate nutes at 3 ba		, followed by a pH of	more than 11, followed by heat treatment at 140 °C		

(2) or	[(b)	in the	case of dicalcium phosphate, has been produced by a process that:		
		(i)	ensures that all Category 3 bonematerial is finely crushed and degreased with hot water and treated with dilute hydrochloric acid (at a minimum concentration of 4 % and a pH of less than 1,5) over a period of at least two days;		
		(ii)	followed by treatment of the obtained phosphoric liquor with lime, resulting in a precipitate of dicalcium phosphate at pH 4 to 7; and		
		(iii)	finally air-dries this precipitate for 15 minutes, with inlet temperature of 270 °C to 325 °C and end temperature between 60 °C and 65 °C;]		
(2) or	[(b)	in the	case of tricalcium phosphate, has been produced by a process ensuring:		
		(i)	that all Category 3 bone-material is finely crushed and degreased in counter-flow with hot water (bone chips less than 14 mm);		
		(ii)	continuous cooking with steam at 145 °C during 30 minutes at 4 bars;		
		(iii)	separation of the protein broth from the hydroxyapatite (tricalcium phosphate) by centrifugation; and		
		(iv)	granulation of the tricalcium phosphate after drying in a fluid bed with air at 200 °C.]		
Notes Part I:					
_			1.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for transit may be filled in if the certificate is for import commodity.		
_	Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses.				
	Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading.				
	Box reference I.19: use the appropriate HS code: 28.35 or 35.04.				
_	Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.				
_	Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.				
_	Box re	eference	1.28: Nature of commodity: specify if hydrolysed protein, dicalcium phosphate or tricalcium phosphate.		
			Manufacturing plant: provide the registration number of treatment/processing establishment.		
Part II:					
(¹)	OJ L	273, 10.1	0.2002, p. 1.		
(2)		e as app	·		
(3)			e means all waste food, including used cooking oils, originating in restaurants, catering facilities and kitchens, al kitchens and household kitchens.		
_	The s	ignature	and the stamp must be in a different colour to that of the printing.		
			erson responsible for the consignment in EU: this certificate is only for veterinary purposes and has to accompany ent until it reaches the border inspection post.		
Official veter	inarian				
	Name	e (in capi	tals):		
			Qualification and title:		
	Date:				
			Signature:		
	Stam	p:			

Health certificate

For apiculture by-products, intended for dispatch to or for transit through (2) the European Community

COUN	TRY	Veterinary certificate to EU
	I.1. Consignor Name	I.2. Certificate reference number I.2.a.
	Address	I.3. Central Competent Authority
Part I: Details of dispatched consignment	Tel. No	I.4. Local Competent Authority
ign	I.5. Consignee	I.6. Person responsible for the consignment in EU
ous	Name	Name
pa	Address	Address
atch	Postal code	Postal code
lisp	Tel. No	Tel. No
ofe	I.7. Country of ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10. Region of Code
tails	origin	destination destination
De	I.11. Place of origin	I.12. Place of destination
ᇤ		Custom warehouse
٦	Name Approval number	Name Approval number
	Address	Address
		Postal code
	I.13. Place of loading	I.14. Date of departure
	I.15. Means of transport	I.16. Entry BIP in EU
	Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐	
	Identification:	1.17.
	Documentary references:	1.17.
	I.18. Description of commodity	I.19. Commodity code (HS code)
		I.20. Quantity
	I.21. Temperature of product	I.22. Number of packages
	Ambient Chilled	Frozen
	I.23. Identification of container/Seal number	I.24. Type of packaging
	I.25. Commodities certified for:	
	Animal feedingstuff	Technical use Other Other
	I.26. For transit to third country vis-à-vis EU	I.27. For import or admission into EU
	3rd country ISO code	
	I.28. Identification of the commodities	
		Approval number of establishments
	Species Nature of commodity (Scientific name)	Manufacturing plant Net weight
	•	

COUN	ITRY					Apiculture by-products
			II.a.	Certificate referen	ce number	II.b.
	11.	Health attestation	L			
	11.		torinarian	declare that I have re	ad and understood	Regulation (EC) No 1774/2002 (1) and in particular
		Article 6 and Annex VIII Cha				
g.	II.1.	come from an area where the	a discasas	mentioned helow are	officially notifiable a	and which is not subject to any restrictions associated
Part II: Certification		with:	, uiscases	mendoned below are	micially notiliable a	and which is not subject to any restrictions associated
ific		(a) American foul brood (F	Paenibacillu	is larvae);		
Seri		(b) Acariosis (Acarapis wo	odi (Renni	e));		
i		(c) Small hive beetle (Aeth	nina tumida); and		
art		(d) Tropilaelaps mites (Tro	pilaelaps s	spp);		
"						
	II.2.	have been				
	(2) either	[subjected to a temperature	of – 12 °C (or lower for at least 24	hours:]	
	(2) or	[in the case of wax refined o			,•	
	, ,					
	Notes					
	Part I:					
	l arei.	D			-11.46.5.6	
	-	it may be filled in if the certifi			EU; this dox is to be	e filled in only if it is a certificate for transit commodity;
	 Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in tra can only be stored in free zones, free warehouses and custom warehouses. 					
		Pov reference I 15: Pogistral	tion numbo	r (railway wagana ar s	antainar and larria	a) flight number (sivereft) or name (ship); information
	-	is to be provided in the even			ontainer and iorne	s), flight number (aircraft) or name (ship); information
		Day reference I 10, use the		UC and at 05 11 00 a	ad an asifutha ass	modify as listed under note Day reference I 20
	 Box reference I.19: use the appropriate HS code: 05.11.99 and specify the commodity as listed under note Box reference I.28. 					
		Box reference I.23: for bulk	containers,	the container number	and the seal num	ber (if applicable) should be given.
		Box reference I.26 and I.27:	fill in accor	ding to whether it is a	transit or an impor	rt certificate.
				_	•	
		Box reference I.28: Nature o	f commodi	ty: means honey, bee	swax, royal jelly, pi	ropolis or pollen used in bee-keeping.
	Part II:	O I 272 10 10 2002 5 1				
	(¹) (²)	OJ L 273, 10.10.2002, p. 1. Delete as appropriate.				
		The signature and the stamp	n must be i	a different colour to	hat of the printing	
	_	- '				ly for veterinary purposes and has to accompany the
		consignment until it reaches				
	Official ve	eterinarian				
		Name (in capitals):			Qua	lification and title:
		Date:			Sinr	nature:
					Sigi	
		Stamp:				

CHAPTER 14(A)

Health certificate

For fat derivatives not intended for human consumption to be used for technical purposes, intended for dispatch to or for transit through (²) the European Community

COUN	TRY				Veterinary certificate to EU
	l.1.	Consignor Name	1.2.	Certificate reference number	1.2.a.
		Address	1.3.	Central Competent Authori	ity
nent		Tel. No	1.4.	Local Competent Authority	,
sign	1.5.	Consignee	1.6.	Person responsible for the	consignment in EU
COU		Name		Name	
ped		Address		Address	
atc		Postal code		Postal code	
dis		Tel. No		Tel. No	
Part I: Details of dispatched consignment	1.7.	Country of ISO code I.8. Region of origin Code origin	1.9.	Country of ISO code destination	1.10.
i. De	l.11.	Place of origin	1.12.	Place of destination	
Part				Custom wareho	_
-		Name Approval number		Name	Approval number
		Address		Address	
				Postal code	
	1.13.	Place of loading	1.14.	Date of departure	
	1.15.	Means of transport	1.16.	Entry BIP in EU	
		Aeroplane Ship Railway wagon			
		Road vehicle Other I	1.17.		
		Documentary references:	1.17.		
	l.18.	Description of commodity	<u></u>	I.19. Commod	dity code (HS code) 15.16.10
					I.20. Quantity
	1.21.	Temperature of product			I.22. Number of packages
		Ambient Chilled Chilled		Frozen	
	1.23.	Identification of container/Seal number			I.24. Type of packaging
	1.25.	Commodities certified for:		L	
		Technical use			
	1.26.	For transit to third country vis-à-vis EU	1.27.	For import or admission into	o EU
		3rd country ISO code			
	1.28.	Identification of the commodities	-		
		Approval number of esta	blishme	ents	
		Species Nature of commodity Manufacturing pla		Number of package	s Net weight Batch number
	(Scie	entific name)			

COUNTRY

For fat derivatives to be used for technical purposes

II.a. Certificate reference number II.b.

II. Health attestation

I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (1) and certify that the fat derivatives described above:

- II.1. consist of fat derivatives that satisfy the health requirements below;
- II.2. consist of fat derivatives containing exclusively fat derivatives not intended for human nor animal consumption;
- II.3. have been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance with Article 14 and where appropriate Article 11 of Regulation (EC) No 1774/2002, in order to kill pathogenic agents;
- II.4. have been prepared from rendered fats exclusively produced from Category 2 and/or Category 3 materials (3);
- II.5. the fat derivatives produced from Category 2 materials:
 - (a) have been produced using the following methods:
 - (²) either [transesterification or hydrolysis at least 200 °C, under corresponding appropriate pressure, for 20 minutes (glycerol, fatty acids and esters); and]
 - (2) or [saponification with NaOH 12 M (glycerol and soap):
 - $(^2)\ either\ [in\ a\ batch\ process\ at\ 95\ ^\circ C$ for three hours; and]
 - (2) or [in a continuous process at 140 °C, 2 bars (2 000 hPa) for eight minutes; and]]
 - (b) are packaged in new containers or in containers that have been cleaned, and all precautions are taken to prevent its contamination which bear labels indicating "NOT FOR HUMAN OR ANIMAL COSUMPTION".

Notes

Part I

- Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for transit commodity;
 it may be filled in if the certificate is for import commodity.
- Box reference I.12: Place of destination; this box is to be filled in only if it is a certificate for transit commodity. The products in transit
 can only be stored in free zones, free warehouses and custom warehouses.
- -- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: Manufacturing plant: provide the registration number of treatment/processing establishment.

Part II:

- (1) OJ L 273, 10.10.2002, p. 1.
- (2) Delete as appropriate.
- (3) List of Category 2 materials:
 - (a) all animal materials collected when treating waste water from slaughterhouses other than slaughterhouses covered by Article 4(1)(d) or from Category 2 processing plants, including screenings, materials from de-sanding, grease and oil mixtures, sludge and materials removed from drains from those premises;
 - (b) products of animal origin containing residues of veterinary drugs and contaminants listed in Group B (1) and (2) of Annex I to Directive 96/23/EC, if such residues exceed the permitted level laid down by Community legislation;
 - (c) products of animal origin, other than Category 1 material, that are imported from third countries and, in the course of the inspections provided for in Community legislation, fail to comply with the veterinary requirements for their importation into the Community, unless they are returned or their importation is accepted under restrictions laid down under Community legislation;
 - (d) animals and parts of animals, other than those referred to in Article 4, that die other than by being slaughtered for human consumption, including animals killed to eradicate an epizootic disease;
 - (e) mixtures of Category 2 material with Category 3 material, including any material destined for processing in a Category 2 processing plant; and
 - (f) animal by-products other than Category 1 material or Category 3 material.

Part II: Certification

_	The signature and the stamp must be in a different colour to that of the pri Note for the person responsible for the consignment in EU: this certificate consignment until it reaches the border inspection post.	· ·				
Official ve	Official veterinarian					
	Name (in capitals):	Qualification and title:				
	Date:	Signature:				
:	Stamp:					

CHAPTER 14(B)

Health certificate

For fat derivatives not intended for human consumption to be used as feed or for technical purposes, intended for dispatch to or for transit through (²) the European Community

COUN	TRY	Veterinary certificate to EU
	I.1. Consignor Name	I.2. Certificate reference number I.2.a.
	Address	I.3. Central Competent Authority
ment	Tel. No	I.4. Local Competent Authority
sign	I.5. Consignee	I.6. Person responsible for the consignment in EU
con	Name Address	Name Address
hed	/ Mulicas	Address
spate	Postal code	Postal code
of dis	Tel. No I.7. Country of ISO code I.8. Region of origin Code	Tel. No le I.9. Country of ISO code I.10.
Part I: Details of dispatched consignment	I.7. Country of ISO code I.8. Region of origin Coo	le I.9. Country of ISO code I.10. destination
ŭ	I.11. Place of origin	I.12. Place of destination
Part		Custom warehouse
	Name Approval number Address	Name Approval number Address
	, adriese	Addioso
		Postal code
	I.13. Place of loading	I.14. Date of departure
	I.15. Means of transport	I.16. Entry BIP in EU
	Aeroplane Ship Railway wagon Road vehicle Other	-
	Identification:	1.17.
	Documentary references:	
	I.18. Description of commodity	I.19. Commodity code (HS code) 15.16.10
		I.20. Quantity
	I.21. Temperature of product	I.22. Number of packages
	Ambient Chilled	· -
	I.23. Identification of container/Seal number	I.24. Type of packaging
	I.25. Commodities certified for:	7 Table 1
	Animal feedingstuff	
	I.26. For transit to third country vis-à-vis EU	I.27. For import or admission into EU
	ISO code 3rd country	
	I.28. Identification of the commodities	
	Approval number of e	stablishments
	Species Nature of commodity Manufacturing (Scientific name)	plant Number of packages Net weight Batch number

COUNTRY

Fat derivatives to be used as feed or for technical purposes

		II.a. Certificate reference number II.b.						
	II.	Health attestation I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (¹) and certify that						
		the fat derivatives described above:						
	II.1.	consist of fat derivatives that satisfy the health requirements below;						
tion	II.2.	consist of fat derivatives containing exclusively fat derivatives not intended for human consumption;						
Part II: Certification	II.3.	have been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance with Article 14 and where appropriate Article 11 of Regulation (EC) No 1774/2002, in order to kill pathogenic agents;						
Ö	11.4.	have been prepared from rendered fats exclusively produced from the following Category 3 materials:						
Part I	(²) either	 parts of slaughtered animals, which are fit for human consumption in accordance with Community legislation, but are not intended for human consumption for commercial reasons.] 						
	(²) and/or	[— parts of slaughtered animals, which are rejected as unfit for human consumption but are not affected by any signs of diseases communicable to humans or animals and derive from carcasses that are fit for human consumption in accordance with Community legislation,]						
	(²) and/or	[— hides and skins, hooves and horns, pig bristles and feathers originating from animals that were slaughtered in a slaughterhouse, after undergoing ante-mortem inspection, and were fit, as a result of such inspection, for slaughter for human consumption in accordance with Community legislation,]						
	(²) and/or	[— blood obtained from animals other than ruminants that are slaughtered in a slaughterhouse, after undergoing ante-mortem inspection, and were fit, as a result of such inspection, for slaughter for human consumption in accordance with Community legislation.]						
	(²) and/or	 animal by-products derived from the production of products intended for human consumption, including degreased bones and greaves,] 						
	(²) and/or	[— former foodstuffs of animal origin, or former foodstuffs containing products of animal origin, other than catering waste (3), which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects which do not present any risk to humans or animals,]						
	(²) and/or	[— milk originating from animals which do not show any clinical signs of any disease communicable through that product to humans or animals.]						
	(²) and/or	[— fish or other sea animals, except sea mammals, caught in the open sea for the purposes of fishmeal production,]						
	(²) and/or	r [by-products from fish from plants manufacturing fish products for human consumption,]						
	(²) and/or	 shells, hatchery by-products and cracked egg by-products originating from animals which did not show clinical signs of any disease communicable through that product to humans or animals; 						
	II.5.	are packaged in new containers or in containers which bear labels indicating "NOT FOR HUMAN CONSUMPTION", that have been cleaned, and all precautions are taken to prevent its contamination.						
	Notes Part I:							
	_	Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity.						
		Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses.						
	17.75.46	Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading.						
	_	Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.						
		Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.						
	_	Box reference I.28: Manufacturing plant: provide the registration number of treatment/processing establishment.						

Part II:			
(1)	OJ L 273, 10.10.2002, p. 1.		
(2)	Delete as appropriate.		
(3)	Catering waste means all waste food, including us including central kitchens and household kitchens.	ed cooking oils, originating in restaurants, catering facilities and kitchens,	
	The signature and the stamp must be in a different of	olour to that of the printing.	
	Note for the person responsible for the consignment in EU: this certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.		
Official ve	terinarian		
	Name (in capitals):	Qualification and title:	
	Date:	Signature:	
	Stamp:		
1			

Health certificate

For egg products not intended for human consumption that could be used as feed material, intended for dispatch to or for transit through (3) the European Community

COUN	TRY	Veterinary certificate to EU
	I.1. Consignor Name	I.2. Certificate reference number I.2.a.
	Address	I.3. Central Competent Authority
ment	Tel. No	I.4. Local Competent Authority
sign	I.5. Consignee	I.6. Person responsible for the consignment in EU
l con	Name Address	Name Address
tchec		
lispa	Postal code Tel. No	Postal code Tel. No
Part I: Details of dispatched consignment	I.7. Country of ISO code I.8. Region of origin Co origin	de I.9. Country of ISO code I.10. destination
: De	I.11. Place of origin	I.12. Place of destination
Part	Name - America I america	Custom warehouse
	Name Approval number Address	Name Approval number Address
	I.13. Place of loading	Postal code I.14. Date of departure
	<u> </u>	·
	I.15. Means of transport Aeroplane Ship Railway wagon	I.16. Entry BIP in EU
	Road vehicle Other	
	Identification: Documentary references:	1.17.
	I.18. Description of commodity	I.19. Commodity code (HS code) 35.02
		I.20. Quantity
	I.21. Temperature of product	I.22. Number of packages
	Ambient Chilled	Frozen 🗌
	I.23. Identification of container/Seal number	I.24. Type of packaging
	I.25. Commodities certified for: Animal feedingstuff	
	I.26. For transit to third country vis-à-vis EU	I.27. For import or admission into EU
	3rd country ISO code	
	I.28. Identification of the commodities	-
	Approval number of	establishments
	Species Nature of commodity Manufacturii (Scientific name)	ng plant Number of packages Net weight Batch number

COUNTRY

Egg products not intended for human consumption that could be used as feed material

		II.a. Certificate reference number II.b.						
	II.	Health attestation						
		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (¹) and certify that the egg products described above:						
	II.1.	consist of egg products that satisfy the health requirements below;						
tior	II.2.	consist exclusively of egg products not intended for human consumption;						
Part II: Certification	н.з.	have been prepared and stored in a plant, approved, validated and supervised by the competent authority in accordance with Article 17 and where appropriate Article 11 of Regulation (EC) No 1774/2002 or Council Directive 89/437/EEC (2), in order to kill pathogenic agents;						
	11.4.	have been prepared (derived) exclusively with the following animal by-product:						
Par		 eggs originating from animals which did not show clinical signs of any disease communicable through that product to humans or animals; 						
	II.5.	have been subjected to processing:						
	(3) either	[in accordance with processing method (4) as set out in Annex V, Chapter III of Regulation (EC) No 1774/2002;]						
	(3) or	[in accordance to a method and parameters which ensure that the products complies with the microbiological standards set in Chapter I, paragraph 10 of Annex VII to Regulation (EC) No 1774/2002]						
	(3) or	[treated in accordance with Chapter V of the Annex to Council Directive 89/437/EC]						
	II.6.	have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply with the following standards (5):						
		Salmonella: absence in 25 g: $n = 5$, $c = 0$, $m = 0$, $M = 0$;						
		Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram;						
	II.7.	meet Community standards on residues of substances that are harmful or might alter the organoleptic characteristics of the product or make its use as feed dangerous or harmful to animal health;						
	II.8.	the end product was:						
	(3) either [packed in new or sterilized bags;]							
	(3) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfect approved by the competent authority before use,]							
	and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION';							
	II.9.	the end product was stored in enclosed storage;						
	II.10.	the product has undergone all precautions to avoid contamination with pathogenic agents after treatment.						
	Notes							
	Part I:							
		Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity.						
		Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses.						
		Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading.						
		Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.						
	_	Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.						
	L							

Part II: OJ L 273, 10.10.2002, p. 1. (1) (²) OJ L 212, 22.07.1989, p. 89. (3) Delete as appropriate. (4) Insert method 1 to 5 or 7 as applicable. (5) n = number of samples to be tested; m = threshold value for the number of bacteria; the result is considered satisfactory if the number of bacteria in all samples does M = maximum value for the number of bacteria; the result is considered unsatisfactory if the number of bacteria in one or more samples is M or more; and c = number of samples the bacterial count of which may be between m and M, the sample still being considered acceptable if the bacterial count of the other samples is m or less. The signature and the stamp must be in a different colour to that of the printing. Note for the person responsible for the consignment in EU: this certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post. Official veterinarian Qualification and title: Name (in capitals): Date: Signature: Stamp:

Model Declaration

Declaration by the importer of bones and bone products (excluding bone meal), horns and horn products (excluding horn meal) and hooves and hoof products (excluding hoof meal) intended for use other than as feed material, organic fertilisers or soil improvers for dispatch to the European Communities

Note for the importer: This declaration is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.

f, the undersigned, declare that the following products (1):

- (a) bones and bone products (excluding bone meal);
- (b) horns and horn products (excluding horn meal);
- (c) hooves and hoof products (excluding hoof meal);

are intended to be imported by me into the Community, and I declare that these products will not be diverted at any stage for any use in food, feed material, organic fertilizers or soil improvers and will be conveyed directly for the purpose of further processing or treatment to:

Name:	Address:
The importer:	
Name:	Address:
Done at	on
(place)	(date)
Signature	
Regulation (EC) No 136/2004:	eterinary entry document (CVED) provided for in Annex III to Commission
Official stamp of the border inspection post of entry	into the EC (²)
	ial veterinarian of the border inspection post) (²)
Name:	(Name in capital letters)

⁽¹⁾ Delete as appropriate.

The signature and the stamp must be in a different colour to that of the printing.

Health certificate

For processed manure and processed manure products intended for dispatch to or for transit through (²) the European Community

COUN	COUNTRY Veterinary certificate to EU						
	l.1.	Consignor Name	1.2.	Certificate reference number	1.2.a.		
		Address	1.3.	Central Competent Author	ity		
nent		Tel. No	1.4.	Local Competent Authority	,		
ignr	1.5.	Consignee	1.6.	Person responsible for the	consignment in EU		
Suo		Name		Name			
ped		Address		Address			
atch		Postal code		Postal code			
disp		Tel. No		Tel. No			
Part I: Details of dispatched consignment	1.7.	Country of ISO code I.8. Region of origin Code origin	1.9.	Country of ISO code destination	e I.10. Region of Code destination		
. De	l.11.	Place of origin	1.12.	Place of destination	•		
art				Custom wareho	use		
"		Name Approval number		Name	Approval number		
		Address		Address			
				Postal code			
	I.13.	Place of loading	1.14.	Date of departure			
	l.15.	Means of transport	1.16.	Entry BIP in EU			
		Aeroplane Ship Railway wagon					
	Road vehicle Other I		1.17.				
		Documentary references:					
	I.18. Description of commodity		I.19. Commodity code (HS code)				
					I.20. Quantity		
	I.21. Temperature of product			_	I.22. Number of packages		
		Ambient Chilled		Frozen			
	I.23. Identification of container/Seal number				I.24. Type of packaging		
	I.25. Commodities certified for:						
		Animal feedingstuff Technical use		Other 🗌			
	1.26.	For transit to third country vis-à-vis EU	1.27.	For import or admission int	to EU		
		3rd country ISO code					
	I.28. Identification of the commodities						
	Species Nature of commodity (Scientific name)		Approval number of establishments Manufacturing plant Net weight				

COUNTRY Processed manure and processed manure products II.a. Certificate reference number II.b. II. Health attestation I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (1) and in particular Article 5 and Annex VIII Chapter VI thereof and certify that the processed manure or processed manure products described above: come from a technical plant, a biogas plant or a composting plant approved by the competent authority of the third country meeting the special conditions laid down in Regulation (EC) No 1774/2002; 11.1. Part II: Certification II.2. (2) have been subjected to: [a heat treatment process of at least 70 °C for at least 60 minutes]; or [an equivalent treatment validated and authorized by the importing Member State in accordance with the specific conditions laid down in Regulation (EC) No 1774/2002 as follows: II.3. are: (a) free from Salmonella (no salmonella in 25 g treated product); (b) free from Escherichia coli or from enterobacteriacea (based on the aerobic count: less than 1 000 cfu per gram of treated product); and (c) have been subjected to reduction in spore-forming bacteria and toxic formation; 11.4. are securely enclosed in: (a) well-sealed and insulated containers; or (b) properly sealed packs (plastic bags or big bags). Notes

Part I:

- Box reference I.6: Person responsible for the consignment in EU: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in the event of unloading and reloading.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be given.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: Nature of commodity: enter if processed manure or if processed manure products.

Part II:

- OJ L 273, 10.10.2002, p. 1. (¹)
- (²) Delete as appropriate.
- The signature and the stamp must be in a different colour to that of the printing.
- Note for the person responsible for the consignment in EU: This certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.

	Official veterinarian				
	Name (in capitals):	Qualification and title:			
	Date:	Signature:'.			
	Stamp:				
L					

- (6) Annex XI is amended as follows:
 - (a) the title of Part VI is replaced by the following:

'PART VI

List of third countries from which Member States may authorise imports of animal by-products and blood products (with the exception of blood products of equidae) intended for technical purposes including pharmaceuticals (health certificates Chapters 4 (C) and 8);

(b) in Part VII, the title and point A. are replaced by the following:

'PART VII

List of third countries from which Member States may authorise imports of animal by-products for the manufacture of processed petfood (health certificate Chapters 3(B) and 3(F))

A. Animal by-products from bovine, ovine, caprine, porcine and equine animals, including farmed and wild animals:

Third countries or parts of third countries listed in Part 1 of Annex II to Council Decision 79/542/EEC, from which imports of that category of fresh meat of the respective species is authorised and the following countries for the by-products specified:

Southern America and Southern Africa countries or parts thereof where matured and de-boned meat of the corresponding species is authorised, concerning matured and de-boned meat (including diaphragm) and/or matured trimmed offal of bovine, caprine, ovine animals and game (wild or farmed).';

(c) Part IX is replaced by the following:

'PART IX

List of third countries from which Member States may authorise imports of processed manure and processed manure products for the treatment of soil (Health Certificate Chapter 17)

For processed manure and processed manure products, third countries listed in:

- (a) Part 1 of Annex II to Decision 79/542/EEC;
- (b) Annex I to Commission Decision 2004/211/EC; or
- (c) Annex II to Commission Decision 2006/696/EC (OJ L 295, 25.10.2006, p. 1).';
- (d) Part XIII is replaced by the following:

'PART XIII

List of third countries from which Member States may authorise imports of serum of equidae (Health Certificate Chapter 4(A))

Third countries or parts of third countries listed in Annex I to Commission Decision 2004/211/EC (8), from which the importation of equidae for breeding and production is allowed.'.