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(Acts adopted under the EC Treaty/Euratom Treaty whose publication is obligatory)

REGULATIONS

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 by-products 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.

⁽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).

- (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 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.
- Regulation (EC) No 1774/2002 provides that certain ani-(11)mal by-products 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.

⁽¹⁾ 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).

- (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 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 by-products 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.

Article 2

For a transitional period of six months from the date of entry into force of this Regulation, the commercial document set out in Annex II to Regulation (EC) No 1774/2002 and the model health certificates set out in Annex X to that Regulation, which where completed in conformity with the provisions applicable before the date of entry into force of the present Regulation, may continue to accompany the products covered by such documents or certificates.

⁽¹) 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.

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.

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

Done at Brussels, 28 June 2007.

For the Commission Markos KYPRIANOU Member of the Commission

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

- 1. 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 eartag 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.
- (i) 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)

FORO	PEAN	COMMUNITY						Comme	rciai docume	; I I L
	l.1.	Consignor Name			1.2.	Document number	reference	I.2.a. Local refere	nce number:	
		Address			1.3.	Central co	mpetent authorit	ty		
pə		Postal code			1.4.	Local com	petent authority			
sent	1.5.	Consignee			1.6.					_
pres		Name								
ent_		Address								
ignme		Postal code			1.7.					_
ons	1.8.	Country of ISO code	I.9. Region of orig	in Code	1 10	Country of	f ISO code	e I.11. Region of	Cod	40
Part I : Details of consignment presented	1.0.	origin	1.3. Region of ong		1. 10.	destination		destination		10
etail	I.12	. Place of origin			I.13.	Place of de	estination			_
ä		Esta	ablishment 🗌				Establishm	nent 🔲	Other]
핕		Name	Approval num	ber		Name		Approval num	ber	_
9		Address				Address				
		Postal code				Postal cod	de			
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	1.40	Postal code			<u> </u>					_
	l.16.	. Means of transport Aeroplane	☐ Ship ☐		1.17.	Transporte Name	er	Approval number		
		Road vehicle	= '=	way wagon 🔲		Address		Approvai name.		
		Identification:		vay wagon		Postal cod	le	Member State		
	1.18	. Description of commodity					I.19. Commor	dity code (CN code)		
						[
								I.20. Number/quanti	ty	
	l.21.	. Temperature of products	Ambient	Chilled		F	rozen 🔲	I.22. Number of pac	kages	
	1.23.	. Identification of container/Se	eal number					I.24. Type of packag	ging	
	1.25	. Commodities certified for								_
		Animal feedingstuff	Tec	chnical use			Other			
	1.26	. Transit through third country	,		1.27.	Transit thro	ough Member St	ates		_
			SO code			Member St		ISO code		
		•	ode IP unit No:			Member St Member St		ISO code ISO code		
	1.28	. Export			1.29.					_
		Third country IS	SO code							
		Exit point C	ode							
	1.30.									
	I.31.	Identification of the commod	ities							
						Appı	roval number of	establishments		
		Species Nature of	commodity Categ	jory Treatr	nent typ		Manufacturin		atch number	
	(Sc	cientific name)								

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.

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 signature must be in a different colour to that of the printing

	Signature Done aton						
	(date)	(place)					
		(signature of the responsible person/consignor)					
		(name, in capital letters)					
Declara	tion by the transporter						
	I, the undersigned, declare that:						
II.2.1.	in the case where the packaging is done by the transporter	r, the animal by-products and/or processed products are:					
(1) eithe	r [in sealed new packaging;]						
(¹) or		hicles or other means of transport that were clean and dry before use and					
11.2.2.	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 during transportation, 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	inting.					
_	Note for the transporters: This document must accompany the point of destination.	y the consignment (*) from the place of loading for dispatch until it reaches					
(*)		type, which may contain different Categories of animal by-products, coming mercial document conveyed by the same means of transport to the same					
Signatu	re						
Done at		on					
	(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 (*)
 - (*) OJ L 139, 30.4.2004, p. 55; corrected version (OJ L 226, 25.6.2004, p. 22).';
 - (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 (*).
 - (*) OJ L 18, 23.1.2003, p. 11.';
 - (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 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:
 - (i) 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:
 - '5. 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 by-products, 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.
 - 5. 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	IRY			Veterinary certificate	to EU		
	l.1.	Consignor Name	1.2.	Certificate reference I.2.a. number			
		Address	1.3.	Central Competent Authority			
nent		Tel. No	1.4.				
signr	1.5.	Consignee	1.6.	I.6. Person responsible for the consignment in EU			
cons		Name		Name			
peu		Address		Address			
Part I: Details of dispatched consignment		Postal code Tel. No		Postal code Tel. No			
o Jo	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	I.12.	2. Place of destination			
art I				Custom warehouse			
-		Name Approval number		Name Approval number			
		Address		Address			
				Postal code			
	I.13.	Place of loading	l.14.	Date of departure			
	l.15.	Means of transport	I.16.	6. Entry BIP in EU			
		Aeroplane Ship Railway wagon					
		Road vehicle Other I	147				
		Documentary references:	l.17.				
	I.18.	Description of commodity		I.19. Commodity code (HS code)			
				I.20. Quantity			
	1.04	Towns earth use of executed		122 Number of posterior			
	1.21.	Temperature of product Ambient Chilled		I.22. Number of packages 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		1 27	. For import or admission into EU			
	1.20.	For transit to third country vis-à-vis EU	1.27.	. For import of admission into EO			
		3rd country ISO code					
	1.28.	Identification of the commodities					
			establi	il number of lishments acturing plant Net weight Batch number			

	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 VII Chapter II thereof and certify that:

- II.1. the processed animal protein or product described above contains exclusively processed animal protein not intended for human consumption that:
 - (a) has 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 (²) and
 - (b) has been prepared exclusively with the following animal by-products:
 - (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 antemortem 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 [— 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;]

and

- (c) has been subjected to the following processing standard:
- (2) either [heating to a core temperature of more than 133 °C for at least 20 minutes without interruption at a pressure (absolute) of at least 3 bars produced by saturated steam, with a particle size prior to processing of not more than 50 millimetres;]
- (²) or [in the case of non-mammalian protein other than fishmeal, the processing method as set out in Annex V, Chapter III, of Regulation (EC) 1774/2002;]
- (²) or [in the case of porcine blood, the processing method......as set out in Annex V, Chapter III to Regulation (EC) No 1774/2002, where in case of method 7 a heat treatment of at least 80 °C has been applied throughout its substance;]
- II.2. the competent authority examined a random sample immediately prior to dispatch and found it to comply with 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 g;

- II.3. the end product:
 - (2) either [was packed in new or sterilised bags;]
 - (2) or [was 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,]

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 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 aris to be provided in the event of unloading and reloading.	nd 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		
<u> </u>	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)	Where:			
	n = number of samples to be tested;			
	m = threshold value for the number of bacteria; the result is considered not exceed m;	ed satisfactory if the number of bacteria in all samples does		
	M = maximum value for the number of bacteria; the result is conside samples is M or more; and	red unsatisfactory if the number of bacteria in one or more		
	c = number of samples the bacterial count of which may be between bacterial count of the other samples is m or less.	m and M, the sample still being considered acceptable if the		
_	Note for the person responsible for the consignment in EU: This certifithe consignment until it reaches the border inspection post.	cate is only for veterinary purposes and has to accompany		
_	The signature and the stamp must be in a different colour to that of the	printing.		
Official ve	eterinarian			
o moiai vo				
	Name (in capitals):	Qualification and title:		
	Date:	Signature:		
	Stamp:			

ı

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	IKI		veterinary certificate to E		
	l.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	1.5.	Consignee	I.6. Person responsible for the consignment in EU		
suc		Name	Name		
Ö		Address	Address		
che					
pate		Postal code	Postal code		
dis		Tel. No	Tel. No		
o of	1.7.	Country of ISO code I.8. Region of origin Code			
tails		origin	destination		
Def	l.11.	Place of origin	I.12. Place of destination		
"		•	Custom warehouse		
Ра		Name Approval number	Name Approval number		
		Address	Address		
		Address	Addiess		
			Postal code		
	I.13.	Place of loading	I.14. Date of departure		
	l.15.	Means of transport	I.16. Entry BIP in EU		
		Aeroplane Ship Railway wagon			
		Road vehicle Other			
		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 🔲		
	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	I.27. For import or admission into EU		
		3rd country ISO code			
	1.28.	Identification of the commodities	1		
		Annual mush or	of a stab liabour sate		
		Approval number of	oi establistiments		
		Species Nature of Manufacture (Scientific name) commodity	uring plant Number of Net weight Batch number packages		

		II.a. C	ertificate reference number	II.b.					
		II.a. C	Standate reference number	11.50					
	II.	Health attestation							
		that:		ood Regulation (EC) No 1774/2002 (1) and certify					
Part II: Certification	II.1.								
licat	II.2.	the milk and milk-based product referred to i	n this certificate:						
erti		(a) has been prepared from raw milk that of	has been prepared from raw milk that comes from animals:						
ا ≝		 not showing clinical signs of a dis 	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 no 	belonging to holdings that are not under official restriction due to footandmouth disease or rinderpest; and						
_		(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;							
	11.4.	the milk/milk-based product was packed:							
	(3) either	[in new containers;]							
	(3) or	or [in vehicles or bulk containers disinfected prior to loading using a product approved by the competent authority;]							
		and the containers are marked so as to indi HUMAN CONSUMPTION".	cate the nature of the milk/milk-ba	sed product and bear labels indicating "NOT FOR					
	Notes								
	Part I:								
	_	Box reference I.6: Person responsible for t commodity; it may be filled in if the certificate	_	s to be filled in only if it is a certificate for transit					
	_	Box reference I.12: Place of destination: thi transit can only be stored in free zones, free	•	a certificate for transit commodity. The products in ses.					
	_	Box reference I.15: Registration number (information is to be provided in case of unload		lorries), flight number (aircraft) or name (ship);					
	_	Box reference I.19: use the appropriate HS	code: 23.09.10; 23.09.90; 35.01; 35	5.02 or 35.04.					
	_	Box reference I.23: for bulk containers, the o	ontainer number and the seal num	ber (if applicable) should be included.					
	_	Box reference I.26 and I.27: fill in according	to whether it is a transit or an impo	rt certificate.					
	_	Box reference I.28: Manufacturing plant: pro	vide the registration number of trea	tion number of treatment or processing establishment.					
	Part II:								
	(¹)	OJ L 273, 10.10.2002, p. 1.							
	(2)	For completion if the authorisation to import	into the Community is restricted to	certain regions of the third country concerned.					
	(³)	Delete as appropriate.							
	_	The signature and the stamp must be in a di	fferent colour to that of the printing						
	_	Note for the person responsible for the const the consignment until it reaches the border i	9	nly for veterinary purposes and has to accompany					
İ	Official vete	rinarian							
		Name (in capitals):	Qua	alification and title:					
		Date:	Sign	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	TRY	Veterinary certificate to EL
	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
Part I: Details of dispatched consignment	I.5. Consignee Name	Person responsible for the consignment in EU Name
ped c	Address	Address
dispato	Postal code Tel. No	Postal code Tel. No
tails of	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 Aeroplane Ship Railway	I.16. Entry BIP in EU
	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 Ambient 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 Further process	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 establi	
	Species Nature of commodity Manufacturing plan (Scientific name)	t Number of packages Net weight Batch number

			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:						
	II.1.	the milk-based product re	ferred to in this	certificate:				
tion		(a) has been prepared f	rom raw milk t	hat comes from animals:				
ifica		(i) not showing cli	nical signs of a diseases that can be transmitted through the milk to humans or animals; and					
Cert		(ii) belonging to he	oldings that are not under official restriction due to foot-and-mouth disease or rinderpest;					
Part II: Certification			cess involving heating to (temperature) for (time), which ensured a negative reaction to the lowed by, in the case of dried milk or dried milk-based product, a drying process; and					
۵		(c) has undergone an a	cidification pro	cess whereby its pH has been ma	intained at less than 6 for at least one hour;			
	II.2.	every precaution was take	en to avoid con	tamination of the milk-based prod	uct after processing;			
	II.3.	the milk-based product wa	as packed:					
	(²) either	[in new containers;]						
	(²) or	[in vehicles or bulk contain	ners disinfecte	d prior to loading using a product	approved by the competent authority;]			
		and the containers are ma CONSUMPTION".	rked so as to ir	ndicate the nature of the milk-base	d product and bear labels indicating "NOT FOR HUMAN			
	Notes							
	Part I:							
	_			for the consignment in EU: this ticate is for import commodity.	pox is to be filled in only if it is a certificate for transit			
	_			: this box is to be filled in only if free warehouses and custom war	t is a certificate for transit commodity. The products in ehouses.			
	_	Box reference I.15: Reginformation is to be provided			and lorries), flight number (aircraft) or name (ship);			
	_	Box reference I.19: use th	e appropriate	HS code: 23.09.10; 23.09.90; 35.0	01; 35.02 or 35.04.			
	_	Box reference I.23: for bu	lk containers, t	he container number and the sea	number (if applicable) should be included.			
	_	Box reference I.26 and I.2	7: fill in accord	ling to whether it is a transit or an	import certificate.			
	_	Box reference I.28: Manuf	facturing plant:	provide the registration number of	of treatment or processing establishment.			
	Part II:							
	(¹)	OJ L 273, 10.10.2002, p.	1.					
	(2)	Delete as appropriate.						
	_			a different colour to that of the pri				
		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 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	TRY				Veterinary certificate to EU
	I.1.	Consignor	1.2. C	Certificate reference	1.2.a.
		Name	n	umber	
		Address			
			I.3. C	Central Competent Authority	У
Part I: Details of dispatched consignment		Tel. No		ocal Competent Authority	
ign	1.5.	Consignee	I.6. P	erson responsible for the c	onsignment in EU
ous		Name	N	ame	
9		Address	Α	ddress	
che					
pat		Postal code		ostal code	
dis		Tel. No	Te	el. No	
o e	1.7.	Country of ISO code I.8. Region of origin Code		Country of ISO code	1.10.
ails		origin	d	estination	
Def	l.11.	Place of origin	I.12. P	Place of destination	
÷				Custom warehou	sa 🗖
Pai		Name Assessed assessed assessed as			_
		Name Approval number		lame	Approval number
		Address	A	ddress	
			Р	ostal code	
	I 13	Place of loading	I 14 D	ate of departure	
	1.10.	Thuse of feating	1.11. D	vato or dopartaro	
	l.15.	Means of transport	I.16. E	intry BIP in EU	
		Aeroplane Ship Railway Railway Road yehicle Other wagon			
		Identification:	I.17.		
		Documentary references:			
	I.18.	Description of commodity		I.19. Commodi	ty code (HS code)
					I.20. Quantity
	I.21.	Temperature of product			I.22. Number of packages
		Ambient Chilled		Frozen	
		<u> </u>			
	I.23.	Identification of container/Seal number			I.24. Type of packaging
	1.25.	Commodities certified for:		•	
		Animal feedingstuff Further process		Technical use	Other
	1.26	For transit to third country vis-à-vis EU	127 E	or import or admission into	<u></u>
	1.20.	For transit to trind country vis-a-vis EO	1.27. F	or import or admission into	
		3rd country ISO code			
	1.28	Identification of the commodities			
		Approval number of establish	monte		
		Species Nature of commodity Manufacturing plant		umber of packages	Net weight Batch number
	' C	,	IN	ambor or packages	not worght Daton number
	(50	cientific name)			

			La Cortificato refer	onco numbor	Тиь			
			II.a. Certificate refer	ence number	II.b.			
	II.	Health attestation						
		I, the undersigned officithat:	ial veterinarian, declare that I hav	ve read and underst	good Regulation (EC) No 1774/2002 (1) and certify			
	II.1.	the milk/milk-based product referred to in this certificate:						
ion		(a) has been prepare	ed from raw milk that comes from a	animals:				
ficat		(i) not showing clinical signs of a diseases that can be transmitted through the milk to humans or animals; and						
Part II: Certification		(ii) belonging t	o holdings that are not under offici	noldings that are not under official restriction due to footandmouth disease or rinderpest; and				
: C		(b) has undergone:						
art	(²) either	[(i) a sterilisati	ion process whereby an Fc value e	equal to or greater th	at 3 is achieved;]			
_	(²) or	[(ii) an initial process involving heating to (temperature) for (time), which ensured a negative reaction to the phosphatase test, followed by a further process involving heating to (temperature) for (time), which ensured a negative reaction to the phosphatase test, followed, in the case of dried milk, or dried milk-based products, by a drying process;]						
	II.2.	every precaution was ta	ken to avoid contamination of the	milk/milk-based proc	duct after processing;			
	II.3.	the milk/milk-based prod	duct was packed:					
	(²) either	[in new containers;]						
	(²) or	[in vehicles or bulk conta	ainers disinfected prior to loading u	using a product appr	oved 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 "NO HUMAN CONSUMPTION".				ased product and bear labels indicating "NOT FOR			
	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 tran commodity; it may be filled in if the certificate is for import commodity. 						
	_		ce of destination: this box is to be d in free zones, free warehouses		a certificate for transit commodity. The products in uses.			
	_		egistration number (railway wagor vided in case of unloading and relo		d lorries), flight number (aircraft) or name (ship);			
	_	Box reference I.19: use	the appropriate HS code: 23.09.10	0; 23.09.90; 35.01; 3	5.02 or 35.04.			
	_	Box reference I.23: for b	oulk containers, the container numl	ber and the seal nun	nber (if applicable) should be included.			
	_	Box reference I.26 and I	I.27: fill in according to whether it is	7: fill in according to whether it is a transit or an import certificate.				
	_	Box reference I.28: Man	nufacturing plant: provide the regist	tration number of tre	atment or processing establishment.			
	Part II:							
	(¹)	OJ L 273, 10.10.2002, p	o. 1.					
	(2)	Delete as appropriate.						
	_	The signature and the s	tamp 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 veterinary purposes and has to accompan the consignment until it reaches the border inspection post.						
	Official vete	erinarian						
		Name (in capitals):		Qu	alification and title:			
		Date:		Sig	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	
	l.1.	Consignor Name	1.2.	Certificate reference number	1.2.a.	
		Address	1.3.	Central Competent Authorit	У	
ent		Tel. No	1.4.	Local Competent Authority		
Part I: Details of dispatched consignment	1.5.	Consignee	1.6.	I.6. Person responsible for the consignment in EU		
ü		Name		Name		
o pe		Address		Address		
tche						
spa		Postal code Tel. No		Postal code Tel. No		
of di	17		10		140 Parism of Code	
ls o	1.7.	Country of ISO code I.8. Region of origin Code origin	1.9.	Country of ISO code destination	I.10. Region of Code destination	
eta				100 to 200 1 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0		
<u>:</u>	l.11.	Place of origin	I.12.	Place of destination	_	
Part				Custom warehou	se	
-		Name Approval number		Name	Approval number	
		Address		Address		
				B		
				Postal code		
	l.13.	Place of loading	l.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.			
		Documentary references:				
	l.18.	Description of commodity		I.19. Commod	ity code (HS code) 23.09.10	
					I.20. Quantity	
	I.21.	Temperature of product		_	I.22. Number of packages	
		Ambient 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		1.27	_	<u></u>	
	1.20.	For transit to third country vis-à-vis EU	1.27.	For import or admission into	E0	
		3rd country ISO code				
	1.28.	Identification of the commodities				
		Approval principles of actablishment				
		Approval number of establishm	ienis	Not weight	Datah numbar	
		Species Manufacturing plant (Scientific name)		Net weight	Batch number	
		(Colonalio Hamo)				

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 pa Article 6 and Annex VIII Chapter II thereof and certify that the petfood described above:							
cation	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;						
rtifi	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,]						
Pai	(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/E 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 aft		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.

Part II:	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 pr	rinting.			
_	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:				

CHAPTER 3(B)

Health certificate

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

COUNTRY Veterinary certificate to EU 1.2. Certificate Consignor reference I.2.a. number Name Address 1.3. Central Competent Authority Part I: Details of dispatched consignment Tel. No 1.4. Local Competent Authority Consignee 1.6. Person responsible for the consignment in EU Name Name Address Address Postal code Postal code Tel. No Tel. No 1.7. Country of ISO code 1.8. Region of origin Code 1.9. Country of ISO code I.10. Region of Code destination destination origin I.11. Place of origin I.12. Place of destination Custom warehouse Name Approval number Name Approval number Address Address Postal code I.13. Place of loading I.14. Date of departure I.16. Entry BIP in EU I.15. Means of transport Ship 🗌 Railway wagon Aeroplane Other 🔲 Road vehicle Identification: I.17. Documentary references: I.18. Description of commodity I.19. Commodity code (HS code) 23.09.10 I.20. Quantity I.21. Temperature of product I.22. Number of packages Ambient Chilled Frozen I.24. Type of packaging I.23. Identification of container/Seal number I.25. Commodities certified for: Technical use Animal feedingstuff 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 **Species** Approval number of establishments Manufacturing plant Net weight Batch number (Scientific name)

COUR	NIKY	Proces	sed petrood other than canned petrood					
		II.a. Certificate reference number II.b.						
	II.	Health attestation						
		I, the undersigned official veterinarian, declare that I have read and understood Regulation Article 6 and Annex VIII Chapter II thereof and certify that the petfood described above:	(EC) No 1774/2002 (¹) and in particular					
ation	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;						
ertific	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,]						
<u> </u>	(²) and/or	d/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 the underwent ante-mortem inspection and were fit, as a result of such inspection, for sl legislation,] 						
	(²) and/or [— blood obtained from animals other than ruminants that were slaughtered in a slaughterhouse, underwent ante-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 b greaves,]							
	origin, other than catering waste, which roblems of manufacturing or packaging							
	(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 ha	d been					
		(a) in the case of meat or meat products subjected to a heat treatment of at least 90 °C to	hroughout 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 submitted to a pasteurisation treatment sufficient to produce a negative phosph	. ,					

- 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 2004/438/EC, submitted to a sterilisation process or a double heat treatment where each treatment was sufficient to produce a negative 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-and-mouth 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,

or

- 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;
- (i) 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 (4); 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;

	(n) in the case of tricalcium phosphate produced by a process that ensures	5		
	 that all Category 3 bone-material is finely crushed and degreased 14 mm); 	d in counter-flow with hot water (bone chips less than		
	(ii) continuous cooking with steam at 145 °C during 30 minutes at 4 b	par;		
	(iii) separation of the protein broth from the hydroxyapatite (tricalciuur	m phosphate) by centrifugation; and		
	(iv) granulation of the tricalcium phosphate after drying in a fluid bed	with air at 200 °C];		
11.4.	was analysed by a random sampling of at least five samples from each processing plant and complies with the following standards (5):	processed batch taken during or after storage at the		
	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.	has undergone all precautions to avoid contamination with pathogenic agent	s after treatment;		
II.6.	was packed in new packaging, which, if the petfood is not dispatched in ready-to-sale packages on which it is clearly indicated the content is destined for feeding to pets only, bear labels indicating "NOT FOR HUMAN CONSUMPTION".			
Notes				
Part I:	Box reference I.6: Person responsible for the consignment in EU: this box	v is to be filled in only if it is a cortificate for transit		
_	commodity; it may be filled in if the certificate is for import commodity.	k 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 can only be stored in free zones, free warehouses and custom warehouses.	certificate for transit commodity. The products in transit		
_	Box reference I.15: Registration number (railway wagons or container and lorr is to be provided in the event of unloading and reloading.	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.			
Part II:				
(¹)	OJ L 273, 10.10.2002, p. 1.			
(²)	Delete as appropriate.			
(3)	OJ L 139, 30.4.2004, p. 55. Corrected by OJ L 226, 25.6.2004, p. 22.			
(4)	OJ L 226, 25.6.2004, p. 22.			
(⁵)	Where:			
	n = number of samples to be tested;			
	m = threshold value for the number of bacteria; the result is considered sat not exceed m;	isfactory if the number of bacteria in all samples does		
	M = maximum value for the number of bacteria; the result is considered un samples is M or more; and	nsatisfactory if the number of bacteria in one or more		
	c = number of samples the bacterial count of which may be between m and bacterial count of the other samples is m or less.	M, the sample still being considered acceptable if the		
_	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 the consignment until it reaches the border inspection post.	•		
Official ve	veterinarian			
	Name (in capitals):	Qualification and title:		
	Date: S	ignature:		
	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	ΕU	
	l.1.	Consignor Name	1.2.	Certificate reference I.2.a. number	_	
		Address	1.0	Combani Commentant Authority		
↓			1.3.	Central Competent Authority		
Part I: Details of dispatched consignment		Tel. No	1.4.	Local Competent Authority		
sigr	I.5.	Consignee	1.6.	I.6. Person responsible for the consignment in EU		
cou		Name		Name		
eq		Address		Address		
atch		Postal code		Postal code		
isp		Tel. No		Tel. No		
of d	17	Country of ISO code I.8. Region of origin Code	1.9.	Country of ISO code I.10. Region of Cod		
iis		origin	1.0.	destination destination	10	
Deta	1 11	Place of origin	112	Place of destination	-	
‡	1.11.	Flace of origin	1.12.			
Par				Custom warehouse		
		Name Approval number		Name Approval number		
		Address		Address		
				Postal code		
	l.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.			
		Documentary references:				
	l.18.	Description of commodity	I.19. Commodity code (HS code) 42.05.00			
				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		
	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 EU		
		3rd country ISO code				
	1.28.	Identification of the commodities			_	
		Approval number of	of estab	blishments		
		Species Manufactu (Scientific name)	ring pla	ant Net weight Batch number	r	
		(Scientific Harrie)				
					ļ	

COUN	IIKI					Dogcnews	
			II.a.	Certificate reference number	•	II.b.	
		L					
	II.	Health attestation					
I, the undersigned official veterinarian, declare that I have read and understood Regu Article 6 and Annex VIII Chapter II thereof and certify that the dogchews described at						• ', '	
ation	II.1.	have been prepared and sto where appropriate Article 11			the com	petent authority in accordance with Article 18 and	
ifica	II.2.	have been prepared exclusiv	ely with th	e following animal by-products:			
Part II: Certification	(²) either			ich were fit for human consump for commercial reasons,]	tion in ac	ecordance with Community legislation, but are not	
(²) and/or [— parts of slaughtered animals, which were rejected as unfit for human consumption but are diseases communicable to humans or animals and derive from carcasses that were fit for huma with Community legislation,]							
	(²) and/or		0	animals that were slaughtered in ction, for slaughter in accordance	U	nterhouse, underwent ante-mortem inspection and immunity legislation,]	
	(²) and/or	[— animal by-products deri greaves,]	ved from t	he production of products intend	ed for hu	man consumption, including degreased bones and	
	(²) and/or	[— fresh by-products from	fish from p	plants manufacturing fish produc	ts for hun	nan consumption,]	
(²) and/or [— material from animals which have been treated with certain substances which are prohibited pur the import of the material being permitted in accordance with Article 28 of Regulation (EC) No 17							
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];					
(°) or [in the case of dogchews made from animal by-products other than hides and skins of ungulates or from fish, to at least 90 °C throughout their substance;]				ins of ungulates or from fish, to a heat treatment of			
	II.4. were examined by random sampling of at least five samples from each processed batch taken during or after storal processing plant and complies with the following standards (*):				essed batch taken during or after storage at the		
		Salmonella: absence	e in 25 g:	n = 5, $c = 0$, $m = 0$, $M = 0$;			
		Enterobacteriaceae: n = 5, c	= 2, m = 1	10, M = 300 in 1 gram;			
	II.5.	have undergone all precaution	ns to avoi	d contamination with pathogenic	agents a	after treatment;	
	II.6.	were packed in new packagii	ng.				
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 commodity; it may be filled in if the certificate is for import commodity. 				to be filled in only if it is a certificate for transit			
Box reference I.12: Place of destina can only be stored in free zones, free						ificate for transit commodity. The products in transit	
	_	_	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 of	ontainers,	the container number and the s	eal numb	per (if applicable) should be given.	
	_	Box reference I.26 and I.27:	fill in acco	rding to whether it is a transit or	an import	t 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 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 may bacterial count of the other samples is m or less.	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 ve	terinarian			
	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	IKI				veterinary certificate to EU
	l.1.	Consignor Name	1.2.	Certificate reference number	I.2.a.
		Address	1.3.	Central Competent Authority	/
nent		Tel. No	1.4.	Local Competent Authority	
signr	1.5.	Consignee	I.6. Person responsible for the consignment in EU		
ned con		Name Address		Name Address	
dispatch		Postal code Tel. No		Postal code 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 destination ISO code	I.10. Region of Code destination
l: De	l.11.	Place of origin	I.12.	Place of destination	_
Part		Name Amazonal assertan		Custom warehous	_
		Name Approval number Address		Name Address	Approval number
	1.40	Diagonal loading	144	Postal code	
	1.13.	Place of loading	1.14.	Date of departure	
	I.15.	Means of transport Aeroplane Ship Railway	I.16.	Entry BIP in EU	
		Road vehicle Other wagon			
		Identification: Documentary references:	l.17.		
	I.18.	Description of commodity		I.19. Commodi	ty code (HS code)
					I.20. Quantity
					1.20. Quantity
ľ	I.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:		Tankai and una	Ott 11 🗖
}	1.26	Animal feedingstuff For transit to third country vis-à-vis EU	1 27	Technical use For import or admission into	Other _
	1.20.			To import of duminosion into	
		3rd country ISO code			
	1.28.	Identification of the commodities			
				of establishments	
		Species Nature of commodity Nature of commodity (Scientific name)	Manufac	cturing plant N	et weight Batch number

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 raw petfood 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 79/542/EEC (2) and provided the animals from which the meat is derived come from a territory or part of a territory (ISO code) as listed in that Decision which has been free of foot and mouth disease, rinderpest, 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 species),
 - and/or Commission Decision 2006/696/EC (³), and provided the animals from which the meat is derived come from a territory or part of a territory (ISO code) as listed in that Decision which has been free from Newcastle disease and Avian Influenza for the last 12 months,
 - (b) derived from animals that, 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 in the Decisions above for which the animals are susceptible; 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 (5) 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 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 diseases 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 Decisions 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" or "ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS NOT FOR HUMAN CONSUMPTION" and then in leak-proof and 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 18 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) and complies with the following standards (6):

Salmonella: absence in 25 g: n = 5, c = 0, m = 0, M = 0;

Enterobacteriacae: n = 5, c = 2, m = 10, M = 300 in 1 gram.

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 warehouse	• •		
_	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: 05.11.91; 05.11.99 or 23	3.09.90.		
_	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: select raw petfood or animal by-	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 the animal and public health and veterinary certification conditions, for important fresh meat.			
(3)	OJ L 295, 25.10.2006, p. 1.			
(4)	Commission Decision 2000/585/EC of 7 September 2000 laying dow certifications for import of wild and farmed game meat and rabbit meat fr 97/217/EC, 97/218/EC, 97/219/EC and 97/220/EC. OJ L 251, 6.10.2000,	om third countries and repealing Commission Decisions		
(5)	Council Directive 93/119/EC of 22 December 1993 on the protection of animals at the time of slaughter or killing. OJ L 340, 31.12.1993, p. 21.			
(6)	Where:			
	n = number of samples to be tested;			
	m = threshold value for the number of bacteria; the result is considered s 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	unsatisfactory if the number of bacteria in one or more		
	c = number of samples the bacterial count of which may be between m a 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 pri	nting.		
_	Note for the person responsible for the consignment in EU: This certificate the consignment until it reaches the border inspection post.	e is only for veterinary purposes and has to accompany		
Official veterinarian				
	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	TRY	Veterinary certificate to EU				
	I.1. Consignor	I.2. Certificate reference I.2.a. number				
	Name	Harribot				
	Address	I.3. Central Competent Authority				
Part I: Details of dispatched consignment	Tel. No	I.4. Local Competent Authority				
sign	I.5. Consignee	I.6. Person responsible for the consignment in EU				
ő	Name	Name				
ρģ	Address	Address				
tche						
spa	Postal code Tel. No	Postal code Tel. No				
fdi						
<u>s</u>	I.7. Country of ISO code I.8. Region of origin C	Code I.9. Country of ISO code I.10. Region of Code destination				
etai	Shight	dodanation dodanation				
<u></u>	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:	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				
	I.26. For transit to third country vis-à-vis EU	I.27. For import or admission into EU				
	3rd country ISO code					
	1.29 Identification of the commodition					
	I.28. Identification of the commodities					
		Approval number of establishments				
	Species Nature of commodity (Scientific name)	Manufacturing plant Net weight Batch number				
	(Scientific name)					

		II.a.	Certificate reference number	er II.b.		
11.	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 XIV thereof and certify that the flavouring innards products described above:					
II.1.	consist of animal by-products that satisfy the animal health requirement below;					
II.2.	have been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance w Article 18 and where appropriate Article 11 of Regulation (EC) No 1774/2002;					
II.3.	have been prepared including the following animal by-products which are exclusively:					
(²) either	[— parts of slaughtered animals, which were fit for human consumption in accordance with Community legislation, but are r intended for human consumption for commercial reasons,]					
II.2. II.3. (²) either (²) 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 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 Communi legislation,]					
(²) and/or	[— blood obtained from animals other than ruminants that were slaughtered in a slaughterhouse, underwent ante-morte 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 greaves,]					
(²) and/or	 former foodstuffs of animal origin, or former foodstuffs containing products of animal origin, other than catering waste, whi are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packagin 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 huma 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 a 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/E 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 th the following standards (3):		nt authority taking a random san	ple immediately prior to dispatch and found it to comply		
	Salmonella: a	bsence 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	r [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 disinfected approved by the competent authority before use;]					
	and which bear labels indicating "NOT FOR HUMAN CONSUMPTION";					
117	the end product was stored in enclosed storage;					
II.7.			itions to avoid contamination with			

- 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: 05.04 or 05.11.91.

_	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 res not exceed m;	sult 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	esult is considered unsatisfactory if the number of bacteria in one or more				
	c = number of samples the bacterial count of which management bacterial count of the other samples is m or less.	ay be between m and M, the sample still being considered acceptable if the				
_	The signature and the stamp must be in a different colo	our to that of the printing.				
_	Note for the person responsible for the consignment in the consignment until it reaches the border inspection p	EU: This certificate is only for veterinary purposes and has to accompany post.				
Official v	veterinarian					
	Name (in capitals):	Qualification and title:				
	Date:	Signature:				
	Stamp:					

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

COUNT	IRY				Veterinary certificate to EU
	I.1.	Consignor Name	1.2.	Certificate reference number	1.2.a.
		Address	1.3.	Central Competent Authorit	у
nent		Tel. No	1.4.	Local Competent Authority	
Part I: Details of dispatched consignment	I.5.	Consignee Name Address	I.6.	Person responsible for the on Name Address	consignment in EU
f dispatch		Postal code Tel. No		Postal code Tel. No	
tails of	1.7.	Country of ISO code I.8. Region of origin Code origin	1.9.	Country of ISO code destination	1.10.
art I: De	I.11.	Place of origin	I.12.	Place of destination Custom warehou	ise
		Name Approval number Address		Name Address	Approval number
				Postal code	
	I.13.	Place of loading	I.14.	Date of departure	
	I.15.	Means of transport Aeroplane Ship Railway Road vehicle Other wagon		Entry BIP in EU	
		Documentary references:	1.17.		
	I.18.	Description of commodity		I.19. Commod	ity code (HS code)
					I.20. Quantity
	I.21.	Temperature of product Ambient Chilled Chilled		Frozen 🗌	I.22. Number of packages
	1.23.	Identification of container/Seal number			I.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 Species Nature of commodity Manufacturing plar (Scientific name)		s Number of packages	Net weight Batch number

			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 that the animal by-products described above:							
	II.1.1.	consist of animal by-prod	ducts that satisfy the animal health requirements be	elow;				
ion	II.1.2. have been obtained in the territory of:							
ficat	(³) either	[(a) that have remained	I in this territory since birth or for at least the last th	ree months before slaughter;]				
Serti	II.1.2. have been obtained in the territory of:							
<u> </u>	II.1.3.	have been obtained from	n animals:					
Par	(³) either	[(a) coming from holding	gs:					
	(i) where, for the following diseases for which the animals are susceptible, there has been neither case/ou rinderpest, swine vesicular disease, Newcastle disease or highly pathogenic avian influenza during the prior nor of classical or African swine fever during the prior 40 days; nor in the holdings situated in their vicinity with during the prior 30 days; and							
		. ,	nas been neither case/outbreak of foot and mouth eir vicinity within 25 km, during the prior 30 days; at	disease during the prior 60 days, nor in the holdings and				
		(b) which:						
		(i) were not killed	d to eradicate any epizootic disease;					
			d in their holdings of origin for at least forty days beforerhouse without contact with other animals which o	ore departure and which have been transported directly did not comply with the same health conditions;				
(iii) at the slaughterhouse, have passed the ante-mortem health inspendave shown no evidence of the diseases referred to above for which								
			eated in the slaughterhouse before and at the time Council Directive 93/119/EC on animal welfare;]	of slaughter or killing in accordance with the relevant				
	(3) or	[(a) captured and killed	in the wild in an area:					
		susceptible: fo		of the following diseases for which the animals are ease or highly pathogenic avian influenza during the prior 40 days; and				
			ed at a distance that exceeds 20 km from the born is not authorised at these dates for exporting this	ders separating another territory of a country or part material to the European Community; and				
			vere transported within 12 hours for chilling either t nt, or directly to a game establishment;]	to a collection centre and immediately afterwards to a				
	II.1.4.	.4. have been obtained in an establishment around which, within a radius of 10 km, there has been no case/outbreak of diseas 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, t preparation of raw material for exportation to the European Community has been authorised only after removal of all meat, a 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	(3) either [— parts of slaughtered animals, which are fit for human consumption in accordance with Community legislation, but are intended for human consumption for commercial reasons,]						
	(³) and/or							
	(³) and/or [— animal by-products derived from the production of products intended for human consumption, including degreased be and greaves,]							

(3) 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,] (3) and/or fish or other sea animals, except sea mammals, caught in the open sea for the purposes of fishmeal production,] fresh by-products from fish from plants manufacturing fish products for human consumption,] (3) and/or (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.] raw material derived from animals which have been treated with certain substances prohibited in accordance with Directive (3) and/or 96/22/EC for the manufacture of petfood, as referred to in Article 28 of Regulation (EC) No 1774/2002;] II.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 they will not spoil between dispatch and delivery to the plant of destination; II.1.9. in the case of raw material derived from animals which have been treated with certain substances prohibited in accordance with Directive 96/22/EC for the manufacture of petfood, as referred to in Article 28 of Regulation (EC) No 1774/2002: it has been marked in the third country before entry into the territory of the Community by a cross of liquefied charcoal or activated carbon on each outer side of each frozen block in a way that the marking covers at least 70 % of the diagonal length of the frozen block and be of at least 10 cm width; in case of material which is not frozen, the raw material has been marked in the third country before entry into the territory of the Community by spraying it with liquefied charcoal or by applying charcoal powder in a way that the charcoal is clearly visible on the material; and in the case the animal by-products are made up of raw material which has been treated as referred to above and other nontreated raw material, all the raw materials have been marked as laid down in point (a) and (b) above. (3)(6) [II.2. Specific requirements The by-products in this consignment come from animals that have been kept in the territory mentioned under (II.1.2), where (3) (7) II.2.1. vaccination programmes against foot-and-mouth disease are being regularly carried out and officially controlled in domestic bovine animals. The by-products in this consignment consists only of animal by-products derived from trimmed offal of domestic ruminants, which have maturated at an ambient temperature of more than + 2 °C for at least three hours, or in the case of masseter muscles of (3) (8) II.2.2. bovine animals and de-boned meat of domestic animals, for at least 24 hours.] 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.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. Box reference I.28: Manufacturing plant: provide the veterinary control number of the approved establishment.

Part II:					
(*)	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 e	he name and ISO code number of the exporting country as laid down in:			
	 part 1 of Annex II of Council Decision 	n 79/542/EEC,			
	 the Annex to Commission Decision 9 	nnex to Commission Decision 94/984/EC, and			
	 the Annex to Commission Decision 2 	2000/585/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 into the European Community.	intended 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	ncluding used cooking oils, originating in restaurants, catering facilities and kitchens, kitchens.			
(⁶)	Supplementary guarantees to be provided when the material of domestic ruminants originated in the territory of a South American or South African country or part thereof from where only maturated and de-boned fresh meat of domestic ruminants for human consumption is permitted for exportation to the European Community. In the case of offal only trimmed offal of domestic ruminants which must be exclusively offal from which the bones, cartilage, trachea and main bronchi, lymphatic glands, adhering connective tissue, fat and mucus have been completely removed is permitted. The whole masseter muscles of bovine animals, incised in 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	S.			
(8)	Only for certain South American and Sou	th African countries.			
_	The signature and the stamp must be in a	a different colour to that of the printing.			
_	Note for the person responsible for the cothe consignment until it reaches the border	onsignment in EU: this certificate is only for veterinary purposes and has to accompany er inspection post.			
Official ve	eterinarian				
	Name (in capitals):	Qualification and title:			
	Date:	Signature:			
	Stamp:				

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	
	l.1.	Consignor Name	1.2.	Certificate reference number	1.2.a.	
		Address	1.3.	Central Competent Authorit	ty	
nent		Tel. No	1.4.	Local Competent Authority		
signı	1.5.	Consignee	1.6.	I.6. Person responsible for the consignment in EU		
COD		Name Address		Name Address		
hed		Address		Address		
Part I: Details of dispatched consignment		Postal code Tel. No		Postal code Tel. No		
s of	1.7.	Country of ISO code I.8. Region of origin Code	1.9.		1.10.	
etails		origin		destination		
Ξ	l.11.	Place of origin	1.12.	Place of destination	_	
Part				Custom warehou	_	
		Name Approval number Address		Name	Approval number	
		Address		Address		
				Postal code		
	l.13.	Place of loading	1.14.	Date of departure		
	l.15.	Means of transport	I.16.	Entry BIP in EU		
	1	Aeroplane Ship Railway wagon Road vehicle Other				
	1	Identification:	1.17.			
	1	Documentary references:				
	I.18.	Description of commodity		I.19. Commod	lity code (HS code) 30.02	
					I.20. Quantity	
	<u> </u>					
	I.21.	Temperature of product			I.22. Number of packages	
		Ambient Chilled		Frozen		
	1.23.	Identification of container/Seal number			I.24. Type of packaging	
	1.25.	Commodities certified for:	1			
	1.26	Technical use	1.07	For import or admission into	. Fu	
	1.26.	For transit to third country vis-à-vis EU	1.27.	For import or admission into	DEU	
		3rd country ISO code				
	1.28.	Identification of the commodities				
		Species (Scientific name)	Approv	al number of establishments Manufacturing plant		

Part II: Certification

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;
- II.2. consist exclusively of serum of equidae not intended for human nor animal consumption;
- 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;
- 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;
- 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) either [(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 p	rinting.			
_	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:				

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 EU
	I.1.	Consignor	1.2.	Certificate reference number	I.2.a.
		Name Address			
		, addiese	1.3.	Central Competent Authorit	у
Part I: Details of dispatched consignment		Tel. No	1.4.	Local Competent Authority	
sign	1.5.	Consignee	I.6.	Person responsible for the	consignment in EU
cou		Name		Name	
pei		Address		Address	
atch		Postal code		Postal code	
disp		Tel. No		Tel. No	
of	1.7.	Country of ISO code I.8. Region of origin Code	1.9.	Country of ISO code	1.10.
tails		origin		destination	
Det	l.11.	Place of origin	I.12.	Place of destination	
빌				Custom warehou	se \square
Pg		Name Approval number		Name	Approval number
		Address		Address	Trivi
				Postal code	
	I.13. Place of loading			Date of departure	
	I.15. Means of transport		I.16.	Entry BIP in EU	
		Aeroplane Ship Railway wagon			
		Road vehicle Other			
		Identification: Documentary references:	I.17.		
	<u> </u>				
	l.18.	Description of commodity		I.19. Commod	ity code (HS code)
					I.20. Quantity
	I.21.	Temperature of product			I.22. Number of packages
		Ambient Chilled		Frozen	
	1.23.	Identification of container/Seal number			I.24. Type of packaging
	1.20.	- Contained of Contained Court III.			Ti.24. Typo of paolaging
	1.25.	Commodities certified for:			
		Animal feedingstuff			
	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			
	1.20.	identification of the commodities			
				Approval number of establishment	
		Species Nature of commodity (Scientific name)		Manufacturing plan	t Batch number
		,			

						·			
			II.a.	Certificate refer	ence number	II.b.			
						L			
	11.	Health attestation							
		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (¹) and certify that the blood products described above:							
uo	II.1.	consist of blood products that satisfy the health requirements below;							
cati	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 wi Article 17 and where appropriate Article 11 of Regulation (EC) No 1774/2002;								
art II	II.4.	have been prepared (deriv	/ed) exclusiv	ely with the followin	g animal by-products	s:			
Pe	(²) either	[blood of slaughtered aning for human consumption for			sumption in accordar	nce with Community legislation, but is not intended			
	(²) and/or) 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 No 1774/2002;]	ance with p	processing method	(3) as set	out in Annex V, Chapter III of Regulation (EC)			
	(²) or	[to a method and param paragraph 10 of Annex VI				n the microbiological standards set in Chapter I,			
		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 the following standards (4):				ediately prior to dispatch and found it to comply with				
		Salmonella:	absence in :	25 g: n = 5, c = 0, m	i = 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 sterilise	d bags;]						
	(²) or	approved by the competer	nt authority b	pefore use,]		oughly cleaned and disinfected with a disinfectant			
		and which bear labels indi	cating "NOT	FOR HUMAN CON	ISUMPTION";				
	II.8.	the end product was store	d in enclose	d storage;					
	II.9.	the product has undergon	e all precaut	ions to avoid contar	nination with pathoge	enic 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 tran commodity; it may be filled in if the certificate is for import commodity. 				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 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. 				I lorries), flight number (aircraft) or name (ship);			
	_	Box reference I.19: use th	e appropriat	e HS code: 05.11.9	1 or 05.11.99.				
	_	Box reference I.23: for bul	k containers	s, the container num	ber and the seal num	nber (if applicable) should be included.			
	_	Box reference I.26 and I.2	7: fill in accc	ording to whether it i	s a transit or an impo	ort certificate.			

Part II:		
(¹)	OJ L 273, 10.10.2002, p. 1.	
(²)	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 bacteria; the result is con not exceed m;	sidered satisfactory if the number of bacteria in all samples does
	M = maximum value for the number of bacteria; the result is co samples is M or more; and	insidered unsatisfactory if the number of bacteria in one or more
	c = number of samples the bacterial count of which may be beto bacterial count of the other samples is m or less.	ween m 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 printing.
_	Note for the person responsible for the consignment in EU: this the consignment until it reaches the border inspection post.	certificate is only for veterinary purposes and has to accompany
Official ve	terinarian	
	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	ΓRY				Veterinary certificate to EU
	I.1. Coi Nai	nsignor	1.2.	Certificate reference number	1.2.a.
		dress	1.0	Cantral Cananatant Authorit	
			1.3.	Central Competent Authorit	у
Part I: Details of dispatched consignment	Tel	I. No	1.4.	Local Competent Authority	
sign	I.5. Coi	nsignee	I.6.	Person responsible for the o	consignment in EU
Sions	Nai			Name	
ed o	Add	dress		Address	
atch	Pos	stal code		Postal code	
isp		I. No		Tel. No	
of d	I.7. Coi	untry of ISO code I.8. Region of origin Code	1.9.	Country of ISO code	1.10.
ails	orig			destination	
Det	I 11 Pla	ace of origin	I 12	Place of destination	
± ±		ace of ongin		Custom warehou	se 🗖
Ра	Naı	me Approval number		Name	Approval number
		dress		Address	Approval named
				Postal code	
	I.13. Pla	ace of loading	I.14.	Date of departure	
	I.15. Me	eans of transport	I.16.	Entry BIP in EU	
		Aeroplane Ship Railway wagon			
	lde	Road vehicle Other characteristics:	1.47		
		cumentary references:	l.17.		
	I.18. Des	scription of commodity		I.19. Commod	ity code (HS code)
		•			30.02
					I.20. Quantity
	I.21. Ter	mperature of product		_	I.22. Number of packages
		Ambient Chilled Chilled		Frozen	
	I.23. Ide	entification of container/Seal number			I.24. Type of packaging
	1.25. Co	mmodities certified for:	ı		
		Technical use			
	I.26. For	r transit to third country vis-à-vis EU	1.27.	For import or admission into	EU
	3rd	d country ISO code			
	I.28. Ide	entification of the commodities			
				Approval number of establish	
		Species Nature of commodity (Scientific name)		Manufacturing plan	t Batch number
		,			

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			II.a. Certificate reference number	II.b.					
	II.	Health attestation							
		I, the undersigned office that the blood products	al veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (¹) and certify described above:						
	II.1.	consist of blood produc	s that satisfy the health requirements below;						
tion	II.2.	consist exclusively of b	od products not intended for human or animal consumption;						
ifica	II.3.	have been prepared ex	cclusively with the following animal by-products:						
Part II: Certification	(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,]							
Part	(²) and/or		humans or animals, derived from carcasses that are	it for human consumption but is not affected by any signs of diseases carcasses that are fit for human consumption in accordance with					
	(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	[— blood and blood p	roducts derived from the production of products intende	ed for human consumption,]					
	(²) and/or	[— blood and blood products originating from animals that did not show clinical signs of any disease communicable through that product to humans or animals;]							
(2) either [II.4. in the case of blood products derived from ruminant animals they originate				a third country or regions, where:					
		(2) either [the animals and products come from a region where no case of foot-and-mouth disease, vesicular st rinderpest, peste des petits ruminants, Rift Valley fever and bluetongue (3) has been recorded for 12 month which vaccination has not been carried out against those diseases for at least 12 months and from which of ruminant animals are authorized pursuant to Community legislation. The blood from which such prod manufactured must have been collected:							
		(2) either [in slaughte	rhouses approved in accordance with Community legis	slation;]					
		(2) or [from live as	nimals in facilities approved in accordance with Commo	unity legislation;]					
		Commissio	erhouses approved and supervised by the competer in and Member States must be notified of the address te shall indicate this information;]]						
			s have undergone one of the following treatments, guar ot-and-mouth disease, vesicular stomatitis, rinderpest (3):						
		(2) either [heat treatm	ent 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 effe	ectiveness check;]					
		(²) or [change in	pH to pH 5 for two hours, followed by an effectiveness	check;]					
		(²) or [heat treatm	ent of at least 90°C throughout their substance, follow	ed by an effectiveness check;]]					
			ve bluetongue animals are present, and the blood prodicals, in vitro diagnosis and laboratory reagents, to be p State] (4);]						
	(2) or [II.4.	in the case of blood pr	oducts derived from animals excluding ruminants they	originate in a third country or regions where:					
		classical sw for 12 mon for at least	and the products come from a region where no case o ine fever, African swine fever, Newcastle disease or highs highest in the susceptible species and in which vaccination 12 months. The health certificate shall follow the model cts are derived;	ghly pathogenic avian influenza has been recorded n has not been carried out against those diseases					
		effectivenes swine vesion	ts have undergone a heat treatment at a temperature is check, guaranteeing the absence of pathogens of ular disease, classical swine fever, African swine few the susceptible species;]]	the following diseases: foot-and-mouth disease,					

II.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 approved by the competent authority before use;]	thoroughly cleaned and disinfected with a disinfectant				
	and which bear labels indicating 'NOT FOR HUMAN OR ANIMAL CONSUMPTION';					
II.6.	the end product was stored in enclosed storage;					
II.7.	the product has undergone all precautions to avoid contamination with pa	athogenic agents after treatment.				
Notos						
Notes Part I:						
— —	Box reference I.6: Person responsible for the consignment in EU: this b	pox 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.	son to to so miss in only in the a commente for manon				
_	Box reference L12: Place of destination: this box is to be filled in only if i	t is a certificate for transit commodity. The products in				
	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 ir 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 1010100 1.20. 101 bulk containers, the container number and the sear number (ii 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:	OLL 272 40 40 2002 p. 4					
(¹) (²)	OJ L 273, 10.10.2002, p. 1. Delete as appropriate.					
(3)	In the case of countries in which bluetongue sero-positive ruminant anima	Is are present, blood products have been treated or the				
()	animals have been tested sero-negative.					
(4)	This must be the same Member State of first entry of the products into the	e Community.				
_	The signature and the stamp must be in a different colour to that of the pr	rinting.				
_	Note for the person responsible for the consignment in EU: this certificate the consignment until it reaches the border inspection post.	is only for veterinary purposes and has to accompany				
Official veter	narian					
	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	IKI				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.	I.4. Local Competent Authority				
signr	1.5.	Consignee	1.6.	Person responsible for the	consignment in EU			
cons		Name		Name Address				
hed		Address		Address				
Part I: Details of dispatched consignment		Postal code Tel. No		Postal code Tel. No				
s of	1.7.	Country of ISO code I.8. Region of origin Code	1.9.	Country of ISO code				
etail		origin		destination	destination			
i.	I.11.	Place of origin	1.12	Place of destination	-			
Parí		Approval number		Custom wareho				
		Name Approval number Address		Name Address	Approval number			
				Postal code				
	I.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.	No(s) of CITES				
		Documentary references:		<u>-</u>				
	I.18.	Description of commodity		I.19. Commod	dity 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			
	1.25.	Commodities certified for:						
	1.00	Animal feedingstuff			nical use Other Other			
	1.26.	For transit to third country vis-à-vis EU	1.27.	For import or admission int	0 EU			
		3rd country ISO code						
	1.28.	Identification of the commodities						
		Approval numb	er of es	stablishments				
		·	facturin	g plant	Net weight			
		(Scientific name)						

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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 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):
 - (a) were slaughtered and their carcasses are fit for human consumption in accordance with Community legislation; or
 - (b) 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;
- II.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 (3):
 - [— classical swine fever, and African swine fever,]
 - [— rinderpest,]

and

- (b) 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 (');
- II.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;]

II.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:							
(1)	OJ L 273, 10.10.2002, p. 1.						
(²)	Delete as appropriate.						
(3)	Delete diseases not applicable to the species concerned.						
_	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 ve	terinarian						
	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	IRY				Veterinary certificate to EU			
	l.1.	Consignor Name	1.2.	Certificate reference number	1.2.a.			
		Address	1.3.	Central Competent Authorit	у			
Part I: Details of dispatched consignment		Tel. No	1.4.					
ignr	1.5.	Consignee	1.6.	Person responsible for the o	consignment in EU			
Sons		Name		Name				
ed c		Address		Address				
tch		Postal code		Postal code				
ispa		Tel. No		Tel. No				
οf σ	1.7.	Country of ISO code I.8. Region of origin Code	1.9.		I.10. Region of Code			
ails		origin		destination	destination			
Det	l.11.	Place of origin	1.12.	Place of destination	1			
벁				Custom warehou	se \square			
9		Name Approval number		Name	Approval number			
		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 wagon Road vehicle Other						
		Identification:	1 17	No(s) of CITES				
		Documentary references:	1.17.	110(3) 01 011 20				
	I.18.	Description of commodity	•	I.19. Commod	ity code (HS code)			
					I.20. Quantity			
	I.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		Techni	cal 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			No.4			
		Species Manufa (Scientific name)	acturing	g piant	Net weight			
		,						

COU	NTRY					Treated hides and skins of Ungulates			
			II.a.	Certificate reference num	ber	II.b.			
	l _{II} .	Health attestation				L			
	".	I, the undersigned officia		n, declare that I have read and u thereof and certify that the hides		Regulation (EC) No 1774/2002 (¹) and in particular s described above:			
uo	II.1.	have been obtained from	n animals th	at (²):					
ficati		(a) were slaughtered	and their car	casses are fit for human consul	mption in a	accordance with Community legislation; or			
Part II: Certification	(b) 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								
Part		(c) did not show any epizootic disease;	clinical signs	s of any disease communicable	to human	s or animals, and were not killed to eradicate any			
	(²) either [II.2		ountry listed	in part 1 of Annex II to Decis		isation in accordance with Community legislation, 2/EEC (3) from which imports of fresh meat of the			
	(²) either	[dried;]							
	(²) or	[dry-salted or wet-salted	for at least	14 days prior to dispatch;]					
	(²) or		by ship and	the duration of transport will be		to the declaration of the transporter, the hides and they will have undergone a minimum of 14 days of			
	(²) or	[salted for seven days in	sea salt wit	h the addition of 2 % of sodium	carbonate	5]			
	(²) or	declaration of the transp	orter, the hi		d by ship	owing date and according to the and the duration of transport will be such that they border inspection post;]			
	(²) or [II.2.		isted in part	1 of Annex II to Decision 79/54		ion in accordance with Community legislation, from m which imports of fresh meat of the corresponding			
	(²) either	[salted for seven days in	sea salt wit	h the addition of 2 % of sodium	carbonate	5]			
	(²) or	declaration of the transp	orter, the hi		d by ship	wing date and according to the and the duration of transport will be such that they border inspection post;]			
	(2) or	[dried for 42 days at a te	mperature o	of at least 20 °C;]					
	II.3.	the consignment has no transmissible disease.	t been in co	ntact with other animal products	or with liv	ve animals presenting a risk of spreading a serious			
	Notes								
	Part I:	Part I:							
	_		and the second second	ble for the consignment in EU: certificate is for import commodi		is to be filled in only if it is a certificate for transit			
	-			ation: this box is to be filled in cases, free warehouses and custo	•	a certificate for transit commodity. The products in uses.			
	-			mber (railway wagons or conta event of unloading and reloading		orries), flight number (aircraft) or name (ship) and			
	-	Box reference I.19: use	the appropri	ate HS code: 41.01; 41.02 or 4	1.03.				
	-	Box reference I.23: for b	ulk containe	ers, the container number and th	e seal nur	nber (if applicable) should be given.			
	-	Box reference I.26 and	.27: fill in ac	cording to whether it is a transit	or an imp	ort 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.						
l –	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 vet	erinarian						
	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 (¹) 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 EL
	I.1.	Consignor Name	1.2.	Certificate reference number	1.2.a.
		Address	1.3.	Central Competent Authority	
ŧ		Tal Na		·	/
Part I: Details of dispatched consignment		Tel. No	1.4.	Local Competent Authority	
sigr	1.5.	Consignee	1.6.	Person responsible for the c	onsignment in EU
cor		Name Address		Name Address	
hed				, , , , , , , , , , , , , , , , , , , ,	
pato		Postal code		Postal code	
f dis	1.7	Tel. No	1.0	Tel. No	Lian Basis of Octo
ils o	1.7.	Country of ISO code I.8. Region of origin Code origin	1.9.	Country of ISO code destination	I.10. Region of Code destination
Deta	I 11	Place of origin	112	. Place of destination	
r. ::		That of origin	""	Custom warehous	se \square
Pa		Name Approval number		Name	Approval number
		Address		Address	
				Postal code	
	I.13.	Place of loading	1.14	. Date of departure	
			1 140	E 1 - DID : - EII	
	1.15	Means of transport Aeroplane Ship Railway wagon	1.16	. Entry BIP in EU	
		Road vehicle Other	1		
		Identification: Documentary references:	1.17	. No(s) of CITES	
	I.18.	Description of commodity		I.19. Commodi	ty 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		Techni	cal 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	<u> </u>		
		Approval numl	ner of ex	etahliehmente	
			ıfacturin		Net weight
		(Scientific name)			-
	l				

				II.a.	Certificate reference	enumber	II.b.			
	II.		Declaration I, the undersigned declare that the hides and skins described above:							
	II.1.									
	11.1.			, ,		onsumption in acc	cordance with Community legislation; or			
_		(α)	word diaugntored and	trion daroado	oo aro ne for naman o	onoumption in doc	ordanies was community regionation, or			
Part II: Certification			(b) 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							
II: Cert		(c) did not show any clinical signs of any disease communicable to humans or animals, and were not killed to eradicate a epizootic disease;								
ar	II.2.	have	been:							
ا ۾	(1) either	[dried	;]							
	(1) or	[dry-s	alted or wet-salted for	at least 14 da	ays prior to dispatch;]					
	(1) or	[salte	d for seven days in sea	a salt with the	addition of 2 % of so	dium carbonate;]				
	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	er [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	[11.4.	following the declara	ition of the tra	ansporter, the duratior	n of the transport p	period is foreseen to be at least 21 days.]			
	Notes									
	_		eference I.6: Person nodity; it may be filled i		-		to be filled in only if it is a certificate for transit			
	_		eference I.12: Place of nly be stored in free zo				ficate for transit commodity. The products in transit			
	_		eference I.15: Registra be provided in the ever			ntainer and lorries)	, flight number (aircraft) or name (ship); information			
	_	Box re	eference I.19: use the	appropriate F	IS code: 41.01; 41.02	or 41.03.				
	_	Box re	eference I.23: for bulk	containers, th	ne container number a	and the seal numb	er (if applicable) should be given.			
	_	Box re	eference I.26 and I.27:	fill in accordi	ing to whether it is a tr	ransit or an import	certificate.			
	Part II:									
	(¹)	Delete	e as appropriate.							
	_	The s	ignature and the stam	p must be in a	a different colour to th	at of the printing.				
	_		for the person respons onsignment until it read			is declaration is o	nly for veterinary purposes and has to accompany			
	Official vet	terinaria	n							
		Name	e (in capitals):			Qual	ification and title:			
		Date:				Signa	ature:			
		Stam	p:							

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.	COUNTRY Veterinary certificate to EU								
	I.1. Consignor Name		I.2. Certificate number	reference	I.2.a.				
	Address	-	I.3. Central Co	ompetent Authority					
ent	Tel. No	<u> </u>		npetent Authority	'				
u u	LE Consigned				anaignment in EU				
nsiç	I.5. Consignee Name	'	I.6. Person res Name	sponsible for the c	onsignment in EU				
8	Address		Address						
chec									
pate	Postal code		Postal cod	е					
fdis	Tel. No		Tel. No						
Part I: Details of dispatched consignment	I.7. Country of ISO code I.8. Region of origin	Code	I.9. Country of destination		1.10.				
: De	I.11. Place of origin	'	I.12. Place of d	estination					
art I				Custom warehous	se 🔲				
۵	Name Approval numbe	er	Name		Approval number				
	Address		Address						
			Postal cod	le					
	I.13. Place of loading		I.14. Date of de	parture					
	I.15. Means of transport		I.16. Entry BIP	in FU					
	Aeroplane Ship Railwa	ay wagon 🔲	1. TO. Entry Dir	20					
	Road vehicle Other								
	Identification: Documentary references:		I.17. No.(s) of 0	CITES					
	I.18. Description of commodity			I.19. Commodi	ty code (HS code)				
			l		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:			L	Oth or \square				
	I.26. For transit to third country vis-à-vis EU		127 Foriman	or admission into	Other _				
		_ '	1.27. For import	or admission into					
	3rd country ISO code								
	I.28. Identification of the commodities								
	Species (Scientific name)	Nature of comr	modity	Num	ber of packages				
	(Goleriulic Hairle)								

			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 certi that the game trophies described above:								
tion	II.1.	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;							
fica	(2) either [II.2	. in the case of game tr	ophies consi	sting solely of hides or skin:					
erti	(²) either	[have been dried;]							
<u>:</u>	(2) or	[have been dry-salted	or wet-salted	d for a minimum of 14 days before di	spatch;]				
Part II: Certification	(²) or		d the duration	n of the transport will be such that th	according to the declaration of the transporter, will be ey will have undergone a minimum of 14 days salting				
	(2) or [II.2.	in the case of game tr	ophies consi	sting solely of bone, horns, hooves, o	claws, antlers or teeth:				
				ng water for an appropriate time so th is removed; and	as to ensure that any matter other than bone, horns,				
		(b) have been disinf parts consisting			authority, in particular with hydrogen peroxide where				
	Notes								
	Part I:								
	_			ible for the consignment in EU: this 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 a certificate for transit commodity. The products 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 (shi information is to be provided in case of unloading and reloading.								
	_	Box reference I.23: fo	or bulk conta	iners, the container number and the	e seal number (if applicable) should be				
	_	Box reference I.19: us	se the approp	priate HS code: 05.05; 05.06; 05.07	or				
	_	Box reference I.26 ar	nd 1.27: fill in	according to whether it is a transi	t or an import				
	_	Box reference I.28: for [hooves], [claws], [ant			ore possibilities among the following: [bones], [horns],				
	Part II:								
	(¹)	OJ L 273, 10.10.2002	, p. 1.						
	(2)	Delete as appropriate							
	_	•		be in a different colour to that of the					
	 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. 								
	Official veteri	narian							
	1	Name (in capitals):		(Qualification and title:				
		Date:		9	Signature:				
		Stamp:							
ı									

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	IKI					Veterinary certificate to EU
	l.1.	Consignor Name	1.2.	Certificate referer number	nce	1.2.a.
		Address	1.3.	Central Competer	nt Authorit	у
nent		Tel. No	1.4.	Local Competent	Authority	
nsignr	1.5.	Consignee Name	I.6.	Person responsib	le for the o	consignment in EU
oo pa		Address		Address		
Part I: Details of dispatched consignment	 	Postal code Tel. No		Postal code Tel. No		
etails of	1.7.	Country of ISO code I.8. Region of origin Code origin	1.9.	Country of destination	ISO code	1.10.
: De	l.11.	. Place of origin	l.12.	Place of destination		
Part		Approval number			n warehou	_
		Name Approval number Address		Name Address		Approval number
				Postal code		
	l.13.	. Place of loading	1.14.	Date of departure)	
	I.15. Means of transport			I.16. Entry BIP in EU		
	Aeroplane Ship Railway wagon Road vehicle Other					
		Identification: Documentary references:	l.17.	I.17. No(s) of CITES		
	I.18.	. Description of commodity		I.19. Commodity code (HS code)		
						I.20. Quantity
	I.21.					I.22. Number of packages
	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 adm	nission into	EU
		3rd country ISO code				
	1.28.	Identification of the commodities				
		Species (Scientific name)	N	umber of package	es	

		II.a. Certificate reference number II.b.						
		II.a. Certificate reference number III.D.						
	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:						
ا ہ ا	(²) either [II.1.	II.1. with respect to game trophies of cloven-hoofed animals, excluding swine:						
fication		(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						
ertil		(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 count part of a third country not authorised to export untreated game trophies of wild swine to the Community;]							
	(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.						
	 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: 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.						
	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 colour to that of the pr	inting.			
_	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	inarian				
	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 Africans wine fever, intended for dispatch to or for transit through $(^2)$ the European Community

COUN	IRY				Veterinary certificate to EU
	l.1.	Consignor Name	1.2.	Certificate reference number	I.2.a.
		Address	1.3.	Central Competent Authori	ty
ment		Tel. No	1.4.	Local Competent Authority	
onsign	1.5.	Consignee Name	I.6.	Person responsible for the Name	consignment in EU
hed co		Address		Address	
Part I: Details of dispatched consignment		Postal code Tel. No		Postal code Tel. No	
tails of	1.7.	Country of ISO code I.8. Region of origin Code origin	1.9.	Country of destination ISO code	1.10.
t I: De	l.11.	Place of origin	I.12.	Place of destination	🗆
Par				Custom warehou	— I
		Name Approval number Address		Name Address	Approval number
				Postal code	
	I.13.	Place of loading	I.14.	Date of departure	
	l.15.	Means of transport	I.16.	Entry BIP in EU	
		Aeroplane Ship Railway wagon Road vehicle Other			
		Identification: Documentary references:	l.17.		
	I.18.	Description of commodity		I.19. Commod	lity code (HS code) 05.02
					I.20. Quantity
	I.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	I.27. For import or admission into EU		
		3rd country ISO code			
	1.28.	Identification of the commodities			
		Approval number of establishments			
		Manufacturing plant Numb	er of pa	ackages	Net weight

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	II.a.	Certificate reference number	II.b.
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II. Health attestation

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

- II.1. the pig bristles described above have been obtained from pigs originating, and slaughtered in a slaughterhouse, in the country of origin:
- 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;
- 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;
- II.4. the pig bristles are dry and securely enclosed 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.
- 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.

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:		Date:	Signature:					
		Stamp:						

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	IRY		Veterinary certificate to
	l.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
nsign	1.5.	Consignee Name	Person responsible for the consignment in EU Name
oo pər		Address	Address
Part I: Details of dispatched consignment		Postal code Tel. No	Postal code Tel. No
tails of	l.7.	Country of ISO code I.8. Region of origin Code origin	I.9. Country of destination ISO code I.10.
i: De	l.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 wagon	I.16. Entry BIP in EU
		Road vehicle Other	
		Identification: Documentary references:	1.17.
	1.18.	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
	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	I.27. For import or admission into EU
		3rd country ISO code	
	1.28.	Identification of the commodities	,
		Approval number of establishments	
	Manufacturing plant Nu		umber of packages Net weight

ı			II.a.	Certificate refe	rence number	II.b.		
ı								
ı	II.	Health attestation						
I, the undersigned official veterinarian, declare that I have read and u						tood Regulation (EC) No 1774/2002 (¹) and cer		
l		that:						
II.1. the pig bristles described above have been obtained from pigs originating, and slaughtered in a slaughterhouse, in origin; II.2. the pigs from which the pig bristles have been obtained did not show during inspection, carried out at the time of signs of diseases communicable to humans or animals and were not killed to eradicate any epizootic disease; II.3. the pig bristles mentioned above have been:								
١		origin;						
١		2 the pige from which the pig briefles have been obtained did not show during inspection, carried out at the time of eleventoring						
١	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;						
ı								
١	II.3.	the pig bristles mentioned above have been:						
ı	(²) either (²) or	[boiled;] [dyed;]						
ı	(²) or	[bleached;]						
١								
١	11.4.	the pig bristles are dry a	nd securely er	nclosed in packagir	ng.			
١								
╛								
	Notes							
	Part I:	Pov reference LG: Derec	n roonanaihl	o for the consignm	ant in Elle this have	is to be filled in only if it is a continuete for trav		
	_	commodity; it may be fill				is to be filled in only if it is a certificate for trar		
	_	Box reference I.12: Plac transit can only be stored			•	a certificate for transit commodity. The products ouses.		
	_	Box reference I.15: Reginformation is to be provi				d lorries), flight number (aircraft) or name (shi		
	_	Box reference I.23: for b	ulk containers	, the container nun	nber and the seal nu	mber (if applicable) should be included.		
	_	Box reference I.26 and I	.27: fill in accc	ording to whether it	is a transit or an imp	port certificate.		
	_	Box reference I.28: Man	ufacturing plar	nt: provide the vete	rinary control numbe	er of the registered establishment.		
	Dort II.							
	Part II:	OJ L 273, 10.10.2002, p	. 1.					
	(2)	Delete as appropriate.						
1	_	The signature and the st	amp must be	in a different colou	r to that of the printir	ng.		
	_			-		only for veterinary purposes and has to accompa		
the consignment until it reaches the border inspection post.								
	Official veterinarian							
	Official veter				0	ualification and title:		
-	Official veter	Name (in capitals):			Q	dallication and title.		
-	Official veter	Name (in capitals):						
-	Official veter					ignature:		

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	IKI				veterinary certificate to EU	
	l.1.	Consignor Name	l.2.	Certificate reference number	1.2.a.	
		Address	1.3.	Central Competent Autho	rity	
ment		Tel. No	1.4.	Local Competent Authorit	у	
sign	I.5.	Consignee	1.6.	I.6. Person responsible for the consignment in EU		
Part I: Details of dispatched consignment		Name Address		Name Address		
tchec						
lispa		Postal code Tel. No		Postal code Tel. No		
s of c	1.7.	Country of ISO code I.8. Region of origin Code	1.9.	Country of ISO coo	le I.10.	
etail		origin		destination		
٦ : D	l.11.	Place of origin	l.12.	Place of destination Custom wareho	nuse \square	
Pal		Name Approval number		Name	Address	
		Address		Address		
				Postal code		
	l.13.	Place of loading	1.14.	Date of departure		
	L15.	Means of transport	L16.	Entry BIP in EU		
	Aeroplane Ship Railway wagon Road vehicle Other					
		Identification: Documentary references:				
	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 Chilled		Frozen		
	I.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 in	to EU	
		3rd country ISO code				
	1.28.	Identification of the commodities				
		Annual gumb on of order	. la l' a la .ea			
	(Sc	Approval number of esta Species Nature of commodity Manufacturing plientific name)		ents Number of packag	es Net weight Batch number	

		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;							
L C	II.1.2.	have been obtained in the territory of:(²) from animals:							
Part II: Certification	(³) either	[(a) that have remained in this territory since birth or for at least the last three months before slaughter;]							
rtifi	(³) or	[(b) killed in the wild in this territory (4);]							
ပ္ပ	II.1.3.	have been obtained from animals:							
= t	(³) either	[(a) coming from holdings:							
Pa		(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 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 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,]							
	(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 manufacturing 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 significance communicable through that product to humans or animals,]	ins of any			
(3) and/or	[— fur originating from animals that did not show clinical signs of any disease communicable through that product to hanimals,]	umans or			
II.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 th will not spoil between dispatch and delivery to the plant of destination.				
(3)(6) [II.2.	Specific requirements				
(³) (⁷) II.2.1.	the by-products in this consignment come from animals that have been obtained in the territory mentioned under (II.1 vaccination programmes against foot-and-mouth disease are being regularly carried out and officially controlled in bovine animals.				
(³) (8) II.2.2.	. The by-products in this consignment consists of animal by-products derived from offal or de-boned meat.]				
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 commodity; it may be filled in if the certificate is for import commodity.	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 p transit can only be stored in free zones, free warehouses and custom warehouses.	roducts in			
_	Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or nar information is to be provided in case of unloading and reloading.	ne (ship);			
_	Box reference I.19: use the appropriate HS code: 05.11.91; 05.11.99 or 30.01.				
_	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 approved establishment.				
Part II:					
(*)	Excluding raw blood, raw milk, hides and skins of ungulates or ruminants and pig bristles (see relevant specific certifica import of these products) as well as wool, hair, feathers or parts of feathers.	tes for the			
(¹)	OJ L 273, 10.10.2002, p. 1.				
(2)	The name and ISO code number of the exporting country as laid down in:				
	— part 1 of Annex II of Council Decision 79/542/EEC,				
	— the Annex to Commission Decision 94/984/EC, and				
	 the Annex to Commission Decision 2000/585/EC. In addition the ISO code of regionalisation in this Annex (where applicable for the susceptible species concerned) 	should be			
	included.	siloulu be			
(3)	Delete as appropriate.				
(4)	Only for countries from where game meat intended for human consumption of the same animal species is auth importation into the European Community.	orised for			
(5)	Catering waste means all waste food, including used cooking oils, originating in restaurants, catering facilities and including central kitchens and household kitchens.	kitchens,			
(6)	Supplementary guarantees to be provided when the material of domestic ruminants originated in the territory of a South Americar or South African country or part thereof from where only maturated and de-boned fresh meat of domestic ruminants for humar consumption is permitted for exportation to the European Community. In the case of offal only trimmed offal of domestic ruminants which must be exclusively offal from which the bones, cartilage, trachea and main bronchi, lymphatic glands adhering connective tissue, fat and mucus have been completely removed is permitted. The whole masseter muscles of bovine animals, incised in 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.				
(8)	Only for certain South American and South 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 consignment in EU: this certificate is only for veterinary purposes and has to a the consignment until it reaches the border inspection post.	ccompany			
Official veter	erinarian	_			
	Name (in capitals): Qualification and title:				
	Date: Signature:				
	Stamp:				

CHAPTER 9

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	IKI				veterinary certificate to EU	
	l.1.	Consignor Name	1.2.	Certificate reference number	l.2.a.	
		Address	1.3.	Central Competent Authorit	у	
nent		Tel. No	1.4.	Local Competent Authority		
signr	1.5.	Consignee	1.6.	I.6. Person responsible for the consignment in EU		
con		Name Address		Name Address		
ched		Addiess		Addiess		
Part I: Details of dispatched consignment		Postal code Tel. No		Postal code Tel. No		
s of c	1.7.	Country of ISO code I.8.	1.9.	Country of ISO code	1.10.	
etail		origin		destination		
 	l.11.	Place of origin	l.12	. Place of destination Custom warehou	uso	
Par		Name Approval number		Name	Approval number	
		Address		Address	4,6	
				Postal code		
	l.13.	Place of loading	1.14.	Date of departure		
	l.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. Commod	ity code (HS code)	
					I.20. Quantity	
	1.21	Temperature of product			I.22. Number of packages	
	1.21.	Ambient Chilled		Frozen	1.22. Number of packages	
	I.23. Identification of container/Seal number				I.24. Type of packaging	
	1.25.	Commodities certified for: Animal feedingstuff		Techn	ical use	
	I.26. For transit to third country vis-à-vis EU		1.27.	For import or admission into		
		3rd country ISO code				
	1.28.	Identification of the commodities				
	Approval number of establishments Nature of commodity Manufacturing plant		Ni	umber of packages	Net weight Batch number	

		II.a.	Certificate reference num	ber II	b.	
l						
II.	Health attestation					
	I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1774/2002 (¹) and certify the fish oil described above:					
II.1.	consists of fish oil the	at satisfy the healt	n requirements below;			
II.2.	contains exclusively fish oil not intended for human consumption;					
II.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;					
II.4.	has been prepared e	exclusively with the	following animal by-products	3:		
(²) either		asons or due to pro	other than catering waste (³ blems of manufacturing or pa			
(²) and/or	[— fish or other se	a animals, except	sea mammals, caught in the	open sea for the	purposes of fishmeal produ	ıction,]
(²) and/or	[— fresh by-produc	cts from fish from p	plants manufacturing fish pro	ducts for human	consumption;]	
II.5.	the fish oil:					
	(a) has been subjected to processing in accordance with Annex VII, Chapter IV of Regulation 1774/2002/EC, in order to k pathogenic agents;					
	(b) has not been in	n contact with othe	r types of oils including rende	ered fats from oth	er animal species; and	
(²) either	[(c) is packaged in	n new containers	or in containers that have	been cleaned	and all precautions taker	n to prevent th
	contamination;]				
(²) or	transportation of	of the product from	the pipe, pumps and bulk tan the manufacturing plant eith be clean before use;]			
1						
	and which bear labe	ls indicating "NOT	FOR HUMAN CONSUMPTION	ON".		
	and which bear labe	ls indicating "NOT	FOR HUMAN CONSUMPTIO	DN".		
	and which bear labe	ls indicating "NOT	FOR HUMAN CONSUMPTIO	DN".		
Notes	and which bear labe	ls indicating "NOT	FOR HUMAN CONSUMPTIO	DN".		
Notes Part I:	Box reference I.6: F	Person responsible	e for the consignment in EU	: this box is to b	pe filled in only if it is a ce	ertificate for tran
	Box reference I.6: F commodity; it may be Box reference I.12: F	Person responsible e filled in if the cert Place of destination		: this box is to t y. rifit is a certificat		
	Box reference I.6: For commodity; it may be Box reference I.12: For can only be stored in	Person responsible e filled in if the cer Place of destinatior i free zones, free v Registration numbe	e for the consignment in EU ificate is for import commodi : this box is to be filled in only varehouses and custom ware r (railway wagons or containe	: this box is to to ty. y. if it is a certificat	e for transit commodity. The	products in tran
	Box reference I.6: For commodity; it may be Box reference I.12: For can only be stored in Box reference I.15: For is to be provided in commodity.	Person responsible e filled in if the cert Place of destinatior of free zones, free v Registration numbe case of unloading a	e for the consignment in EU ificate is for import commodi : this box is to be filled in only varehouses and custom ware r (railway wagons or containe	: this box is to to ty. y. if it is a certificat	e for transit commodity. The	products in tran
	Box reference I.6: For commodity; it may be Box reference I.12: For can only be stored in Box reference I.15: For is to be provided in a Box reference I.19: U	Person responsible e filled in if the ceri Place of destination if free zones, free vegistration numbersase of unloading asse the appropriate	e for the consignment in EU ificate is for import commodit : this box is to be filled in only varehouses and custom ware r (railway wagons or containe and reloading.	: this box is to by. If it is a certificathouses. If and lorries), flig	e for transit commodity. The ht number (aircraft) or name	products in tran
	Box reference I.6: For commodity; it may be Box reference I.12: For can only be stored in Box reference I.15: For is to be provided in a Box reference I.19: Leaver the Box reference I.23: for Box reference I.26 and a store of the box reference I.26 and a store	Person responsible e filled in if the cert Place of destination free zones, free values ase of unloading a use the appropriate or bulk containers, and 1.27: fill in acco	e for the consignment in EU ificate is for import commodit this box is to be filled in only varehouses and custom ware r (railway wagons or containe and reloading. HS code: 15.04 or 15.18. the container number and the roling to whether it is a transit	: this box is to by. If it is a certificatehouses. If and lorries), fligue seal number (if or an import cert	e for transit commodity. The ht number (aircraft) or name f applicable) should be inclu ificate.	products in tran (ship); informati
	Box reference I.6: For commodity; it may be Box reference I.12: For can only be stored in Box reference I.15: For is to be provided in a Box reference I.19: Leaver the Box reference I.23: for Box reference I.26 and a store of the box reference I.26 and a store	Person responsible e filled in if the cert Place of destination free zones, free values ase of unloading a use the appropriate or bulk containers, and 1.27: fill in acco	e for the consignment in EU ificate is for import commodit It this box is to be filled in only varehouses and custom ware Ir (railway wagons or containe and reloading. In the container number and the	: this box is to by. If it is a certificatehouses. If and lorries), fligue seal number (if or an import cert	e for transit commodity. The ht number (aircraft) or name f applicable) should be inclu ificate.	products in tran (ship); informati
Part I:	Box reference I.6: For commodity; it may be Box reference I.12: For can only be stored in Box reference I.15: For is to be provided in Box reference I.19: UBox reference I.23: for Box reference I.26 a Box reference I.28: May be a supplemental box reference II.28: May be a suppl	Person responsible e filled in if the cert Place of destination if free zones, free values as e of unloading a use the appropriate for bulk containers, and 1.27: fill in acco	e for the consignment in EU ificate is for import commodit this box is to be filled in only varehouses and custom ware r (railway wagons or containe and reloading. HS code: 15.04 or 15.18. the container number and the roling to whether it is a transit	: this box is to by. If it is a certificatehouses. If and lorries), fligue seal number (if or an import cert	e for transit commodity. The ht number (aircraft) or name f applicable) should be inclu ificate.	products in tran (ship); informati
Part I:	Box reference I.6: For commodity; it may be Box reference I.12: For can only be stored in Box reference I.15: For is to be provided in a Box reference I.23: for Box reference I.26 a Box reference I.28: MOJ L 273, 10.10.200	Person responsible e filled in if the cere Place of destination if the example of the containers or bulk containers, and I.27: fill in acco Manufacturing plant 2, p. 1.	e for the consignment in EU ificate is for import commodit this box is to be filled in only varehouses and custom ware r (railway wagons or containe and reloading. HS code: 15.04 or 15.18. the container number and the roling to whether it is a transit	: this box is to by. If it is a certificatehouses. If and lorries), fligue seal number (if or an import cert	e for transit commodity. The ht number (aircraft) or name f applicable) should be inclu ificate.	products in tran (ship); informati
Part I: (1) (2)	Box reference I.6: For commodity; it may be Box reference I.12: For can only be stored in Box reference I.15: For is to be provided in a Box reference I.23: For ference I.26 a Box reference I.28: For ference I.28: For ference I.28: For control of the III and III are the	Person responsible e filled in if the cert Place of destination of the expression numbers as e of unloading a use the appropriate or bulk containers, and I.27: fill in accommunication plant 2, p. 1.	e for the consignment in EU ificate is for import commodit this box is to be filled in only varehouses and custom ware r (railway wagons or containe and reloading. HS code: 15.04 or 15.18. the container number and the reding to whether it is a transit try provide the registration number.	: this box is to by. If it is a certificatehouses. If and lorries), fligue seal number (if or an import certimber of the treati	e for transit commodity. The ht number (aircraft) or name f applicable) should be inclu ifficate. ment/processing establishm	e products in tran (ship); informati uded.
Part I:	Box reference I.6: For commodity; it may be Box reference I.12: For can only be stored in Box reference I.15: For is to be provided in a Box reference I.23: For ference I.26 a Box reference I.28: For ference I.28: For ference I.28: For control of the III and III are the	Person responsible e filled in if the cere place of destination free zones, free values of unloading a use the appropriate or bulk containers, and I.27: fill in accontainers plants of the properties of the plants	e for the consignment in EU ificate is for import commodit this box is to be filled in only varehouses and custom ware r (railway wagons or containe and reloading. HS code: 15.04 or 15.18. the container number and the right to whether it is a transit to provide the registration number coulding used cooking oils, oricluding used cooking oils, oriclinication in the registration of the coulding used cooking oils, oriclinication in the registration of the coulding used cooking oils, oriclinication in the registration of the	: this box is to by. If it is a certificatehouses. If and lorries), fligue seal number (if or an import certimber of the treati	e for transit commodity. The ht number (aircraft) or name f applicable) should be inclu ifficate. ment/processing establishm	e products in tran (ship); informati ided.
Part I: (1) (2)	Box reference I.6: Frommodity; it may be Box reference I.12: From only be stored in Box reference I.15: From the Box reference I.19: Use Box reference I.26: Box reference I.26: Modulate Box reference I.28: Modulate Box	Person responsible of illed in if the cert Place of destination of the expension of the exp	e for the consignment in EU ificate is for import commodit this box is to be filled in only varehouses and custom ware r (railway wagons or containe and reloading. HS code: 15.04 or 15.18. the container number and the right to whether it is a transit to provide the registration number coulding used cooking oils, oricluding used cooking oils, oriclinication in the registration of the coulding used cooking oils, oriclinication in the registration of the coulding used cooking oils, oriclinication in the registration of the	: this box is to by. If it is a certificatehouses. If and lorries), fligue seal number (if or an import certimber of the treate	e for transit commodity. The ht number (aircraft) or name f applicable) should be inclu ifficate. ment/processing establishm	e products in tran (ship); informati uded.
Part I: (1) (2)	Box reference I.6: For commodity; it may be Box reference I.12: For can only be stored in Box reference I.15: For it is to be provided in the Box reference I.23: For IIII Box reference I.26: For IIII Box reference I.28: For IIII Box reference III Box reference II Box refer	Person responsible e filled in if the cert Place of destination of the content of	e for the consignment in EU ificate is for import commodifultion if this box is to be filled in only varehouses and custom ware represent (railway wagons or contained and reloading. HS code: 15.04 or 15.18. The container number and the reding to whether it is a transite to provide the registration number and the container of the registration	: this box is to by. If it is a certificatehouses. If and lorries), fligue seal number (if or an import certimber of the treateful or an import certification or an import certification or an import certification or an important certification or an import	e for transit commodity. The ht number (aircraft) or name f applicable) should be inclu ifficate. ment/processing establishm rants, catering facilities and	e products in tran (ship); information ided. ent. kitchens, includio
Part I: (1) (2)	Box reference I.6: For commodity; it may be Box reference I.12: For can only be stored in Box reference I.15: For is to be provided in the Box reference I.23: If Box reference I.23: If Box reference I.26: If Box reference I.28: I	Person responsible e filled in if the cert Place of destination of the content of	e for the consignment in EU ificate is for import commodifultion if this box is to be filled in only varehouses and custom ware represent (railway wagons or contained and reloading. HS code: 15.04 or 15.18. The container number and the reding to whether it is a transite to provide the registration number and the container of the registration	: this box is to by. If it is a certificatehouses. If and lorries), fligue seal number (if or an import certimber of the treateful or an import certification or an import certification or an import certification or an important certification or an import	e for transit commodity. The ht number (aircraft) or name f applicable) should be inclu ifficate. ment/processing establishm rants, catering facilities and	e products in tran (ship); informati ided. ent. kitchens, includi
Part I: (1) (2) (3)	Box reference I.6: For commodity; it may be Box reference I.12: For can only be stored in Box reference I.15: For is to be provided in the Box reference I.23: If Box reference I.23: If Box reference I.26: If Box reference I.28: I	Person responsible e filled in if the cert Place of destination of the content of	e for the consignment in EU ificate is for import commodifultion if this box is to be filled in only varehouses and custom ware represent (railway wagons or contained and reloading. HS code: 15.04 or 15.18. The container number and the reding to whether it is a transite to provide the registration number and the container of the registration	: this box is to by. If it is a certificatehouses. If and lorries), fligue seal number (if or an import certificate of the treatember of	e for transit commodity. The ht number (aircraft) or name f applicable) should be inclu ifficate. ment/processing establishm rants, catering facilities and	e products in tran (ship); information ided. ent. kitchens, includio
Part I: (1) (2) (3)	Box reference I.6: For commodity; it may be Box reference I.12: For can only be stored in Box reference I.15: For is to be provided in a Box reference I.23: for Box reference I.26: for Box reference I.28: for Modern I.28: for	Person responsible e filled in if the cert Place of destination of the content of	e for the consignment in EU ificate is for import commodifultion if this box is to be filled in only varehouses and custom ware represent (railway wagons or contained and reloading. HS code: 15.04 or 15.18. The container number and the reding to whether it is a transite to provide the registration number and the container of the registration	: this box is to by. If it is a certificatehouses. If and lorries), fligue seal number (if or an import certificate of the treatember of	e for transit commodity. The ht number (aircraft) or name f applicable) should be incluificate. ment/processing establishments, catering facilities and reterinary purposes and has tion and title:	e products in tran (ship); informati ided. ent. kitchens, includi

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 (4) the European Community

COUN	IKI					veterinary certificate to EU
	l.1.	Consignor Name	1.2.	Certificate reference number)	1.2.a.
		Address	1.3.	Central Competent	Authority	У
nent		Tel. No	1.4.	Local Competent Au	ıthority	
signr	I.5.	Consignee	1.6.	Person responsible	for the c	onsignment in EU
con		Name Address		Name Address		
ched						
Part I: Details of dispatched consignment		Postal code Tel. No		Postal code Tel. No		
s of c	1.7.	Country of ISO code I.8.	1.9.		O code	1.10.
etail		origin		destination		
1 1:0	l.11.	Place of origin	1.12.	Place of destination Custom v	zrehou	sa 🗖
Pai		Name Approval number		Name	varenou.	Approval number
		Address		Address		
				Postal code		
	I.13.	Place of loading	l.14.	Date of departure		
	l 15	Means of transport	I 16	Entry BIP in EU		
	1.10.	Aeroplane Ship Railway wagon	1.10.	Linky Bit iii Lo		
		Road vehicle Other I				
		Documentary references:	l.17.			
	I.18.	Description of commodity		I.19. C	ommodi	ty code (HS code)
					1.20). Quantity
	I.21.	Temperature of product			1.22	2. Number of packages
		Ambient Chilled Chilled		Frozen		
	I.23.	Identification of container/Seal number			1.24	l. Type of packaging
	1.25.	Commodities certified for: Animal feedingstuff			Techni	cal use
	1.26.	For transit to third country vis-à-vis EU	1.27.	For import or admiss		
		3rd country ISO code				
	1.28.	Identification of the commodities				
		Approval number of 6	establis	hments		
		Species Nature of commodity Manufacturing (Scientific name)	g plant	Number of	packag	es Net weight Batch number

		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 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	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 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 an 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;]					
	1						

and which bear labels indicating "NOT FOR HUMAN CONSUMPTION".

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 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: 15.02; 15.03; 15.04; 15	5.05; 15.06; 15.16.10; 15.17 or 15.18.				
_	Box reference I.23: for bulk containers, the container number and the sea	al number (if applicable) should be included.				
_	Box reference I.26 and I.27: fill in according to whether it is a transit or an	n 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, originatic central kitchens and household kitchens.	ng in restaurants, catering facilities and kitchens, including				
_	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 consignment until it reaches the border inspection post.	is only for veterinary purposes and has to accompany the				
Official ve	terinarian					
	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	IKI				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	
ignn	1.5.	Consignee	1.6.	Person responsible for the	consignment in EU
cons		Name		Name	
peq (Address		Address	
Part I: Details of dispatched consignment		Postal code Tel. No		Postal code Tel. No	
of c	1.7.	Country of ISO code I.8.	1.9.	Country of ISO code	e I.10.
tails		origin		destination	
i: De	I.11.	. Place of origin	I.12.	Place of destination	
Part				Custom wareho	_
		Name Approval number		Name	Approval number
		Address		Address	
				Postal code	
	l.13.	. Place of loading	1.14.	Date of departure	
	l.15.	. Means of transport	I.16.	Entry BIP in EU	
		Aeroplane Ship Railway wagon Road vehicle Other			
		Identification:	1.17.		
		Documentary references:			
	l.18.	. Description of commodity		I.19. Commod	dity code (HS code)
					I.20. Quantity
	l.21.	. Temperature of product Ambient Chilled		Frozon 🗖	I.22. Number of packages
				Frozen	
	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	o EU
		3rd country ISO code			
	1.28.	. Identification of the commodities	•		
		Approval number of e	establis	hments	
		Species Nature Manufactu	ıring	Number of	Net Batch
		(Scientific name) of commodity plant		packages	weight number
	l				

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 rendered fats described above:

- II.1. consist of rendered fats that satisfy the health requirements below;
- 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;
- II.4. have been prepared exclusively with the following animal by-products:
- (2) either [Category 2 materials (3);]
- (2) or [a mixture of Category 2 materials with Category 3 materials (4);]
- 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 XII of Regulation (EC) No 1774/2002/EC, in order to kill pathogenic agents; and
- (2) either [(b) are packaged in new containers or in containers that have been cleaned and all precautions taken to prevent their contamination;]
- (2) 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 OR ANIMAL 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 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.19: use the appropriate HS code: 15.01; 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 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 the 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.
- (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;
 - 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.

	consignment until it reaches the border inspection post.					
Official veterinarian						
	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	IKI		veterinary certificate to i
	l.1.	Consignor Name	I.2. Certificate reference number
		Address	I.3. Central Competent Authority
nent		Tel. No	I.4. Local Competent Authority
signn	1.5.	Consignee	I.6. Person responsible for the consignment in EU
con		Name	Name
hed		Address	Address
Part I: Details of dispatched consignment		Postal code Tel. No	Postal code Tel. No
s of	I.7.	Country of ISO code I.8.	I.9. Country of ISO code I.10.
etail		origin	destination
<u>:</u>	l.11.	. Place of origin	I.12. Place of destination
Parl		Approval number	Custom warehouse
		Name Approval number Address	Name Approval number Address
		Address	Audioss
			Postal code
	l.13.	. Place of loading	I.14. Date of departure
	l.15.	. Means of transport Aeroplane Ship Railway wagon	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)
			I.20. Quantity
	I.21.	Temperature of product	I.22. Number of packages
		Ambient Chilled	Frozen
	1.23.	Identification of container/Seal number	I.24. Type of packaging
	1.25.	Commodities certified for:	•
		Animal feedingstuff	Technical use
	1.26.	For transit to third country vis-à-vis EU	I.27. For import or admission into EU
		3rd country ISO code	
	1.28.	Identification of the commodities	
		Approval number of e	establishments
	(Sci	Species Nature of commodity Manufacturin entific name)	

					or for technical purposes		
			II.a.	Certificate reference number	II.b.		
	11.	Health attestation	al votorinarion	declare that I have read and understood	Regulation (EC) No 1774/2002 (1) and certify that		
		the gelatine/collagen (2)	,		Regulation (EC) No 1774/2002 (*) and certify that		
	II.1.	consists of gelatine/coll	agen (2) that sa	atisfy the health requirements below;			
ıtion	II.2.	consist exclusively of ge	elatine/collager	n (²) not intended for human consumption	on;		
Part II: Certification	II.3.			lant approved, validated and supervise le 11 of Regulation (EC) No 1774/2002,	ed by the competent authority in accordance with in order to kill pathogenic agents;		
3	II.4.	has been prepared exc	usively with the	e following animal by-products:			
Part	(²) either			which are fit for human consumption in a ion for commercial reasons,]	accordance with Community legislation, but are not		
	(²) and/or	[— parts of slaugh	tered animals,	which are rejected as unfit for human 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	slaughterhouse	, after undergo		inating from animals that were slaughtered in a fit, as a result of such inspection, for slaughter for		
	(²) and/or	[— animal by-prode	ucts derived fro	om 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 wast which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing packaging defects or other defects which do not present any risk to humans or animals,]						
	(²) and/or	[— fish or other sea	a animals, exce	ept sea mammals, caught in the open so	ea for the purposes of fishmeal production,]		
	(²) and/or	[— fresh by-produc	ts from fish fro	m plants manufacturing fish products fo	or human consumption;]		
	II.5.	the gelatine/collagen (2)	:				
			-		nygiene conditions, and in particular wrapping and mitted under Community legislation were used.		
		Wrappings and ANIMAL CONS			ords "GELATINE/COLLAGEN (2) SUITABLE FOR		
	(²) either	to a treatment v	vith acid or alk	ali, followed by one or more rinses, invo	that unprocessed Category 3 material is subjected olving pH adjustment, extraction by heating one or tration and sterilisation, in order to kill pathogenic		
	1 1 7			ıg, pH adjustment using acid or alkali follo	that unprocessed Category 3 material is subjected owed by one or more rinses, filtration and extrusion,		
	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 commodity; it may be filled in if the certificate is for import commodity. 						
	_			on: this box is to be filled in only if it is a s, free warehouses and custom wareho	a certificate for transit commodity. The products in uses.		
	_			nber (railway wagons or container and funloading and reloading.	I lorries), flight number (aircraft) or name (ship);		
	_	Box reference I.19: use	the appropriat	e HS code: 35.03 or 35.04.			
	_	Box reference I.23: for I	oulk containers	, the container number and the seal nur	mber (if applicable) should be included.		
	_	Box reference I.26 and	I.27: fill in acco	ording to whether it is a transit or an imp	ort certificate.		
	Box reference I.28: Nature of commodity: select gelatine or collagen. 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 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 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 vete	rinarian					
	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	IKT					veterinary certificate to EU
	l.1.	Consignor Name	1.2.	Certificate reference number		1.2.a.
		Address	1.3.	Central Competent A	uthority	1
nent		Tel. No	1.4.	Local Competent Aut	hority	
signr	I.5.	Consignee	1.6.	Person responsible fo	or the c	onsignment in EU
con		Name Address		Name Address		
ched		Addiess		Address		
Part I: Details of dispatched consignment		Postal code Tel. No		Postal code Tel. No		
ils of	1.7.	Country of ISO code I.8.	1.9.	Country of ISC destination) code	1.10.
Deta	1.11.	Place of origin	1.12.	Place of destination		
art I:		3		Custom wa	arehous	se 🔲
^		Name Approval number		Name		Approval number
		Address		Address		
			Postal code			
	I.13.	Place of loading	l.14.	Date of departure		
	l.15.	. Means of transport Aeroplane Ship Railway wagon	I.16.	Entry BIP in EU		
		Road vehicle Other Railway wagon				
		Identification:	I.17.			
	1 10	Documentary references:		110.00		
	1. 10.	Description of commodity		1.19. Coi		ty code (HS code)
					1.20). Quantity
	1 21	. Temperature of product			1 122	2. Number of packages
	I.∠ ı.	Ambient Chilled		Frozen	1.22	Nulliber of packages
	1.23.	Identification of container/Seal number			1.24	. Type of packaging
	1.25.	Commodities certified for: Animal feedingstuff			Technic	cal use
	1.26.	For transit to third country vis-à-vis EU	1.27.	For import or admission		
	3rd country ISO code					_
	1.28.	Identification of the commodities				
		Approval number of esta	ıblishme	ents		
		Species Nature of commodity Manufacturing plentific name)		Number of pac	ckages	Net weight Batch number
	í					

					I					
			II.a.	Certificate reference number	II.b.					
	II.	Health attestation								
				declare that I have read and understor phate/tricalcium phosphate (²) describ	od Regulation (EC) No 1774/2002 (¹) and certify that ed above:					
Part II: Certification	II.1.	consists of hydrolys	ed protein/dicalciu	nm phosphate/tricalcium phosphate (²)	that satisfy the health requirements below;					
I: Certif	II.2.	consists exclusively	of hydrolysed pro	tein/dicalcium phosphate/tricalcium ph	osphate (²) not intended for human consumption;					
Part	II.3.			olant approved, validated and supervisule 11 of Regulation (EC) No 1774/2002	sed by the competent authority in accordance with 2, in order to kill pathogenic agents;					
	II.4.	has been prepared	exclusively with th	e following animal by-products:						
	(²) either	[— parts of sla	ightered animals,		accordance with Community legislation, but are not					
	(²) and/or	of diseases		humans or animals and derive from	an consumption but are not affected by any signs a carcasses that are fit for human consumption in					
	(²) and/or	slaughterho	use, after underg		ginating from animals that were slaughtered in a fit, as a result of such inspection, for slaughter for					
	(²) and/or	mortem ins		mals other than ruminants that are slaughtered in a slaughterhouse, after undergoing antewere fit, as a result of such inspection, for slaughter for human consumption in accordance with						
	(²) and/or	[— animal by-p	animal by-products derived from the production of products intended for human consumption;]							
(2) and/or [— former foodstuffs of animal origin, or former foodstuffs containing products or during the foodstuffs containing products or former foodstuffs containing products or food				ial reasons or due to problems of manufacturing or						
	(²) and/or	[— raw milk or humans or		nals that do not show clinical signs of	any disease communicable through that product to					
	(²) and/or	[— fish or othe	sea animals, exc	ept sea mammals, caught in the open	sea for the purposes of fishmeal production;]					
	(²) and/or	[— fresh by-pro	resh by-products from fish from plants manufacturing fish products for human consumption;]							
	(²) and/or			and cracked egg by-products originatir rough that product to humans or anima	ng from animals which did not show clinical signs of als;]					
	II.5.	the hydrolysed prot	ein/dicalcium phos	phate/tricalcium phosphate (2):						
(a) was wrapped and packaged in packaging which bear labels indicating "NC stored and transported under satisfactory hygiene conditions, and in particular dedicated room, and only preservatives permitted under Community legislation"				n particular wrapping and packaging took place in a						
	(²) either	-, ,	[(b) in the case of hydrolysed protein, has been produced by a process involving appropriate measures to minimise contamination of raw Category 3 material.							
		processing	plant dedicated or		uminants hides and skins, has been produced in a using a process involving the preparation of the raw d by:					
				rial to a pH of more than 11 for more the treatment at more than 140 °C for 30 in	han 3 hours at temperature of more than 80 °C and minutes at more than 3,6 bar; and					
			osure of the mater 30 minutes at 3 ba		f more than 11, followed by heat treatment at 140 °C					

(0)			
(²) or	[(b)		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 $^{\circ}$ C to 325 $^{\circ}$ C and end temperature between 60 $^{\circ}$ C and 65 $^{\circ}$ 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:			
_			e I.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.
-			1.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in y be stored in free zones, free warehouses and custom warehouses.
_			e I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship);
	inform	ation is	to be provided in case of unloading and reloading.
-	Box re	eference	I.19: use the appropriate HS code: 28.35 or 35.04.
_	Box re	eference	I.23: for bulk containers, the container number and the seal number (if applicable) should be included.
_	Box re	eference	I.26 and I.27: fill in according to whether it is a transit or an import certificate.
_	Box re	eference	I.28: Nature of commodity: specify if hydrolysed protein, dicalcium phosphate or tricalcium phosphate.
			Manufacturing plant: provide the registration number of treatment/processing establishment.
D4 II-			
Part II:	011.0	22 40	40,000 - 4
(¹)			10.2002, p. 1.
(2)			ropriate.
(3)	includ	ing cent	re means all waste food, including used cooking oils, originating in restaurants, catering facilities and kitchens, ral kitchens and household kitchens.
_		•	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	(in capi	itals):
			Qualification and title:
	Date:		
			Signature:
	Stamp	o:	g
	Clairi		

Health certificate

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

COUN	IRY				Veterinary certificate to EU
	l.1.	Consignor Name	1.2.	Certificate reference number	1.2.a.
		Address	1.3.	Central Competent Authority	
nent		Tel. No	1.4.	Local Competent Authority	
ısignr	1.5.	Consignee	1.6.	Person responsible for the co	nsignment in EU
oo pə		Name Address		Name Address	
Part I: Details of dispatched consignment		Postal code Tel. No		Postal code Tel. No	
tails of	1.7.	Country of ISO code I.8. Region of origin Code origin	1.9.	Country of destination ISO code	I.10. Region of Code destination
: I: De	l.11.	Place of origin	I.12.	Place of destination]
Part		Name Approval number		Custom warehouse	Approval number
		Address		Address	Approvar nambor
				Postal code	
	I.13.	Place of loading	l.14.	Date of departure	
	l.15.	Means of transport	I.16.	Entry BIP in EU	
		Aeroplane Ship Railway wagon Road vehicle Other			
		Identification: Documentary references:	l.17.		
	I.18.	Description of commodity		I.19. Commodity	code (HS code)
				ı	.20. Quantity
	l.21.	Temperature of product Ambient Chilled Chilled		Frozen 🔲	.22. Number of packages
	1.23.	Identification of container/Seal number		- 1	.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 E	EU
		3rd country ISO code			
	1.28.	Identification of the commodities			
		Species Nature of commodity (Scientific name)	Approva	al number of establishments Manufacturing plant	Net weight

COUNTRY Apiculture by-products II.a. Certificate reference number II.b. П. 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 IX thereof and certify that the apiculture by-products described above: II.1. come from an area where the diseases mentioned below are officially notifiable and which is not subject to any restrictions associated Part II: Certification (a) American foul brood (Paenibacillus larvae); (b) Acariosis (Acarapis woodi (Rennie)); Small hive beetle (Aethina tumida); and (c) (d) Tropilaelaps mites (Tropilaelaps spp); 11.2. have been (2) either [subjected to a temperature of -12 °C or lower for at least 24 hours;] [in the case of wax refined or rendered.] (2) or 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: 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 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: means honey, beeswax, royal jelly, propolis or pollen used in bee-keeping. 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:

Signature:

Date:

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	ΓRY			Veterinary certificate to EU
	I.1. Consignor Name	1.2	. Certificate reference number	1.2.a.
	Address	1.3	. Central Competent Author	ity
nent	Tel. No		. Local Competent Authority	
nsignı	I.5. Consignee I. Name		I.6. Person responsible for the consignment in EU	
oo pə	Address		Name Address	
Part I: Details of dispatched consignment	Postal code Tel. No		Postal code Tel. No	
tails of	I.7. Country of ISO code I.8. Reg	gion of origin Code I.9). Country of ISO code destination	1.10.
t I: De	I.11. Place of origin	l.1	2. Place of destination	
Par	Name App	proval number	Custom wareho Name	Approval number
	Address		Address	
			Postal code	
	I.13. Place of loading	1.1	4. Date of departure	
	I.15. Means of transport Aeroplane Ship		6. Entry BIP in EU	
	Road vehicle Other			
	Identification: Documentary references:	1.1	7.	
	I.18. Description of commodity		I.19. Commod	dity code (HS code) 15.16.10
				I.20. Quantity
	I.21. Temperature of product	Chilled	Frozon 🗖	I.22. Number of packages
	Ambient 1.23. Identification of container/Seal number		Frozen	I.24. Type of packaging
				1.24. Type of packaging
	I.25. Commodities certified for:	Technical use		
	I.26. For transit to third country vis-à-vis EU		7. For import or admission int	o EU
	3rd country ISO co	de		
	I.28. Identification of the commodities	<u>'</u>		
	Species Nature of commodity (Scientific name)	Approval number of establishr Manufacturing plant	ments Number of package	s Net weight Batch number

Part II: Certification

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 °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.

_	 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	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 El
	I.1. Consignor Name	I.2. Certificate reference number
	Address	I.3. Central Competent Authority
nent	Tel. No	I.4. Local Competent Authority
Part I: Details of dispatched consignment	I.5. Consignee Name Address	I.6. Person responsible for the consignment in EU Name Address
dispatche	Postal code Tel. No	Postal code Tel. No
tails of	I.7. Country of ISO code I.8. Region of origin Code origin	I.9. Country of destination ISO code I.10.
ırt I: De	I.11. Place of origin	I.12. Place of destination Custom warehouse
Ра	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 Road vehicle Other	I.16. Entry BIP in EU
	Identification: Documentary references:	1.17.
	I.18. Description of commodity	I.19. Commodity code (HS code) 15.16.10
		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
	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 es Species Nature of commodity Manufacturing (Scientific name)	

			II.a. Certificate reference number	II.b.				
	II.	Health attestation						
		I, the undersigned officia the fat derivatives descri	l veterinarian, declare that I have read and understood bed above:	d Regulation (EC) No 1774/2002 (1) and certify that				
	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;						
ပ္ပ	II.4.	have been prepared from rendered fats exclusively produced from the following Category 3 materials:						
Part	(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	diseases communi	[— 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	slaughterhouse, af	hooves and horns, pig bristles and feathers origing ter undergoing ante-mortem inspection, and were fith In 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 and greaves,]	derived from the production of products intended for	r human consumption, including degreased bones				
	(²) and/or	which are no long	animal origin, or former foodstuffs containing products of animal origin, other than catering waste (3), or intended for human consumption for commercial reasons or due to problems of manufacturing or or other defects which do not present any risk to humans or animals,]					
	(²) and/or	[— milk originating fro humans or animals	m animals which do not show any clinical signs of any disease communicable through that product to					
	(²) and/or	[— fish or other sea ar	nimals, except sea mammals, caught in the open sea	for the purposes of fishmeal production,]				
	(²) and/or	[— by-products from fi	sh from plants manufacturing fish products for human	consumption,]				
	(²) and/or		-products and cracked egg by-products originating fro able through that product to humans or animals;]	m animals which did not show clinical signs of any				
	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:							
	_		on responsible for the consignment in EU: this box it add in if the certificate is for import commodity.	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: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (sinformation 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 imp	ort 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 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 different colour to that of the printing.		
_	Note for the person responsible for the consignment in EU: this certificate the consignment until it reaches the border inspection post.	e is only for veterinary purposes and has to accompany	
Official veter	inarian		
	Name (in capitals):	Qualification and title:	
	Date:	Signature:	
	Stamp:		

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	IKI			veterinary certificate to) EU
	l.1.	Consignor Name	l.2.	Certificate reference I.2.a. number	
		Address	1.3.	Central Competent Authority	
nent		Tel. No	1.4.	Local Competent Authority	
signr	I.5.	Consignee	1.6.	I.6. Person responsible for the consignment in EU	
con		Name Address		Name Address	
hed		Addiess		Address	
Part I: Details of dispatched consignment		Postal code Tel. No		Postal code Tel. No	
s of c	1.7.	Country of ISO code I.8. Region of origin Code	1.9.	Country of ISO code I.10.	_
etails		origin		destination	
. I: D	I.11.	Place of origin	I.12.	. Place of destination	
Parl		Name Approval number		Custom warehouse	
		Address		Address	
				Postal code	
	I.13. Place of loading			. Date of departure	
	I.15. Means of transport		I.16.	. Entry BIP in EU	
		Aeroplane Ship Railway wagon Road vehicle Other			
		Identification: Documentary references:			_
	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	
	1.23.	Identification of container/Seal number		I.24. Type of packaging	
	I.25.	Commodities certified for: Animal feedingstuff			
	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			
	0.		مطمئلطم	wante.	
	(Sci	Approval number of est Species Nature of commodity Manufacturing plentific name)		пепts Number of packages Net weight Batch numb	er

			II.a. Certificate reference number	II.b.				
		l						
	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;						
atio	II.2.	consist exclusively of egg products not intended for human consumption;						
art 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 or Council Directive 89/437/EEC (2), in order to kill pathogenic agents;						
∺	II.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 pro-	cessing:					
	(³) either	[in accordance with process	sing method (4) as set out in Annex V, Chap	pter 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	7/EC]				
	II.6.	have been examined by the the following standards (5):	competent authority taking a random sample imme	ediately prior to dispatch and found it to comply with				
		Salmonella: ab	osence 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 or make its use as feed dangerous or harmful to animal health;				at alter the organoleptic characteristics of the product				
	II.8.	the end product was:						
	(³) either	[packed in new or sterilized	bags;]					
	(³) or	[transported in bulk in cont approved by the competent		roughly cleaned and disinfected with a disinfectant				
		and which bear labels indica	ating '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:							
	_		responsible for the consignment in EU: this box in if the certificate is for import commodity.	is to be filled in only if it is a certificate for transit				
	_		f destination: this box is to be filled in only if it is a cerones, free warehouses and custom warehouses.	rtificate for transit commodity. The products in transit				
	_	Box reference I.15: Registra is to be provided in case of		s), flight number (aircraft) or name (ship); information				
	–	Box reference I.23: for bulk	containers, the container number and the seal num	nber (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)	OJ L 212, 22.07.1989, p. 89.			
(3)	Delete as appropriate.			
(4)	Insert method 1 to 5 or 7 as applicable.			
(5)	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 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 bacterial count of the other samples is m or less	may be between m and M, the sample still being considered acceptable if the s.		
-	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 ve	eterinarian			
	Name (in capitals):	Qualification and title:		
	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.

- I, 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:				
	on				
(place)	(date)				
Signature					
Regulation (EC) No 136/2004:	veterinary entry document (CVED) provided for in Annex III to Commission				
Official stamp of the border inspection post of entry into the EC (²)					
	fficial veterinarian of the border inspection post) (²)				
Name:					

(Name in capital letters)

Delete as appropriate.

⁽²⁾ 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	IRY				Veterinary certificate to EU
	l.1.	Consignor Name	1.2.	Certificate reference number	1.2.a.
		Address	1.3.	Central Competent Authorit	ty
nent		Tel. No	1.4.	Local Competent Authority	
nsignr	1.5.	Consignee Name	1.6.	Person responsible for the o	consignment in EU
ned co		Address		Address	
Part I: Details of dispatched consignment		Postal code Tel. No		Postal code Tel. No	
tails of	l.7.	Country of origin ISO code I.8. Region of origin Code origin	1.9.	Country of destination ISO code	I.10. Region of Code destination
t I: De	l.11.	Place of origin	I.12.	Place of destination	
Par		Name Approval number		Custom warehou	Approval number
		Address		Address	
				Postal code	
	I.13.	Place of loading	l.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:	l.17.		
	I.18. Description of commodity			I.19. Commod	ity code (HS code)
					I.20. Quantity
	I.21.	Temperature of product			I.22. Number of packages
		Ambient 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.	For import or admission into	EU
		3rd country ISO code			
	1.28.	Identification of the commodities			
		Species Nature of commodity (Scientific name)		al number of establishments Manufacturing plant	Net weight

			II.a. Certificate reference number	II.b.				
	II.	Health attestation		•				
				erstood Regulation (EC) No 1774/2002 (1) and in particular				
		Article 5 and Annex VIII Chapte	er vi thereof and certify that the processed	manure or processed manure products described above:				
on	II.1. come from a technical plant, a biogas plant or a composting plant approved by the competent authority of the third country meeting special conditions laid down in Regulation (EC) No 1774/2002;							
II.2. (²) have been subjected to:								
ırtif	special conditions laid down in Regulation (EC) No 1774/2002; 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 in Regulation (EC) No 1774/2002 as follows:							
ა 								
art II								
_								
	1000							
	II.3.	are:	colmonally in 25 a treated and dust).					
			salmonella in 25 g treated product);	ne aerobic count: less than 1 000 cfu per gram of treated				
		product); and	in or nom emeropacienacea (pased on the	le deloble count. less than 1 000 clu per gram of freated				
		(c) have been subjected to re	eduction in spore-forming bacteria and toxi	c formation;				
	II.4.	are securely enclosed in:						
		(a) well-sealed and insulated	containers; or					
		(b) properly sealed packs (pl	astic 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 com 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 product can only be stored in free zones, free warehouses and custom warehouses. 						
	_	Box reference I.15: Registratio	n number (railway wagons or container an	d lorries), flight number (aircraft) or name (ship); information				
		is to be provided in the event o	· ·					
	_	Box reference I.23: for bulk cor	ntainers, the container number and the sea	I 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 c	ommodity: enter if processed manure or if	processed manure products.				
	Part II:							
	(¹)	OJ L 273, 10.10.2002, p. 1.						
	(²)	Delete as appropriate.						
	_		nust be in a different colour to that of the pr	_				
		Note for the person responsible for the consignment in EU: This certificate is only for veterinary purposes and has to accompany consignment until it reaches the border inspection post.						
	Official \	veterinarian						
		Name (in capitals):		Qualification and title:				
		Date:		Signature:'.				
		Stamn:						

- (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.'.