COMMISSION IMPLEMENTING REGULATION (EU) 2016/759

of 28 April 2016

drawing up lists of third countries, parts of third countries and territories from which Member States are to authorise the introduction into the Union of certain products of animal origin intended for human consumption, laying down certificate requirements, amending Regulation (EC) No 2074/2005 and repealing Decision 2003/812/EC

(Text with EEA relevance)

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard 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 (1) and in particular Articles 8(1) and 9(4) thereof,

Having regard to Regulation (EC) No 854/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific rules for the organisation of official controls on products of animal origin intended for human consumption (2), and in particular Article 11(1) thereof,

Whereas:

- (1)Regulation (EC) No 854/2004 requires products of animal origin to be imported only from a third country or a part of third country that appears on a list drawn up in accordance with that Regulation.
- Commission Decision 2003/812/EC (3) draws up lists of third countries from which Member States are to (2)authorise imports of certain products for human consumption subject to Council Directive 92/118/EEC (*). Those lists include a list of third countries or parts of third countries from which imports of gelatine intended for human consumption are authorised. However, there is no list which covers collagen, or raw materials for the production of gelatine and collagen, for human consumption. It is appropriate to draw up such lists.
- (3) In accordance with Regulation (EC) No 853/2004 of the European Parliament and of the Council (5), food business operators importing products of animal origin are to ensure that the documents accompanying the consignment meet the requirements of Article 14 of Regulation (EC) No 854/2004. Commission Regulation (EC) No 2074/2005 (6) lays down model certificates for imports of certain products of animal origin intended for human consumption. Those model certificates include outdated references to previous legislation that need to be updated.

⁽¹⁾ OJ L 18, 23.1.2003, p. 11.

OJ L 139, 30.4.2004, p. 206.

Commission Decision 2003/812/EC of 17 November 2003 drawing up lists of third countries from which Member States are to authorise imports of certain products for human consumption subject to Council Directive 92/118/EEC (OJ L 305, 22.11.2003, p. 17). Council Directive 92/118/EEC of 17 December 1992 laying down animal health and public health requirements governing trade in

and import into the Community of products not subject to the said requirements laid down in specific Community rules referred to in Annex A(I) to Directive 89/662/EEC, and, as regards pathogens, Directive 90/425/EEC (OJ L 62, 15.3.1993, p. 49).

Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific hygiene rules for

food of animal origin (OJ L 139, 30.4.2004, p. 55). Commission Regulation (EC) No 2074/2005 of 5 December 2005 laying down implementing measures for certain products under Regulation (EC) No 853/2004 of the European Parliament and of the Council and for the organisation of official controls under (⁶) Regulation (EC) No 854/2004 of the European Parliament and of the Council and Regulation (EC) No 882/2004 of the European Parliament and of the Council, derogating from Regulation (EC) No 852/2004 of the European Parliament and of the Council and amending Regulations (EC) No 853/2004 and (EC) No 854/2004 (OJ L 338, 22.12.2005, p. 27).

- (4) Third countries, parts of third countries and territories listed in Annex II to Commission Decision 2006/766/EC (¹), in Part 1 of Annex I to Commission Regulation (EC) No 798/2008 (²), in Part 1 of Annex I to Commission Regulation (EC) No 119/2009 (³) or in Part 1 of Annex II to Commission Regulation (EU) No 206/2010 (⁴) meet the Union requirements with regard to imports of fresh meat and certain fishery products. Those lists could also be used for imports of raw materials for the production of gelatine and collagen. However, less strict requirements should apply if those raw materials have been subjected to certain treatments as provided for in Sections XIV and XV of Annex III to Regulation (EC) No 853/2004.
- (5) Raw materials for the production of gelatine and collagen, whether or not treated, introduced into the Union for transit to a third country, pose a negligible risk to public health. Such raw materials, even when treated, should, however, comply with the relevant animal health requirements. Accordingly, a list of third countries, parts of third countries and territories should be drawn up and model certificates for transit, and storage before transit, of raw materials and treated raw materials for the production of gelatine and collagen should be laid down.
- (6) Due to the geographical situation of Kaliningrad, specific animal health conditions should be laid down for transit via the Union of consignments of raw materials and treated raw materials for the production of gelatine or collagen to and from Russia, which only concern transit through Latvia, Lithuania and Poland.
- (7) In the interest of clarity and simplification of Union legislation, and without prejudice to Commission Decision 2003/863/EC (⁵), the lists of third countries, parts of third countries and territories from which Member States are to authorise the introduction of frogs' legs, snails, gelatine, collagen, raw materials and treated raw materials for the production of gelatine and collagen, and honey, royal jelly and other products of apiculture for human consumption, and the model certificates for those products, should be set out in an Annex to this Regulation. Consequently, the corresponding existing certificates should be deleted from Annex VI to Regulation (EC) No 2074/2005.
- (8) In order to ensure the safety of certain highly refined products of animal origin, specific requirements have been inserted in Annex III to Regulation (EC) No 853/2004. Therefore it is appropriate to draw up the list of countries from which those products may be imported and lay down a model certificate for those products.
- (9) As the lists of third countries, parts of third countries and territories from which Member States are to authorise imports of furred farm game meat products and feathered farm game meat products and leporidae (rabbit and hare) meat and their meat products have been laid down in Commission Decision 2007/777/EC (⁶) and in Regulation (EC) No 119/2009 respectively, Decision 2003/812/EC becomes redundant and should be repealed.
- (10) It is appropriate to introduce a transitional period to allow Member States and food business operators to adapt to the new requirements laid down in this Regulation.
- (11) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on Plants, Animals, Food and Feed,

^{(&}lt;sup>1</sup>) Commission Decision 2006/766/EC of 6 November 2006 establishing the lists of third countries and territories from which imports of bivalve molluscs, echinoderms, tunicates, marine gastropods and fishery products are permitted (OJ L 320, 18.11.2006, p. 53).

 ^{(&}lt;sup>2</sup>) Commission Regulation (EC) No 798/2008 of 8 August 2008 laying down a list of third countries, territories, zones or compartments from which poultry and poultry products may be imported into and transit through the Community and the veterinary certification requirements (OJ L 226, 23.8.2008, p. 1).
 (³) Commission Regulation (EC) No 119/2009 of 9 February 2009 laying down a list of third countries or parts thereof, for import into, or

^{(&}lt;sup>3</sup>) Commission Regulation (EC) No 119/2009 of 9 February 2009 laying down a list of third countries or parts thereof, for import into, or transit through, the Community of meat and wild leporidae, of certain wild land mammals and farmed rabbits and the veterinary certification requirements (OJ L 39, 10.2.2009, p. 12).

^(*) Commission Regulation (EU) No 206/2010 of 12 March 2010 laying down lists of third countries, territories or parts thereof authorised for the introduction into the European Union of certain animals and fresh meat and the veterinary certification requirements (OJ L 73, 20.3.2010, p. 1).

 ^{(&}lt;sup>5</sup>) Commission Decision 2003/863/EC of 2 December 2003 on health certificates for the importation of animal products from the United States of America (OJ L 325, 12.12.2003, p. 46).
 (⁶) Commission Decision 2007/777/EC of 29 November 2007 laying down the animal and public health conditions and model certificates

⁽⁶⁾ Commission Decision 2007/777/EC of 29 November 2007 laying down the animal and public health conditions and model certificates for imports of certain meat products and treated stomachs, bladders and intestines for human consumption from third countries and repealing Decision 2005/432/EC (OJ L 312, 30.11.2007, p. 49).

HAS ADOPTED THIS REGULATION:

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CHAPTER 1

IMPORTS OF CERTAIN PRODUCTS OF ANIMAL ORIGIN

Article 1

Lists of third countries, parts of third countries and territories

The third countries, parts of third countries and territories from which Member States are to authorise the import of the following products of animal origin intended for human consumption are set out in the relevant Parts of Annex I:

- (a) frogs' legs, Part I;
- (b) snails, Part II;
- (c) gelatine and collagen, Part III;
- (d) raw materials for the production of gelatine and collagen, Part IV;
- (e) treated raw materials for the production of gelatine and collagen, Part V;
- (f) honey, royal jelly and other products of apiculture, Part VI;
- (g) the following highly refined products, Part VII:
 - (i) chondroitin sulphate;
 - (ii) hyaluronic acid;
 - (iii) other hydrolysed cartilage products;
 - (iv) chitosan;
 - (v) glucosamine;
 - (vi) rennet;
 - (vii) isinglass;
 - (viii) amino acids that are authorised as food additives in accordance with Regulation (EC) No 1333/2008 of the European Parliament and of the Council (¹).

Article 2

Model certificates

1. The model certificates for imports into the Union of the products referred to in Article 1 are set out in Annex II as follows:

- (a) frogs' legs, Part I;
- (b) snails, Part II;
- (c) gelatine, Part III;
- (d) collagen, Part IV;
- (e) raw materials for the production of gelatine and collagen, Part V;
- (f) treated raw materials for the production of gelatine and collagen, Part VI;
- (g) honey, royal jelly and other products of apiculture, Part VII;
- (h) the following highly refined products, Part VIII:
 - (i) chondroitin sulphate;
 - (ii) hyaluronic acid;

^{(&}lt;sup>1</sup>) Regulation (EC) No 1333/2008 of the European Parliament and of the Council of 16 December 2008 on food additives (OJ L 354, 31.12.2008, p. 16).

(iii) other hydrolysed cartilage products;

- (iv) chitosan;
- (v) glucosamine;
- (vi) rennet;
- (vii) isinglass;
- (viii) amino acids that are authorised as food additives in accordance with Regulation (EC) No 1333/2008.

Those certificates must be completed in accordance with the explanatory notes set out in Annex IV and the notes in the relevant certificate.

2. Electronic certification and other systems agreed between the Union and the third country concerned may be used.

CHAPTER 2

TRANSIT OF CERTAIN PRODUCTS OF ANIMAL ORIGIN

Article 3

Lists of third countries, parts of third countries and territories

The third countries, parts of third countries and territories from which Member States are to authorise the transit through the Union of raw materials and treated raw materials for the production of gelatine and collagen intended for human consumption bound for a third country, either by immediate transit or after storage in the Union in accordance with Article 12(4) and Article 13 of Council Directive 97/78/EC (¹), are set out in Parts IV and V of Annex I to this Regulation, respectively.

Article 4

Model certificate

1. The model certificate for the transit through the Union of the raw materials and treated raw materials referred to in Article 3 is set out in Annex III.

That certificate must be completed in accordance with the notes set out in Annex IV and in the relevant model certificate.

2. Electronic certification and other systems harmonised at Union level may be used.

Article 5

Derogation for transit through Latvia, Lithuania and Poland

1. By way of derogation from Article 3, transit by road or by rail between the specific, designated border inspection posts in Latvia, Lithuania and Poland, listed and marked with special remark 13 in Annex I to Commission Decision 2009/821/EC (²), of consignments of the raw materials or treated raw materials referred to in Article 3 of this Regulation coming from and bound for Russia, directly or via another third country, shall be authorised where the following conditions are met:

(a) the consignment is sealed with a serially numbered seal by the official veterinarian at the border inspection post of entry;

 ^{(&}lt;sup>1</sup>) Council Directive 97/78/EC of 18 December 1997 laying down the principles governing the organisation of veterinary checks on products entering the Community from third countries (OJ L 24, 30.1.1998, p. 9).
 (²) Commission Decision 2009/821/EC of 28 September 2009 drawing up a list of approved border inspection posts, laying down certain

^{(&}lt;sup>2</sup>) Commission Decision 2009/821/EC of 28 September 2009 drawing up a list of approved border inspection posts, laying down certain rules on the inspections carried out by Commission veterinary experts and laying down the veterinary units in Traces (OJ L 296, 12.11.2009, p. 1).

- (b) the documents accompanying the consignment, as provided for in Article 7 of Directive 97/78/EC, are stamped with the words 'Only for transit to Russia via the EU' on each page by the official veterinarian at the border inspection post of entry;
- (c) the procedural requirements provided for in Article 11 of Directive 97/78/EC are complied with;
- (d) the consignment is certified as acceptable for transit on the common veterinary entry document issued by the official veterinarian at the border inspection post of entry.

2. The consignments referred to in paragraph 1 shall not be unloaded or put into storage, as referred to in Article 12(4) or in Article 13 of Directive 97/78/EC, within the Union.

3. Regular audits shall be conducted by the competent authority to ensure that the number of consignments referred to in paragraph 1 and the corresponding quantities of products leaving the Union correspond with the number and quantities which have been introduced in the Union.

CHAPTER 3

FINAL PROVISIONS

Article 6

Amendment

Annex VI to Regulation (EC) No 2074/2005 is amended as follows:

(1) in Section I, Chapters I, II, III and VI are deleted;

(2) Appendices I, II, III and VI are deleted.

Article 7

Repeal

Decision 2003/812/EC is repealed.

Article 8

Transitional provisions

Consignments of products of animal origin in respect of which the relevant certificates have been issued in accordance with Regulation (EC) No 2074/2005 may continue to be introduced into the Union provided that the certificate was signed before 3 December 2016.

Article 9

Entry into force

This Regulation shall enter into force on the twentieth day following that of 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 April 2016.

For the Commission The President Jean-Claude JUNCKER

ANNEX I

Lists of third countries, parts of third countries and territories as referred to in Article 1

PART I

FROGS' LEGS

Third countries and territories listed in the column 'Countries' of Annex II to Decision 2006/766/EC, except those for which a restriction is mentioned in the column 'Restrictions' of that Annex, and the following countries or territories:

COUNTRY ISO CODE	COUNTRY/TERRITORY
MK (*)	former Yugoslav Republic of Macedonia

(*) The former Yugoslav Republic of Macedonia; provisional code that does not prejudge in any way the definitive nomenclature for this country, which will be agreed following the conclusion of negotiations currently taking place on this subject in the United Nations.

PART II

SNAILS

Third countries and territories listed in the column 'Countries' of Annex II to Decision 2006/766/EC, except those for which a restriction is mentioned in the column 'Restrictions' of that Annex, and the following countries/territories:

COUNTRY ISO CODE	COUNTRY/TERRITORY
MD	Moldova
MK (*)	former Yugoslav Republic of Macedonia
SY	Syria

(*) The former Yugoslav Republic of Macedonia; provisional code that does not prejudge in any way the definitive nomenclature for this country, which will be agreed following the conclusion of negotiations currently taking place on this subject in the United Nations.

PART III

GELATINE AND COLLAGEN INTENDED FOR HUMAN CONSUMPTION

SECTION A

Gelatine and collagen derived from bovine, ovine, caprine, porcine and equine animals, both farmed and wild

Third countries and territories listed in column 1 of Part 1 of Annex II to Regulation (EU) No 206/2010 and the following countries or territories:

COUNTRY ISO CODE	COUNTRY/TERRITORY			
KR Republic of Korea				
МҮ	Malaysia			
РК	Pakistan			
TW	Taiwan			

SECTION B

Gelatine and collagen derived from poultry including ratites and feathered game

Third countries and territories listed in column 1 of Part 1 of Annex I to Regulation (EC) No 798/2008.

SECTION C

Gelatine and collagen derived from fishery products

All third countries and territories listed in the column 'Countries' of Annex II to Decision 2006/766/EC, regardless of whether a restriction is mentioned in the column 'Restrictions' of that Annex.

SECTION D

Gelatine and collagen derived from leporidae and from wild land mammals not referred to in Section A

Third countries listed in column 1 of Part 1 of Annex I to Regulation (EC) No 119/2009.

PART IV

RAW MATERIALS FOR THE PRODUCTION OF GELATINE AND COLLAGEN INTENDED FOR HUMAN CONSUMPTION

SECTION A

Raw materials from bovine, ovine, caprine, porcine and equine animals, both farmed and wild

Third countries, territories and parts thereof listed in Part 1 of Annex II to Regulation (EU) No 206/2010 from which introduction into the Union of that category of fresh meat of the respective species is authorised as specified in that Part of that Annex, unless such introduction is limited by supplementary guarantees A or F as indicated in column 5.

SECTION B

Raw materials from poultry including ratites and feathered game

Third countries, parts of third countries and territories listed in Part 1 of Annex I to Regulation (EC) No 798/2008 from which imports of fresh poultry meat of the respective species is authorised as specified in that Part of that Annex.

SECTION C

Raw materials from fishery products

Third countries and territories listed in the column 'Countries' of Annex II to Decision 2006/766/EC, subject to the restrictions mentioned in the column 'Restrictions' of that Annex.

SECTION D

Raw materials from leporidae and from wild land mammals not referred to in Section A

Third countries listed in column 1 of Part 1 of Annex I to Regulation (EC) No 119/2009 from which imports of fresh meat of the respective species is authorised as specified in that Part of that Annex.

PART V

TREATED RAW MATERIALS FOR THE PRODUCTION OF GELATINE AND COLLAGEN INTENDED FOR HUMAN CONSUMPTION

SECTION A

Treated raw materials from bovine, ovine, caprine, porcine and equine animals, both farmed and wild

Third countries and territories and parts thereof listed in column 1 of Part 1 of Annex II to Regulation (EU) No 206/2010 and the following countries or territories:

COUNTRY ISO CODE	COUNTRY/TERRITORY
KR	Republic of Korea
МҮ	Malaysia
РК	Pakistan
TW	Taiwan

SECTION B

Treated raw materials from poultry including ratites and feathered game

Third countries and territories listed in column 1 of Part 1 of Annex I to Regulation (EC) No 798/2008.

SECTION C

Treated raw materials from fishery products

All third countries and territories listed in the column 'Countries' of Annex II to Decision 2006/766/EC regardless of whether a restriction is mentioned in the column 'Restrictions' of that Annex.

SECTION D

Treated raw materials from leporidae and wild land mammals not referred to in Section A

Third countries listed in column 1 of Part 1 of Annex I to Regulation (EC) No 119/2009.

SECTION E

Treated raw materials referred to in Annex III to Regulation (EC) No 853/2004, Section XIV, Chapter I point 4(b)(iii) and Section XV, Chapter I, point 4(b)(iii)

Third countries, parts of third countries and territories referred to in Part IV of this Annex.

PART VI

HONEY, ROYAL JELLY AND OTHER PRODUCTS OF APICULTURE INTENDED FOR HUMAN CONSUMPTION

Third countries and territories listed in the column 'Country' in the Annex to Commission Decision 2011/163/EU (¹) and marked with an 'X' in the column 'Honey' in that Annex.

^{(&}lt;sup>1</sup>) Commission Decision 2011/163/EU of 16 March 2011 on the approval of plans submitted by third countries in accordance with Article 29 of Council Directive 96/23/EC (OJ L 70, 17.3.2011, p. 40).

PART VII

HIGHLY REFINED CHONDROITIN SULPHATE, HYALURONIC ACID, OTHER HYDROLYSED CARTILAGE PRODUCTS, CHITOSAN, GLUCOSAMINE, RENNET, ISINGLASS AND AMINO ACIDS FOR HUMAN CONSUMPTION

(a) In the case of raw materials derived from ungulates including equidae, third countries and territories listed in column 1 of Part 1 of Annex II to Regulation (EU) No 206/2010 and the following countries or territories:

COUNTRY ISO CODE	COUNTRY/TERRITORY			
KR	Republic of Korea			
МҮ	Malaysia			
РК	Pakistan			
TW	Taiwan			

- (b) In the case of the raw materials derived from fishery products, all third countries and territories listed in the column 'Countries' in Annex II to Decision 2006/766/EC, regardless of whether a restriction is mentioned in the column 'Restrictions' of that Annex.
- (c) In the case of raw materials derived from poultry, third countries and territories listed in column 1 of Part 1 of Annex I to Regulation (EC) No 798/2008.

ANNEX II

Model certificates as referred to in Article 2

PART I

MODEL CERTIFICATE FOR IMPORTS OF CHILLED, FROZEN OR PREPARED FROGS' LEGS INTENDED FOR HUMAN CONSUMPTION

COUNTRY:

Veterinary certificate to EU

	l.1.	Consignor		I.2. Certificate reference No I.2.a.			
		Name	I.3. Central competent authority				
		Address	I.4. Local competent authority				
		Tel.					
t.	1.5.	Consignee	1.6.				
nen		Name					
ign		Address			/		
ons							
s c		Postcode					
tche		Tel.					
Part I: Details of dispatched consignment				<u> </u>			
of d	I.7.	Country of ISO code I.8. origin	1.9.	Country of destination	ISO cod	e I.10.	
ails							
Det							
ü t	I.11.	Place of origin	I.12.				
ъ В							
		Name Approval number					
		Address					
	113	Place of loading	114	Date of depar	ture		
	1.10.		1. 1 7.	Date of depar			
	I.15.	Means of transport	I.16.	Entry BIP in E	U		
		Aeroplane 🛛 Ship 🖾 Railway wagon 🗆					
		Road vehicle	I.17.				
		Identification					
		Documentation references					
	l.18.	Description of commodity	_		I.19. Commo	dity code (HS code)	
						02.08.90	
					· · · · · · · · · · · · · · · · · · ·	.20. Quantity	
	121	Temperature of product				.22. Number of	
	1.21.	Ambient Chilled	1	Frozen 🗖		packages	
	I.23.	Seal/Container No				.24. Type of packaging	

I.25. Commodities c	ertified for:				
Human consum	nption 🗖				
1.26.			I.27. For i	mport or admission into	EU 🗖
I.28. Identification of	the commodities				
Species (scientific name)	Treatment type	Approval r establish Manufactu	nments	Number of packages	Net weight

EN COUNTRY Model FRG Frogs' legs II. Health information II.a. Certificate reference No II.b. II.1. **Public Health Attestation** I, the undersigned, declare that I am aware of the relevant provisions of Regulation (EC) No 178/2002 of the European Parliament and of the Council of 28 January 2002 laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety (OJ L 31, 1.2.2002, p. 1), Regulation (EC) No 852/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs (OJ L 139, 30.4.2004, p. 1) and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying Part II: Certification down specific hygiene rules for food of animal origin (OJ L 139, 30.4.2004, p. 55) and certify that the frogs' legs described above were produced in accordance with those requirements, in particular that they: come from (an) establishment(s) implementing a programme based on the HACCP principles in accordance with Article 5 of Regulation (EC) No 852/2004; and originate from frogs that have been bled, prepared and, where appropriate, chilled, frozen or processed, packaged and stored in a hygienic manner in accordance with the requirements of Section XI of Annex III to Regulation (EC) No 853/2004. Notes Part I: Box reference I.11: Place of origin: name and address of the dispatch establishment. Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship). Separate information is to be provided in the event of unloading and reloading. Box reference I.20: Indicate total gross weight and total net weight. - Box reference I.23: Identification of container/Seal number: only where applicable. - Box reference I.28: Treatment type: fresh, treated. Part II: The colour of the stamp and signature must be different from that of the other particulars in the certificate. Official inspector Name (in capital letters): Qualification and title: Date: Signature: Stamp:

PART II

MODEL CERTIFICATE FOR IMPORTS OF CHILLED, FROZEN, SHELLED, COOKED, PREPARED OR PRESERVED SNAILS INTENDED FOR HUMAN CONSUMPTION

COUNTRY:

Veterinary certificate to EU

	l.1.	Consignor	I.2. Certificate reference No I.2.a.		
		Name Address	I.3. Central competent authority		
		Address	I.4. Local competent authority		
		Tel.			
	1.5.	Consignee	1.6.		
lent		Name			
nsignn		Address			
ioo be		Postcode			
atche		Tel.			
Part I: Details of dispatched consignment	1.7.	Country of ISO code I.8. origin	I.9. Country of ISO code I.10. destination		
Detai					
Part I:	l.11.	Place of origin	1.12.		
		Name Approval number			
		Address			
	l.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 D Other D	l.17.		
		Identification			
	Documentation references				
	l.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		
	1.23.	Seal/Container No	I.24. Type of packaging		

I.25. Commodities certified for:		
Human consumption 🗖		
1.26.		I.27. For import or admission into EU
I.28. Identification of the commoditie	S	
Species (scientific name)	Treatment type	Approval number of establishments
Number of packages	Net weight	
Manufacturing plant		

Official Journal of the European Union EN COUNTRY Model SNS Snails II. Health information II.a. Certificate reference No II.b II.1. **Public Health Attestation** I, the undersigned, declare that I am aware of the relevant provisions of Regulation (EC) No 178/2002 of the European Parliament and of the Council of 28 January 2002 laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety (OJ L 31, 1.2.2002, p. 1), Regulation (EC) No 852/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs (OJ L 139, 30.4.2004, p. 1) and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying Part II: Certification down specific hygiene rules for food of animal origin (OJ L 139, 30.4.2004, p. 55) and certify that the snails described above were produced in accordance with those requirements, in particular that they: come from (an) establishment(s) implementing a programme based on the HACCP principles in accordance with Article 5 of Regulation (EC) No 852/2004; and have been handled and, where appropriate, shelled, cooked, prepared, preserved, frozen, packaged and stored in a hygienic manner in accordance with the requirements of Section XI of Annex III to Regulation (EC) No 853/2004. Notes Part I: Box reference I.11: Place of origin: name and address of the dispatch establishment. Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship). Separate information is to be provided in case of unloading and reloading. Box reference I.19: Use the appropriate Harmonised System (HS) code under the following headings: 03.07, 16.05. Box reference I.20: Indicate total gross weight and total net weight. - Box reference I.23: Identification of container/Seal number: only where applicable. - Box reference I.28: Treatment type: fresh, treated. Part II: The colour of the stamp and signature must be different from that of the other particulars in the certificate. Official inspector Name (in capital letters): Qualification and title: Date: Signature: Stamp:

PART III

MODEL CERTIFICATE FOR IMPORTS OF GELATINE INTENDED FOR HUMAN CONSUMPTION

COUNTRY:

Veterinary certificate to EU

	l.1.	Consignor	I.2. Ce	rtificate refe	erence No	l.2.a.	
		Name Address	I.3. Central competent authority				
				I.4. Local competent authority			
		Tel.					
	1.5.	Consignee	1.6.				
Jen		Name					
Isignn		Address					
COL							
hed		Postcode					
patc		Tel.					
Part I: Details of dispatched consignment	I.7.	Country of ISO code I.8. origin	I.9. Co des	untry of stination	ISO co	de I.10.	
: Deta							
art I	I.11.	Place of origin	l.12.				
e							
		Name Approval number			_		
		Address					
	I.13.	Place of loading	l.14. Da	te of depart	ure		
	l.15.	Means of transport	I.16. Ent	ry BIP in El	U		
		Aeroplane □ Ship □ Railway wagon □					
		Road vehicle Other	l.17.				
		Identification					
		Documentation references					
	l.18.	Description of commodity			I.19. Comm	odity code (HS code)	
						I.20. Quantity	
	I.21.	Temperature of product			T	I.22. Number of	
		Ambient Chilled	Froz	en 🗖		packages	
	I.23.	Seal/Container No				I.24. Type of packaging	

I.25.	Commodities cert	tified for:					
	Human consump	tion 🗖					
1.26.				1.27.	For import or admissior	n into EU 🛛 🗖	
1.28.	Identification of th	ne commodities					
(sc	Species cientific name)	Date production (dd/mm/yyyy)	Approval r establisł Manufactu	nments	Number of packages	Net weight	

	Model GE Gelatine intended for human consumptio									
	П.	Heal	th information	II.a.	Certificate reference No	II.b.				
	II.1.									
Part II: Certification		I, the undersigned, declare that I am aware of the relevant provisions of Regulation (EC) No 178/200 the European Parliament and of the Council of 28 January 2002 laying down the general principles requirements of food law, establishing the European Food Safety Authority and laying down proced in matters of food safety (OJ L 31, 1.2.2002, p. 1), Regulation (EC) No 852/2004 of the Europ Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs (OJ L 139, 30.4.2004, and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 la down specific hygiene rules for food of animal origin (OJ L 139, 30.4.2004, p. 55) and certify that gelatine described above was produced in accordance with those requirements, in particular that:								
Part II: C		d on the HACCP principles in								
-		—	it has been produced from Section XIV of Annex III to R		naterials that met the requireme n (EC) No 853/2004;	nts of Chapters I and II of				
		_	it has been manufactured in Annex III to Regulation (EC)	complia No 853/	ance with the conditions set out in 2004;	Chapter III of Section XIV of				
		_		EC) No	of Section XIV of Annex III to Regu 2073/2005 of 15 November 2005 1);					
		and i	f from ruminant origin, except	for gela	tine derived from hides and skins o	of ruminants,				
		(1) eit	ther							
		_	No 999/2001 of the Europea the prevention, control and	n Parlia d eradio	n classified in accordance with A ment and of the Council of 22 Ma cation of certain transmissible s ntry or region posing a negligible B	ay 2001 laying down rules for pongiform encephalopathies				
		—			was derived were born, continuou and passed ante-mortem and post-					
		—	if in the country or region the	re have	been BSE indigenous cases:					
					vere born after the date from whi meal and greaves derived from r					
(ii) the products of bovine, ovine and caprine animal origin do not contain and are no from specified risk material as defined in Annex V to Regulation (EC) No 999, mechanically separated meat obtained from bones of bovine, ovine or caprine animal										
		(¹) or								
		 [it comes from a country or a region classified in accordance with Article 5(2) of Regulation (E No 999/2001 as a country or region posing a controlled BSE risk; 								
		— the animals from which the gelatine was derived passed ante-mortem and post-mortem inspection								
		_	stunning by means of gas	injecte ter stun	destined for export was derived ha d into the cranial cavity or kille ning of central nervous tissue by e cranial cavity;	ed by the same method or				
		_			not derived from specified risk mat nechanically separated meat obta					

со	UNTRY				Gelatine i	ntended for h	Model GEL numan consumption
П.	Hea	th inform	ation	II.a.	Certificate reference No	II.b.	
	(¹) 0	r				-	
	—				n classified in accordance v posing a negligible BSE risk;		2) of Regulation (EC)
	_	the gelati	ne is derived fr	rom animals v	which passed ante-mortem a	and post-morte	em inspections;
	_	region wi if there h the ban o had been Article 5(and whic or killed b	th negligible B ave been BSE on the feeding n enforced, an 2) of Regulation h have not bee by the same mo	SE risk in acc i indigenous of ruminants d from anim on (EC) No s in slaughtere ethod or slau	mals born, continuously real cordance with Article 5(2) of cases in the country or reg with meat-and-bone meal a als born in a country or re 099/2001 as a country or re d after stunning by means o ghtered by laceration after s instrument introduced into the	Regulation (Ē ion, born after nd greaves de gion classified gion posing a f gas injected tunning of cen	C) No 999/2001, and, the date from which erived from ruminants d in accordance with controlled BSE risk, into the cranial cavity tral nervous tissue by
	_	Regulatio		9/2001, or r	not derived from specified ris nechanically separated mea		
	(¹) 01	-					
	—				n classified in accordance v vith an undetermined BSE ri		2) of Regulation (EC)
	—				was derived have not been f ante-mortem and post-morte		
	_	means o laceration	f gas injected	into the cra of central no	e was derived have not b nial cavity or killed by the ervous tissue by means of a	same metho	od or slaughtered by
	—	the gelati	ne is not derive	ed from:			
		(i) spe	cified risk mate	erial as define	ed in Annex V to Regulation	(EC) No 999/2	2001;
		(ii) ner	vous and lymp	hatic tissues	exposed during the debonin	g process;	
		(iii) me	chanically sepa	arated meat c	btained from bones of bovin	le, ovine or ca	prine animals.]
Not	es						
Par	t I:						
—	Box refe	erence I.11	: Place of origi	n: name and	address of the dispatch esta	ablishment.	
					vay wagons or container an vided in case of unloading a		nt number (aircraft) or
—	Box refe	erence I.19	: Use the appr	opriate Harm	onised System (HS) code ur	nder the headi	ng of 35.03.
—	Box refe	erence I.20	: Indicate total	gross weight	and total net weight.		
_	Box refe	erence I.23	: Identification	of container/	Seal number: only where ap	plicable.	

со	UNTRY		Gelatine inte	Model GEL ended for human consumption
П.	Health information	II.a.	Certificate reference No	II.b.
Par	t II:			
(1)	Delete as appropriate.			
-	The colour of the stamp and signature	e must be	e different from that of the other	r particulars in the certificate.
Offi	cial veterinarian			Name (in capital letters):
	Qualification and title:			
	Date:			Signature:
	Stamp:			

PART IV

MODEL CERTIFICATE FOR IMPORTS OF COLLAGEN INTENDED FOR HUMAN CONSUMPTION

COUNTRY:

Veterinary certificate to EU

	l.1.	Consignor	1.2.	Certificate refe	erence No	l.2.a.
		Name Address	1.3.	Central compe	etent authority	
		Address	I.4.	Local compete	ent authority	
		Tel.				
	1.5.	Consignee	I.6.			
lent		Name				
nsignm		Address				
ed cor		Postcode				
atche		Tel.				
Part I: Details of dispatched consignment	I.7.	Country of ISO code I.8. origin	I.9.	Country of destination	ISO co	de I.10.
: Deta						
Part	l.11.	Place of origin	I.12.			
		Name Approval number				
		Address				
			/			
	I.13.	Place of loading	I.14.	Date of depar	ture	
	l.15.	Means of transport	I.16.	Entry BIP in E	U	
		Aeroplane □ Ship □ Railway wagon □				
		Road vehicle	I.17.			
		Identification				
		Documentation references				
	l.18.	Description of commodity			I.19. Comm	odity code (HS code)
				l		I.20. Quantity
	l.21.	Temperature of product Ambient Chilled	I	=rozen 🗖		I.22. Number of packages
	I.23.	Seal/Container No				I.24. Type of packaging

I.25. Commodities certified for:		
Human consumption 🗖		
1.26.	I.27. For import o	or admission into EU 🛛 🗖
I.28. Identification of the commodities		
(scientific name) (dd/mm/uuuu)	establistifients	lumber of Net weight backages

	COUN	TRY				Collagen inten	Mode ded for human consur	el COL mption
	II.	Heal	lth inf	ormation	II.a.	Certificate reference No	II.b.	
	II.1.	Pub	lic He	alth Attestation				
Part II: Certification		the E requ in m Parli and dow	Europ iireme natters lamen Regul n spe	ean Parliament and of nts of food law, estal of food safety (OJ t and of the Council lation (EC) No 853/20 cific hygiene rules fo	of the Cour blishing the L 31, 1.2. of 29 April 004 of the E r food of a	e of the relevant provisions of Re icil of 28 January 2002 laying do European Food Safety Authority 2002, p. 1), Regulation (EC) N 2004 on the hygiene of foodstuff European Parliament and of the C nimal origin (OJ L 139, 30.4.200 accordance with those requirem	wn the general principl y and laying down proc o 852/2004 of the Eu fs (OJ L 139, 30.4.2004 Council of 29 April 2004 04, p. 55) and certify tl	es and edures ropean 4, p. 1) Haying
art II: C		—				implementing a programme base on (EC) No 852/2004;	d on the HACCP princ	iples in
a .		—		as been produced fi ion XV of Annex III to		naterials that met the requirement (EC) No 853/2004;	ents of Chapters I an	d II of
		—		s been manufactured ex III to Regulation (E		ance with the conditions set out i 2004;	n Chapter III of Sectior	ı XV of
		_	of C		n (ĖC) No :	of Section XV of Annex III to Reg 2073/2005 of 15 November 2005 1);		
		and	if fron	n ruminant origin, exc	ept for colla	igen derived from hides and skins	of ruminants,	
		(1) ei	ither					
		_	No 9 the	99/2001 of the Europ prevention, control	bean Parlia and eradic	n classified in accordance with A ment and of the Council of 22 M ation of certain transmissible ntry or region posing a negligible l	ay 2001 laying down ru spongiform encephalor	ules for
		—				was derived were born, continuo and passed ante-mortem and pos		ered in
		—	if in t	the country or region	there have	been BSE indigenous cases:		
			(i)			vere born after the date from wh meal and greaves derived from		
			(ii)	from specified risk	material a	and caprine animal origin do no is defined in Annex V to Regu ibtained from bones of bovine, ov	lation (EC) No 999/20	001, or
		(¹) or	r					
		—				n classified in accordance with A posing a controlled BSE risk;	rticle 5(2) of Regulatio	n (EC)
		—		animals from whic ections;	n the coll	agen was derived passed an	te-mortem and post-r	nortem
		_	after slau	stunning by means	of gas inje after stun	n destined for export were derive ected into the cranial cavity or k ning of central nervous tissue by e cranial cavity;	villed by the same met	thod or
		—	to R		9/2001, or	not derived from specified risk n mechanically separated meat of		

CO	UNTRY				Collagen intend	Model COL ed for human consumption
II.	Heal	th inf	ormation	II.a.	Certificate reference No	ll.b.
	(¹) 0I	ſ				
	—				n classified in accordance with Ar osing a negligible BSE risk;	ticle 5(2) of Regulation (EC)
	_	the o	collagen is derived from	animals	which passed ante-mortem and po	ost-mortem inspections;
	_	region if the the l had Artic and or ki	on with negligible BSE ri ere have been BSE indi ban on the feeding of ru been enforced, and fro ble 5(2) of Regulation (E which have not been sla lled by the same method	sk in acc igenous minants om anima EC) No 9 aughtered d or slaug	mals born, continuously reared an ordance with Article 5(2) of Regul cases in the country or region, bo with meat-and-bone meal and gre als born in a country or region o 99/2001 as a country or region p d after stunning by means of gas i ghtered by laceration after stunnin instrument introduced into the crani	ation (EC) No 999/2001, and, orn after the date from which eaves derived from ruminants classified in accordance with posing a controlled BSE risk, njected into the cranial cavity g of central nervous tissue by
	—	to R			not derived from specified risk m mechanically separated meat ob	
	(1) or	•				
	_				n classified in accordance with Ar vith an undetermined BSE risk;	ticle 5(2) of Regulation (EC)
	—				en was derived have not been passed ante-mortem and post-mo	
	_	mea lace	ns of gas injected into	the cra	n was derived have not been s nial cavity or killed by the same ervous tissue by means of an elon	e method or slaughtered by
	—	the o	collagen was not derived	l from:		
		(i)	specified risk material	as define	d in Annex V to Regulation (EC) N	lo 999/2001;
		(ii)	nervous and lymphatic	tissues e	exposed during the deboning proc	ess;
		(iii)	mechanically separate	d meat o	btained from bones of bovine, ovir	ne or caprine animals.]
Not	es					
Par						
_			-		address of the dispatch establishn	
_					vay wagons or container and lorrie vided in case of unloading and rel	
—	Box refe	erence	e I.18: This certificate ma	ay also b	e used for import of collagen casir	igs.
—	Box refe	erence	e I.19: Use the appropria	ite Harmo	onised System (HS) code under th	he heading of 35.04 or 39.17.
—	Box refe	erence	e I.20: Indicate total gros	s weight	and total net weight.	
_	Box refe	erence	e I.23: Identification of co	ontainer/s	Seal number: only where applicabl	e.

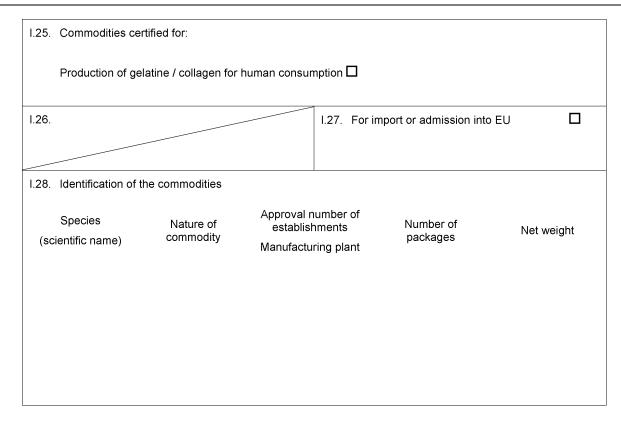
COL	INTRY		Collagen int	Model COL ended for human consumption
11.	Health information	II.a.	Certificate reference No	II.b.
Part	11:			
(¹)	Delete as appropriate.			
_	The colour of the stamp and signature	e must be	e different from that of the othe	er particulars in the certificate.
Offic	ial veterinarian			
	Name (in capital letters):			Qualification and title:
	Date:			Signature:
	Stamp:			

PART V

MODEL CERTIFICATE FOR IMPORTS OF RAW MATERIALS FOR THE PRODUCTION OF GELATINE/ COLLAGEN INTENDED FOR HUMAN CONSUMPTION $(^{\rm n})$

COL	INTRY	1				Veterinary certificate to EU			
	l.1.	Consignor Name Address				I.2. I.3.	Certificate re	eference No petent autho	I.2.a.
		Tel.				1.4.	Local compe	etent authorit	у
Part I: Details of dispatched consignment	I.5.	Consignee Name Address Postcode Tel.				1.6.			
Details of dis	I.7.	Country of origin	ISO code	I.8. Region of origin	Code	1.9.	Country of destination	ISO code	1.10.
Part I: I	I.11.	Place of or Name Address	igin	Approval nur	nber	I.12.			
	l.13.	Place of loa	ading			I.14.	Date of depa	arture	
	I.15.	Means of the Aeroplane		Ship 🗖 🛛 Ri	ailway	I.16.	Entry BIP in	EU	
		wagon D Road vehic Identificatio Documenta		Other 🗖		1.17.			
	l.18.	Description	of commo	dity				I.19. Comn	nodity code (HS code)
									I.20. Quantity
	I.21.	Temperatu Ambient 🗖		ct Chilled 🗖	l		Frozen 🗖		I.22. Number of packages
	I.23.	Seal/Conta	iner No						I.24. Type of packaging

⁽¹⁾ Unless covered by Part VI.



COUNTRY

	П.	Healt	h information	II.a.	Certificate reference No	ll.b.
	II.1.	Publi	c Health Attestation			
Part II: Certification		the Eu requir in ma Parlia Regul specif No 85 for the (OJ L requir	uropean Parliament and of the ements of food law, establis itters of food safety (OJ L ment and of the Council of 2 lation (EC) No 853/2004 of the ic hygiene rules for food of 4/2004 of the European Par e organisation of official cou- 139, 30.4.2004, p. 206) and ements, in particular that: (¹) [bones, hides and skins of and sinews described above	ne Counc hing the E 31, 1.2.2 9 April 20 e Europe of animal liament a ntrols on d certify f domestic derive fro	of the relevant provisions of Reg il of 28 January 2002 laying dow European Food Safety Authority 002, p. 1), Regulation (EC) No 004 on the hygiene of foodstuffs an Parliament and of the Council origin (OJ L 139, 30.4.2004,] nd of the Council of 29 April 200 products of animal origin inten- that the raw materials describe c and farmed ruminant animals,] om animals which have been slate of found fit for human consumpt	n the general principles and and laying down procedures \$52/2004 of the European (OJ L 139, 30.4.2004, p. 1), of 29 April 2004 laying down p. 55) and Regulation (EC) 04 laying down specific rules ded for human consumption ad above comply with those pigs and poultry and tendons ughtered in a slaughterhouse
			and/or			
					described above derive from kill ption following post-mortem insp	
			and/or			
			(¹) [fish skins and bones de human consumption authoris		above derive from plants manuf port,]	acturing fishery products for
		(¹) and	b			
		[if fror	n ruminant origin, except for	hides and	skins of ruminants,	
		(¹) eith	ner:			
			No 999/2001 of the Europea the prevention, control and	n Parliam 1 eradica	a classified in accordance with A nent and of the Council of 22 Ma tion of certain transmissible s ry or region posing a negligible B	y 2001 laying down rules for pongiform encephalopathies
				ed and sl	rials of bovine, ovine and caprin aughtered in the country with ne ions;	
		—	if in the country or region the	re have b	een BSE indigenous cases:	
					e date from which the ban on th s derived from ruminants had be	
			derived from specified	risk mater	vine and caprine animal origin ial as defined in Annex V to Reg tained from bones of bovine, ovir	ulation (EC) No 999/2001, or
		(1) or:				
			[they come from a country o No 999/2001 as a country or		a classified in accordance with A using a controlled BSE risk;	rticle 5(2) of Regulation (EC)
			the animals from which the passed ante-mortem and pos		rials of bovine, ovine and caprin inspections;	e animal origin were derived

COUNTRY

Health Information II.a. Certificate reference No ILb - animals from which the raw materials of bovine, ovine and caprine animal origin intended for export were derived have not been slaughtered after stunning by means of gas injected into the cranal cavity or killed by the same method or slaughtered by laceration after stunning of certral nervous tissue by means of an elongade tord-shaped instrument introduced into the cranal cavity. - the raw materials of bovine, ovine and caprine animal origin do not contain and are not derived from specified ram at obtained from bones of bovine, ovine or caprine animals.] (!) or - [they come from a country or a region classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region posing a negligible BSE risk: - the animals from which the raw materials of bovine, ovine and caprine animal origin intended for export are derived both from animals which were born, continuously reared and slaughtered in a country or region with negligible BSE risk in accordance with Article 5(2) of Regulation (EC) No 999/2001, and, if there have been BSE indigenous cases in the country or region, bom after the date from which the ban on the feeding of runniants with meat-and-bone meal and greaxes derived form which the ban on the feeding of runniants with meat-and-bone meal and greaxes derived from envous tissue by means of an elongade rod-shaped instrument introduced into the cranial cavity or will dive by as method or slaughtered by laceration after thurning of eartral nervous tissue by means of an elongade rod-shaped instrument introduced into the cranial cavity. - the raw materials of bovine, ovine art caprine				conagen intend	
 were derived have not been slaughtered after stuming by means of gas injected into the cranial cavity. the raw materials of bovine, ovine and caprine animal origin do not contain and are not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001, or mechanically separated meat obtained from bones of bovine, ovine or caprine animals.] (!) or [(!) or the animals from which the raw materials of bovine, ovine and caprine animal origin were derived passed ante-mortem and post-mortem inspections; the animals from which the raw materials of bovine, ovine and caprine animal origin intended for export are derived both rong animals which were born, continuously reared and slaughtered in a country or region with negligible BSE risk in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or normal may which were born, continuously reared and slaughtered in a country or region with negligible BSE risk in accordance with Article 5(2) of Regulation (EC) No 999/2001, and, if there have been BSE indigenous cases in the country or region posing a controled BSE risk, and which have not been slaughtered for subming by means of gas inglected into the cranial cavity or killed by the same method or slaughtered by laceration after stuming of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity; the raw materials of bovine, ovine and caprine animal origin do not contain and are not derived from twich they not been slaughtered and slaughtered the scale in accordance with Article 5(2) of Regulation (EC) No 999/2001, or mechanically separated meat obtained from bones of bovine, ovine or caprine paing a controled BSE risk, and which have not been slaughtered after stuming by means of gas inglected into the cranial cavity or killed by the same method or slaughtered at the cannial acting in the row materials of bovine, ovine and caprine animal origin were derived frawe not b	II.	Hea	Ith information	II.a. Certificate reference No	II.b.
 specified risk material as defined in Annex V to Regulation (EC) No 999/2001, or mechanically separated meat obtained from bones of bovine, ovine or caprine animals.] (1) or [(1) or [(1) the provided regulation (EC) No 999/2001 as a country or region posing a negligible BSE risk; the animals from which the raw materials of bovine, ovine and caprine animal origin were derived passed ante-mortem and post-mortem inspections; the raw materials of bovine, ovine and caprine animal origin intended for export are derived both from animals which were born, continuously reared and slaughtered in a country or region with hergulation (EC) No 999/2001, and, if there have been BSE indigenous cases in the country or region, born after the date from which the base of the same method or slaughtered after stunning by means of gas injected into the cranial cavity or region of slaughtered after stunning by means of gas injected in the caralial cavity or willed by the same method or slaughtered by laceration after stunning of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity; the raw materials of bovine, ovine and caprine animal origin do not contain and are not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001, or mechanically separated meat obtained from bones of bovine, ovine or caprine animals.] (1) or [(1) or [1] they come from a country or a region classified in accordance with Article 5(2) of Regulation (EC) No 999/2001, as a country or region which the raw materials of bovine, ovine and caprine animal origin were derived have not been faculated from bones of bovine, ovine and caprine animals.] (1) or the raw materials of bovine, origin were derived from ruminants and passed ante-mortem and post-mortem inspections. the animals from which the raw materials of bovine, ovine and caprine animal origin were derived have n		_	were derived have not been cavity or killed by the same	n slaughtered after stunning by means of method or slaughtered by laceration afte	gas injected into the cranial r stunning of central nervous
 [they come from a country or a region classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region posing a negligible BSE risk; the animals from which the raw materials of bovine, ovine and caprine animal origin were derived passed ante-mortem and post-mortem inspections; the raw materials of bovine, ovine and caprine animal origin intended for export are derived both from animals which were born, continuously reared and slaughtered in a country or region, and, if there have been BSE indigenous cases in the country or region, born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants had been enforced, and from animals born in a country or region posing a controlled BSE risk, and which have not been slaughtered after stunning by means of gas injected into the cranial cavity or willed by the same method or slaughtered by laceration after stunning of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity separated meat obtained from bones of bovine, ovine and caprine animals.] (1) or (1) or (1) or (1) the animals from which the raw materials of bovine, ovine and caprine animal origin were derived from specified risk material as defined in a greaves derived from ruminants and passed ante-mortem and post-mortem inspections; the animals from which the raw materials of bovine, ovine and caprine animal origin were derived from specified risk material as defined after stunning by means of gas injected into the cranial cavity or No 999/2001 as a country or region with an undetermined BSE risk; the animals from which the raw materials of bovine, ovine and caprine animal origin were derived have not been fed meat-and-bone meal or greaves derived from ruminants and passed ante-mortem and post-mortem inspections; the animals from which the raw m		_	specified risk material as d	efined in Annex V to Regulation (EC) N	o 999/2001, or mechanically
 No 999/2001 as a country or region posing a negligible BSE risk; the animals from which the raw materials of bovine, ovine and caprine animal origin were derived passed ante-mortem and post-mortem inspections; the raw materials of bovine, ovine and caprine animal origin intended for export are derived both from animals which were born, continuously reared and slaughtered in a country or region with negligible BSE risk in accordance with Article 5(2) of Regulation (EC) No 999/2001, and, if there have been BSE indigenous cases in the country or region, born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants had been enforced, and from animals born in a country or region classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region posing a controlled BSE risk, and which have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration after stunning of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity. the raw materials of bovine, ovine and caprine animal origin do not contain and are not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001, or mechanically separated meat obtained from bones of bovine, ovine and caprine animal.] (!) or [(they come from a country or a region classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region with an undetermined BSE risk; the animals from which the raw materials of bovine, ovine and caprine animal origin were derived have not been field meat-and-bone meal or greaves derived from ruminants and passed antemortem and post-mortem inspections; the animals from which the raw materials of bovine, ovine and caprine animal origin were derived have not been fed meat-and-bone meal or greaves derived fr		(¹) 0	r		
 passed ante-mortem and post-mortem inspections; the raw materials of bovine, ovine and caprine animal origin intended for export are derived both from animals which were born, continuously reared and slaughtered in a country or region with negligible BSE risk in accordance with Article 5(2) of Regulation (EC) No 999/2001, and, if there have been BSE indigenous cases in the country or region, born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants had been enforced, and from animals born in a country or region posing a controlled ESE risk, and which have not been slaughtered after stunning by means of gas injected into the cranial cavity or relied plaves derived from ruminants had been enforced, and from an animals thered by laceration after stunning of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity; the raw materials of bovine, ovine and caprine animal origin do not contain and are not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001, or mechanically separated meat obtained from bones of bovine, ovine or caprine animals.] (!) or [(!) or the animals from which the raw materials of bovine, ovine and caprine animal origin were derived have not been fed meat-and-bone meal or greaves derived from ruminants and passed antemortem and post-mortem inspections; the animals from which the raw materials of bovine, ovine and caprine animal origin were derived have not been fed meat-and-bone meal or greaves derived from ruminants and passed antemortem and post-mortem inspections; the animals from which the raw materials of bovine, ovine and caprine animal origin were derived have not been fed meat-and-bone meal or greaves derived from ruminants and passed antemortem and post-mortem inspections; the animals from which the raw materials of bovine, ovine and caprine animal origin we		—			rticle 5(2) of Regulation (EC)
 from animals which were born, continuously reared and slaughtered in a country or region with negligible BSE risk in accordance with Article 5(2) of Regulation (EC) No 999/2001, and, if there have been BSE indigenous cases in the country or region, born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants had been enforced, and from animals born in a country or region classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region posing a controlled BSE risk, and which have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered after stunning of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity; the raw materials of bovine, ovine and caprine animal origin do not contain and are not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001, or mechanically separated meat obtained from bones of bovine, ovine or caprine animals.] (¹) or [they come from a country or a region classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region with an undetermined BSE risk; the animals from which the raw materials of bovine, ovine and caprine animal origin were derived have not been fed meat-and-bone meal or greaves derived from ruminants and passed antemortem and post-mortem inspections; the animals from which the raw materials of bovine, ovine and caprine animal origin were derived have not been fad meat-and-bone meal or greaves derived from ruminants and passed antemortem and post-mortem inspections; the animals from which the raw materials of bovine, ovine and caprine animal origin were derived have not been fad meat-and-bone meal or greaves derived from ruminants and passed of an elongated rod-shaped instrument introduced into the cranial cavity; the anim		_			ne animal origin were derived
 specified risk material as defined in Annex V to Regulation (EC) No 999/2001, or mechanically separated meat obtained from bones of bovine, ovine or caprine animals.] (¹) or [they come from a country or a region classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region with an undetermined BSE risk; the animals from which the raw materials of bovine, ovine and caprine animal origin were derived have not been fed meat-and-bone meal or greaves derived from ruminants and passed antemortem and post-mortem inspections; the animals from which the raw materials of bovine, ovine and caprine animal origin were derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration after stunning of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity; the raw materials of bovine, ovine and caprine animal origin are not derived from: specified risk material as defined in Annex V to Regulation (EC) No 999/2001; nervous and lymphatic tissues exposed during the de-boning process; 		_	from animals which were b negligible BSE risk in accor have been BSE indigenous on the feeding of ruminants been enforced, and from Article 5(2) of Regulation (E and which have not been sla or killed by the same method	orn, continuously reared and slaughtered dance with Article 5(2) of Regulation (EC cases in the country or region, born after s with meat-and-bone meal and greaves animals born in a country or region c EC) No 999/2001 as a country or region p aughtered after stunning by means of gas d or slaughtered by laceration after stunning	d in a country or region with C) No 999/2001, and, if there the date from which the ban derived from ruminants had lassified in accordance with posing a controlled BSE risk, injected into the cranial cavity g of central nervous tissue by
 [they come from a country or a region classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region with an undetermined BSE risk; the animals from which the raw materials of bovine, ovine and caprine animal origin were derived have not been fed meat-and-bone meal or greaves derived from ruminants and passed antemortem and post-mortem inspections; the animals from which the raw materials of bovine, ovine and caprine animal origin were derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration after stunning of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity; the raw materials of bovine, ovine and caprine animal origin are not derived from: specified risk material as defined in Annex V to Regulation (EC) No 999/2001; nervous and lymphatic tissues exposed during the de-boning process; 		_	specified risk material as d	efined in Annex V to Regulation (EC) N	o 999/2001, or mechanically
 No 999/2001 as a country or region with an undetermined BSE risk; the animals from which the raw materials of bovine, ovine and caprine animal origin were derived have not been fed meat-and-bone meal or greaves derived from ruminants and passed antemortem and post-mortem inspections; the animals from which the raw materials of bovine, ovine and caprine animal origin were derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration after stunning of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity; the raw materials of bovine, ovine and caprine animal origin are not derived from: specified risk material as defined in Annex V to Regulation (EC) No 999/2001; nervous and lymphatic tissues exposed during the de-boning process; 		(¹) 0	r		
 have not been fed meat-and-bone meal or greaves derived from ruminants and passed ante-mortem and post-mortem inspections; the animals from which the raw materials of bovine, ovine and caprine animal origin were derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration after stunning of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity; the raw materials of bovine, ovine and caprine animal origin are not derived from: specified risk material as defined in Annex V to Regulation (EC) No 999/2001; nervous and lymphatic tissues exposed during the de-boning process; 		—			rticle 5(2) of Regulation (EC)
 have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration after stunning of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity; the raw materials of bovine, ovine and caprine animal origin are not derived from: (i) specified risk material as defined in Annex V to Regulation (EC) No 999/2001; (ii) nervous and lymphatic tissues exposed during the de-boning process; 		_	have not been fed meat-a	nd-bone meal or greaves derived from	
 (i) specified risk material as defined in Annex V to Regulation (EC) No 999/2001; (ii) nervous and lymphatic tissues exposed during the de-boning process; 		—	have not been slaughtered a the same method or slaught	after stunning by means of gas injected into ered by laceration after stunning of centra	o the cranial cavity or killed by
(ii) nervous and lymphatic tissues exposed during the de-boning process;		—	the raw materials of bovine,	ovine and caprine animal origin are not de	rived from:
			(i) specified risk material a	as defined in Annex V to Regulation (EC) N	No 999/2001;
(iii) mechanically separated meat obtained from bones of bovine, ovine or caprine animals.]]			(ii) nervous and lymphatic	tissues exposed during the de-boning pro	cess;
			(iii) mechanically separate	d meat obtained from bones of bovine, ovi	ne or caprine animals.]]

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COUNTRY

П.	Health in	formation	II.a.	Certificate reference No	II.b.
(¹) [II.2.	Animal H	lealth Attestation			
	I, the und	lersigned official vete	rinarian,	certify that the raw materials o	described above:
II.2.1.	consist of	f animal products that	t satisfy t	he animal health requirement	ts below;
II.2.2.	have bee from:	n obtained in the terr	itory of (¹) either [:]	(¹) or [] (²) (³) (⁴)
(¹) either	[11.2.2.1	animals that come f the last 3 months be			nat territory since birth or for at leas
	(¹) either	12 March 20 authorised for meat and the all the relevan that were slav European Uni	10 laying the intro veterinar nt animal ughtered on of fre	g down lists of third coun oduction into the European L ry certification requirements (I health import requirements for human consumption on sh meat from animals of thos	n Regulation (EU) No 206/2010 o htries, territories or parts thereo Union of certain animals and fresh (OJ L 73, 20.3.2010, p. 1), fulfilling laid down in that Regulation, and n a date for which import into the se species was authorised from the umn 8 of Part 1 of Annex II to tha
	(¹) or	9 February 20 or transit thro mammals and	09 laying ough, the of farme 12), fulfill	g down a list of third countrie e Community of meat of wi ed rabbits and the veterinary o	n Regulation (EC) No 119/2009 o es or parts thereof, for imports into ild leporidae, of certain wild land certification requirements (OJ L 39 ealth import requirements laid dowr
(¹) or	[II.2.2.1	chicks or slaughter Annex I to Commiss third countries, terri may be imported in requirements (OJ L Regulation, and co animal health impo human consumptio animals of those sp	poultry f sion Reg tories, zo nto and 226, 23.1 nsist of rt require n on a o ecies wa	from (a) third country(ies) list ulation (EC) No 798/2008 of 8 ones or compartments from y transit through the Commun 8.2008, p. 1) under conditions species referred to in that Re ements laid down in that Re date for which import into the	g or have been imported as day-old ted for that commodity in Part 1 o 8 August 2008 laying down a list o which poultry and poultry products hity and the veterinary certification s at least equivalent to those in tha Regulation, fulfilling all the relevan gulation, and were slaughtered fo he European Union of meat from ry or territory thereof in accordance
(¹) or	[11.2.2.1	animals that have area:	been kill	ed in the wild in that territor	$\mathbf{y}^{(5)}$; and captured and killed in ar
		diseases for v Newcastle dis	which the ease or h	e animals are susceptible: fo	voutbreak of any of the following bot and mouth disease, rinderpest enza during the prior 30 days, nor o lays and
		territory of a	country		rom the borders separating anothen not authorised at these dates for n, and
					n 12 hours for chilling either to a jame establishment, or directly to a

COUNTRY

						ded for human consumptio		
II.	Health inform	ation	ll.a.	Certificate refere	nce No	II.b.		
II.2.3.	3. have been obtained in an establishment around which, within a radius of 10 km, there has been no case/outbreak of the following diseases for which the animals are susceptible: foot and mouth disease, rinderpest, Newcastle disease or highly pathogenic avian influenza, classical or African swine fever during the prior 30 days or, in the event of a case of one of those diseases, the preparation of raw materials for exportation to the European Union has been authorised only after the removal of all meat and the total cleaning and disinfection of the establishment under the control of an official veterinarian; and:							
II.2.4.		have been obtained and prepared without contact with other materials not complying with the conditions required above, and it has been handled so as to avoid contamination with pathogenic agents; and						
II.2.5.	have been trar	nsported in clean ar	nd seal	ed containers or lor	ries.]			
Notes								
Part I:								
20 biv 18	06/766/EC of 6 No alve molluscs, ec .11.2006, p. 53) a	ovember 2006 estal hinoderms, tunicate	blishing es, mar nnex l	the lists of third co ine gastropods and to Regulation (EC	untries and te fishery prod) 798/2008 a	II to Commission Decision prritories from which imports ucts are permitted (OJ L 32 nd/or in Part 1 of Annex I 06/2010.		
	x reference l.11: l mber as appropria		ne and	address of the disp	atch establish	ment; registration or approv		
		Registration numbe te information is to				ies), flight number (aircraft) loading.		
		Jse the appropriate 05.11.91, 05.11.99		•) code under	the following headings: 02.0		
— Во	x reference I.20: I	ndicate total gross	weight	and total net weigh	t.			
— Во	x reference I.23: I	dentification of cont	tainer/S	Seal number: only w	here applicat	le.		
— Во	x reference I.28:	Nature of commo	<i>dity</i> : hio	des, skins, bones, t	endons and s	inews;		
		Approval number	of esta	ablishments: registra	ation or appro	val number as appropriate;		
				cludes slaughterho and processing plar		vessel, cutting plant, gan		
Part II:								
. ,	lete as appropria leted.	te. In case of prod	ucts de	erived from fishery	products, the	whole section II.2 should I		
(²) Th	e name and ISO o	ode number of the	exporti	ing country or territo	ory or zone as	laid down in:		
_	the Annexes to	Decision 2006/766	6/EC;					
_	Annex I to Reg	ulation (EC) 798/20	08;					

COUNTRY

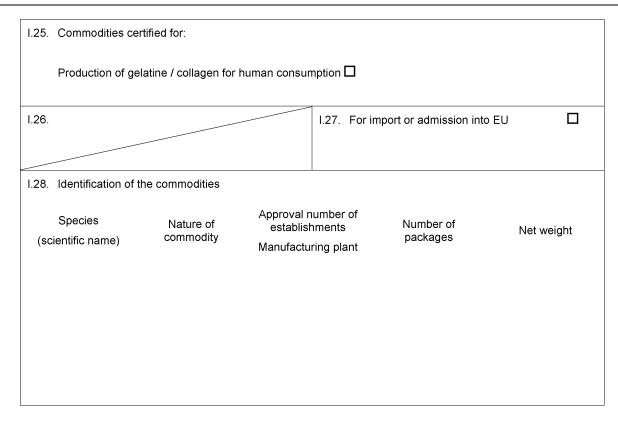
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П.	Health information	II.a.	Certificate reference No	II.b.				
	 Part 1 of Annex II to Regulation (EC) No 119/2009; 							
	— Part 1 of Annex II to Regulation (EU) No 206/2010.							
(3)	If parts of the materials were derived from animals originating from an(other) third country(ies) listed in Annex II to Regulation (EU) No 206/2010 for import of that commodity into the Union, then the code(s) of country(ies) or territory(ies) and of the third country slaughtering the animals shall be indicated (the material cannot come from a country or territory that has supplementary guarantees A or F as indicated in column 5 of that Annex).							
(4)) If the materials were derived from slaughter poultry originating from an(other) third country(ies) listed in Part 1 of Annex I to Regulation (EC) No 798/2008 for imports of that commodity into the Union, then the code(s) of country(ies) or territory(ies) and of the third country slaughtering the poultry shall be indicated.							
(⁵)	Only for countries from where game meat intended for human consumption of the same animal species is authorised for importation into the European Union.							
_	The signature and the stamp must be in a different colour to that of the printing.							
NB Note for the person responsible for the consignment in the EU: this certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post. The consignment must be transported directly to the manufacturing plant of destination.								
Official veterinarian								
	Name (in capital letters): Qualification and title:							
Date: Signature:								
	Stamp:							

PART VI

MODEL CERTIFICATE FOR IMPORTS OF TREATED RAW MATERIALS FOR THE PRODUCTION OF GELATINE/COLLAGEN INTENDED FOR HUMAN CONSUMPTION

COL	DUNTRY:						Veterinary certificate to EU			
	I.1. Consignor				1.2.	Certificate r	reference No	l.2.a.	\geq	
		Name Address				I.3. Central competent authority				
						I.4. Local competent authority				
	Tel.									
ment	1.5.	Consignee				1.6.				
		Name								
nsign		Address								
d co		Postcode								
tche		Tel.								
spat										
s of di	1.7.	Country of origin	ISO code	I.8. Region of origin	Code	1.9.	Country of destination	ISO code	l.10.	
Part I: Details of dispatched consignment										
oart I:	I.11.	Place of or	igin			I.12.				
-										
		Name Approval number			nber					
	Address									
	I.13.	Place of lo	ading			I.14.	Date of dep	parture		
	l.15.	Means of t	ransport			I.16.	Entry BIP in	n EU		
		Aeropla <u>n</u> e	-	Ship 🗖 🛛 R	ailway					
		wagon 🛯			anway					
		Road vehic		Other 🗖		I.17.				
	Identification Documentation references									
		I.18. Description of commodity								
	I.18.	Descriptior	n of commo	dity				I.19. Comn	nodity code (HS code)	
									I.20. Quantity	
	I.21.	Temperatu Ambient 🗖		ct Chilled 🛙]		Frozen 🗖		I.22. Number of packages	
	1.23.	Seal/Conta	ainer No						I.24. Type of packag	jing



COUNTRY

Model TCG Treated raw materials for the production of gelatine and collagen

	П.	. Health information		II.a.	Certificate reference No	II.b.					
	II.1.	Public H	ealth Attestation								
Part II: Certification		I, the undersigned, certify that the treated raw materials described above comply with the following requirements:									
		 they have been derived from establishments under the control of and listed by the competent authority 									
		and,									
		als, pigs and poultry described in a slaughterhouse and the lowing ante- and post-mortem									
		(¹) and/or									
		 [wild game hides, skins and bones described above are derived from killed animals whose carcasses have been found fit for human consumption following post-mortem inspection,] 									
		(¹) and/or									
		 [fish skins and bones described above are derived from plants manufacturing fishery products for human consumption authorised for export,] 									
		and									
	(¹) either	[they are dried bones of species from bovine, ovine, caprine, porcine and equine animals, including farmed and wild animals, poultry including ratites and feathered game for the production of collagen or gelatine, they derived from healthy animals slaughtered in a slaughterhouse, and they have been treated as follows:									
		(¹) either	(1) either [crushed to pieces of approximately 15 mm and degreased with hot water at a temperature of minimum 70 °C for at least 30 minutes, minimum 80 °C for at least 15 minutes, or minimum 90 °C for at least 10 minutes, and then separated and subsequently washed and dried for at least 20 minutes in a stream of hot air with an initial temperature of minimum 350 °C, or for 15 minutes in a stream of hot air with an initial temperature of more than 700 °C.]								
		(1) or	r [sun-dried for a minimum of 42 days at an average temperature of at least 20 °C.]								
		(¹) or	[acid treatment such before drying.]]	that the p	H is maintained at less than 6 to	the core for at least one hour					
	(¹) or	[they are hides and skins of farmed ruminant animals, pig skins, poultry skins or wild game hides and skins, they derived from healthy animals and they:									
		(¹) <i>either</i> [have undergone an alkali treatment which ensures a PH>12 to the core followed by salti for at least 7 days]									
		(¹) or) or [were dried for at least 42 days at a temperature of at least 20 °C.]								
		(¹) or	(¹) or [have undergone an acid treatment that provides at least a pH of less than 5 to the core t minimum of one hour.]								
		(¹) or	[have undergone an 8 hours.]]	alkali tre	eatment which ensures a pH >	12 to the core for at least					
	(¹) or	[they are bones, hides or skins of farmed ruminant animals, pig skins, poultry skins, fish skins and wild game hides and skins from third countries, parts of third countries and territories referred to in Part IV of Annex I to this Regulation that have undergone any other treatment than those listed above, and that come from establishments registered or approved pursuant to Regulation (EC) No 852/2004 or in accordance with Regulation (EC) No 853/2004									
		and									
		(¹) [if from ruminant origin, except for hides and skins of ruminants,									

COUNTRY

Model TCG Treated raw materials for the production of gelatine and collagen

II.	Healt	h information	II.a.	Certificate reference No	ll.b.
	(1) eit	her:			
	۔ ۲	No 999/2001 of the Europe he prevention, control ar	an Parliar nd eradica	n classified in accordance with A nent and of the Council of 22 Ma ation of certain transmissible s ntry or region posing a negligible	ay 2001 laying down rules for pongiform encephalopathies
	C		lously rea	aw materials of bovine, ovine ar red and slaughtered in the cou ortem inspections;	
	— i	f in the country or region th	nere have	been BSE indigenous cases:	
	(e date from which the ban on the sen on the sen on the sen on the sen of the	
	(not derived from sp	pecified r	vine, ovine and caprine animal o isk material as defined in An separated meat obtained from	inex V to Regulation (EC)
	(1) or:				
				n classified in accordance with A posing a controlled BSE risk;	rticle 5(2) of Regulation (EC)
		he animals from which the derived passed ante-morte		aw materials of bovine, ovine ar st-mortem inspections;	d caprine animal origin were
	f t c	or export were derived ha he cranial cavity or killed	ive not be by the s	materials of bovine, ovine and c en slaughtered after stunning b ame method or slaughtered by f an elongated rod-shaped ins	y means of gas injected into laceration after stunning of
	C	derived from specified risk	material	ovine and caprine animal origir as defined in Annex V to Regu ed from bones of bovine, ovine o	lation (EC) No 999/2001, or
	(¹) or			
	— [1	they come from a country No 999/2001 as a country o	or a regio or region p	n classified in accordance with A posing a negligible BSE risk;	rticle 5(2) of Regulation (EC)
		he animals from which the derived passed ante-morte		aw materials of bovine, ovine ar st-mortem inspections;	d caprine animal origin were
	r a v r a c i c	derived both from animals region with negligible BSE and, if there have been BS which the ban on the feed ruminants had been enfo accordance with Article 5(controlled BSE risk, and wh nto the cranial cavity or kil	which we risk in ac SE indiger ling of rur prced, and 2) of Reg nich have lled by the	e, ovine and caprine animal or re born, continuously reared and coordance with Article 5(2) of Re- nous cases in the country or reg- ninants with meat-and-bone me d from animals born in a cou- julation (EC) No 999/2001 as a not been slaughtered after stunn e same method or slaughtered b of an elongated rod-shaped ins	d slaughtered in a country or egulation (EC) No 999/2001, ion, born after the date from al and greaves derived from intry or region classified in a country or region posing a ing by means of gas injected y laceration after stunning of

COUNTRY

Model TCG Treated raw materials for the production of gelatine and collagen

	of gelatine and collagen						
II.	Health information II.a. Certificate reference No II.b.						
	 the treated raw materials of bovine, ovine and caprine animal origin do not contain and are not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001, or mechanically separated meat obtained from bones of bovine, ovine or caprine animals.] 						
	(¹) or						
	 [they come from a country or a region classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region with an undetermined BSE risk; 						
	 the animals from which the treated raw materials of bovine, ovine and caprine animal origin were derived have not been fed meat-and-bone meal or greaves derived from ruminants and passed ante-mortem and post-mortem inspections; 						
	 the animals from which the treated raw materials of bovine, ovine and caprine animal origin were derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration after stunning of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity; 						
	— the treated raw materials of bovine, ovine and caprine animal origin are not derived from:						
	(i) specified risk material as defined in Annex V to Regulation (EC) No 999/2001;						
	(ii) nervous and lymphatic tissues exposed during the de-boning process;						
	(iii) mechanically separated meat obtained from bones of bovine, ovine or caprine animals.]]]						
(¹) [II.2.	Animal Health Attestation						
	I, the undersigned official veterinarian, certify that the treated raw materials described above:						
II.2.1.	consist of animal products that satisfy the animal health requirements below;						
II.2.2.	have been obtained in the territory(ies) of: (1) [] (1) or [] (2) (3)						
II.2.3.	have been obtained and prepared without contact with other materials not complying with the conditions required above, and have been handled so as to avoid contamination with pathogenic agents;						
II.2.4.	have been transported in clean and sealed containers or lorries.]						
Notes							
Part I:							
— Box reference I.8: Provide the code of territory as appearing in Annex II to Commission Decision 2006/766/EC of 6 November 2006 establishing the lists of third countries and territories from which imports of bivalve molluscs, echinoderms, tunicates, marine gastropods and fishery products are permitted (OJ L 320, 18.11.2006, p. 53) or in Part 1 of Annex I to Commission Regulation (EC) 798/2008 of 8 August 2008 laying down a list of third countries, territories, zones or compartments from which poultry and poultry products may be imported into and transit through the Community and the veterinary certification requirements (OJ L 226, 23.8.2008, p. 1) or in Part 1 of Annex I to Commission Regulation (EC) No 119/2009 of 9 February 2009 laying down a list of third countries or parts thereof, for imports into, or transit through, the Community of meat of wild leporidae, of certain wild land mammals and of farmed rabbits and the veterinary certification requirements (OJ L 39, 10.2.2009, p. 12) or in Part 1 of Annex II to Commission Regulation (EU) No 206/2010 of 12 March 2010 laying down lists of third countries, territories or parts thereof, the introduction into the European Union of certain animals and fresh meat and the veterinary certification requirements (OJ L 73, 20.3.2010, p. 1).							
	x reference I.11: Place of origin: name and address of the dispatch establishment and approval number or npetent authority identification number as appropriate.						

 Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship). Separate information is to be provided in case of unloading and reloading.

COUNTRY

Model TCG Treated raw materials for the production of gelatine and collagen

				of gelatine and collagen			
II.	Health information	n II.a.	Certificate reference No	II.b.			
_	Box reference I.19: Use the appropriate Harmonised System (HS) code under the following headings: 03.05, 05.05, 05.06, 05.11.91, 05.11.99, 41.01, 41.02, 41.03.						
_	Box reference I.20: Indic	ate total gross weig	ht and total net weight.				
—	Box reference I.23: Iden	ification of containe	er/Seal number: only where applic	cable.			
—	Box reference I.28:	Nature of commodit	<i>ty:</i> hides, skins, bones, tendons a	and sinews;			
		Approval number identification numbe		number or competent authority			
			<i>t:</i> includes slaughterhouse, fact nent and processing plant;	tory vessel, cutting plant, game			
		Approval number: w	vhen applicable.				
Pai	t II:						
(¹)	Delete as appropriate. I deleted.	n case of products	derived from fishery products, t	the whole section II.2 should be			
(²)	The name and ISO code	number of the expo	orting country or territory or zone	as laid down in:			
	— Part 1 of Annex II to	o Regulation (EU) N	lo 206/2010;				
	 Annex I to Regulati 	on (EC) 798/2008;					
	— Part 1 of Annex II to	o Regulation (EC) N	lo 119/2009.				
(3)	Annex I to Commission countries, parts of third into the Union of certain requirements, amending	Implementing Regr countries and territo products of anima Regulation (EC) N	ulation (EU) 2016/759 of 28 App pries from which Member States I origin intended for human cons	other) third country(ies) listed in ril 2016 drawing up lists of third are to authorise the introduction sumption, laying down certificate ecision 2003/812/EC (OJ L 126, ed.			
_	The signature and the st	amp must be in a di	ifferent colour to that of the printi	ng.			
NB		consignment until it	reaches the border inspection	only for veterinary purposes and post. The consignment must be			
—	The time of transportatio	n may be included i	in the duration of treatment.				
Off	Official veterinarian						
	Name (in capital lette	rs):		Qualification and title:			
	Date:			Signature:			
	Stamp:						

PART VII

MODEL CERTIFICATE FOR IMPORTS OF HONEY, ROYAL JELLY AND OTHER APICULTURE PRODUCTS INTENDED FOR HUMAN CONSUMPTION

cou	INTRY	:	Veterinary certificate to EU			
	I.1.	Consignor	I.2. Certificate reference No I.2.a.			
		Name Address	I.3. Central competent authority			
		Address	I.4. Local competent authority			
		Tel.				
¥	1.5.	Consignee	1.6.			
mei		Name				
nsign		Address				
ed co		Postcode				
batch		Tel.				
Part I: Details of dispatched consignment	1.7.	Country of ISO code I.8. origin	I.9. Country of ISO code I.10. destination			
art I: D	I.11.	Place of origin	1.12.			
		Name Approval number				
		Address				
	I.13.	Place of loading	I.14. Date of departure			
	I.15.	Means of transport	I.16. Entry BIP in EU			
		Aeroplane □ Ship □ Railway wagon □				
		Road vehicle	1.17.			
		Identification				
		Documentation 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.	Seal/Container No	I.24. Type of packaging			

I.25. Commodities certi	fied for:				
Human consumpti	ion 🗖				
1.26.			I.27. For im	nport or admission into EU	
I.28. Identification of the	e commodities				
Species (scientific name)	Treatment type	Approval n establish Manufactu	nments	Number of packages	Net weight

COUNTRY Model HON Honey, royal jelly and other apiculture products II. Health information II.a. Certificate reference No II.b II.1. **Public Health Attestation** I, the undersigned, declare that I am aware of the relevant provisions of Regulation (EC) No 178/2002 of the European Parliament and of the Council of 28 January 2002 laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety (OJ L 31, 1.2.2002, p. 1), Regulation (EC) No 852/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs (OJ L 139, 30.4.2004, p. 1) and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying Part II: Certification down specific hygiene rules for food of animal origin (OJ L 139, 30.4.2004, p. 55) and certify that honey, royal jelly and other apiculture products described above were produced in accordance with those requirements, in particular that they: come from (an) establishment(s) implementing a programme based on the HACCP principles in accordance with Regulation (EC) No 852/2004; have been handled and, where appropriate, prepared, packaged and stored in a hygienic manner in accordance with the requirements of Annex II to Regulation (EC) No 852/2004; and the guarantees covering live animals and products thereof provided by the residue plans submitted in accordance with Council Directive 96/23/EC of 29 April 1996 on measures to monitor certain substances and residues thereof in live animals and animal products and repealing Directives 85/358/EEC and 86/469/EEC and Decisions 89/187/EEC and 91/664/EEC (OJ L 125, 23.5.1996, p. 10), and in particular Article 29 thereof, are fulfilled. Notes Part I: - Box reference I.11: Place of origin: name and address of the dispatch establishment. Approval number means registration number. Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship). Separate information is to be provided in case of unloading and reloading. Box reference I.19: Use the appropriate Harmonised System (HS) code under the following headings: 04.09, 04.10. Box reference I.20: Indicate total gross weight and total net weight. Box reference I.23: Identification of container/Seal number: only where applicable. Box reference I.28: Treatment type: Indicate 'ultrasonication', 'homogenisation', ultrafiltration'. 'pasteurisation', 'no thermal treatment'. Approval number of establishments: approval number or competent authority identification number as appropriate Part II: The colour of the stamp and signature must be different from that of the other particulars in the certificate. Official inspector Name (in capital letters): Qualification and title: Date: Signature: Stamp:

PART VIII

MODEL CERTIFICATE FOR IMPORTS OF HIGHLY REFINED CHONDROITIN SULPHATE, HYALURONIC ACID, OTHER HYDROLYSED CARTILAGE PRODUCTS, CHITOSAN, GLUCOSAMINE, RENNET, ISINGLASS AND AMINO ACIDS INTENDED FOR HUMAN CONSUMPTION

COL	INTRY	:				Ve	terinary	certificate to EU
	l.1.	Consignor		I.2.	Certificate ref	erence No	I.2.a.	
		Name Address		I.3. Central competent authority				
				I.4.	Local compet	ent authority		
		Tel.						
	I.5.	Consignee		I.6.				
ment		Name						
nsign		Address						
ed co		Postcode						
oatche		Tel.						
Part I: Details of dispatched consignment	I.7.	Country of ISO code origin	1.9.	Country of destination	ISO co	de	l.10.	
: Deta								
Part I	l.11.	Place of origin				I		
		Name App						
		Address						
	l.13.	Place of loading		l.14.	Date of depar	ture		
	l.15.	Means of transport		l.16.	Entry BIP in E	U		
		Aeroplane 🛛 Ship 🗖 wagon 🗖	Railway					
		-	er 🗖	I.17.			_	
		Identification						
		Documentation references						
	I.18. Description of commodity				I.19. Comm	odity co	ode (HS code)	
							1.20. (Quantity
	I.21.	Temperature of product						Number of
		Ambient 🗖	Chilled 🗖	I	Frozen 🗖		þ	backages
	1.23.	Seal/Container No					1.24.	Type of packaging

I.25.	I.25. Commodities certified for:						
	Human consump	tion 🗖					
1.26.				1.27.	For impor	t or admission into EU	
1.28.	Identification of th	ne commodities		1			
(sc	Species cientific name)	Date production (dd/mm/yyyy)	Approval r establisł Manufactu	nments		Number of packages	Net weight

COUNTRY

Model HRP Highly refined chondroitin sulphate, hyaluronic acid, other hydrolysed cartilage products, chitosan, glucosamine, rennet, isinglass and amino acids for human consumption

ſ										
	П.	Health information		II.a.	Certificate reference No	II.b.				
	II.1.	Public Health Attestation								
 I, the undersigned, declare that I am aware of the relevant provisions of Regulation (EC) No the European Parliament and of the Council of 28 January 2002 laying down the general prequirements of food law, establishing the European Food Safety Authority and laying down in matters of food safety (OJ L 31, 1.2.2002, p. 1), Regulation (EC) No 852/2004 of the Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs (OJ L 139, 30.4 and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April down specific hygiene rules for food of animal origin (OJ L 139, 30.4.2004, p. 55) and ce highly refined products described above were produced in accordance with those requirements. — that they come from (an) establishment(s) implementing a programme based on principles in accordance with Regulation (EC) No 852/2004; — that they have been handled and, where appropriate, prepared, packaged and stored in the provinciples in accordance with the provinciples of the requirements of forms with the provinciples of the provinciples in accordance with Regulation (EC) No 852/2004; 										
л II: С		k	orino	ciples in accordance with	n Regulat	nment(s) implementing a progr tion (EC) No 852/2004; here appropriate, prepared, pac				
Ра						ements of Annex II to Regulation				
		— t and	hat	they comply with the red	quiremen	ts of Section XVI of Annex III to	Regulation (EC) No 853/2004;			
			¹) [it	amino acids, that						
		(i)	human hair was not us	ed as a s	source for their manufacture; and	1			
	 they comply with Regulation (EC) No 1333/2008 of the European Parliament and of the Council of 16 December 2008 on food additives ((OJ L 354, 31.12.2008, p. 16)] 									
	Notes									
	Part I:									
	— Вс	ox refere	ence	e I.11: Place of origin: na	me and a	address of the dispatch establish	iment.			
						/ay wagons or container and lor vided in case of unloading and r				
				e I.19: Use the approp 29.32, 35.07, 35.03 or 3		rmonised System (HS) code u	nder the heading of 21.06.90,			
	— Вс	ox refere	ence	e I.20: Indicate total gros	s weight	and total net weight.				
	— Во	ox refere	ence	e I.23: Identification of co	ontainer/S	Seal number: only where applica	ble.			
	Part II	:								
	(1) De	elete as	app	propriate.						
	— The colour of the stamp and signature must be different from that of the other particulars in the certificate.									
	Official veterinarian Name (in capital letters):									
		Qualification and title:								
		Date:				S	ignature:			
		Stamp	:							

ANNEX III

MODEL CERTIFICATE FOR THE TRANSIT THROUGH THE UNION, IMMEDIATE TRANSIT OR AFTER STORAGE, FOR RAW MATERIALS OR TREATED RAW MATERIALS FOR THE PRODUCTION OF GELATINE/COLLAGEN FOR HUMAN CONSUMPTION

COL	JNTRY	1						Ve	terinary certificate to EU
	l.1.	Consignor				1.2.	Certificate re	eference No	l.2.a.
		Name Address				I.3. Central competent authority		ity	
						I.4.	Local compe	etent authority	4
		Tel.							
	1.5.	Consignee				I.6. Person responsible for the consignment in EU			
nent		Name Address					Name Address		
signn									
con		Postcode					Postcode		
ched		Tel.					Tel.		
dispat	1.7.		ISO code	I.8. Region of origin	Code	1.9.	Country of destination	ISO code	l.10.
ls of c									
Part I: Details of dispatched consignment	l.11.	11. Place of origin					Place of des	stination	
art								_	_
		Name Address					Customs warehouse \Box		Ship supplier 🗖
		Audress				Name		Approval number	
							Address		
							Postal code		
	I.13.	Place of load	ling			l.14.	Date of depa	arture	
		Means of trai					Entry BIP in		
	1.10.		noport			1.10.			
		Aeroplane 🗖 wagon 🗖]	Ship 🗖 🛛 Ra	ailway				
		Road vehicle		Other 🗖		I.17.			
		Identification							
		Documentatio	on referen	ces					
	l.18.	I.18. Description of commodity				I.19. Comm	odity code (HS code)		
							l		I.20. Quantity
	I.21.	Temperature	of produc	t					I.22. Number of
		Ambient 🗖		Chilled 🗖]		Frozen 🗖		packages
	1.23.	Seal/Contain	er No						I.24. Type of packaging

I.25.	Commodities certified for:						
	Production of gelat	ine / collagen for human consur	nption 🗖				
I.26.	For transit through	EU to third country	1.27.				
	Third country	ISO code					
l.28.	Identification of the	commodities					
	Species (scientific name)	Manufacturing plant	Number of packages	Net weight			

Part II: Certification

EN

COUN	IIKI				WOULD TRANSITISTORAGE			
11.	Health	information	II.a.	Certificate reference number	II.b.			
II.1.	Public Health Attestation							
	•	indersigned official veterin ed in Part I:	narian, h	nereby certify, that the raw mater	ials or treated raw materials			
	II.1.1.	Annex I to Commission	Regulat	authorised for imports into the E ion (EC) 798/2008 of 8 August 20	008 laying down a list of third			

- countries, territories, zones or compartments from which poultry and poultry products may be imported into and transit through the Community and the veterinary certification requirements (OJ L 226, 23.8.2008, p. 1) or in Commission Regulation (EC) No 119/2009 of 9 February 2009 laying down a list of third countries or parts thereof, for imports into, or transit through, the Community of meat of wild leporidae, of certain wild land mammals and of farmed rabbits and the veterinary certification requirements (OJ L 39, 10.2.2009, p. 12) or in Part 1 of Annex II to Commission Regulation (EU) No 206/2010 of 12 March 2010 laying down lists of third countries, territories or parts thereof authorised for the introduction into the European Union of certain animals and fresh meat and the veterinary certification requirements (OJ L 73, 20.3.2010, p. 1), and
- II.1.2. comply with the relevant animal health conditions as laid down in the animal health attestation in the model certificate in Part V or VI of Annex II to Commission Implementing Regulation (EU) 2016/759 of 28 April 2016 drawing up lists of third countries, parts of third countries and territories from which Member States are to authorise the introduction into the Union of certain products of animal origin intended for human consumption, laying down certificate requirements, amending Regulation (EC) No 2074/2005 and repealing Decision 2003/812/EC (OJ L 126, 14.5.2016, p. 13).

Notes

This certificate is meant for transit and storage in accordance with Article 12(4) or Article 13 of Council Directive 97/78/EC of 18 December 1997 laying down the principles governing the organisation of veterinary checks on products entering the Community from third countries (OJ L 24, 30.1.1998, p. 9) of raw materials or treated raw materials for the production of gelatine/collagen for human consumption of:

- (1) domestic bovine animals (including *Bubalus* and *Bison* species and their cross-breeds);
- (2) domestic ovine animals (*Ovis aries*) or domestic caprine animals (*Capra hircus*);
- (3) domestic porcine animals (Sus scrofa);
- (4) domestic solipeds (*Equus caballus*, *Equus asinus* and their cross-breeds);
- (5) farmed non-domestic animals of the order Artiodactyla (excluding bovine animals (including *Bison* and *Bubalus* species and their cross-breeds), *Ovis aries, Capra hircus*, Suidae and Tayassuidae), and of the families Rhinocerotidae and Elephantidae;
- (6) wild non-domestic animals of the order Artiodactyla (excluding bovine animals (including *Bison* and *Bubalus* species and their cross-breeds), *Ovis aries*, *Capra hircus*, Suidae and Tayassuidae), and of the families Rhinocerotidae and Elephantidae;
- (7) farmed non-domestic animals belonging to the Suidae, Tayassuidae, or Tapiridae families;
- (8) wild non-domestic animals belonging to the Suidae, Tayassuidae, or Tapiridae families;
- (9) wild solipeds belonging to the subgenus *Hippotigris* (zebra);
- (10) wild leporidae (rabbits and hares);
- (11) wild land mammals other than ungulates and leporidae;
- (12) farmed rabbits;
- (13) poultry;
- (14) farmed ratites;
- (15) wild game;
- (16) fish.

co	UNTRY			Model TRANSIT/STORAGE					
II.	Health information	II.a.	Certificate reference number	II.b.					
Pa	Part I:								
_	— Box reference I.8: Provide the code of territory as appearing in Part 1 of Annex II to Regulation (EU) No 206/2010 or in Part 1 of Annex I to Regulation (EC) No 119/2009 or Part 1 of Annex I to Regulation (EC) No 798/2008 or Annex II to Commission Decision 2006/766/EC of 6 November 2006 establishing the lists of third countries and territories from which imports of bivalve molluscs, echinoderms, tunicates, marine gastropods and fishery products are permitted (OJ L 320, 18.11.2006, p. 53).								
—	Box reference I.11: Place of origin: na	ame an	d address of the dispatch establish	ment.					
-	Box reference I.12: Address (and warehouse, customs warehouse or sh			ehouse in a free zone, free					
-	Box reference I.15: Registration num name (ship) is to be provided. In case into the EU.								
-	Box reference I.19: Use the appropri 05.04, 05.05, 05.06, 05.11.91, 05.11.			r the heading of 02.08, 03.05,					
-	Box reference I.20: Indicate total gros	s weigl	nt and total net weight.						
-	Box reference I.23: For containers or be included.	boxes,	the container number and the sea	l number (if applicable) should					
_	 Box reference I.28: Manufacturing plant: provide the registration number, approval number or competent authority identification number of establishment as appropriate. It includes slaughterhouse, factory vessel, cutting plant, game handling establishment and processing plant. 								
Off	Official veterinarian or Official inspector								
	Name (in capital letters): Qualification and title:								
	Date: Signature:								
	Stamp:								

ANNEX IV

EXPLANATORY NOTES FOR COMPLETING THE CERTIFICATES

(referred to in Articles 2(1) and 4(1))

(a) Certificates shall be issued by the exporting third country, based on the models set out in Annexes II and III according to the layout of the model that corresponds to the products of animal origin 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.

If the Member State of destination imposes, for the products of animal origin concerned, additional certification requirements, attestations to certify that those requirements are fulfilled shall also be incorporated in the original form of the certificate.

- (b) Where the model certificate states that certain statements shall be kept as appropriate, statements which are not relevant, may be crossed out nd initialled and stamped by the certifying officer, or completely deleted from the certificate.
- (c) A separate and unique certificate must be provided for the products of animal origin that are exported from a territory or territories or zones of the same exporting country listed or referred to in Annex I which are consigned to the same destination and transported in the same railway wagon, lorry, aircraft or ship.
- (d) The original of each certificate shall consist of a single sheet of paper, or, where more text is required it must be in such a form that all sheets of paper required are part of an integrated whole and indivisible.
- (e) The certificate shall be drawn up in at least one of the official languages of the Member State of the border inspection post of entry of the consignment into the EU and of the Member State of destination. However, those Member States may authorise the certificate to be drawn up in the official language of another Member State, and accompanied, if necessary, by an official translation.
- (f) If for reasons of identification of the items of the consignment (schedule in point I.28 of the model certificate), additional sheets of paper are attached to the certificate, those sheets of paper shall also be considered as forming part of the original of the certificate by the application of the signature and stamp of the certifying officer, on each of the pages.
- (g) When the certificate, including additional sheets of paper referred to in (f), comprises more than one page, each page shall be numbered, (page number) of (total number of pages), at the end of the page and shall bear the certificate reference number that has been designated by the competent authority at the top of the pages.
- (h) The original of the certificate must be completed and signed by an official veterinarian or by another designated official inspector where this is provided for in the model certificate. The competent authorities of the exporting third country shall ensure that rules of certification equivalent to those laid down in Council Directive 96/93/EC (¹) are followed.

The colour of the signature shall be different from that of the printing. This requirement also applies to stamps other than those embossed or watermarked.

(i) The certificate reference number referred to in boxes I.2 and II.a. must be issued by the competent authority.

⁽¹⁾ Council Directive 96/93/EC of 17 December 1996 on the certification of animals and animal products (OJ L 13, 16.1.1997, p. 28).