II

(Non-legislative acts)

REGULATIONS

COMMISSION REGULATION (EU) 2019/319

of 6 February 2019

amending Annex IX to Regulation (EC) No 999/2001 of the European Parliament and of the Council and Annex XV to Commission Regulation (EU) No 142/2011 as regards health certification at import into the Union concerning transmissible spongiform encephalopathies

(Text with EEA relevance)

THE EUROPEAN COMMISSION.

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

Having regard to Regulation (EC) No 999/2001 of the European Parliament and of the Council of 22 May 2001 laying down rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies (1), and in particular the first paragraph of Article 23 and the introductory phrase and point (m) of Article 23a thereof,

Having regard to Regulation (EC) No 1069/2009 of the European Parliament and of the Council of 21 October 2009 laying down health rules as regards animal by-products and derived products not intended for human consumption and repealing Regulation (EC) No 1774/2002 (Animal by-products Regulation) (2), and in particular the introductory phrase and point (d) and the final paragraph of Article 42(2) thereof,

Whereas:

- Regulation (EC) No 999/2001 lays down rules for the prevention, control and eradication of transmissible spongiform encephalopathies ('TSEs') in bovine, ovine and caprine animals. It applies to the production and placing on the market of live animals and products of animal origin and in certain specific cases to exports thereof. That Regulation also provides a legal basis for the classification, as laid down in Commission Decision 2007/453/EC (3), of Member States and third countries or regions thereof according to their disease status for bovine spongiform encephalopathy (BSE) into those with a negligible BSE risk, a controlled BSE risk and an undetermined BSE risk.
- (2)Annex IX to Regulation (EC) No 999/2001 set outs the requirements for the importation into the Union of live animals, embryos, ova and products of animal origin. More particularly, Chapter B of that Annex sets out the requirements for imports of bovine animals, which takes into account the BSE status of the third countries or regions. In addition, Chapter D of that Annex lays down requirements for the provision of an attestation concerning the TSE related risk in the health certificate required for the importation into the Union of certain animal by-products and derived products, including, inter alia, processed animal protein.
- Chapter B of Annex IX to Regulation (EC) No 999/2001, as amended by Commission Regulation (EU) (3) 2016/1396 (4), requires that live bovine animals imported into the Union must not have been exposed to BSE

⁽¹) OJ L 147, 31.5.2001, p. 1. (²) OJ L 300, 14.11.2009, p. 1.

Commission Decision 2007/453/EC of 29 June 2007 establishing the BSE status of Member States or third countries or regions thereof according to their BSE risk (OJ L 172, 30.6.2007, p. 84).

Commission Regulation (EU) 2016/1396 of 18 August 2016 amending certain Annexes to Regulation (EC) No 999/2001 of the European Parliament and of the Council laying down rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies (OJ L 225, 19.8.2016, p. 76).

cases or their cohort. Taking into account the fact that the main transmission route of BSE is through feed contaminated with the BSE prion, that requirement should be amended to provide that live bovine animals imported into the Union may not be BSE cases or their cohort. Chapter B of Annex IX to Regulation (EC) No 999/2001 should therefore be amended accordingly.

- (4) Regulation (EC) No 1069/2009 lays down public health and animal health rules for animal by-products and derived products in order to prevent and minimise risks to public and animal health arising from those products. Commission Regulation (EU) No 142/2011 (5) lays down implementing measures for the public and animal health rules for animal by-products and derived products laid down in Regulation (EC) No 1069/2009, including certain requirements for the importation of animal by-products and derived products from third countries.
- (5) Annex I to Regulation (EU) No 142/2011 lists certain definitions to be used for the purposes of that Regulation. Article 31 of Regulation (EU) No 142/2011 provides that consignments of animal by-products and derived products for importation into or transit through the Union are to be accompanied by health certificates and declarations, in accordance with the models set out in Annex XV thereto.
- (6) Point 1 of Article 11.4.13 of the Terrestrial Animal Health Code of the World Organisation for Animal Health ('OIE Code') (6) recommends that meat-and-bone meal and greaves derived from ruminants, as defined in the OIE Code, and commodities containing such products, which originate from countries or zones with a negligible BSE risk status in which there has been an indigenous BSE case, may enter international trade only if the products were derived from cattle born after the date of the effective implementation, in the country, of the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants, as defined in the OIE Code. Point 2 of that Article recommends that meat-and-bone meal and greaves derived from ruminants, as defined in the OIE Code, and commodities containing such products, should not enter international trade if they originate from countries or zones with a controlled or undetermined BSE risk status.
- (7) The OIE Code defines meat-and-bone meal as the solid protein products obtained when animal tissues are rendered, including any intermediate protein product other than peptides of a molecular weight less than 10 000 daltons and amino-acids. Thus, meat-and-bone meal as defined in the OIE Code covers both the definition of meat-and-bone meal set out in point 27 of Annex I to Regulation (EU) No 142/2011 and the definition of processed animal protein set out in point 5 of that Annex.
- (8) In accordance with Article 41(2)(c) of Regulation (EC) No 1069/2009, imports into the Union of meat-and-bone meal, as defined in Union legislation, may only take place if implementing rules have been adopted setting out the conditions for such importation. Since no such implementing rules have been adopted, the importation into the Union of meat-and-bone meal, derived from Category 1 or Category 2 material, is currently not allowed. However, imports into the Union of processed animal protein, as defined in Union legislation, may take place, subject to compliance with the TSE related import conditions laid down in Section B of Chapter D of Annex IX to Regulation (EC) No 999/2001, as well as with the conditions on the import of processed animal protein laid down in Regulation (EU) No 142/2011.
- (9) In order to align the TSE conditions for imports into the Union, laid down in Regulation (EC) No 999/2001, with the recommendations included in the BSE Chapter of the OIE Code, it is appropriate to amend Section B of Chapter D of Annex IX to Regulation (EC) No 999/2001 so that the requirement laid down in that Section take account of the recommendations of Article 11.4.13 of the OIE Code. However, since the use of processed animal protein derived from ruminants in the manufacturing of petfood is authorised in the Union, in order not to apply a discriminatory treatment towards imports compared to European Union production, the recommendations of Article 11.4.13 of the OIE Code should not be followed for the importation of petfood containing processed animal protein derived from ruminants, provided that such petfood is processed and labelled in accordance with Union legislation.
- (10) Section B of Chapter D of Annex IX to Regulation (EC) No 999/2001 should therefore be amended accordingly.

⁽⁵⁾ Commission Regulation (EU) No 142/2011 of 25 February 2011 implementing Regulation (EC) No 1069/2009 of the European Parliament and of the Council laying down health rules as regards animal by-products and derived products not intended for human consumption and implementing Council Directive 97/78/EC as regards certain samples and items exempt from veterinary checks at the border under that Directive (OJ L 54, 26.2.2011, p. 1).

⁽⁶⁾ http://www.oie.int/international-standard-setting/terrestrial-code/access-online/

- (11) Products of animal origin may be required to be declared animal by-products by Union law, or by the decision of the responsible operator. When an operator decides that products of animal origin are to be declared as animal by-products, that decision is irreversible. Such animal by-products are excluded from use for human consumption. Certain animal by-products have the same Combined Nomenclature (CN) customs codes as animal products intended for human consumption which are laid down in Annex I to Council Regulation (EEC) No 2658/87 (?). For the classification in the CN customs codes the customs authorities in Member States need to be able to clearly differentiate between products which are fit for human consumption and those which are unfit for human consumption. In order to avoid any confusion for the purpose of that classification, the health guarantees referred to in the import certificates of unprocessed animal by-products should clarify that, although the animal by-products originate from animal products which were fit for human consumption at a former stage, they are now classified and treated as animal by-products which are permanently excluded from the food chain. The model health certificates set out in Chapters 3(D), 3(F) and 8 of Annex XV to Regulation (EU) No 142/2011 should therefore be amended accordingly.
- In addition, the TSE attestation in the model certificates for imports of and transit through the Union of certain animal by-products set out in Chapters 1, 1a, 2(A), 2(B), 3(A), 3(B), 3(C), 3(D), 3(E), 3(F), 4(B), 4(C), 4(D), 6(B), 8, 10(A), 10(B), 11, 12 and 18 of Annex XV to Regulation (EU) No 142/2011 should be amended to take account of the requirements of Chapter D of Annex IX to Regulation (EC) No 999/2001, as amended by Commission Regulation (EU) No 630/2013 (*), by Regulation (EU) 2016/1396 and by this Regulation.
- (13) The import conditions for processed animal protein referred to in the model health certificate set out in Chapter 1 of Annex XV to Regulation (EU) No 142/2011 require the absence of blood from ruminants in processed animal proteins imported from third countries. However, the new TSE attestation set out in point II.7. of that model health certificate, as amended by this Regulation, provides for adequate guarantees to mitigate the TSE risk in such products. Therefore, the wording 'other than ruminants' should be deleted in all the model health certificates set out in Annex XV to Regulation (EU) No 142/2011 that are to be amended by this Regulation.
- (14) Chapters 1, 1a, 2(A), 2(B), 3(A), 3(B), 3(C), 3(D), 3(E), 3(F), 4(B), 4(C), 4(D), 6(B), 8, 10(A), 10(B), 11, 12 and 18 of Annex XV to Regulation (EU) No 142/2011 should therefore be amended accordingly.
- In addition, consignments of intermediate products intended for the manufacturing of cosmetic and pharmaceutical products are required to be accompanied by a declaration completed in accordance with the model set out in Chapter 20 of Annex XV to Regulation (EU) No 142/2011 when presented at a border inspection post ('BIP') for the purpose of veterinary checks. Intermediate products may consist of or may contain animal by-products. The existing model declaration indicates only a limited number of appropriate HS codes which are to be used by the operator to notify the product to the customs authorities in the Member States. It is not possible to set out an exhaustive list of HS codes in advance in the model declaration which would cover all combinations of animal by-products in the intermediate products. Therefore, it is appropriate to replace the existing HS codes in order that the person responsible for the consignment may declare intermediate products to the BIP by an appropriate HS code in accordance with Commission Decision 2007/275/EC (°). Chapter 20 of Annex XV to Regulation (EU) No 142/2011 should be amended accordingly.
- (16) In order to avoid any disruption of trade, this Regulation should provide for a transitional period during which time the commodities concerned by the amendments made to Regulation (EU) No 142/2011 should continue to be accepted for importation into and transit through the Union, provided that those commodities comply with the requirements laid down in Regulation (EU) No 142/2011 before they were amended by this Regulation.
- (17) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on Plants, Animals, Food and Feed,

HAS ADOPTED THIS REGULATION:

Article 1

Annex IX to Regulation (EC) No 999/2001 is amended in accordance with Annex I to this Regulation.

⁽⁷⁾ Council Regulation (EEC) No 2658/87 of 23 July 1987 on the tariff and statistical nomenclature and on the Common Customs Tariff (OJ L 256, 7.9.1987, p. 1).

⁽⁸⁾ Commission Regulation (EU) No 630/2013 of 28 June 2013 amending the Annexes to Regulation (EC) No 999/2001 of the European Parliament and of the Council laying down rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies (OJ L 179, 29.6.2013, p. 60).

lopathies (OJ L 179, 29.6.2013, p. 60).

(°) Commission Decision 2007/275/EC of 17 April 2007 concerning lists of animals and products to be subject to controls at border inspection posts under Council Directives 91/496/EEC and 97/78/EC (OJ L 116, 4.5.2007, p. 9).

Article 2

Annex XV to Regulation (EU) No 142/2011 is amended in accordance with Annex II to this Regulation.

Article 3

For a transitional period until 30 September 2019, consignments of animal by-products and of derived products accompanied by a health certificate duly completed and signed in accordance with the appropriate model health certificate set out in Chapters 1, 1a, 2(A), 2(B), 3(A), 3(B), 3(C), 3(D), 3(E), 3(F), 4(B), 4(C), 4(D), 6(B), 8, 10(A), 10(B), 11, 12 and 18 of Annex XV to Regulation (EU) No 142/2011 in the version applicable before the amendments provided for by Article 2 of this Regulation, and, where applicable, by a declaration, which has been duly completed and signed in accordance with the model declaration set out in Chapter 20 of that Annex in its version applicable before the amendments provided for by Article 2 of this Regulation, shall continue to be accepted for importation into and transit though the Union, provided that such health certificates or declarations were duly completed and signed no later than 31 July 2019.

Article 4

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, 6 February 2019.

For the Commission
The President
Jean-Claude JUNCKER

ANNEX I

Annex IX to Regulation (EC) No 999/2001 is amended as follows:

- (1) in Chapter B:
 - (i) in Section A, the introductory phrase of point (b) is replaced by the following:
 - '(b) the animals are identified by a permanent identification system enabling them to be traced back to the dam and herd of origin, and are not the following bovine animals:'
 - (ii) in Section B, the introductory phrase of point (b) is replaced by the following:
 - '(b) the animals are identified by a permanent identification system enabling them to be traced back to the dam and herd of origin, and are not the following bovine animals:'
 - (iii) in Section C, the introductory phrase of point (c) is replaced by the following:
 - '(c) the animals are identified by a permanent identification system enabling them to be traced back to the dam and herd of origin, and are not the following bovine animals:'
- (2) in Chapter D, Section B is replaced by the following:

SECTION B

Health certificate requirements

- 1. Imports of the animal by-products and derived products of bovine, ovine and caprine origin referred to in Section A shall be subject to the presentation of a health certificate which has been completed with the following attestation:
 - (a) the animal by-product or derived product:
 - (i) does not contain and is not derived from specified risk material as defined in point 1 of Annex V to this Regulation; and
 - (ii) does not contain and is not derived from mechanically separated meat obtained from bones of bovine, ovine or caprine animals, except if the animals, from which the animal by-product or derived product are derived, were born, continuously reared and slaughtered in a country or region classified in accordance with Decision 2007/453/EC as a country or region posing a negligible BSE risk, in which there has been no BSE indigenous cases; and
 - (iii) is derived from animals which have not been killed, after stunning, by laceration of the central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, or by means of gas injected into the cranial cavity, except for animals born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC;

or

- (b) the animal by-product or derived product does not contain and is not derived from bovine, ovine and caprine materials other than those derived from animals born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.
- 2. In addition to the requirements of point 1 of this Section, imports of the animal by-products and derived products referred to in points (d) and (f) of Section A shall be subject to the presentation of a health certificate which has been completed with the following attestation:
 - (a) the animal by-product or derived product originates from a country or region, which is classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC, and in which there has been no BSE indigenous case;

or

(b) the animal by-product or derived product originates from a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC in which there has been a BSE indigenous case, and the animal by-product or derived product was derived from animals born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants, as defined in the OIE Terrestrial Animal Health Code, has been effectively enforced in that country or region.

By way of derogation from the preceding paragraph, the attestation referred to in points (a) and (b) shall not be required for the importation of processed petfood, which is packaged and labelled in accordance with Union legislation.

- 3. In addition to the requirements of points 1 and 2 of this Section, imports of the animal by-products and derived products referred to in Section A, containing milk or milk products of ovine or caprine animal origin and intended for feed, shall be subject to the presentation of a health certificate which has been completed with the following attestation:
 - (a) the ovine and caprine animals from which those animal by-products or derived products have been derived have been kept continuously since birth in a country where the following conditions are fulfilled:
 - (i) classical scrapie is compulsorily notifiable;
 - (ii) an awareness, surveillance and monitoring system is in place;
 - (iii) official restrictions apply to holdings of ovine or caprine animals in the case of a suspicion of TSE or a confirmation of classical scrapie;
 - (iv) ovine and caprine animals affected with classical scrapie are killed and completely destroyed;
 - (v) the feeding to ovine and caprine animals of meat-and-bone meal or greaves of ruminant origin, as defined in the OIE Terrestrial Animal Health Code, has been banned and effectively enforced in the whole country for a period of at least the preceding seven years;
 - (b) the milk and milk products of ovine or caprine animals originate from holdings where no official restrictions are imposed due to a suspicion of TSE;
 - (c) the milk and milk products of ovine or caprine animals originate from holdings where no case of classical scrapie has been diagnosed during a period of at least the preceding seven years or, following the confirmation of a case of classical scrapie:
 - (i) all ovine and caprine animals on the holding have been killed and destroyed or slaughtered, except for breeding rams of the ARR/ARR genotype, breeding ewes carrying at least one ARR allele and no VRQ allele and other ovine animals carrying at least one ARR allele;

or

- (ii) all animals in which classical scrapie was confirmed have been killed and destroyed, and the holding has been subjected for a period of at least two years since the date of confirmation of the last classical scrapie case to intensified TSE monitoring, including testing with negative results for the presence of TSE in accordance with the laboratory methods set out in point 3.2 of Chapter C of Annex X, of all of the following animals which are over the age of 18 months, except ovine animals of the ARR/ARR genotype:
 - animals which have been slaughtered for human consumption; and
 - animals which have died or been killed on the holding but which were not killed in the framework of a disease eradication campaign.'

ANNEX II

Annex XV to Regulation (EU) No 142/2011 is amended as follows:

(1) Chapters 1 to 3(F) are replaced by the following:

'CHAPTER 1

Health certificate

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

COUNTRY: Veterinary certificate to EU

	l.1.	Consignor					1.2.	Certificate refere	ence No	1.2	2.a.	
		Name Address					1.3.	Central compete	nt authority			
		Tel.					1.4.	Local competent	authority			
	1.5.	Consignee					1.6.	Person responsi	ble for the loa	ad in E	U	
		Name						Name				
nent		Address						Address				
ignr												
suo:		Postcode						Postcode				
o pau		Tel.						Tel.				
Part I : Details of dispatched consignment	1.7.	Country of origin	ISO code	1.8.	Region of origin	Code	1.9.	Country of destination	ISO code	I.10.	Region of destination	Code
ls of												
: Detai	l.11.	Place of orig	in				I.12.	Place of destinat	tion			
artı		Mana	Δ		l				0			
		Name Address	Aţ	oprova	l number			Name		om war oval nu	rehouse	Ц
		Name	Δη	nrova	l number			Address	Appro	oval III	iiiibei	
		Address	~⊦	prova	i namber			Addices				
		Name	Ar	prova	l number			Postcode				
		Address										
	I.13.	Place of load	ling				1.14.	Date of departure	e			
	I.15.	Means of tra	nsport				I.16.	Entry BIP in EU				
		Aeroplane 🗖	•		Railway wa	gon 🛘						
		Road vehicle					1.17.					
		Identification						_				
		Documentati	on references	;								



I.18.	Description of commodity	/			I.19. Commo	odity c	ode (HS code)
						1.20.	Quantity
I.21.	Temperature of product Ambient □	Chilled □		Frozen 🗆]	I.22.	Number of packages
1.23.	Seal/Container No					1.24.	Type of packaging
1.25.	Commodities certified for	. .					
	Animal feedingstuff \square	Technic	cal use 🛘	Manufacture of	petfood \square		
1.26.	For transit through EU to	third country		I.27. For import of	or admission in	to EU	
	Third country	ISO code					
1.28.	Identification of the comm		oval number	of establishments			
Sp	ecies (Scientific Nat name)	ture of commodity	Manufactı	uring plant	Net weight		Batch number

II.

II.1.

(2) and/or

[-

Part II: Certification

COUNTRY

Processed animal protein, other than those derived from farmed insects, not intended for human consumption including mixtures and products other than petfood containing such protein

						includir	ng mixtures and		other than petfood ining such protein		
Heal	th informatio	n		II.a.	Certifica	te reference N	lo	II.b.			
the E	European Par	liamen	t and of the	Counci	I (^{1a}) and	in particular	Article 10 thereof	f, and Com	C) No 1069/2009 of imission Regulation nex XIV thereto and		
	processed ar ided for huma				bedirozek	above conta	ins exclusively	processed	animal protein not		
(a)	authority in accordance with Article 24 of Regulation (EC) No 1069/2009, and										
(b)	(b) has been prepared exclusively with the following animal by-products:										
	(²) either	[-	animals k	illed, an	d which	are fit for hu		on in acco	, bodies or parts of ordance with Union al reasons;]		
	(²) and/or	[-	carcases and the following parts originating either from animals that has slaughtered in a slaughterhouse and were considered fit for slaughter for consumption following an ante-mortem inspection or bodies and the following animals from game killed for human consumption in accordance with Union legisless								
			con	sumptior	ses or bodies and parts of animals which are rejected as unfit for human mption in accordance with Union legislation, but which did not show any of disease communicable to humans or animals;						
			(ii) hea	ds of pou	ıltry;						
			` '	phalang					and feet, including us and metatarsus		
			(iv) pig	bristles;							
			(v) feat	hers;]							
	(²) and/or	[-	to humar slaughterl	ns or ai nouse af	nimals, o ter having	btained from g been consid	animals that I	nave been ghter for h	cable through blood slaughtered in a uman consumption ion;]		
	(²) and/or	[-		ion, incli	uding deg				tended for human or separator sludge		
	(²) and/or	[-	longer int	ended fo uring or	r human packagin	consumption	for commercial r	easons or	origin, which are no due to problems of no risk to public or		
	(²) and/or	[-		nat did n	ot show				originating from live ugh that product to		
	(²) and/or	[-					als, except sea r umans or animals		which did not show		

animal by-products from aquatic animals originating from establishments or plants manufacturing products for human consumption;]

Processed animal protein, other than those derived from farmed insects, not intended for human consumption including mixtures and products other than petfood containing such protein

					,			containing such protein
II.	Health in	formatio	า		1	l.a.	Certificate reference No	II.b.
	(2)	and/or	[-				rial originating from animals which dic ough that material to humans or animal	
				(i)	shells fro	om s	hellfish with soft tissue or flesh;	
				(ii)	the follo	wing	originating from terrestrial animals:	
					— hat	cher	ry by-products,	
					— egç	gs,		
					— egg	g by-	products, including egg shells;	
				(iii)	day-old (chicl	ks killed for commercial reasons;]	
	(2)	and/or	[-		c and te her than		trial invertebrates other than species p ects;]	athogenic to humans or animals
	(2)	and/or	[-	Categ	ory 1 ma	ateri	thereof of the zoological orders of Ro al as referred to in Article 8(a)(iii), (iv) ticle 9(a) to (g) of Regulation (EC) No	and (v) and Category 2 material
	and							
	(c) ha	s been su	bjecte	d to the	followin	g pr	ocessing standard:	
	(2)	either	at a	pressu	ire (abso	olute	erature of more than 133°C for at leas) of at least 3 bars produced by satur ot more than 50 millimetres;]	
	(2)	or				ndica	nmalian protein other than fishmeal, the ate the processing method) as set ou //2011;]	
	(2)	or	(ind		e proces		the processing method 1-2-3-4-5-6-7 g method) as set out in Chapter III o	
	(2)	or	(ind No	icate th 142/201	e proces	ssin e in	olood, the processing method 1-2-3-4-5 g method) as set out in Chapter III of case of method 7 a heat treatment of e;]	of Annex IV to Regulation (EU)
II.2.	the comp following			examine	ed a ran	dom	sample immediately prior to dispatch	and found it to comply with the
	Salmonel	la:			Absence	e in 2	25 g: n = 5, c = 0, m = 0, M = 0	
	Enteroba	cteriaceae) :		n = 5, c	= 2,	m = 10, M = 300 in 1g;	
II.3.	the produ	ct has un	dergo	ne all pr	ecaution	ıs to	avoid recontamination with pathogenic	agents after treatment;
II.4.	the end p	roduct:						
	(²) either	[was pa	acked	in new	or sterilis	sed l	bags,]	

Processed animal protein, other than those derived from farmed insects, not intended for human consumption including mixtures and products other than petfood containing such protein

						containing such protein
II.	Health informati	ion		II.a. Certificate reference No		II.b.
	•	transported in fected before us		n containers or other means of transpo	rt that	were thoroughly cleaned and
	which bear labels	s indicating 'NO	T FOI	R HUMAN CONSUMPTION';		
II.5.	the end product v	was stored in en	close	d storage;		
(²) [II.6.	the processed a ruminant origin a	•	or pr	duct described above contains or is	derived	d from animal-by products of
	(²) either		with	a country or region, which is classified Decision 2007/453/EC, and in which t		
	(²) or	with Decision by-product ban on the	on 20 or de e fee as de	a country or region classified as posing 07/453/EC in which there has been an irived product were derived from animalading of ruminants with meat-and-bone ined in the OIE Terrestrial Animal Health region, and]	ndigeno s born e mea	ous BSE case, and the animal after the date from which the al and greaves derived from
	(²) either	[is derived f	from o	ther ruminants than bovine, ovine or cap	rine ar	nimals.]
	(²) or	[is derived f	from b	ovine, ovine or caprine animals and does	s not c	ontain and is not derived from:
			contir	e, ovine and caprine materials other th uously reared and slaughtered in a cou ible BSE risk in accordance with Decisio	ıntry oı	r region classified as posing a
		(²) or	[(a)	specified risk material as defined in po No 999/2001 of the European Parliame		
			(b)	mechanically separated meat obtained caprine animals, except from those a reared and slaughtered in a country negligible BSE risk in accorda 2007/453/EC (5), in which there has be	nimals y or r nce	s that were born, continuously egion classified as posing a with Commission Decision
			(c)	animal by-product or derived producaprine animals which have been killed central nervous tissue by means of a introduced into the cranial cavity, or cranial cavity, except for those animals and slaughtered in a country or region risk in accordance with Decision 2007/	d, after an elor by mo s that v classif	r stunning, by laceration of the ngated rod-shaped instrument eans of gas injected into the were born, continuously reared ied as posing a negligible BSE
II.7.	the processed an	nimal protein or p	produ	ct described above:		
		s not contain mil ed animals, othe		nilk products of ovine or caprine animal fur animals.]	origin	or is not intended for feed for
				ducts of ovine or caprine animal origin mals, and the milk or milk products:	and is	s intended for feed for farmed
				ne and caprine animals which have bed lowing conditions are fulfilled:	en kep	ot continuously since birth in a
		(i) cla	assica	scrapie is compulsorily notifiable;		

Processed animal protein, other than those derived from farmed insects, not intended for human consumption including mixtures and products other than petfood containing such protein

			-				
II.	Health info	ormation		II.a.	Certificate reference No	_	II.b.
		(ii)	an aware	eness,	surveillance and monitoring sy	stem is in	place for classical scrapie;
		(iii)			ons apply to holdings of ovin SE or the confirmation of classion		
		(iv)	ovine an	d capr	ine animals affected with class	ical scrapi	e are killed and destroyed;
		(v)	defined i Health (0	n the OIE),	ovine and caprine animals of Terrestrial Animal Health Cod of ruminant origin has been l for a period of at least the prec	le of the V banned ar	Norld Organisation for Animal and effectively enforced in the
		(b) originate fr	rom holding	s whe	re no official restrictions are im	posed due	e to a suspicion of TSE;
					ere no case of classical scrapie n years or, following the confirm		
		(²) either	slaughte carrying	red, e at lea	caprine animals on the holdi xcept for breeding rams of th st one ARR allele and no VRQ RR allele;]	ne ARR/AI	RR genotype, breeding ewes
		(²) or	and the I of confir including laborator No 999/2	noldin mation testir y met 2001,	which classical scrapie was cog has been subjected for a pen of the last classical scraping with negative results for the hods set out in point 3.2 of Crof all of the following animals nimals of the ARR/ARR genoty	riod of at l e case to presence napter C o which are	least two years since the date intensified TSE monitoring, of TSE in accordance with the f Annex X to Regulation (EC)
			— anim	ıals wl	nich have been slaughtered for	human co	onsumption; and
					nich have died or been killed or work of a disease eradication ca		
II.8.					scribed above contains or is d ment of the Consignor referred		
	(²) either	[not intended for	r the produc	ction o	f feed for farmed animals, othe	r than fur	animals.]
	(²) (⁶) or	[intended for the	e productio	n of fe	eed for non-ruminant farmed a	nimals, ot	her than fur animals, and the

Notes

Part I:

Box reference I.6: Person responsible for the consignment in the European Union: this box is required to be filled in only if
it is a certificate for a commodity to be transited through the European Union; it may be filled in if the certificate is for a
commodity that is to be imported into the European Union.

Regulation (EC) No 152/2009 (7).]

Consignor has undertaken to ensure that the Border Inspection Post of entry will be provided with the results of the analyses carried out in accordance with the methods set out in Annex VI to Commission

- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. Products in transit may only be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in the event of unloading and reloading.

COUNTRY

Processed animal protein, other than those derived from farmed insects, not intended for human consumption including mixtures and products other than petfood containing such protein

			i	ncluding mixtures and	products other than petfood containing such protein			
II.		Health information	II.a. Certificate refe	rence No	II.b.			
_	Вох і	reference I.19: use the appropriate HS o	ode: 05.05; 05.06; 05.0	07; 05.11; 23.01 or 23.09).			
_		reference I.25: technical use: any us uction or manufacturing of pet food.	e other than feeding	of farmed animals, oth	er than fur animals, and the			
_	Вох і	reference I.26 and I.27: fill in according	o whether it is a transit	or an import certificate.				
_	Suida	reference I.28: Species: select from th ae, Pesca, Mollusca, Crustacea, inverte cientific name of the fish.						
Part	: II:							
(^{1a})	OJ L	300, 14.11.2009, p. 1.						
(1b)	OJ L	54, 26.2.2011, p. 1.						
(²)	Delete as appropriate.							
(3)	Whe	re:						
	n =	number of samples to be tested;						
	m = threshold value for the number of bacteria; the result is considered satisfactory if the number of bacteria in all samples does not exceed m;							
	M = maximum value for the number of bacteria; the result is considered unsatisfactory if the number of bacteria in one or more samples is M or more; and							
	c =	number of samples the bacterial cou acceptable if the bacterial count of the			sample still being considered			
(4)	OJ L	147, 31.5.2001, p. 1.						
(⁵)	OJ L	172, 30.6.2007, p. 84.						
(⁶)	desc than (EC) resul	Person responsible for the load referreribed in this health certificate is intended fur animals, the consignment must be No 152/2009, in order to verify the abt of such analysis must be attached to ection post.	I to be used for the pro analysed, in accordand sence of unauthorised	duction of feed for non-rice with the methods set I constituents of animal	uminant farmed animals, other out in Annex VI to Regulation origin. The information on the			
(⁷)	OJ L	54, 26.2.2009, p. 1.						
_	The	signature and the stamp must be in a di	ferent colour to that of	the printing.				
_		for the person responsible for the consi must accompany the consignment until i			is only for veterinary purposes			
Offic	cial ve	terinarian/Official inspector						
	Nam	e (in capital letters):		Qualification a	and title:			
	Date	:		Signature:				
	Stam	np:						

CHAPTER 1a

Health certificate

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

COL	JNTRY	/ :				Veterinary certific	ate to EU	
	l.1.	Consignor	1.2.	Certificate refer	ence No	I.2.a.		
		Name	1.3.	Central compete	ent authority			
		Address	1.4.	Local competen	nt authority			
		Tel.						
	1.5.	Consignee	1.6.	Person respons	sible for the lo	ad in EU		
nent		Name		Name				
ignn		Address		Address				
cons		Postcode		Postcode				
pet		Tel.		Tel.				
oatch	1.7.	Country ISO code I.8. Region of Code	1.9.	Country of	ISO	I.10. Region of	Code	
dist		of origin origin		destination	code	destination	1	
s of								
Part I : Details of dispatched consignment	l.11.	Place of origin	I.12.	Place of destina	ation			
<u> </u>							_	
Part		Name Approval number				om warehouse		
		Address		Name	Appr	oval number		
		Name Approval number		Address				
		Address						
		Name Approval number		Postcode				
		Address	—					
	1.13.	Place of loading	1.14.	Date of departu	re			
	l.15.	Means of transport	I.16.	Entry BIP in EU				
		Aeroplane Ship Railway wagon Railway wagon						
		Road vehicle Other	l.17.					
		Identification						
		Documentation references						
	l.18.	Description of commodity			I.19. Comr	modity code (HS code	r)	
						I.20. Quantity		
	I.21.	Temperature of product				I.22. Number of p	ackages	
		Ambient ☐ Chilled ☐	Frozen 🗖					
	1.23.	Seal/Container No				I.24. Type of pack	kaging	



1.25.	Commodities certi-	fied for:				
	Animal feedingstut	ff 🗆	Technical use \square		Manufacture of pe	tfood □
1.26.	For transit through	EU to third country		I.27. For imp	port or admission into EU	
	Third country	ISO cod	e			
1.28.	Identification of the	e commodities	Approval number	of establishme	ents	
Sp	ecies (Scientific name)	Nature of commo	dity Manufactu	uring plant	Net weight	Batch number

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

			and products other than petfood	containing such protein
	II.	Health information	II.a. Certificate reference No	II.b.
		the European Parliament and of the Council	lare that I have read and understood Regulati cil (¹a) and in particular Article 10 thereof, and ction 1 of Chapter II of Annex X, and Chapter I	d Commission Regulation
ation	II.1.	the processed animal protein derived from processed animal protein not intended for his	om farmed insects or product described ab uman consumption that:	ove contains exclusively
Part II: Certification			in establishment or plant approved and supe 4 of Regulation (EC) No 1069/2009, and	ervised by the competent
Part II:		(b) has been prepared exclusively from f	armed insects of the following species:	
		(²) either [- Black Soldier Fly	(Hermetia illucens);]	
		(²) and/or [- Common Housef	y (Musca domestica);]	
	_	(²) and/or [- Yellow Mealworm	n (Tenebrio molitor);]	
		(²) and/or [- Lesser Mealworm	n (Alphitobius diaperinus);]	
		(²) and/or [- House cricket (Ad	cheta domesticus);]	
		(²) and/or [- Banded cricket (0	Gryllodes sigillatus);]	
		(²) and/or [- Field Cricket (<i>Gry</i>	rllus assimilis).]	
		and		
		(c) has been processed by method [1]- (EU) No 142/2011;	[2]-[3]-[4]-[5]-[7] (²) as set out in Chapter III c	of Annex IV to Regulation
		and		
		(d) the substrate for the feeding of fa following products of animal origin of	rmed insects may only contain products of Category 3 material:	non-animal origin or the
		— fishmeal;		
		 blood products from non-rumina 	ants;	
		 — di and tricalcium phosphate of a 	nimal origin;	
		 hydrolysed proteins from non-ru 	ıminants;	
		 hydrolysed proteins from hides 	and skins of ruminants;	
		 gelatine and collagen from non- 	ruminants;	
		 eggs and egg products; 		
		 milk, milk based-products, milk- 	derived products, and colostrum;	
		— honey;		
		— rendered fats;		

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

						and pro	ducts othe	r than petfoo	d containing such	protein
II.	Health inf	formatio	า		II.a.	Certificate	reference N	lo	II.b.	
	and									
	ma	terials of		in than tho					en in contact with ar e did not contain n	•
II.2.	the compe following s			ned a rando	om samp	ole immedia	tely prior to	dispatch and	found it to comply v	with the
	Salmonell	a:		Absence i	n 25 g: n	= 5, c = 0, ı	m = 0, M =	0		
	Enterobac	cteriaceae) :	n = 5, c = 2	2, m = 1	0, M = 300 ii	n 1g;			
II.3.	the produc	ct has und	dergone all p	recautions	to avoid	recontamina	ation with p	athogenic age	nts after treatment;	
II.4.	the end pr	oduct:								
	(²) either	[was pa	acked in new	or sterilise	ed bags,]					
	(²) or		ansported in cted before u		ontainers	or other m	eans of tra	nsport that we	ere thoroughly clean	ed and
								ESSED INSEC AND FUR ANI	T PROTEIN – SHAL MALS';	_L NOT
II.5.	the end product was stored in enclosed storage;									
(²) [II.6.	the proce			or product	t describ	ped above	contains or	is derived f	om animal-by prod	ucts of
	(2)	either		e with Dec					ng a negligible BSE been no indigenou	
	(2)	or	with Decis by-product ban on th ruminants,	ion 2007/45 or derived ne feeding	53/EC in d produc of rum I in the C	which there t were deriv inants with DIE Terrestri	e has been red from an meat-and	an indigenous imals born aft -bone meal a	ble BSE risk in acco BSE case, and the er the date from whand greaves derive as been effectively en	animal nich the ed from
	(2)	either	[is derived	from other	ruminan	ts than bovi	ne, ovine oi	caprine anim	als.]]	
	(2)	or	[is derived	from bovin	e, ovine	or caprine a	inimals and	does not cont	ain and is not derive	d from:
			(²) either	continuous	sly reare	d and slaug	ghtered in a		derived from animal gion classified as p 33/EC.]]	
			(²) or						Annex V to Regulation the Council (4);	on (EC)
				ca rea ne	prine an ared and gligible	imals, exce d slaughtere BSE risk	pt from tho ed in a co c in acc	se animals th ountry or regi ordance wit	bones of bovine, o at were born, contin on classified as po n Commission D igenous BSE case,	nuously

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

						and products other than petfood containing such protein
II.	Health in	format	ion			II.a. Certificate reference No II.b.
				cap cer intr cra and	rine tral odu nial I sla	al by-product or derived product obtained from bovine, ovine or ne animals which have been killed, after stunning, by laceration of the ral nervous tissue by means of an elongated rod-shaped instrument duced into the cranial cavity, or by means of gas injected into the al cavity, except for those animals that were born, continuously reared slaughtered in a country or region classified as posing a negligible BSE in accordance with Decision 2007/453/EC.]]]
II.7.	the proces	ssed ar	nimal proteir	n or product des	scrit	ribed above:
	(²) either			n milk or milk p other than fur a		oducts of ovine or caprine animal origin or is not intended for feed for imals.]
	(²) or	-		•		of ovine or caprine animal origin and is intended for feed for farmed and the milk or milk products:
		(a)				d caprine animals which have been kept continuously since birth in a conditions are fulfilled:
			(i)	classical scra	pie	ie is compulsorily notifiable;
			(ii)	an awarenes	3, SI	surveillance and monitoring system is in place for classical scrapie;
			(iii)			ons apply to holdings of ovine or caprine animals in the case of a SE or the confirmation of classical scrapie;
			(iv)	ovine and cap	rin [,]	ine animals affected with classical scrapie are killed and destroyed;
			(v)	defined in the Health (OIE)	e Te	ovine and caprine animals of meat-and-bone meal or greaves, as Terrestrial Animal Health Code of the World Organisation for Animal of ruminant origin has been banned and effectively enforced in the for a period of at least the preceding seven years;
		(b)	originate fr	om holdings wh	ere	re no official restrictions are imposed due to a suspicion of TSE;
		(c)	•	•		ere no case of classical scrapie has been diagnosed during a period of a years or, following the confirmation of a case of classical scrapie:
			(²) either	slaughtered,	exc ast	caprine animals on the holding have been killed and destroyed or xcept for breeding rams of the ARR/ARR genotype, breeding ewes st one ARR allele and no VRQ allele and other ovine animals carrying RR allele;]
			(²) or	and the holdi of confirmation including test laboratory me No 999/2001	ng l on ing etho , of	which classical scrapie was confirmed have been killed and destroyed, g has been subjected for a period of at least two years since the date of the last classical scrapie case to intensified TSE monitoring, g with negative results for the presence of TSE in accordance with the hods set out in point 3.2 of Chapter C of Annex X to Regulation (EC) of all of the following animals which are over the age of 18 months, nimals of the ARR/ARR genotype:
				— animals v	vhic	nich have been slaughtered for human consumption; and
						nich have died or been killed on the holding but which were not killed in work of a disease eradication campaign.]]

II.8. [the processed animal protein or product described above contains or is derived from animal-by products of non-ruminant origin and is, according to the statement of the Consignor referred to in Box I.1,

COUNTRY

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

II.	Health information	II.a. Certificate reference No	II.b.
	(2) attended for the conduction of		-1- 1

- (2) either [not intended for the production of feed for farmed animals, other than fur animals.]
- (²) (6) or [intended for the production of feed for non-ruminant farmed animals, other than fur animals, and the Consignor has undertaken to ensure that the border inspection post of entry into the European Union will be provided with the results of the analyses carried out in accordance with the methods set out in Annex VI to Commission Regulation (EC) No 152/2009 (7).]

Notes

Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is required to be filled in only if
 it is a certificate for a commodity to be transited through the European Union; it may be filled in if the certificate is for an a
 commodity to be imported into the European Union.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for a transit commodity. Products
 in transit may only be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in the event of unloading and reloading.
- Box reference I.19: use the appropriate HS code: 05.11, 23.01 or 23.09.
- Box reference I.25: technical use: any use other than feeding of farmed animals, other than fur animals, and the
 production or manufacturing of pet food..
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: Species: insects, specify its scientific name.

Part II:

- (1a) OJ L 300, 14.11.2009, p. 1.
- $(^{1b})$ OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.
- (3) Where:
 - n = number of samples to be tested;
 - m = threshold value for the number of bacteria; the result is considered satisfactory if the number of bacteria in all samples does not exceed m;
 - M = maximum value for the number of bacteria; the result is considered unsatisfactory if the number of bacteria in one or more samples is M or more; and
 - c = number of samples the bacterial count of which may be between m and M, the sample still being considered acceptable if the bacterial count of the other samples is m or less.
- (4) OJ L 147, 31.5.2001, p. 1.
- (5) OJ L 172 30.6.2007, p. 84.

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

II.	Health information	II.a. Certificate referenc	e No	II.b.		
(⁶)	(6) The Person responsible for the load referred to in Box I.6 must ensure that, if the processed animal protein or product described in this health certificate is intended to be used for the production of feed for non-ruminant farmed animals, other than fur animals, the consignment must be analysed, in accordance with the methods set out in Annex VI to Regulation (EC) No 152/2009, in order to verify the absence of unauthorised constituents of animal origin. The information on the result of such analysis must be attached to this health certificate when presenting the consignment at an EU Border Inspection Post.					
(7)	(⁷) OJ L 54, 26.2.2009, p. 1.					
_	The signature and the stamp must be in a different colour to that of the printing.					
_	 Note for the person responsible for the consignment in the European Union: This certificate is only for veterinary purposes and must accompany the consignment until it reaches the border inspection post. 					
Offic	cial veterinarian/Official inspector					
	Name (in capital letters):		Qualification and t	itle:		
	Date: Signature:					
	Stamp:					
1						

CHAPTER 2(A)

Health certificate

For milk, milk-based products and milk-derived products not intended for human consumption for dispatch to or transit through (2) the European Union

COL	JNTRY	/ :				Veterinary certificat	e to EU	
	l.1.	Consignor	1.2.	I.2. Certificate reference No I.2.a.				
		Name	1.3.	I.3. Central competent authority				
		Address	1.4.	I.4. Local competent authority				
		Tel.	ļ					
	1.5.	Consignee	I.6.		r the load	d in EU		
nen		Name		I.3. Central competent authority I.4. Local competent authority I.6. Person responsible for the load in EU Name Address Postcode Tel. I.9. Country of destination code destination I.12. Place of destination Custom warehouse Approval number Address Postcode				
ignr		Address		Address				
cons		Postcode		Postcode				
ped		Tel.						
Part I : Details of dispatched consignment	1.7.	Country ISO code I.8. Region of Code origin	1.9.	Country of ISC			Code	
Details	l.11.	Place of origin	I.12.	Place of destination				
# ::		Name Approval number				Custom warehouse		
Ъа		Address		Name		Approval number		
		Name Approval number		Address				
		Address						
		Name Approval number	Postcode					
		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 Other	1.17.	Number(s) of CITES				
		Identification						
		Documentation references						
	I.18.	Description of commodity	•	I.19.	Comm	odity code (HS code)		
						I.20. Quantity		
	I.21.	Temperature of product				I.22. Number of pac	kages	
		Ambient ☐ Chilled ☐		Frozen \square				
	1.23.	Seal/Container No				I.24. Type of packa	ging	

1.25.	Commodities certified for:			
	Animal feedingstuff ☐ Technical use ☐	Further process D	Production of per	tfood 🗖
1.26.	For transit through EU to thir	d country	I.27. For import or admission into EU	
	Third country	ISO code		
1.28.	Identification of the commodi	ities		
		Approval number	of establishments	
	Species (Scientific name)	Manufacturing plant	Net weight	Batch number

Milk, milk-based products and milk-derived products not for human consumption

Γ										
	II.	Health inforn	nation		II.a.	Certificate reference No		II.b.		
		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council (¹a¹), and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011 (¹b¹), and in particular Section 4 of Chapter II of Annex X, and Chapter I of Annex XIV thereto, and certify that the milk (²), the milk-based products (²) and milk-derived products (²) referred to in box I.28 comply with the following conditions:								
u	II.1.							name of exporting country) (3),		
they were produced and derived in							nich has been free from foot-and-			
Part	II.2. they were produced from raw milk derived from animals which at the time of milking did not show clinical signs of any disease transmissible through milk to humans or animals, and which had been kept for a period of at least 30 days prior to production on holdings that were not subject to official restrictions due to foot-and-mouth disease or rinderpest;							een kept for a period of at least		
	II.3.	they are milk	or milk products	that:						
		(²) either	[have undergo	ne one c	f the ti	reatments or combinations the	ereof des	cribed in point II.4;]		
		(²) or				animals of species suscept subjected to one of the treatn		pot-and-mouth disease, and that scribed in point II.4 and:		
			(²) either	[the whe	ey was	collected at least 16 hours af	ter clottin	g and has a pH below 6;]		
						s been produced at least 21 es of FMD have been detected		ore the shipping and during that xporting country;]		
			(²) (⁵) or	voyage	durati		efore the	in consideration of the foreseen consignment is presented to a		
	II.4.	they have bee	en subject to one	e of the f	ollowin	g treatments:				
		(²) either						t 15 seconds, or an equivalent st in bovine milk, in combination		
			(²) either	15 seco	nds or			pasteurisation at 72°C for at least self achieves a negative reaction		
			(²) or			t drying process that in the additional heating to 72°C or		of milk intended for feeding is		
		(²) or	[a subse level be		educed a	nd kept for at least one hour at a				
	the date							roduced at least 21 days prior to s of FMD have been detected in		
			(²) (⁵) or	conside	ration	of the foreseen voyage duration	on, being	(insert the date), this date, in at least 21 days prior to the date aspection post of the European		
			(²) or	[sterilisa	ition at	t a level of at least F ₀ 3;]]				

Milk, milk-based products and milk-derived products not for human consumption

		4*		0 15 1 5	nor numan consumption			
II.	Health inforr	mation	II.	a. Certificate reference No	II.b.			
	(²) or	[ultra high tempera	ture tre	atment at 132°C for at least one second i	n combination with:			
				uent drying process that in the case with additional heating to 72°C or higher;]				
			ubsequ el below	lent process by which the pH is reduced a $(6;]$	and kept for at least one hour at a			
		the	date of	condition that the milk/milk product has been produced at least 21 days prior to late of shipping and during that period no cases of FMD has been detected in the rting country;]				
		con that	[the milk/milk product has been produced on// (insert the date), this date, in consideration of the foreseen voyage duration, being at least 21 days prior to the date that the consignment is presented to a border inspection post of the European Union;]]					
II.5.	every precau processing;	ution was taken to	avoid	contamination of the milk/milk-based pr	roduct/milk-derived product after			
II.6.	the milk/milk-	based product/milk-d	lerived	product was packed:				
	(²) either	[in new containers;]					
	(²) or	[in vehicles or bulk containers disinfected prior to loading using a product approved by the competent authority;]						
	and	and the containers are marked so as to indicate the nature of the milk/milk-based product/milk-derived product and bear labels indicating that the product is Category 3 material and not intended for human consumption;						
II.7.	the milk, milk	-based products and	milk-de	erived products described above:				
	(²) either	[does not contain n farmed animals, ot		milk products of ovine or caprine animal on n fur animals.]	rigin or is not intended for feed for			
	(²) or	-		ducts of ovine or caprine animal origin ar imals, and the milk or milk products:	nd is intended for feed for farmed			
				d from ovine and caprine animals which hountry where the following conditions are				
		(i)		classical scrapie is compulsorily notifia	ble;			
		(ii)		an awareness, surveillance and mo classical scrapie;	enitoring system is in place for			
		(iii)		official restrictions apply to holdings o case of a suspicion of TSE or the confi				
		(iv)		ovine and caprine animals affected widestroyed;	th classical scrapie are killed and			
		(v)		the feeding to ovine and caprine and greaves, as defined in the Terrestrial organisation for Animal Health (OIE banned and effectively enforced in the least the preceding seven years;	Animal Health Code of the World E), of ruminant origin has been			
		· ,	inate fr SE;	om holdings where no official restrictions	s are imposed due to a suspicion			

COUNTRY

Milk, milk-based products and milk-derived products not for human consumption

			T
II.	Health information		II.a. Certificate reference No II.b.
	(c)	during a	te from holdings where no case of classical scrapie has been diagnosed a period of at least the preceding seven years or, following the confirmation of of classical scrapie:
		(²) eithe	[all ovine and caprine animals on the holding have been killed and destroyed or slaughtered, except for breeding rams of the ARR/ARR genotype, breeding ewes carrying at least one ARR allele and no VRQ allele and other ovine animals carrying at least one ARR allele;]
		(²) or	[all animals in which classical scrapie was confirmed have been killed and destroyed, and the holding has been subjected for a period of at least two years since the date of confirmation of the last classical scrapie case to intensified TSE monitoring, including testing with negative results for the presence of TSE in accordance with the laboratory methods set out in point 3.2 of Chapter C of Annex X to Regulation (EC) No 999/2001 (⁶), of all of the following animals which are over the age of 18 months, except ovine animals of the ARR/ARR genotype:
			 animals which have been slaughtered for human consumption; and
			 animals which have died or been killed on the holding but which were not killed in the framework of a disease eradication campaign.]]
1			

Notes

Part I:

- Box reference I.6: Person responsible for the load in the European Union: this box is required to be filled in only if it is a
 certificate for a commodity to be transited through the European union; it may be filled in if the certificate is for a
 commodity to be imported into the European Union.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to be provided. In the case of unloading and reloading, the consignor must inform the border inspection post of the European Union.
- Box reference I.19: use the appropriate Harmonised System (HS) code of the World Customs Organisation: 04.01; 04.02; 04.03; 04.04; 23.09.10, 23.09.90, 35.01, 35.02 or 35.04.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) must be included.
- Box reference I.25: technical use: any use other than feeding of farmed animals, other than fur animals, and the
 production or manufacturing of pet food.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: 'Manufacturing plant': provide the registration number of treatment or processing establishment.

Part II:

- (1a) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.

COUNTRY

Milk, milk-based products and milk-derived products not for human consumption

II.	Health information	II.a.	Certificate reference No		II.b.				
(2)	Delete as appropriate.								
(3)	For completion if the authorisation to import into or transit through the European Union is restricted to certain regions of the third country concerned.								
(4)	OJ L 175, 10.7.2010, p. 1.								
(5)	this condition applies only to third countries listed in column 'A' of Annex I to Regulation (EU) No 605/2010.								
(6)	OJ L 147, 31.5.2001, p. 1.								
_	The signature and the stamp must be in a d	ifferen	t colour to that of the printin	ng.					
_	Note for the person responsible for the cons and must accompany the consignment until				ate is only for veterinary purposes				
0	ficial veterinarian/Official inspector								
	Name (in capital letters):			Qualification	on and title:				
	Date: Signature:								
	Stamp:								

CHAPTER 2(B)

Health certificate

For colostrum and colostrum products from bovine animals not intended for human consumption for dispatch to or transit through (2) the European Union

COL	JNTRY	′ :								Vete	erinary certifica	te to EU			
	l.1.	Consignor					1.2.	Certificate referer	ice No	1.2	2.a.				
		Name					I.3. Central competent authority								
		Address					I.4. Local competent authority								
											in EU				
		Tel.													
	1.5.	Consignee					I.6. Person responsible for the load in EU								
nent		Name						Name							
ignr		Address						Address			.10. Region of Code destination				
cons		Postcode					Postcode								
ped		Tel.						Tel.							
Part I : Details of dispatched consignment	1.7.	Country	ISO code	1.8.	Region of	Code	1.9.	Country of	ISO	I.10.		Code			
f dis		of origin	I.	I	origin	I		destination	code	1	destination	I			
ils o															
Deta	1.11.	Place of origin Name Approval number Address						Place of destination	on						
<u> </u>										Cust	tom warehouse				
- a								Name		Арр	roval number				
		Name Approval number						Address							
		Address													
		Name		Appro	val number			Postcode							
		Address													
	I.13.	Place of lo	ading				I.14. Date of departure								
	I.15.	Means of t	ransport				I.16. Entry BIP in EU								
		Aeroplane	☐ Ship		Railway wa	agon 🗖									
		Road vehic	cle 🔲 Othe	r 🗆			I.17. Number(s) of CITES								
		Identification	on												
	Documentation references														
	I.18. Description of commodity							I.19. Comr	modity (code (HS code)					
								1.20	. Quantity						
	I.21.	Temperatu	re of product							1.22	. Number of page	ckages			
		Ambient 🗆	1		Chilled]		Frozen \square							
	1.23.	Seal/Conta	iner No				I.24. Type of packaging			ging					

28.2.2019

1.25.	Commodities certified for:			
	Animal feedingstuff ☐ Technical use ☐	Further process C	Production of pe	tfood 🗖
1.26.	For transit through EU to third	d country	I.27. For import or admission into EU	
	Third country	ISO code		
1.28.	Identification of the commodi	ties		
		Approval number	of establishments	
	Species (Scientific name)	Manufacturing plant	Net weight	Batch number

Colostrum and colostrum products from bovine animals not for human consumption

	II.	Health informati	on	II.a. Certificate reference No	II.b.					
		the European Pa (EU) No 142/201	rliament and of the 1 (^{1b}), and in particu	n, declare that I have read and understo Council (^{1a}), and in particular Article 10 lar Section 4 of Chapter II of Annex X and lostrum products (²) referred to in box I.28	thereof, and Commission Regulation d Chapter I of Annex XIV thereto, and					
	II.1.	they were produc	ed and derived in		(insert name of exporting country) (3),					
ill carlo		listed in Annex I to Commission Regulation (EU) No 605/2010 (4), and which has been free from foot-and-mout disease (FMD) and rinderpest for a period of 12 months immediately prior to export and has not practise vaccination against rinderpest during that period;								
rait III. Ceitillication	II.2.	any disease trans	smissible through co ne date of productio	derived from animals which at the time of lostrum to humans or animals, and which n on holdings that were not subject to offic	had been kept for a period of at least					
	II.3.	they are colostrum or colostrum products of bovine animals that have been subject to high temperature short time pasteurisation at 72°C for at least 15 seconds, or an equivalent pasteurisation achieving a negative reaction to a phosphatase test in bovine colostrum, in combination with:								
		(²) (⁵) either [the condition that the colostrum or colostrum products have been produced during a period at least 21 days before the date of shipping and during this period no cases of FMD have been detected in the exporting country,]								
		(²) (⁵) or [the condition that the colostrum or colostrum products have been produced on// (inset the date), this date, in consideration of the foreseen voyage duration, being at least 21 day before the consignment is presented to a border inspection post of the European Union,]								
		and		ed from animals subject to regular veter is on which all bovine herds are:	inary inspections to ensure that they					
			(²) (⁵) either	[recognised as officially tuberculosis and	brucellosis free (6),]					
			(²) (⁵) or	[not restricted under the national legislation of the third country of origin for the eradication of tuberculosis and brucellosis,]						
		and	(²) (⁵) either	[recognised as official enzootic-bovine-le	ukosis-free (⁶),]					
			(²) (⁵) or	[included in an official system for the co there has been no evidence as result of disease in the herd during the period of t	clinical and laboratory testing of this					
	II.4.	every precaution	has been taken to a	void contamination of the colostrum/colos	strum product after processing;					
	II.5.	the colostrum or	colostrum product w	as packed:						
		(²) either	[in new containers	,]						
		(²) or	[in vehicles or bu	ilk containers disinfected prior to loadin ty,]	g using a product approved by the					
	and the containers are marked so as to indicate the nature of the colostrum/colostrum product a bear labels indicating that the product is Category 3 material and not intended for hum consumption;									
	II.6.	the colostrum or o	colostrum product d	oes not contain milk or milk products of ov	vine or caprine animal origin.					
	N. 4									
	Notes									
	Part I:									

Box reference I.6: Person responsible for the load in the European Union: this box is required to be filled in only if it is a certificate for a commodity to be transited through the European Union; it may be filled in if the certificate is for a commodity to be imported into the European Union.

COUNTRY

Colostrum and colostrum products from bovine animals not for human consumption

					not for manian concampaon				
II.	Health information	II.a.	Certificate reference No		II.b.				
_	Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity.								
_	Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to be provided. In the case of unloading and reloading in the European Union, the consignor must inform the border inspection post of the European Union.								
_	Box reference I.19: use the appropriate Harmonised System (HS) code of the World Customs Organisation: 04.04.90; 23.09.10, 23.09.90, 35.01, 35.02 or 35.04.								
_	Box reference I.23: for bulk containers, the	contai	iner number and the seal nu	mber (if app	olicable) must be included.				
_	Box reference I.25: technical use: any use other than feeding of farmed animals, other than fur animals, and the production or manufacturing of pet food								
_	Box reference I.26 and I.27: fill in according	to wh	nether it is a transit or an imp	ort certifica	te.				
_	Box reference I.28: 'Manufacturing plant': p	rovide	the registration number of t	he treatmer	nt or processing establishment.				
Part	II:								
(^{1a})	OJ L 300, 14.11.2009, p. 1.								
(1b)	OJ L 54, 26.2.2011, p. 1.								
(²)	Delete as appropriate.								
(3)	For completion if the authorisation for introduction into the European Union is restricted to certain regions of the third country concerned.								
(4)	OJ L 175, 10.7.2010, p. 1.								
(⁵)	This condition applies only to third coun No 605/2010 (OJ L 175, 10.7.2010, p. 1).	tries	authorised in column 'A' o	f Annex I	to Commission Regulation (EU)				
(⁶)	Officially tuberculosis-free and brucellosis-29.7.1964, p. 1977/64) and officially enzo Directive.								
_	The signature and the seal must be in a diff	erent	colour from that of the printil	ng.					
_	Note for the importer: this certificate is only the border inspection post of the European			accompan	y the consignment until it reaches				
Offic	Official veterinarian/Official inspector								
	Name (in capital letters):			Qualification	on and title:				
	Date:			Signature:					
	Stamp:								

Veterinary certificate to EU

COUNTRY:

CHAPTER 3(A)

Health certificate

For canned petfood intended for dispatch to or for transit through (2) the European Union

	l.1.	Consignor Name						Certificate refere	ence No	I.2.a.	
								Central competent authority			
		Address					1.4.	Local competent authority			
		Tel.									
	1.5.	Consignee						6. Person responsible for the load in EU			
ent		Name						Name			
gnr		Address						Address			
onsi											
ğ Ç		Postcode						Postcode			
tche		Tel.						Tel.			
ispa	1.7.	Country of origin	ISO code	1.8.	Region of origin	Code	1.9.	Country of destination	ISO code	I.10. Region of destination	Code
of d		3			g						
Part I : Details of dispatched consignment	l.11.	Place of or	igin					I.12. Place of destination			
: Del											
T.		Name		Appro	val number					Custom warehouse	
ď		Address						Name		Approval number	
		Name		Appro	val number			Address			
		Address									
		Name		Appro	val number			Postcode			
		Address									
	l.13.	Place of loa	ading				1.14.	Date of departur	re ·		
	115	Means of tr	ransnort				116	Entry BIP in EU			
	1. 10.	Wodilo of a	апорот				1.10.	Linky Dir iii Lo			
		Aeroplane		Railway wa	agon 🗖						
		Road vehic	•		·		1.17.				
		Identification	on								
		Documenta	ation referenc	ces							
	I.18.	I.18. Description of commodity				I.19. Commodity code (HS code)					
										23.09	
										I.20. Quantity	
	l.21.	Temperatu	re of product							I.22. Number of pa	ckages
		Ambient \square			Chilled]		Frozen []		
	123	Seal/Conta	iner No							124 Type of packa	aging

1.25.	Commodities certified for:				
	Petfood \square		Technical use □		
1.26.	For transit through EU to the	hird country	I.27. For import or admission into EU		
	Third country	ISO code			
1.28.	Identification of the commo	odities			
		Approval number	of establishments		
	Species (Scientific name)	Manufacturing plant	Net weight	Batch number	

	COUNTR	Y				Canned Petfood			
	II. Health informa		natio	on	II.a. Certificate reference No	II.b.			
		the European Regulation (E	Pa U) N	rliament and of the	n, declare that I have read and understood Regue Council (^{1a}), and in particular Articles 8 and in and in particular Chapter II of Annex XIII and Cha above:	10 thereof, and Commission			
tion	II.1.	.1. has been prepared and stored in an establishment or plant approved and supervised by the competent author accordance with Article 24 of Regulation (EC) No 1069/2009;							
rtifica	II.2.	has been prep	are	d exclusively with th	e following animal by-products:				
Part II: Certification		(²) either	[-	killed, and which a	d parts of animals slaughtered or, in the case of game, bodies or parts of animals hich are fit for human consumption in accordance with Union legislation, but are not human consumption for commercial reasons;]				
		(²) and/or	[-	slaughterhouse an mortem inspection	rcases and the following parts originating either from animals that have been slaughtered in a aughterhouse and were considered fit for slaughter for human consumption following an ante-ortem inspection or bodies and the following parts of animals from game killed for human insumption in accordance with Union legislation:				
				co	carcases or bodies and parts of animals which are rejected as unfit for human consumption in accordance with Union legislation, but which did not show any signs of disease communicable to humans or animals;				
	(ii)				ads of poultry;				
	ind				hides and skins, including trimmings and splitting thereof, horns and feet, including the phalanges and the carpus and metacarpus bones, tarsus and metatarsus bones;				
				(iv) pig	pristles;				
				(v) fea	athers;]				
		(²) and/or	[-	Article 1(3)(d) of	s from poultry and lagomorphs slaughtered or Regulation (EC) No 853/2004 of the Europe did not show any signs of disease communicable	ean Parliament and of the			
		(²) and/or	[-	[- blood of animals which did not show any signs of disease communicable throu humans or animals, obtained from animals that have been slaughtered in a slaughter having been considered fit for slaughter for human consumption following an inspection in accordance with Union legislation;]					
		(²) and/or	[-		s arising from the production of products intend d bone, greaves and centrifuge or separator slud				
		(²) and/or	[-	intended for huma	origin, or foodstuffs containing products of animal consumption for commercial reasons or due to or other defects from which no risk to public or ar	problems of manufacturing or			
		(²) and/or	[-	[- petfood and feedingstuffs of animal origin, or feedingstuffs containing animal by-performed products, which are no longer intended for feeding for commercial reasons problems of manufacturing or packaging defects or other defects from which no risk animal health arise;]					
		(²) and/or	[-		originating from live animals that product to humans or				
		(²) and/or	[-		nd parts of such animals, except sea mammals, varicable to humans or animals;]	which did not show any signs			
		(²) and/or	[-	animal by-products products for human	from aquatic animals originating from plants or en consumption;]	establishments manufacturing			

II.	Health infor	mation	II.a. Certificate reference No II.b.							
	(²) and/or	[- the following	g material originating from animals which did not show any signs of diseas ele through that material to humans or animals:							
		(i)	shells from shellfish with soft tissue or flesh;							
		(ii)	the following originating from terrestrial animals:							
			 hatchery by-products, 							
			— eggs,							
			egg by-products, including egg shells;							
		(iii)	day-old chicks killed for commercial reasons;]							
	(²) and/or	[- animal by-pi humans or a	roducts from aquatic or terrestrial invertebrates other than species pathogenic t nimals;]							
	(²) and/or	Category 1 r	I parts thereof of the zoological orders of Rodentia and Lagomorpha, except material as referred to in Article 8(a)(iii), (iv) and (v) of Regulation (EC) No 1069/200 y 2 material as referred to in Article 9(a) to (g) of that Regulation;]							
	(²) and/or	Council Dire	n animals which have been treated with certain substances which are prohibited bective 96/22/EC (2b), the import of the material being permitted in accordance wit (ii) of Regulation (EC) No 1069/2009;]							
1.3.	has been su	bjected to heat trea	cted to heat treatment to a minimum Fc value of 3 in hermetically sealed containers;							
1.4.		analysed by a random sampling of at least five samples from each processed batch by laboratory diagnostic hod to ensure adequate heat treatment of the whole consignment as foreseen under point II.3;								
1.5.	has undergone all precautions to avoid contamination with pathogenic agents after treatment.									
²) [II.6.	the petfood described above									
	(²) either	[is derived from other ruminants than bovine, ovine or caprine animals.]								
	(²) or	[is derived from	s derived from bovine, ovine or caprine animals and does not contain and is not derived from:							
		(²) either	bovine, ovine and caprine materials other than those derived from animals bor continuously reared and slaughtered in a country or region classified as posing negligible BSE risk in accordance with Decision 2007/453/EC.]]							
		(²) or	[(a) specified risk material as defined in point 1 of Annex V to Regulation (EC No 999/2001 of the European Parliament and of the Council (³);							
			(b) mechanically separated meat obtained from bones of bovine, ovine of caprine animals, except from those animals that were born, continuous reared and slaughtered in a country or region classified as posing negligible BSE risk in accordance with Commission Decisio 2007/453/EC (4), in which there has been no indigenous BSE case,							
			(c) animal by-product or derived product obtained from bovine, ovine of caprine animals which have been killed, after stunning, by laceration of the central nervous tissue by means of an elongated rod-shape instrument introduced into the cranial cavity, or by means of gas injected into the cranial cavity, except for those animals that were born continuously reared and slaughtered in a country or region classified a posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]							

COUNTRY Canned Petfoo									
II.	Health information	II.a.	Certificate reference No	II.b.					
Note	es								
Part	Part I:								
_	Box reference I.6: Person responsible for the consignment in the European Union: this box is required to be filled in only if it is a certificate for a commodity to be transited through the European Union; it may be filled in if the certificate is for a commodity to be imported into the European Union.								
_	Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. Products in transit may only be stored in free zones, free warehouses and custom warehouses.								
_	Box reference I.15: Registration number (rail information is to be provided in the event of u								
_	Box reference I.23: for bulk containers, the co	ontaine	er number and the seal number (if app	olicable) must be given.					
_	Box reference I.25: technical use: any use production or manufacturing of pet food	e othe	er than feeding of farmed animals,	other than fur animals, and the					
_	Box reference I.26 and I.27: fill in according t	o whe	ther it is a transit or an import certifica	te.					
_	Box reference I.28: Species: select from the Suidae, Pesca, Mollusca, Crustacea, invertel			mmalia other than Ruminantia or					
Part	:11:								
(^{1a})	OJ L 300, 14.11.2009, p. 1.								
(^{1b})	OJ L 54, 26.2.2011, p. 1.								
(²)	Delete as appropriate.								
(^{2a})	OJ L 139, 30.4.2004, p. 55.								
(^{2b})	OJ L 125, 23.5.1996, p. 3.								
(³)	OJ L 147, 31.5.2001, p. 1.								
(⁴)	OJ L 172, 30.6.2007, p. 84.								
_	The signature and the stamp must be in a dif	ferent	colour to that of the printing.						
_	Note for the person responsible for the consi and must accompany the consignment until i			ate is only for veterinary purposes					
Official veterinarian/Official inspector									
	Name (in capital letters):		Qualification	on and title:					
	Date:		Signature:						
	Stamp:								

Veterinary certificate to EU

(CHAPTER 3(B)

Health certificate

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

	l.1.	Consignor	1.2.	I.2. Certificate reference No I.2.a.			
		Name	1.3.	Central competent authority			
		Address	1.4.	Local competent authority			
		Tel.					
	1.5.	Consignee	1.6.	·			
nent		Name		Name Address			
ignr		Address					
suo:		Postcode	Destands				
eq		Tel.		Postcode Tel.			
atch	1.7.	Country ISO code I.8. Region of Code	1.9.	Country of ISO	I.10. Region of Code		
disp	1.7.	of origin origin	1.9.	destination code	destination		
o o o							
Part I : Details of dispatched consignment	l.11.	Place of origin	I.12.	12. Place of destination			
art		Name Approval number			Custom warehouse		
		Address		Name	Approval number		
		Name Approval number		Address			
		Address					
		Name Approval number		Postcode			
		Address					
	I.13.	Place of loading	l.14.	Date of departure			
	I.15.	Means of transport	I.16.	Entry BIP in EU			
		Aeroplane ☐ Ship ☐ Railway wagon ☐					
		Road vehicle Other	l.17.				
		Identification					
		Documentation references					
	I.18.	Description of commodity		I.19. Comm	nodity code (HS code)		
					I.20. Quantity		
	I.21.	Temperature of product			I.22. Number of packages		
		Ambient ☐ Chilled ☐		Frozen			
	1.23.	Seal/Container No			I.24. Type of packaging		



1.25.	Commodities certified for:						
	Petfood		Technical use □				
1.26.	For transit through EU to thi	rd country	I.27. For import or admission into EU				
	Third country	ISO code					
1.28.	Identification of the commod	lities					
		Approval number	of establishments				
	Species (Scientific name)	Manufacturing plant	Net weight	Batch number			

COUNTRY Processed petfood other than canned petfood II. **Health information** II.a. Certificate reference No II.b. I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council (1a), and in particular Articles 8 and 10 thereof, and Commission Regulation (EU) No 142/2011 (1b), and in particular Chapter II of Annex XIII and Chapter II of Annex XIV thereto, and certify that the petfood described above: II.1. has been prepared and stored in a plant approved and supervised by the competent authority in accordance with Part II: Certification Article 24 of Regulation (EC) No 1069/2009; II.2. has been prepared exclusively with the following animal by-products: (2) either carcases and parts of animals slaughtered or, in the case of game, bodies or parts of animals killed, and which are fit for human consumption in accordance with Union legislation, but are not intended for human consumption for commercial reasons;] (2) and/or carcases and the following parts originating either from animals that have been slaughtered in a slaughterhouse and were considered fit for slaughter for human consumption following an antemortem inspection or bodies and the following parts of animals from game killed for human consumption in accordance with Union legislation: (i) carcases or bodies and parts of animals which are rejected as unfit for human consumption in accordance with Union legislation, but which did not show any signs of disease communicable to humans or animals; (ii) heads of poultry; (iii) hides and skins, including trimmings and splitting thereof, horns and feet, including the phalanges and the carpus and metacarpus bones, tarsus and metatarsus bones; (iv) pig bristles; (v) feathers;] (2) and/or animal by-products from poultry and lagomorphs slaughtered on the farm as referred to in [-Article 1(3)(d) of Regulation (EC) No 853/2004 of the European Parliament and of the Council (^{2a}), which did not show any signs of disease communicable to humans or animals] (2) and/or blood of animals which did not show any signs of disease communicable through blood to humans or animals, obtained from animals that have been slaughtered in a slaughterhouse after having been considered fit for slaughter for human consumption following an ante-mortem inspection in accordance with Union legislation;] (2) and/or [animal by-products arising from the production of products intended for human consumption, including degreased bone, greaves and centrifuge or separator sludge from milk processing;] products of animal origin, or foodstuffs containing products of animal origin, which are no longer (2) and/or [intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects from which no risk to public or animal health arise;] petfood and feedingstuffs of animal origin, or feedingstuffs containing animal by-products or (2) and/or derived products, which are no longer intended for feeding for commercial reasons or due to problems of manufacturing or packaging defects or other defects from which no risk to public or animal health arise;] (2) and/or [blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk originating from live animals that did not show signs of any disease communicable through that product to humans or animals;] (2) and/or aquatic animals, and parts of such animals, except sea mammals, which did not show any signs ſ-

of diseases communicable to humans or animals;]

COUNTRY Processed petfood other than canned petfood II. **Health information** Certificate reference No II.b. (2) and/or animal by-products from aquatic animals originating from plants or establishments manufacturing products for human consumption;] (2) and/or the following material originating from animals which did not show any signs of disease communicable through that material to humans or animals: (i) shells from shellfish with soft tissue or flesh; the following originating from terrestrial animals: (ii) hatchery by-products, eggs, egg by-products, including egg shells, (iii) day-old chicks killed for commercial reasons:] (2) and/or animal by-products from aquatic or terrestrial invertebrates other than species pathogenic to [humans or animals;] (2) and/or animals and parts thereof of the zoological orders of Rodentia and Lagomorpha, except ſ-Category 1 material as referred to in Article 8(a)(iii), (iv) and (v) of Regulation (EC) No 1069/2009 and Category 2 material as referred to in Article 9(a) to (g) of that Regulation;] (2) and/or material from animals which have been treated with certain substances which are prohibited by Council Directive 96/22/EC (2b), the import of the material being permitted in accordance with Article 35(a)(ii) of Regulation (EC) No 1069/2009;] II.3. (2) either [was subjected to a heat treatment of at least 90 °C throughout its substance;] (2) or [was produced as regards ingredients of animal origin using exclusively products which had been: in the case of animal by-products or derived products from meat or meat products subjected to a heat treatment of at least 90 °C throughout its substance; in the case of milk and milk based products, (i) if they are from third countries or parts of third countries listed in column B of Annex I to Commission Regulation (EU) No 605/2010 (3) submitted to a pasteurisation treatment sufficient to produce a negative phosphatase test; with a pH reduced to less than 6 from third countries or parts of third countries listed in (ii) column C of Annex I to Regulation (EU) No 605/2010, first submitted to a pasteurisation treatment sufficient to produce a negative phosphatase test; (iii) if they are from third countries or parts of third countries listed in column C of Annex I to Regulation (EU) No 605/2010, submitted to a sterilisation process or a double heat treatment where each treatment was sufficient to produce a negative phosphatase test on its own; (iv) if they are from third countries or parts of third countries listed in column C of Annex I to Regulation (EU) No 605/2010, where there has been an outbreak of foot-and-mouth disease in the preceding 12 months or where vaccination against foot-and-mouth disease has been carried out in the preceding 12 months, submitted to either a sterilisation process whereby an Fc value equal or greater than 3 is achieved or an initial heat treatment with a heating effect at least equal to that achieved by a pasteurisation process of at least 72 °C for at least 15 seconds and sufficient to produce a negative reaction to a phosphatase test, followed by

COUNTRY Processed petfood other than canned petfood

II. Health information II.a. Certificate reference No II.b.

either

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

or

- an acidification process such that the pH has been maintained at less than 6 for at least one hour;
- (c) in the case of gelatine, produced using a process that ensures that unprocessed Category 3 material is subjected to a treatment with acid or alkali, followed by one or more rinses with subsequent adjustment of the pH and subsequent, if necessary repeated, extraction by heat, followed by purification by means of filtration and sterilisation;
- (d) in the case of hydrolysed protein produced using a production process involving appropriate measures to minimise contamination of raw Category 3 material, and, in the case of hydrolysed protein entirely or partly derived from ruminant hides and skins produced in a processing plant dedicated only to hydrolysed protein production, using only material with a molecular weight below 10000 Dalton and a process involving the preparation of raw Category 3 material by brining, liming and intensive washing followed by:
 - (i) exposure of the material to a pH of more than 11 for more than three hours at a temperature of more than 80 °C and subsequently by heat treatment at more than 140 °C for 30 minutes at more than 3,6 bar; or
 - (ii) exposure of the material to a pH of 1 to 2, followed by a pH of more than 11, followed by heat treatment at 140 °C for 30 minutes at 3 bar;
- (e) in the case of egg products submitted to any of the processing methods 1 to 5 or 7, as referred to in Chapter III of Annex IV to Regulation (EU) No 142/2011; or treated in accordance with Chapter II of Section X of Annex III to Regulation (EC) No 853/2004;
- (f) in the case of collagen submitted to a process ensuring that unprocessed Category 3 material is subjected to a treatment involving washing, pH adjustment using acid or alkali followed by one or more rinses, filtration and extrusion, the use of preservatives other than those permitted by Union legislation being prohibited;
- (g) in the case of blood products, produced using any of the processing methods 1 to 5 or 7, as referred to in Chapter III of Annex IV to Regulation (EU) No 142/2011;
- (h) in the case of mammalian processed animal protein submitted to any of the processing methods 1 to 5 or 7 and, in the case of porcine blood, submitted to any of the processing methods 1 to 5 or 7 provided that in the case of method 7 a heat treatment throughout its substance at a minimum temperature of 80 °C has been applied;
- in the case of non-mammalian processed protein with the exclusion of fishmeal submitted to any
 of the processing methods 1 to 5 or 7 as referred to in Chapter III of Annex IV to Regulation (EU)
 No 142/2011;
- (j) in the case of fishmeal submitted to any of the processing methods 1 to 7 as referred to in Chapter III of Annex IV to Regulation (EU) No 142/2011 or to a method and parameters which ensure that the product complies with the microbiological standards for derived products set out in Chapter I of Annex X to Regulation (EU) No 142/2011;
- (k) in the case of rendered fat, including fish oils, submitted to any of the processing methods 1 to 5 or 7 (and method 6 in the case of fish oil) as referred to in Chapter III of Annex IV to Regulation (EU) No 142/2011 or produced in accordance with Chapter II of Section XII of Annex III to Regulation (EC) No 853/2004; rendered fats from ruminant animals must be purified in such a way that the maximum level of the remaining total insoluble impurities does not excess 0,15 % in weight;

COUNTRY Processed petfood other than canned petfood II. **Health information** II.a. Certificate reference No II.b. in the case of dicalcium phosphate produced by a process that (i) ensures that all Category 3 bone-material is finely crushed and degreased with hot water and treated with dilute hydrochloric acid (at a minimum concentration of 4 % and a pH of less than 1,5) over a period of at least two days; following the procedure referred to in (i), applies a treatment of the obtained phosphoric (ii) liquor with lime, resulting in a precipitate of dicalcium phosphate at pH 4 to 7; and (iii) finally, air dries the precipitate of dicalcium phosphate with inlet temperature of 65 °C to 325 °C and end temperature between 30 °C and 65 °C; (m) in the case of tricalcium phosphate produced by a process that ensures that all Category 3 bone-material is finely crushed and degreased in counter-flow with (i) hot water (bone chips less than 14 mm); (ii) continuous cooking with steam at 145 °C during 30 minutes at 4 bar; (iii) separation of the protein broth from the hydroxyapatite (tricalcium phosphate) by centrifugation; and granulation of the tricalcium phosphate after drying in a fluid bed with air at 200 °C; (iv) in the case of flavouring innards, produced according to a treatment method and parameters, which ensure that the product complies with the microbiological standards referred to in point II.4.] (2) or [was subject to a treatment such as drying or fermentation, which has been authorised by the competent authority;] [in the case of aquatic and terrestrial invertebrates other than species pathogenic to humans or (2) or animals, has been subject to a treatment which has been authorised by the competent authority and which ensures that the petfood poses no unacceptable risks to public and animal health;] 11.4. was analysed by a random sampling of at least five samples from each processed batch taken during or after storage at the processing plant and complies with the following standards (4): Salmonella: absence in 25g: n = 5, c = 0, m = 0, M = 0, n = 5, c = 2, m = 10, M = 300 in 1 gramme; Enterobacteriaceae: 11.5. has undergone all precautions to avoid contamination with pathogenic agents after treatment; II.6. was packed in new packaging, which, if the petfood is not dispatched in ready-to-sell packages on which it is clearly indicated that the content is destined for feeding to pets only, bear labels indicating "NOT FOR HUMAN CONSUMPTION"; $(^{2})$ [II.7. the petfood described above (2) either [is derived from other ruminants than bovine, ovine or caprine animals.] (2) or lis derived from bovine, ovine or caprine animals and does not contain and is not derived from: (2) either [bovine, ovine and caprine materials other than those derived from animals born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]] (2) or specified risk material as defined in point 1 of Annex V to Regulation (EC) [(a)

No 999/2001 of the European Parliament and of the Council (5);

COUNTRY Processed petfood other than canned petfood

II.	Health information		II.a. Certificate reference No II.b.						
		anii slai acc	mechanically separated meat obtained from bones of bovine, ovine or caprine animals, except from those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Commission Decision 2007/453/EC (⁶), in which there has been no indigenous BSE case,						
		anii ner the tho or r	imal by-product or derived product obtained from bovine, ovine or caprine imals which have been killed, after stunning, by laceration of the central rvous tissue by means of an elongated rod-shaped instrument introduced into e cranial cavity, or by means of gas injected into the cranial cavity, except for use animals that were born, continuously reared and slaughtered in a country region classified as posing a negligible BSE risk in accordance with Decision 07/453/EC.]]]						

Notes

Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is required to be filled in only if
 it is a certificate for a commodity to be transited through the European Union; it may be filled in if the certificate is for a
 commodity to be imported into the European Union.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for a transit commodity. Products intransit may only be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to be provided. In case of unloading and reloading, the consignor must inform the border inspection post of entry into the European Union.
- Box reference I.19: use the appropriate Harmonized System (HS) code under the following headings: 04.01; 04.02; 04.03; 04.04; 04.08, 05.04, 05.05, 05.06; 05.11, 15.01, 15.02, 15.03, 15.04, 23.01, 23.09; 28.35.25; 28.35.26; 35.01; 35.02; 35.03 or 35.04.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) must be given.
- Box reference I.25: technical use: any use other than feeding of farmed animals, other than fur animals, and the
 production or manufacturing of pet food.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: Species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or Suidae, Pesca, Mollusca, Crustacea, Invertebrates other than Mollusca and crustacea.

Part II:

- (1a) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (²) Delete as appropriate.
- (2a) OJ L 139, 30.4.2004, p. 55.
- (2b) OJ L 125, 23.5.1996, p. 3.
- (3) OJ L 175, 10.7.2010, p. 1.

COUNTRY Processed petfood other than canned pet											
	Health information	II.a.	Certificate reference No		II.b.						
Whe	re:										
n =	number of samples to be tested;										
m =	threshold value for the number of bacteria; the result is considered satisfactory if the number of bacteria in all samples does not exceed m;										
M = maximum value for the number of bacteria; the result is considered unsatisfactory if the number of bacteria in one or more samples is M or more; and											
c =				m and M, the	sample still being considered						
OJ L	147, 31.5.2001, p. 1.										
OJ L	172, 30.6.2007, p. 84.										
The	signature and the stamp must be in a di	fferent	colour to that of the printing	ng.							
cial vet	erinarian/Official inspector										
Name (in capital letters): Qualification and title:											
Date				Signature:							
Stam	p:										
	When n = m = M = OJ L OJ L The se and r Cial vet	Health information Where: n = number of samples to be tested; m = threshold value for the number of backamples does not exceed m; M = maximum value for the number of backamore samples is M or more; and c = number of samples the bacterial count of the OJ L 147, 31.5.2001, p. 1. OJ L 172, 30.6.2007, p. 84. The signature and the stamp must be in a diffusion of the person responsible for the consistent must accompany the consignment until interpretation.	Health information Where: n = number of samples to be tested; m = threshold value for the number of bacteria; samples does not exceed m; M = maximum value for the number of bacteria; the more samples is M or more; and c = number of samples the bacterial count of acceptable if the bacterial count of the other; OJ L 147, 31.5.2001, p. 1. OJ L 172, 30.6.2007, p. 84. The signature and the stamp must be in a different and must accompany the consignment until it reactions are consignment and must accompany the consignment until it reactions. Date:	Health information Where: n = number of samples to be tested; m = threshold value for the number of bacteria; the result is considered samples does not exceed m; M = maximum value for the number of bacteria; the result is considered unimore samples is M or more; and c = number of samples the bacterial count of which may be between acceptable if the bacterial count of the other samples is m or less. OJ L 147, 31.5.2001, p. 1. OJ L 172, 30.6.2007, p. 84. The signature and the stamp must be in a different colour to that of the printing that the person responsible for the consignment in the European Union: and must accompany the consignment until it reaches the border inspection cital veterinarian/Official inspector Name (in capital letters):	Health information II.a. Certificate reference No Where: n = number of samples to be tested; m = threshold value for the number of bacteria; the result is considered satisfactory if samples does not exceed m; M = maximum value for the number of bacteria; the result is considered unsatisfactory if the more samples is M or more; and c = number of samples the bacterial count of which may be between m and M, the acceptable if the bacterial count of the other samples is m or less. OJ L 147, 31.5.2001, p. 1. OJ L 172, 30.6.2007, p. 84. The signature and the stamp must be in a different colour to that of the printing. Note for the person responsible for the consignment in the European Union: This certificate and must accompany the consignment until it reaches the border inspection post of entry in the consignment in						

Veterinary certificate to EU

CHAPTER 3(C)

Health certificate

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

	l.1.	Consignor					1.2.	Certificate refere	ence No	I.2.a.		
		Name					1.3.	Central compete	ent authority			
		Address					1.4.	Local competen	t authority			
		- .										
		Tel.										
	1.5.	Consignee					1.6.	I.6. Person responsible for the load in EU				
nen		Name						Name				
signr		Address						Address				
cons		Postcode						Postcode				
;hed		Tel.						Tel.				
Part I : Details of dispatched consignment	1.7.	Country of origin	ISO code	1.8.	Region of origin	Code	1.9.	Country of destination	ISO code	I.10. Region of destination	Code	
of di		or origin			Origin			destination				
tails	I.11.	Place of or	igin				1.12.	Place of destina	ıtion		<u> </u>	
: De												
art		Name		Appro	val number					Custom warehouse		
Δ.		Address						Name		Approval number		
		Name		Appro	val number			Address				
		Address										
		Name		Appro	val number			Postcode				
		Address										
	I.13.	Place of lo	ading				I.14.	Date of departur	re			
	I.15.	Means of t	ransport				I.16.	Entry BIP in EU				
		Aeroplane	☐ Ship		Railway wa	agon 🗖						
		Road vehic	cle 🔲 Othe	er 🔲			I.17.					
		Identification	on									
		Documenta	ation referen	ces					ı			
	I.18.	Description	n of commod	ity					I.19. Comr	modity code (HS code)		
										I.20. Quantity		
	I.21.	Temperatu	re of product	t						I.22. Number of page	ckages	
		Ambient 🗆]		Chilled D]		Frozen [
	1.23.	Seal/Conta	ainer No							I.24. Type of packa	ging	



1.25.	Commodities certified for:						
	Petfood \square		Technical use □				
1.26.	For transit through EU to the	hird country	I.27. For import or admission into EU				
	Third country	ISO code					
1.28.	Identification of the commo	odities					
		Approval number	of establishments				
	Species (Scientific name)	Manufacturing plant	Net weight	Batch number			

	COUNT	RY				_												Dogo	hews
	II.	Health info	rmati	on		II.a	a. C	ertific	ate re	feren	ce No)			.b.				
	_	the Europe Regulation	an Pa (EU)	d official vet rliament and No 142/2011 e dogchews	ıncil (n part	^{1a}), a icular	nd in	partic	cular ,	Articl	e 10 c	of that	Regul	ation	, and	Comm	ission		
_	II.1.	have been	ave been prepared exclusively with the following animal by-products:																
Part II: Certification		(²) either	[-	carcases a killed, and intended fo	which	are fi	t for h	numai	n con	sump	tion in	n acc	ordand						
Part II: C		(²) and/or	[-	carcases a slaughterho mortem ins consumption	use a pectio	nd w n or	ere co	onside es an	ered t	it for follo	slaug wing	hter parts	for hur	nan c	onsum	ption	follow	ing an	ante-
	_			cor		ion ii	n acc	ordan	ice wi	ith Ur	nion le	egisla	which ition, b		-				numan gns of
				(ii) hea	ids of p	ooulti	γ;												
													ting th						ng the
				(iv) pig	bristle	s;													
				(v) fea	(v) feathers;]														
		(²) and/or	[-	humans or having bee	blood of animals which did not shourned from a naving been considered fit for slanspection in accordance with Union						hat hat ha	ave b	een sl	aught	ered ir	n a sla	aughte	rhouse	e after
		(²) and/or	[-	animal by- including d			-												
		(²) and/or	[-	aquatic ani of disease								ct se	a mam	mals,	which	did n	ot sho	ow any	signs
		(²) and/or	[-	animal by-p products fo						ıls ori	ginatir	ng fro	om plai	nts or	establi	ishme	ents m	anufac	turing
		(²) and/or	[-	material fro Council Di Article 35(a	ective	96/2	2/EC	$(^{2a}),$	the ir	nport	of the	e ma							-
	II.2.	have been	subjed	cted															
		(²) either		he case of d estroy patho														ent suf	ficient
		(²) and/or		the case of on fish, to a he												d skii	ns of	ungula	tes or
	II.3.			oy random s ocessing plar										cesse	d batc	h tak	en du	ring oi	r after
		Salmonella	:	а	bsence	e in 2	5g: n	= 5, 0	c = 0,	m = 0), M =	€ 0,							
		Enterobacte	eriace	ae: n	= 5, c	= 2,	m = 1	0, M	= 300	in 1 (gramr	ne;							



COUNTRY **Dogchews** II. **Health information** II.a. Certificate reference No II.b. have undergone all precautions to avoid contamination with pathogenic agents after treatment; 11.4. 11.5. were packed in new packaging; (2) [II.6. the dogchews described above (2) either [is derived from other ruminants than bovine, ovine or caprine animals.]] (2) or [is derived from bovine, ovine or caprine animals and does not contain and is not derived from: (2) either [bovine, ovine and caprine materials other than those derived from animals born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]] (2) or specified risk material as defined in point 1 of Annex V to Regulation (EC) [(a) No 999/2001 of the European Parliament and of the Council (4); (b) mechanically separated meat obtained from bones of bovine, ovine or caprine animals, except from those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Commission Decision 2007/453/EC (5), in which there has been no indigenous BSE case, animal by-product or derived product obtained from bovine, ovine or caprine (c) animals which have been killed, after stunning, by laceration of the central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, or by means of gas injected into the cranial cavity, except for those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]]

Notes

Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a
 certificate for transit commodity; it may be filled in if the certificate is for a commodity to be imported into the European
 Union.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for a transit commodity. Products
 in transit may only be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship);
 the information is to be provided in the event of unloading and reloading in the European Union.
- Box reference I.19: 05.11, 23.09, 41.01 or 42.05.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) must be given.
- Box reference I.25: technical use: any use other than feeding of farmed animals, other than fur animals, and the
 production or manufacturing of pet food.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: Species: select from the following: Aves, Ruminantia, Suidae, Mammalia Other Than Ruminantia or Suidae, Pesca, Mollusca, Crustacea, Invertebrates Other Than Mollusca And Crustacea.

Part II:

- (1a) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.

COL	INTRY				Dogchews						
II.	Health information	II.a.	Certificate reference No		II.b.						
(²)	Delete as appropriate.										
(^{2a})	OJ L 125, 23.5.1996, p. 3.										
(3)	Where:										
_	n = number of samples to be tested;										
_	m = threshold value for the number of bacteria; the result is considered satisfactory if the number of bacteria in all samples does not exceed m;										
_	M = maximum value for the number of bac more samples is M or more; and	xteria;	the result is considered uns	atisfactory if	the number of bacteria in one or						
_	c = number of samples the bacterial co acceptable if the bacterial count of the			n and M, th	e sample still being considered						
(4)	OJ L 147, 31.5.2001, p. 1.										
(⁵)	OJ L 172, 30.6.2007, p. 84.										
_	The signature and the stamp must be in a d	ifferer	nt colour to that of the printin	ıg.							
_	Note for the person responsible for the consand must accompany the consignment until										
Offic	ial veterinarian/Official inspector										
2											
	Name (in capital letters):			Qualification	n and title:						
	Date:			Signature:							
	Stamp:										

I.24. Type of packaging

I.23. Seal/Container No

CHAPTER 3(D)

Health certificate

For raw petfood for direct sale or animal by-products to be fed to fur animals, intended for dispatch to or for transit through (2) the European Union

COL	JNTRY	/ :								Vete	rinary certificat	te to EU
	l.1.	Consignor					1.2.	Certificate referen	nce No	1.2	2.a	
		Name					1.3.	Central competer	nt authority			
		Address					1.4.	Local competent	authority			
		Tel.										
	1.5.	Consignee					I.6. Person responsible for the load in EU					
ent		Name						Name				
ignm		Address						Address				
cons		Postcode						Postcode				
)ed		Tel.						Tel.				
Part I : Details of dispatched consignment	l.7.	Country of origin	ISO code	I.8.	Region of origin	Code	1.9.	Country of destination	ISO code	I.10.	Region of destination	Code
<u>s</u> of												
etai	l.11.	I.11. Place of origin						Place of destinati	on			
												_
Part		Name Approval number								Cust	om warehouse	
		Address						Name		Аррі	roval number	
		Name Approval number						Address				
		Address										
		Name		Appro	val number			Postcode				
		Address										
	I.13.	Place of lo	ading				l.14.	Date of departure	!			
	I.15.	Means of t	ransport				I.16.	Entry BIP in EU				
		Aeroplane	•		Railway wa	agon 🗖						
		Road vehic	cle 🔲 Othe	er 🗆			1.17.					
	Identification											
	Documentation references											
	I.18. Description of commodity								I.19. Comn	nodity o	code (HS code)	
								L		1.20	Quantity	
	I.21.	I.21. Temperature of product								1.22	Number of pag	kages
		Ambient □]		Chilled []		Frozen 🗆				

1.25.	Commodities certifi	ied for:					
	Petfood			Technical use □			
1.26.	For transit through	EU to third country		I.27. For imp	port or admission into EU		
	Third country	ISO code					
1.28.	Identification of the		oval number	of establishme	nts		
		Λρριν	oval Hambel	or catabilatine	110		
(8	Species Scientific name)	Nature of commodity	Manufactu	uring plant	Net weight	Batch number	

Part II: Certification

COUNTRY

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

II. Health information II.a. Certificate reference No II.b.

I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council (¹a) and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011 (¹b), and in particular Chapter II of Annex XIII and Chapter II of Annex XIV thereto, and certify that the raw petfood or animal by-products described above:

- II.1. consist of animal by-products that satisfy the health requirements below;
- II.2. consist of animal by-products:
 - (a) derived from meat which satisfies the relevant animal and public health requirements laid down in:

 - and/or Commission Regulation (EC) No 798/2008 (4), and provided that the animals from which the
 meat is derived come from the third countries, territories or parts thereof(ISO code in the
 case of a country, or codes in the case of territories or parts thereof) as listed in that Regulation which
 has been free from Newcastle disease and avian influenza for the last 12 months;
 - and/or Commission Regulation (EC) No 119/2009 (5), and provided that the animals from which the meat is derived come from the third countries, territories or parts thereof (ISO code in the case of a country, or codes in the case of territories or parts thereof) as listed in that Regulation which has been free from foot and mouth disease, rinderpest, classical swine fever, African swine fever, swine vesicular disease, Newcastle disease and avian influenza for the preceding 12 months and where no vaccination has taken place during that time (only where relevant for the susceptible species);
 - (b) derived from animals that, at the slaughterhouse, have passed the ante-mortem health inspection during the period of 24 hours before the time of slaughter and have shown no evidence of the diseases referred in the Regulations referred to in point (a) for which the animals are susceptible; and
 - (c) derived from animals that have been handled in the slaughterhouse before and at the time of slaughter or killing in accordance with the relevant provisions of Union legislation and have met requirements at least equivalent to those laid down in Chapters II and III of Council Regulation (EC) No 1099/2009 (⁶); or
 - (d) in the case of feed for fur animals, are derived from aquatic animals which satisfy the relevant animal and public health requirements laid down in Commission Decision 2006/766/EC (⁷), and come from countries or territories thereof(ISO code of the country) as listed in Annex II to that Decision;
- II.3.1. consist only of the following animal by-products:
 - (a) carcases and parts of animals slaughtered or, in the case of game, bodies or parts of animals killed which
 were deemed fit for human consumption in accordance with Union legislation until irreversibly declared as
 animal by-products for commercial reasons;
 - (b) parts of slaughtered animals, which are rejected as unfit for human consumption but are not affected by any signs of diseases communicable to humans or animals and derived from carcases that are fit for human consumption in accordance with Union legislation;
- II.3.2. in the case of feed for fur animals in addition to II.3.1. consist also of the following animal by-products:
 - (²) either [- animal by-products from poultry and lagomorphs slaughtered on the farm as referred to in Article 1(3)(d) of Regulation (EC) No 853/2004 of the European Parliament and of the Council (²a), which did not show any signs of disease communicable to humans or animals;]
 - (2) and/or [- blood of animals which did not show any signs of disease communicable through blood to humans or animals, obtained from animals that have been slaughtered in a slaughterhouse after having been considered fit for slaughter for human consumption following an ante-mortem inspection in accordance with Union legislation;]
 - (2) and/or [- animal by-products arising from the production of products intended for human consumption, including degreased bone, greaves and centrifuge or separator sludge from milk processing;]

COUNTRY

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

II.	Health info	rmati	on		II.a. Certificate reference No	II.b.								
	(²) and/or	[-	intended t	lucts of animal origin, or foodstuffs containing products of animal origin, which are no longer nded for human consumption for commercial reasons or due to problems of manufacturing or caging defects or other defects from which no risk to public or animal health arises;]										
	(²) and/or	[-	derived p	roducts, which a	of animal origin, or feedingstuffs contain are no longer intended for feeding for com g or packaging defects or other defects fron	mercial reasons or due to								
	(²) and/or	[-			thers, hair, horns, hoof cuts and raw milk o of any disease communicable through th	•								
	(²) and/or	[-		atic animals, and parts of such animals, except sea mammals, which did not show any signs seases communicable to humans or animals;]										
	(²) and/or	[-	-	nal by-products from aquatic animals originating from plants or establishments manufacturing ucts for human consumption;]										
	(²) and/or	[-			iginating from animals which did not sh at material to humans or animals:	ow any signs of disease								
			(i) sł	nells from shellfis	h with soft tissue or flesh;									
			(ii) th	e following origir	nating from terrestrial animals:									
			_	hatchery b	y-products,									
			_	eggs,										
			_	egg by-pro	ducts, including egg shells,									
			(iii) da	ay-old chicks kille	ed for commercial reasons;]									
	(²) and/or	[-		r-products from r animals;]	aquatic or terrestrial invertebrates other t	nan species pathogenic to								
	(²) and/or	[-	Category	1 material as ref	of of the zoological orders of Rodentia erred to in Article 8(a)(iii), (iv) and (v) of Reg s referred to in Article 9(a) to (g) of that Reg	gulation (EC) No 1069/2009								
II.4.					ntact with other material which does not con and it has been handled so as to avoid con									
II.5.	CONSUMF CONSUMF preventing NOT FOR	TION TION any le HUMA	or 'ANIM' and then eakage and AN CONSU	d in final packaging which bear labels indicating 'RAW PET FOOD — NOT FOR HUMAN or 'ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS — NOT FOR HUMAN and then placed in leak-proof and officially sealed boxes/containers or in new packaging kage and officially sealed boxes/containers which bear labels indicating 'RAW PET FOOD — I CONSUMPTION' or 'ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS — NOT FOR PTION', and the name and the address of the establishment of destination;										
II.6.	in the case	of rav	v petfood:											
				d stored in a pla lation (EC) No 10	nt approved and supervised by the compet 069/2009 and	ent authority in accordance								
				dom sampling o	f at least five samples from each batch tang standards (8):	ken during storage (before								

COUNTRY

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

					fed to fur animals
II.	Health informati	on		II.a. Certificate reference No	II.b.
	Salmonella:	abse	ence in 25	g: n=5, c=0, m=0, M=0	
	Enterobacteriacea	ae: n=5,	c=2, m=10	0, M=5000 in 1 gram;	
(²) [II.7.	[the petfood or ar products of rumin		ts to be fe	d to fur animals described above contains o	r is derived from animal-by
	(²) either			try or region, which is classified as posing n 2007/453/EC, and in which there has beer	
	(²) or	Decision 2007 product or der the feeding or	7/453/EC ii ived produ f ruminant OIE Terre	y or region classified as posing a negligible En which there has been an indigenous BSE ict were derived from animals born after the cost with meat-and-bone meal and greaves distrial Animal Health Code, has been effective	case, and the animal by- date from which the ban on erived from ruminants, as
	(²) either	[is derived fror	n other rur	ninants than bovine, ovine or caprine animals	.]]
	(²) or	[is derived fror	n bovine, d	ovine or caprine animals and does not contain	and is not derived from:
		(²) either	continuo	ovine and caprine materials other than those usly reared and slaughtered in a country or resease. BSE risk in accordance with Decision 2007/	egion classified as posing a
		(²) or		pecified risk material as defined in point 1 of A o 999/2001 of the European Parliament and o	
			ca aı B	echanically separated meat obtained from aprine animals, except from animals that were nd slaughtered in a country or region classi SE risk in accordance with Commission De hich there has been no indigenous BSE case	e born, continuously reared fied as posing a negligible cision 2007/453/EC (10), in
			ca th in in ca	nimal by-product or derived product obtain aprine animals which have been killed, after se central nervous tissue by means of strument introduced into the cranial cavity, of the cranial cavity, except for those ontinuously reared and slaughtered in a coupsing a negligible BSE risk in accordance with	stunning, by laceration of an elongated rod-shaped r by means of gas injected animals that were born, ntry or region classified as
Notes					

Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is required to be filled in only if
 it is a certificate for a commodity to be transited through the European Union; it may be filled in if the certificate is for a
 commodity to be imported into the European Union.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. Products in transit may only be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to be provided. In case of unloading and reloading, the consignor must inform the border inspection post of entry into the European Union.
- Box I.19: use the appropriate Harmonized System (HS) code under the following heading: 04.08; 05.06; 05.08; 05.11, 23.01 or 23.09.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) must be given.
- Box reference I.25: technical use: any use other than feeding of farmed animals, other than fur animals, and the
 production or manufacturing of pet food.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.

COUNTRY

Raw petfood for direct sale or animal by- products to be

				fed to fur animals						
II.	Health information	II.a.	Certificate reference No	II.b.						
_	Box reference I.28:									
	Nature of commodity: select raw petfood or animal b	y-pro	oduct.							
	In the case of raw material for the manufacture of ra	w pet	food indicate the scientific name of t	he species.						
	In case of raw material for manufacture of feed f Mammalia other than Ruminantia or Suidae, Pe Crustacea.									
Part	II:									
(^{1a})	OJ L 300, 14.11.2009, p. 1.									
(^{1b})	OJ L 54, 26.2.2011, p. 1.									
(²)	Delete as appropriate.									
(^{2a})	OJ L 139, 30.4.2004, p. 55.									
(³)	OJ L 73, 20.3.2010, p. 1.									
(4)	OJ L 226, 23.8.2008, p. 1.									
(⁵)	OJ L 39, 10.2.2009, p. 12.									
(⁶)	OJ L 303, 18.11.2009, p. 1.									
(⁷)	OJ L 320, 18.11.2006, p. 53.									
(8)	Where:									
	n = number of samples to be tested;									
	m = threshold value for the number of bacteria; samples does not exceed m;	the r	esult is considered satisfactory if th	e number of bacteria in all						
	M = maximum value for the number of bacteria; the or more samples is M or more; and	ne res	sult is considered unsatisfactory if the	e number of bacteria in one						
	c = number of samples the bacterial count of w acceptable if the bacterial count of the other s			mple still being considered						
(⁹)	OJ L 147, 31.5.2001, p. 1.									
(10)	OJ L 172, 30.6.2007, p. 84.									
_	The signature and the stamp must be in a different of	colour	to that of the printing.							
_	Note for the person responsible for the consignment and must accompany the consignment until it reach									
Offic	cial veterinarian/Official inspector									
	Name (in capital letters):		Qualification and	d title:						
	Date:		Signature:							
	Stamp:									

I.24. Type of packaging

I.23. Seal/Container No

CHAPTER 3(E)

Health certificate

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

COL	JNTRY	′ :				Veterinary certificat	te to EU			
	l.1.	Consignor	1.2.	Certificate referen	ice No	I.2.a.				
		Name	1.3.	Central competen	t authority					
		Address	1.4.	Local competent a	authority					
				·						
		Tel.								
	1.5.	Consignee	1.6.	I.6. Person responsible for the load in EU						
ent		Name		Name						
gnm		Address	Address							
isuc										
oo pa		Postcode		Postcode						
tche		Tel.		Tel.						
ispa	1.7.	Country ISO code I.8. Region of Code of origin origin	1.9.	Country of destination	ISO code	I.10. Region of destination	Code			
of d										
Part I : Details of dispatched consignment	l.11.	Place of origin	1.12.	Place of destination	⊥ on					
: Del		•								
art I		Name Approval number				Custom warehouse				
ď		Address		Name		Approval number				
		Name Approval number		Address						
		Address								
		Name Approval number		Postcode						
		Address								
	l.13.	Place of loading	I.14.	Date of departure						
	I.15.	Means of transport	I.16.	Entry BIP in EU						
		·		•						
		Aeroplane ☐ Ship ☐ Railway wagon ☐								
		Road vehicle Other	1.17.							
		Identification		_						
		Documentation references								
	I.18.	Description of commodity			I.19. Comm	nodity code (HS code)				
						I.20. Quantity				
	I.21.	Temperature of product				I.22. Number of page	ckages			
		Ambient ☐ Chilled ☐		Frozen 🗖						

1.25.	Commodities certifi	ed for:				
	Petfood			Technical use □		
1.26.	For transit through	EU to third country		I.27. For imp	ort or admission into EU	
	Third country	ISO code				
1.28.	Identification of the		aval numbar	of octoblishme	nto.	
		Аррго	ovai number	of establishmer	its	
(\$	Species Scientific name)	Nature of commodity	Manufactu	uring plant	Net weight	Batch number

Flavouring innards for use in the manufacture of petfood

	II.	Health information		on	II.a. Certificate reference No	II.b.					
		the Europe Regulation	an Pa (EU) N	rliament and of the (No 142/2011 (^{1b}), and	declare that I have read and understood Regulation (EC) No 1069/2009 of Council (1a), and in particular Article 8 and 10 thereof, and Commission in particular Chapter III of Annex XIII and Chapter II of Annex XIV thereto, products described above:						
_	II.1.	consist of a	nimal I	oy-products that satisf	y the animal health requirements below;						
catior	II.2.	have been ¡	prepar	ed and include the foll	owing animal by-products which are exclusive	ly:					
Part II: Certification		(²) either	[-	killed, and which are	of animals slaughtered or, in the case of gan fit for human consumption in accordance with consumption for commercial reasons;]						
Par		(²) and/or [- carcases and the following parts originating either from animals that have been sl slaughterhouse and were considered fit for slaughter for human consumption following mortem inspection or bodies and the following parts of animals from game kill consumption in accordance with Union legislation:									
				consumption	bodies and parts of animals which are in accordance with Union legislation, but wh municable to humans or animals;						
				(ii) heads of pou	Itry;						
				• ,	ins, including trimmings and splitting thereof, horns and feet, including the d the carpus and metacarpus bones, tarsus and metatarsus bones;						
				(iv) pig bristles;							
				(v) feathers;]							
		(²) and/or	[-	humans or animals, of having been consider	rhich did not show any signs of disease communicable through blood to obtained from animals that have been slaughtered in a slaughterhouse after dered fit for slaughter for human consumption following an ante-mortem ance with Union legislation;]						
		(²) and/or	[-		arising from the production of products inter bone, greaves and centrifuge or separator slu						
		(²) and/or	[-	intended for human of	origin, or foodstuffs containing products of animal origin, which are no longer consumption for commercial reasons or due to problems of manufacturing or r other defects from which no risk to public or animal health arise;]						
		(²) and/or	[-	derived products, wh	stuffs of animal origin, or feedingstuffs con nich are no longer intended for feeding for o cturing or packaging defects or other defects	commercial reasons or due to					
		(²) and/or	[-		l, feathers, hair, horns, hoof cuts and raw mil signs of any disease communicable through						
		(²) and/or	[-		parts of such animals, except sea mammals, icable to humans or animals;]	which did not show any signs					
		(²) and/or	[-	animal by-products fr products for human c	rom aquatic animals originating from plants or consumption;]	establishments manufacturing					
		(²) and/or	[-	_	al originating from animals which did not gh that material to humans or animals:	show any signs of disease					
				(i) shells from sl	hellfish with soft tissue or flesh;						

Flavouring innards for use in the manufacture of petfood

							of petfood
II.	Health info	rmation		II.a.	Certificate reference No		II.b.
		(ii)	the follo	owing origina	ating from terrestrial anima	ıls:	
			_	hatchery by-	-products,		
			_	eggs,			
			_	egg by-prod	ucts, including egg shells;		
		(iii)	day-old	d chicks killed	d for commercial reasons;]		
	(²) and/or		al by-proc ans or anir		quatic or terrestrial inver	tebrates othe	er than species pathogenic to
	(²) and/or	Cate	gory 1 ma	terial as refe		v) and (v) of I	itia and Lagomorpha, except Regulation (EC) No 1069/2009 Regulation;]
	(²) and/or	Cour	ncil Directi	ve 96/22/EC			ances which are prohibited by permitted in accordance with
II.3.		subjected to pathogenic		ig in accorda	nce with Chapter III of An	nex XIII to Re	egulation (EU) No 142/2011, in
II.4.					east five samples from eavith the following standard		ed batch taken during or after
	Salmonella:	:	abse	nce in 25g: n	a = 5, $c = 0$, $m = 0$, $M = 0$,		
	Enterobacte	eriaceae:	n = 5	, c = 2, m = 1	10, M = 300 in 1 gramme;		
II.5.	the end pro	duct was:					
	(²) either	[packed ir	new or st	erilised bags	,]		
	(²) or				ers or other means of traproved by the competent a	•	were thoroughly cleaned and re use,]
	and which b	oear labels in	ndicating 'N	NOT FOR HI	JMAN CONSUMPTION';		
II.6.	the end pro	duct was sto	red in enc	closed storag	e;		
II.7.	the product	has underg	one all pre	cautions to a	void contamination with p	athogenic age	ents after treatment;
(²) [II.8.	the flavouring	ng innards p	roducts de	escribed abov	/e		
	(²) either	[is derived	I from othe	er ruminants t	than bovine, ovine or capr	ine animals.]]	
	(²) or	[is derived	I from bovi	ine, ovine or	caprine animals and does	not contain a	and is not derived from:
		(²) either	continu	iously reared		country or r	derived from animals born, region classified as posing a EC.]]
		(²) or	[(a)		sk material as defined ir 1 of the European Parliam		Annex V to Regulation (EC) e Council (4);
			(b)	animals, ex slaughtered accordance	cept from those animals in a country or region cl	that were b assified as p	es of bovine, ovine or caprine form, continuously reared and osing a negligible BSE risk in EC (5), in which there has been
			(c)	animals wh nervous tiss the cranial of those animal	ich have been killed, af sue by means of an elong cavity, or by means of ga als that were born, continu assified as posing a negli	ter stunning, ated rod-shap s injected into uously reared	rom bovine, ovine or caprine by laceration of the central ped instrument introduced into o the cranial cavity, except for d and slaughtered in a country k in accordance with Decision

Official veterinarian/Official inspector Name (in capital letters):

Date:

Stamp:

COUNTRY

Flavouring innards for use in the manufacture

Qualification and title:

Signature:

		<u> </u>	of petfood							
II.	Health information	II.a. Certificate reference No	II.b.							
Note	es									
Part	l:									
_		e consignment in the European Union: this box sited through the European Union; it may be fi Union.								
_	Box reference I.12: Place of destination: this transit may only be stored in free zones, free	box is to be filled in only if it is a certificate for warehouses and custom warehouses.	transit commodity. Products in							
_		lway wagons or container and lorries), flight nu unloading and reloading in the European Union								
_	Box reference I.19: use the appropriate HS of	code: 05.04; 05.06, 05.11 or 23.09 .								
_	Box reference I.23: for bulk containers, the containers	ontainer number and the seal number (if applic	able) should be given.							
_	Box reference I.25: technical use: any use production or manufacturing of pet food.	e other than feeding of farmed animals, oth	er than fur animals, and the							
_	Box reference I.26 and I.27: fill in according t	to whether it is a transit or an import certificate.								
_	Box reference I.28:									
	 species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or Suidae, Pesca, Mollusca, Crustacea, Invertebrates other than Mollusca and crustacea 									
	 define the innard product. 									
Part	II:									
(^{1a})	OJ L 300, 14.11.2009, p. 1.									
(1b)	OJ L 54, 26.2.2011, p. 1.									
(²)	Delete as appropriate.									
(^{2a})	OJ L 125, 23.5.1996, p. 3.									
(³)	Where:									
	n = number of samples to be tested;									
	m = threshold value for the number of ba samples does not exceed m;	cteria; the result is considered satisfactory if	the number of bacteria in all							
	M = maximum value for the number of bac or more samples is M or more; and	cteria; the result is considered unsatisfactory if	the number of bacteria in one							
	c = number of samples the bacterial cou acceptable if the bacterial count of the	nt of which may be between m and M, the other samples is m or less.	sample still being considered							
(4)	OJ L 147, 31.5.2001, p. 1.									
(⁵)	OJ L 172, 30.6.2007, p. 84.									
_	The signature and the stamp must be in a diff	fferent colour to that of the printing.								
_	Note for the person responsible for the consi and must accompany the consignment until i	gnment in the European Union: This certificate it reaches the border inspection post.	is only for veterinary purposes							

Veterinary certificate to EU

CHAPTER 3(F)

Health certificate

For animal by-products (3) for the manufacture of petfood, intended for dispatch to or for transit through (2) the European Union

	l.1.	Consignor	1.2.	Certificate reference N	0	1.2.a.	
		Name	1.3.	Central competent auth	hority		
		Address	1.4.	Local competent autho	ority		
		Tel.					
	1.5.	Consignee	1.6.	Person responsible for	the load in	n EU	
nent		Name		Name			
ignr		Address		Address			
Part I : Details of dispatched consignment		Postcode		Postcode			
ped (Tel.		Tel.			
atch	1.7.	Country ISO code I.8. Region of Code	1.9.	Country of ISC) I	10. Region of	Code
disp	1.7.	of origin	1.5.	destination cod		destination	Code
s of							
etail	I.11.	Place of origin	I.12.	Place of destination			
<u> </u>							
² art		Name Approval number			C	Custom warehouse	
		Address		Name	A	Approval number	
		Name Approval number		Address			
		Address					
		Name Approval number		Postcode			
		Address					
	I.13. 	Place of loading	1.14.	Date of departure			
	I.15.	Means of transport	I.16.	Entry BIP in EU			
		Aeroplane Ship Railway wagon Railway					
		Road vehicle Other	I.17.				
		Identification					
		Documentation references					
	l.18.	Description of commodity		I.19.	Commod	lity code (HS code)	
					I	.20. Quantity	
	I.21.	Temperature of product			ı	.22. Number of page	ckages
		Ambient ☐ Chilled ☐		Frozen 🗖			
	123	Seal/Container No				24 Type of packa	aina



1.25.	Commodities certified for:										
	Manufacture of pe	etfood 🗖	Further pro	ocess 🗆	Technical use □						
1.26.	For transit through	h EU to third countr	у 🗆	I.27. For import or	admission into EU						
	Third country	ISO co	ode								
1.28.	Identification of th	e commodities									
			Approval number	of establishments							
(Sci	Species entific name)	Nature of commodity	Manufacturing plant	Number of packages	Net weight	Batch number					

Animal by-products for the manufacture of petfood

II. **Health information** II.a. Certificate reference No II b I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council (1a) and Commission Regulation (EU) No 142/2011 (1b), and in particular Chapter II of Annex XIV thereto, and certify that the animal by-products described above: consist of animal by-products that satisfy the animal health requirements below; II.1.1. II.1.2. Part II: Certification [(a) that have remained in this territory since birth or for a period of at least three months preceding (2) either the date of slaughter or production;] (2) or [(b) killed in the wild in this territory (1d);] (2) or [(c) derived from rodents, lagomorphs, aquatic animals or terrestrial or aquatic invertebrates;] II.1.3. have been obtained from or produced by animals: (2) either [(a) coming from holdings:

- (i) where, for the following diseases for which the animals are susceptible, there has been no case/outbreak of rinderpest, swine vesicular disease, Newcastle disease or highly pathogenic avian influenza during the period of the preceding 30 days, nor of classical or African swine fever during the period of the preceding 40 days; nor in the holdings situated in their vicinity within a 10 km radius, during the period of the preceding 30 days; and
- (ii) where there has been no case/outbreak of foot-and-mouth disease during the period of the preceding 60 days, nor in the holdings situated in their vicinity within a 25 km radius, during the period of the preceding 30 days; and
- (b) which:
 - (i) were not killed to eradicate any epizootic disease;
 - (ii) have remained in their holdings of origin for a period of at least 40 days before the date of departure and which have been transported directly to the slaughterhouse without any contact with other animals which did not comply with the same health conditions;
 - (iii) at the slaughterhouse, have passed the ante-mortem health inspection during the period of 24 hours preceding the time of slaughter and have shown no evidence of the diseases referred to above for which the animals are susceptible; and
 - (iv) have been handled in the slaughterhouse before and at the time of slaughter or killing in accordance with the relevant provisions of Union legislation and have met requirements at least equivalent to those laid down in Chapters II and III of Council Regulation (EC) No 1099/2009 (4)]
- (2) or [(a) captured and killed in the wild in an area:
 - (i) in which within a 25 km radius there has been no case/outbreak of any of the following diseases for which the animals are susceptible: foot-and-mouth disease, rinderpest, Newcastle disease or highly pathogenic avian influenza during the period of the preceding 30 days, nor of classical or African swine fever during the period of the preceding 40 days; and
 - (ii) situated at a distance of at least 20 km from any country or part of the territory of a country not authorised for export to the European Union of poultry material during the preceding 30 days or of porcine material during the preceding 40 days; and
 - (b) which after killing were transported within a period of 12 hours following the killing for chilling either to a collection centre and immediately afterwards to a game handling establishment, or directly to a game handling establishment;]

Animal by-products for the manufacture of petfood

						of pet	τοοα			
II.	Health inform	ation	II.a.	Certificate	reference No	II.b.				
II.1.4.	have been obtained in an establishment around which, within a radius of 10 km, there has been no case/outbreak of the diseases referred to in point II.1.3 for which the animals are susceptible during the period of the preceding 30 days or, in the event of a case of disease, the preparation of raw material 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;									
II.1.5.						terial that does not comply with nation with pathogenic agents;	n the			
II.1.6.	have been packed in new packaging preventing any leakage and in officially sealed containers bearing the label indicating 'RAW MATERIAL ONLY FOR THE MANUFACTURE OF PET FOOD' and the name and address of the establishment of destination in the European Union;									
II.1.7.	consist only of the following animal by-products:									
	(²) either [-	er [- carcases and parts of animals slaughtered or, in the case of game, bodies or parts of animals killed which were deemed fit for human consumption in accordance with Union legislation until irreversibly declared as animal by-products for commercial reasons;]								
	(²) and/or [-	d/or [- carcases and the following parts originating either from animals that have been slaughtered in a slaughterhouse and were considered fit for slaughter for human consumption following an antemortem inspection or bodies and the following parts of animals from game killed for human consumption in accordance with Union legislation:								
		 carcases or bodies and parts of animals which are rejected as unfit for human consumption in accordance with Union legislation, but which did not show any signs of disease communicable to humans or animals; 								
		(ii) heads of po	oultry;							
						thereof, horns and feet, including tarsus and metatarsus bones;	g the			
		(iv) pig bristles	y;							
		(v) feathers;]								
	(²) and/or [-					cts intended for human consump rator sludge from milk processing;				
	(²) and/or [-	intended for humar	consu	mption for	commercial reasons	s of animal origin, which are no lo or due to problems of manufacturi ublic or animal health arise;]				
	(²) and/or [-	aquatic animals, ar of diseases commu				ammals, which did not show any :	signs			
	(²) and/or [-	animal by-products products for humar		•	nals originating from բ	olants or establishments manufact	uring			
	(²) and/or [-				rom animals which to humans or animals	did not show any signs of dis :	ease			
		(i) shells from	shellfis	sh with soft	tissue or flesh;					
		(ii) the followin	ng originating from terrestrial animals:							
		— hate	chery by-products,							
		— egg	S,							
		— egg	by-pro	ducts, inclu	uding egg shells;					

Animal by-products for the manufacture of petfood

								_		of petfood
II.	Heal	th informa	ition	II.a.	Certific	ate referenc	e No	II.b.		
			(iii) day-old chic	ks kille	ed for co	mmercial re	easons;]			
	(²) aı	nd/or [-	animal by-products humans or animals;		aquatic	or terrestria	al invertebrat	tes, other th	nan species	pathogenic to
	(²) aı	nd/or [-	animals and parts Category 1 material and Category 2 mat	as ref	erred to	in Article 8(a)(iii), (iv) an	d (v) of Reg	ulation (EC)	
	(²) aı	nd/or [-	Council Directive 9	hals which have been treated with certain substances which are prohibited by $96/22/EC$ (4a), the import of the material being permitted in accordance with Regulation (EC) No 1069/2009;]						
II.1.8.	have been deep-frozen at the plant of origin or have been preserved in accordance with European Union legislation in such a way that they will not spoil between dispatch and delivery to the plant of destination in the European Union or during the transit through the European Union;									
II.1.9.	in the case of raw material derived from animals which have been treated with certain substances prohibited by Directive 96/22/EC for the manufacture of petfood, the import being permitted in accordance with Article 35(a)(ii) of Regulation (EC) No 1069/2009:									
	(a) it has been marked in the third country before entry into the territory of the European Union by a cross of liquefied charcoal or activated carbon on each outer side of each frozen block, or, when the raw material is transported in pallets which are not divided into separate consignments during transport to the petfood plant of destination in the European Union or during the transit through the European Union, on each outer side of each pallet, in a way that the marking covers at least 70 % of the diagonal length of the frozen block and be of at least 10 cm width;								raw material is e petfood plant ach outer side	
	(b) in the case of material which is not frozen, the raw material has been marked in the third country entry into the territory of the European Union by spraying it with liquefied charcoal or by applying ch powder in such a way that the charcoal is clearly visible on the material; and									
	(c)		animal by-products -treated raw materia							
(²) (⁵) [II.2.	Spec	cific require	ments							
(²) (⁶) [II.2.1.	(II.1.:	2), where	s in this consignmen vaccination program led in domestic bovin	mes a	against					
(²) (⁷) [II.2.2.	rumii hour	nants, whic	s in this consignmer ch have maturated a case of masseter mo ours.]]	t an a	mbient	temperature	of more tha	n + 2 °C for	r a period of	at least three
(²) [II.3.		animal by-p n and:	roducts for the manu	facture	e of pett	ood contain	s or is derive	ed from anim	nal-by produc	cts of ruminant
	(²) ei		inate from a country Decision 2007/453/E	-			•			
	(²) OI	Dec or d rum	pinate from a countrision 2007/453/EC in erived product were of inants with meat-and estrial Animal Health	which derived d-bone	n there l d from a e meal	nas been ar nimals born and greave	indigenous after the date s derived fro	BSE case, e from whicl om ruminan	and the anir h the ban on its, as define	nal by-product the feeding of ed in the OIE
	(²) ei	<i>ither</i> [is d	erived from other run	ninants	s than b	ovine, ovine	or caprine a	nimals.]		

COUNTRY

Animal by-products for the manufacture of petfood

							oi petiood
II.	. Health information			II.a.	. Certificate reference No	0	II.b.
	(²) or	[is derived	from bovine	, ovine	or caprine animals and do	oes not co	ontain and is not derived from:
	continuous				·	a countr	those derived from animals born, ry or region classified as posing a 07/453/EC.]]]
		(²) or	• (/		d risk material as defined 2001 of the European Parli	•	t 1 of Annex V to Regulation (EC) d of the Council (8);
			ar sl ac	imals, aughter cordan	except from those animing red in a country or region	nals that v n classifie	m bones of bovine, ovine or caprine were born, continuously reared and at as posing a negligible BSE risk in 7/453/EC(9), in which there has been
	anir nen the thos or r				which have been killed, tissue by means of an eld ial cavity, or by means of himals that were born, cor	after stu ongated ro gas injec ntinuously	ained from bovine, ovine or caprine unning, by laceration of the central od-shaped instrument introduced into sted into the cranial cavity, except for reared and slaughtered in a country as Erisk in accordance with Decision

Notes

Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for a commodity to be imported into the European Union.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for a transit commodity. Products
 in transit may only be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in the case of unloading and reloading in the European Union.
- Box reference I.19: use the appropriate HS code: 05.04; 05.06; 05.07; 05.11.91 or 05.11.99; 23.01; 41.01.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.
- Box reference I.25: technical use: any use other than feeding of farmed animals, other than fur animals, and the
 production or manufacturing of pet food.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28:
 - species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or Suidae, Pesca,
 Mollusca, Crustacea, invertebrates other than Mollusca and Crustacea;
 - Manufacturing plant: provide the veterinary control number of the approved establishment.

Part II:

- (^{1a}) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.

COUNTRY

Animal by-products for the manufacture of petfood

II.	Health information	II.a.	Certificate reference No	II.b.					
(^{1c})	The name and ISO code number of the expo	rting c	country as laid down in:						
	 Part 1 of Annex II to Regulation (EU) N 	lo 206	/2010;						
	Part 1 of Annex I to Regulation (EC) N	o 798/	2008, and						
	Part 1 of Annex I to Regulation (EC) N	o 119/	2009.						
	In addition the ISO code of regionalisation in the abovementioned Annexes (where applicable for the susceptible species concerned) must be included.								
(^{1d})	Only for countries from which game meat intended for human consumption of the same animal species is authorised for importation into the European Union.								
(2)	Delete as appropriate.								
(3)	Excluding raw blood, raw milk, hides and skins, hooves and horn, pig bristles and feathers (see relevant specific certificates in that Annex for the import of these products).								
(4)	OJ L 303, 18.11.2009, p. 1.								
(^{4a})	OJ L 125, 23.5.1996, p. 3.								
(⁵)	Supplementary guarantees to be provided when the material of domestic ruminants originated in the territory of a South American or South African country or part thereof from where only maturated and deboned fresh meat of domestic ruminants for human consumption is permitted for exportation to the European Union. The whole masseter muscles of bovine animals, incised in accordance with Part B.1 of Chapter I of Section IV of Annex I to Regulation (EC) No 854/2004 of the European Parliament and of the Council (OJ L 139, 30.4.2004, p. 206), are also permitted.								
(⁶)	Only for certain South American countries.								
(⁷)	Only for certain South American and South A	African	countries.						
(8)	OJ L 147, 31.5.2001, p. 1.								
(⁹)	OJ L 172, 30.6.2007, p. 84.								
_	The signature and the stamp must be in a dif	ferent	colour to that of the printing.						
_	Note for the person responsible for the cons and must accompany the consignment until i								
Offic	cial veterinarian/Official inspector								
	Name (in capital letters):		Qua	lification and title:					
	Date:		Sign	ature: '					
	Stamp:								

Type of packaging

I.23. Seal/Container No

(2) Chapters 4(B) to 4(D) are replaced by the following:

'CHAPTER 4(B)

Health certificate

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

COUNTRY: Veterinary certificate to EU Consignor 1.2. Certificate reference No I.2.a. Name 1.3. Central competent authority Address 1.4. Local competent authority Tel. 1.5. Consignee 1.6. Person responsible for the load in EU Part I: Details of dispatched consignment Name Name Address Address Postcode Postcode Tel. Tel. 1.7. Country ISO code Region of Code 1.9. Country of ISO I.10. Region of Code destination origin destination of origin code I.11. Place of origin I.12. Place of destination Approval number Name Custom warehouse Address Name Approval number Name Approval number Address Address Name Approval number Postcode Address I.13. Place of loading I.14. Date of departure I.15. Means of transport I.16. Entry BIP in EU Aeroplane \square Ship 🔲 Railway wagon \square Road vehicle Other \square 1.17. Identification Documentation references I.18. Description of commodity I.19. Commodity code (HS code) I.20. Quantity I.21. Temperature of product 1.22. Number of packages Chilled \square Ambient \square Frozen

1.25.	5. Commodities certified for:										
	Animal feedingstuff ☐		Manufactu	ıre of petfood □	Technical	nical use 🛘					
1.26.	For transit through EU to third	d country		I.27. For import or admissio	n into EU						
	Third country	ISO code									
1.28.	. Identification of the commodities Approval number of establishments										
	Species (Scientific name)			Manufacturing plant		Batch number					

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

						Coun	u be used as feed material			
	II. Health inform		nation	II.a.	Certificate reference No	11.	.b.			
		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No the European Parliament and of the Council (^{1a}) and Commission Regulation (EU) No 142/2011 (^{1b}) a the blood products described above:								
	II.1.	consist of blood products that satisfy the health requirements below;								
ıtion	II.2.	consist exclusively of blood products not intended for human consumption;								
Part II: Certification	II.3.	have been prepared and stored in a plant, approved and supervised by the competent authority in accordance with Article 24 of Regulation (EC) No 1069/2009;								
art II:	II. 4 .	have been prepared exclusively with the following animal by-products:								
Ф		(²) either [blood of slaughtered animals, which is fit for human consumption in accordance legislation, but which is not intended for human consumption for commercial reasons;]								
		(²) and/or	[blood of slaughtered animals, which has been rejected as unfit for human consumption in accordance with Union legislation, but which did not show any signs of diseases communicable to humans or animals, which has been derived from carcases that have been slaughtered in a slaughterhouse and which were considered fit for human consumption following an ante-mortem inspection in accordance with Union legislation;]							
	II.5.	in order to inactivate pathogenic agents, have been submitted								
		(²) either	[to processing in accordance with processing method							
		(²) or	[to a method and parameters which ensure that the product complies with the microbiological standards set out in Chapter I of Annex X to Regulation (EU) No 142/2011;]							
		(²) or	[in the case of blood products, including spray dried blood and blood plasma, of porcine origin intended for the feeding of porcine animals, to a heat treatment at a temperature of at least 80°C throughout the substance and the dry blood and blood plasma does not contain more than 8% w/w moisture with a water activity (Aw) of less than 0,60.]							
	II.6.									
		(²) either	[packed in new or sterilised bags;]							
		(²) or	[transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectant approved by the competent authority before use,]							
	and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION';									
II.7. the end product was stored in enclosed storage;										
	II.8.	the product ha	as undergone all precautio	ns to avoid contamination with pathogenic agents after treatment;						
(²) and [in the case of blood products, including spray dried blood and blood intended for the feeding of porcine animals, has been stored in dry room temperature for a period of at least 6 weeks.]										
	II.9.	have been examined prior to dispatch under the responsibility of the competent authority by taking a sample during or on removal from storage which was found to comply with the following standards (4):								
		Salmonella:	absence in 25	5g: n =	5, c = 0, m = 0, M = 0,					
		Enterobacteria	aceae: $n = 5, c = 2, i$	m = 10	, M = 300 in 1 gram;					

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

					buid be used as feed material			
II.	Health information			II.a. Certificate reference No	II.b.			
(²) [II.10.	the blood prod	lucts descri	bed above					
	(²) either	[is derived from other ruminants than bovine, ovine or caprine animals.]]						
	(²) or	[is derived	[is derived from bovine, ovine or caprine animals and does not contain and is not derived from:					
		(²) either	[bovine, ovine and caprine materials other than those derived from animals bor continuously reared and slaughtered in a country or region classified as posing negligible BSE risk in accordance with Decision 2007/453/EC.]]					
		(²) or		of Annex V to Regulation (EC) of the Council (5);				
			; ;	mechanically separated meat obtained from bones of boving animals, except from those animals that were born, continuous slaughtered in a country or region classified as posing a negacordance with Commission Decision 2007/453/EC (6), in been no indigenous BSE case,				
			; ; ;	animal by-product or derived product obtained from bovine, ovine or canimals which have been killed, after stunning, by laceration of the conervous tissue by means of an elongated rod-shaped instrument introfinto the cranial cavity, or by means of gas injected into the cranial except for those animals that were born, continuously reared and slaugin a country or region classified as posing a negligible BSE risk in according with Decision 2007/453/EC.]]]				
II.11.	the blood prod	pod products described above:						
	(²) either		do not contain milk or milk products of ovine or caprine animal origin or is not intended for feed for armed animals, other than fur animals.]					
	(²) or	-	nin milk or milk products of ovine or caprine animal origin and is intended for feed for far als, other than fur animals, which:					
		(a)		d from ovine and caprine animals which have been kept continuously sin country where the following conditions are fulfilled:				
			(i) c	classical scrapie is compulsorily notifiable;				
				an awareness, surveillance and monitoring s scrapie;	ystem is in place for classical			
				official restrictions apply to holdings of ovine or a suspicion of TSE or the confirmation of class				
				ovine and caprine animals affected with cla destroyed;	assical scrapie are killed and			
			; ;	he feeding to ovine and caprine animals of mas defined in the Terrestrial Animal Health Coc Animal Health (OIE), of ruminant origin has enforced in the whole country for a periodeven years;	le of the World Organisation for been banned and effectively			
		(b)	originate fro	m holdings where no official restrictions are	imposed due to a suspicion of			
		(c)		om holdings where no case of classical scrap of at least the preceding seven years or, follow scrapie:				

COUNTRY

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

II.	Health inforn	nation	II.a	ā.	Certificate reference No	II.b.	
		(²) either	[all ovine and caprine animals on the holding have been killed and destroyed or slaughtered, except for breeding rams of the ARR/ARR genotype, breeding ewes carrying at least one ARR allele and no VRQ allele and other ovine animals carrying at least one ARR allele;]				
(2) or [all animals in which classical scrapie was confirm destroyed, and the holding has been subjected two years since the date of confirmation of the last intensified TSE monitoring, including testing with presence of TSE in accordance with the labora point 3.2 of Chapter C of Annex X to Regulation (E the following animals which are over the age of animals of the ARR/ARR genotype:					cted for a period of at least e last classical scrapie case to with negative results for the aboratory methods set out in on (EC) No 999/2001, of all of		
			— a	anir	mals which have been slaughtered for	human consumption; and	
					mals which have died or been killed killed in the framework of a disease e	•	
II.12.	the blood products described above contain or are derived from animal-by products of non-ruminant origin, and are, according to the statement of the Consignor referred to in Box I.1,						
	(²) either	[not intended for the p	roducti	on	of feed for farmed animals, other than	ı fur animals.]	
(²) (7) or [intended for the production of feed for non-ruminant farmed animals, other than Consignor has undertaken to ensure that the border inspection post of entry will I results of the analyses carried out in accordance with the methods set of Commission Regulation (EC) No 152/2009 (8).]					of entry will be provided with the		

Notes

Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is required to be filled in only if
 it is a certificate for a commodity that is to be transited through the European Union; it may be filled in if the certificate is
 for a commodity that is to be imported into the European Union.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for a transit commodity. Products
 in transit may only be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in the case of unloading and reloading in the European Union.
- Box reference I.19: use the appropriate HS code: 05.11.91, 05.11.99, 35.02 or 35.04.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.
- Box reference I.25: technical use: any use other than feeding of farmed animals, other than fur animals, and the
 production or manufacturing of pet food.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: Species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or Suidae, Pesca, Reptilia.

COUNTRY

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

	could be used as feed if							
II.	Health information	II.a. Certificate reference No		II.b.				
Part II:								
(^{1a})	OJ L 300, 14.11.2009, p. 1.							
(1b)	OJ L 54, 26.2.2011, p. 1.							
(²)	Delete as appropriate.							
(3)	Insert method 1 to 5 or method 7 as applicable.							
(4)	Where:							
	n = number of samples to be tested;							
	m = threshold value for the number of bacteria; the result is considered satisfactory if the number of bacteria in a samples does not exceed m;							
	M = maximum value for the number of bacteria; the result is considered unsatisfactory if the number of bacteria in one or more samples is M or more; and							
	c = number of samples the bacterial count of which may be between m and M, the sample still being considered acceptable if the bacterial count of the other samples is m or less.							
(⁵)	OJ L 147, 31.5.2001, p. 1.							
(⁶)	OJ L 172, 30.6.2007, p. 84.							
(7)	The person responsible for the load referred to in Box I.6 must ensure that, if the blood products described in this health certificate are intended to be used for the production of feed for non-ruminant farmed animals, other than fur animals, the consignment must be analysed, in accordance with the methods set out in Annex VI to Regulation (EC) No 152/2009, in order to verify the absence of unauthorised constituents of animal origin. The information on the result of such analysis must be attached to this health certificate when presenting the consignment at a border inspection post of the European Union.							
(8)	OJ L 54, 26.2.2009, p. 1.							
_	The signature and the stamp must be in a different colour to that of the printing.							
_	Note for the person responsible for the consignment in the European Union: this certificate is only for veterinary purposes and must accompany the consignment until it reaches the border inspection post of the point of entry into the European Union.							
Official veterinarian/Official inspector								
	Name (in capital letters):	Quali	nd title:					
	Date:	Signature:						
	Stamp:							

CHAPTER 4(C)

Health certificate

For untreated blood products, excluding those of equidae, for the manufacture of derived products for purposes outside the feed chain for farmed animals, intended for dispatch to or for transit through (2) the European Union

COL	JNTRY	/ :				Veterinary certificat	te to EU	
	l.1.	Consignor	1.2.	Certificate reference	No	I.2.a.		
		Name	1.3.	I.3. Central competent authority				
		Address	1.4.	I.4. Local competent authority				
	Tel.							
	1.5.	Consignee	I.6.	I.6. Person responsible for the load in EU				
nen		Name Address		Name Address				
signi		Address		Address				
con		Postcode		Postcode				
hed		Tel.		Tel.				
Part I : Details of dispatched consignment	1.7.	Country ISO code I.8. Region of Code origin	1.9.		SO code	I.10. Region of destination	Code	
ls of								
)etai	l.11.	Place of origin	I.12.	Place of destination				
t : [Name Approved asserting				0		
Part		Name Approval number Address		Name		Custom warehouse Approval number	Ц	
		Name Approval number		Address		Approvar number		
		Address		Addioso				
		Name Approval number		Postcode				
		Address						
	I.13.	Place of loading	1.14.	Date of departure				
	I.15.	Means of transport	I.16.	Entry BIP in EU				
		·		,				
		Aeroplane ☐ Ship ☐ Railway wagon ☐						
		Road vehicle Other	1.17.					
		Identification						
		Documentation references						
	l.18.	Description of commodity		I.19	9. Commo	odity code (HS code)		
						I.20. Quantity		
	I.21.	Temperature of product				I.22. Number of page	ckages	
		Ambient ☐ Chilled ☐		Frozen 🗖				
	123	Seal/Container No				124 Type of packa	aina	

Γ	EN
	LIN

1.25.	Commodities certified for:	
	Technical use	
1.26.	For transit through EU to third country	I.27. For import or admission into EU
	Third country ISO code	
1.28.		
	Approval numbe	er of establishments
	Species (Scientific name) Manufac	turing plant Batch number

Untreated blood products, excluding those of equidae, for the manufacture of derived products for purposes outside the feed chain for farmed animals

	II.	Health information		II.a. Certificate reference No					
		the European	Parliament and of the C	n, declare that I have read and understood Regulation (EC) No 1069/2009 of council (1a), and in particular Article 8(c) and Article 8(d) and Article 10 thereof, 142/2011 (1b), and in particular Chapter II of Annex XIV thereto, and certify					
II.1. the blood products described above consist of blood products that satisfy the health requirements									
catio	II.2.	they consist exclusively of blood products not intended for human or animal consumption;							
Part II: Certification	II.3.	II.3. they have been prepared and stored in a plant supervised by the competent authority or in the establishment collection, exclusively with the following animal by-products:							
Part II		(²) either		[- blood of slaughtered animals, which is fit for human consumption in accordance with Union legislation, but is not intended for human consumption for commercial reasons;]					
		(²) and/or	[- blood of slaughtered animals, which is rejected as unfit for human consumption in accordar with Union legislation, but which did not show any signs of diseases communicable to humans animals, derived from carcases that have been slaughtered in a slaughterhouse and we considered fit for human consumption following an ante-mortem inspection in accordance we Union legislation;]						
		(2) and/or [- blood of slaughtered animals, which did not show any signs of diseases communicable to humans or animals, obtained from animals that have been slaughtered in a slaughterhouse after having been considered fit for human consumption following an ante-mortem inspection in accordance with Union legislation;]							
		(²) and/or	for [- blood and blood products derived from the production of products intended for huma consumption;]						
		(²) and/or	[- blood and blood products originating from live animals that did not show signs of any dise communicable through that product to humans or animals;]						
		(²) and/or	[- animal by-products derived from animals which have been submitted to illegal treatme defined in Article 1(2)(d) of Council Directive 96/22/EC (^{2a}) or Article 2(b) of Council Directive 96/23/EC (^{2b});]						
		(²) and/or	[- animal by-products containing residues of other substances and environmental con listed in Group B(3) of Annex I to Directive 96/23/EC, if such residues exceed the perm laid down in Union legislation or, in the absence thereof, in national legislation;]						
	II.4.	the blood, that such products were manufactured from, was collected in slaughterhouses approved in accordan with Union legislation, in slaughterhouses approved and supervised by the competent authority of the country collection or from live animals in facilities approved and supervised by the competent authority of the country collection;							
	(²) [II.5.	in the case of blood products obtained from animals belonging to the taxa Artiodactyla, Perissodactyla and Proboscidea, including crossbreds between species of those taxa, the blood was collected in a country or region where no case of rinderpest, peste des petits ruminants and Rift Valley fever has been recorded for a period of at least the preceding 12 months and in which vaccination has not been carried out against those diseases for a period of at least the preceding 12 months, and;							
		(²) either	[in third countries, territories or parts thereof						
		(²) or	country or codes (3) for the been recorded for a programmes against	tories or parts thereof					

Untreated blood products, excluding those of equidae, for the manufacture of derived products for purposes outside the feed chain for farmed animals

			the	feed chain for farmed animals			
II.	Health inforn	nation	II.a. Certificate reference No	II.b.			
(²) [II.5.1.	in the case of	animals other than Sui	lae and Tayassuidae, in third countries or regi	ons in which :			
	(²) either [no case of vesicular stomatitis and bluetongue (²) (including the presence of seropositive animals) has been recorded for a period of at least the preceding 12 months and in which vaccination has not been carried out against those diseases for a period of at least the preceding 12 months;]						
	(²) or	[vesicular stomatitis and bluetongue (²) seropositive animals are present (4);]]					
(²) [II.5.2.	classical swin	e fever and African sv	ae, in third countries or regions in which no ne fever has been recorded for a period of a out against those diseases for a period of at	at least the preceding 12 months			
	(²) either	for a period of at least the preceding 12 months and in which vaccination has not been carried against this disease for a period of at least the preceding 12 months;]]					
	(²) or	[vesicular stomatitis s	eropositive animals are present (4);]]]				
(²) [II.6.		blood products derive the country or region v	I from poultry or other avian species the animith code (5)	nals and the products come from			
		en free from Newcast Code of the OIE,	e disease and highly pathogenic avian influe	nza as defined in the Terrestrial			
	which for a pe	eriod of at least the pred	eding 12 months has not carried out vaccinati	on against avian influenza,			
			oducts are derived, have not been vaccinated disease master strain showing a higher pa				
II.7.	the products v	were:					
	(²) either	[packed in new or ste	ilised bags or bottles,]				
	(²) or		n containers or other means of transport the offectant approved by the competent authority				
	the outer pack	kaging or containers be	ar labels indicating 'NOT FOR HUMAN OR AN	IIMAL CONSUMPTION';			
II.8.	the products v	were stored in enclosed	storage;				
II.9.	all precaution	s were taken to avoid o	ontamination of the products with pathogenic a	gents during transport;			
(²) [II.10.	the untreated	blood products describ	ed above				
	(²) either	[is derived from other	ruminants than bovine, ovine or caprine anima	als.]]			
	(²) or	[is derived from bovir	e, ovine or caprine animals and does not cont	ain and is not derived from:			
		(²) either [bovine, ovine and caprine materials other than those derived from animals born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]					
			ecified risk material as defined in point 1 o 999/2001 of the European Parliament and of				
		ar sla ac	chanically separated meat obtained from bomals, except from those animals that were ughtered in a country or region classified ascordance with Commission Decision 2007/453 indigenous BSE case,	born, continuously reared and posing a negligible BSE risk in			

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COUNTRY

Untreated blood products, excluding those of equidae, for the manufacture of derived products for purposes outside the feed chain for farmed animals

II.	Health information			II.a.	Certificate reference No	II.b.
		(c)	animals nervous the crait those a	s which s tissumial ca nial ca nimal on cla	roduct or derived product obtained ch have been killed, after stunning ue by means of an elongated rod-sha avity, or by means of gas injected in s that were born, continuously reare ssified as posing a negligible BSE rist.	, by laceration of the central ped instrument introduced into to the cranial cavity, except for d and slaughtered in a country

Notes

Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is required to be filled in only if it is a certificate for a commodity that is to be transited through the European Union; it may be filled in if the certificate is for a commodity that is to be imported into the European Union.
- Box reference I.11 and I.12: Approval number: the registration number of the establishment or plant, which has been issued by the competent authority.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for a transit commodity. Products in transit may only be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to be provided. In the case of unloading and reloading in the European Union, the consignor must inform the border inspection post of the point of entry into the European Union.
- Box I.19: use the appropriate Harmonized System (HS) code under the following headings: 05.11; 30.02 or 35.02.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) must be included.
- Box reference I.25: technical use: any use other than feeding of farmed animals, other than fur animals, and the production or manufacturing of pet food.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28 Species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or Suidae, Pesca, Reptilian.

Part II:

- OJ L 300, 14.11.2009, p. 1.
- OJ L 54, 26.2.2011, p. 1.
- Delete as appropriate.
- OJ L 125, 23.5.1996, p. 3.
- OJ L 125, 23.5.1996, p. 10.
- Code of the territory as it appears in Part 1 of Annex II to Regulation (EU) No 206/2010 (OJ L 73, 20.3.2010, p. 1).
- In this case following the veterinary checks provided for in Directive 97/78/EC (OJ L 24, 30.1.1998, p. 9), and in accordance with the conditions laid down in Article 8(4) of that Directive, the products must be transported directly to the establishment at the place of destination.

COUNTRY

Untreated blood products, excluding those of equidae, for the manufacture of derived products for purposes outside the feed chain for farmed animals

II.	Health information	II.a. Certificate reference N	lo	II.b.			
(⁵)	Code of the territory as it appears in Part 1 of Annex I to Commission Regulation (EC) No 798/2008 (OJ L 226, 23.8.2008, p. 1).						
(⁶)	OJ L 147, 31.5.2001, p. 1.						
(7)	OJ L 172, 30.6.2007, p. 84.						
_	The signature and the stamp must be in a diffe	rent colour to that of the printir	ng.				
_	Note for the person responsible for the consignment in the European Union: this certificate is only for veterinary purposes and must accompany the consignment until it reaches the border inspection post of the point of entry into the European Union.						
Offic	cial veterinarian/Official inspector						
	Name (in capital letters):		Qualification a	and title:			
	Date: Signature:						
	Stamp:						

Veterinary certificate to EU

COUNTRY:

CHAPTER 4(D)

Health certificate

For treated blood products, excluding those of equidae, for the manufacture of derived products for purposes outside the feed chain for farmed animals, intended for dispatch to or for transit through (2) the European Union

	l.1.	Consignor	1.2.	Certificate reference No		I.2.a.	
		Name	I.3. Central competent authority				
		Address	1.4.	Local competent authority	•		
		T-1					
		Tel.					
	1.5.	Consignee	1.6.	Person responsible for the	e load in	EU	
ent		Name		Name			
ignr		Address		Address			
Part I : Details of dispatched consignment		Postcode		Postcode			
pet		Tel.		Tel.			
atcl	1.7.	Country ISO code I.8. Region of Code	1.9.	Country of ISO	I 1	0. Region of	Code
disp		of origin origin	1.0.	destination code		destination	
s of							
etail	l.11.	Place of origin	1.12.	Place of destination			
.: O							
art		Name Approval number			С	ustom warehouse	
a.		Address		Name	А	pproval number	
		Name Approval number		Address			
		Address					
		Name Approval number		Postcode			
		Address					
	I.13.	Place of loading	I.14.	Date of departure			
	1.15.	Means of transport	I.16.	Entry BIP in EU			
		,		•			
		Aeroplane ☐ Ship ☐ Railway wagon ☐					
		Road vehicle Other	1.17.				
		Identification					
		Documentation references					
	I.18.	Description of commodity		I.19. Co	ommodit	ty code (HS code)	
					1.3	20. Quantity	
	I.21.	Temperature of product			1.3	22. Number of pac	kages
		Ambient ☐ Chilled ☐		Frozen 🗖			
	1.23.	Seal/Container No			1.	24. Type of packa	ging



1.25.	Commodities certified for:	
	Technical use □	
1.26.	For transit through EU to third country	I.27. For import or admission into EU
	Third country ISO code	
1.28.	Identification of the commodities	
	Approval numb	er of establishments
	Species (Scientific name) Manufa	octuring plant Batch number

Treated blood products, excluding those of equidae, for the manufacture of derived products for purposes outside the feed chain for farmed animals

	II.	Health information		II.a.	Certificate refer	rence No	II.b.			
		the European Pa	rliament and of the Co	uncil (1	a), and in particul	ar Article 8(c) and Artic	ulation (EC) No 1069/2009 of cle 8(d) and Article 10 thereof, Annex XIV thereto, and certify			
u	II.1.	II.1. the blood products described above consist of blood products that satisfy the requirements below;								
ficatio	II.2.	they consist exclusively of blood products not intended for human or animal consumption;								
Part II: Certification	II.3.	they have been prepared and stored in a plant supervised by the competent authority, exclusively with the following animal by-products:								
Part		(²) either [-	(²) either [- blood of slaughtered animals, which is fit for human consumption in accordance with Union legislation, but is not intended for human consumption for commercial reasons;]							
(2) and/or [- blood of slaughtered animals, which is rejected as unfit fo with Union legislation, but which did not show any signs of animals, derived from carcases that have been slaughted considered fit for human consumption following an ante-munion legislation;]						w any signs of disease: been slaughtered in	s communicable to humans or a slaughterhouse and were			
		(²) and/or [-	humans or animals,	obtaine ered fi	d from animals t t for human co	hat have been slaughte	of diseases communicable to ered in a slaughterhouse after n ante-mortem inspection in			
		(²) and/or [-				live animals that did r cts to humans or anima	not show clinical signs of any sls;]			
		(²) and/or [-	blood and blood p consumption;]	roduct	derived from	the production of pr	roducts intended for human			
		(²) and/or [-		in Arti			nave been submitted to illegal C (^{2a}) or Article 2(b) of Council			
	listed in Group B(3)				nex I to Directiv		environmental contaminants esidues exceed the permitted eational legislation;]			
	II.4.	4. the blood that these products were manufactured from was been collected in slaughterhouses a accordance with Union legislation, in slaughterhouses approved and supervised by the competent auth country of collection or from live animals in facilities approved and supervised by the competent auth country of collection.								
	(²) [II.5.	(²) [II.5. In the case of blood products derived from Artiodactyla, Perissodactyla and Proboscidea including crossbreeds, other than Suidae and Tayassuidae, the products have undergone one of the following treatm guaranteeing the absence of pathogens of foot-and-mouth disease, vesicular stomatitis, rinderpest, peste des ruminants, Rift Valley fever and bluetongue:								
(²) either [heat treatment at a temperature of 65 °C for at least three check;]						or at least three hours	followed by an effectiveness			
		(²) and/or	[irradiation at 25 kGy	by gar	mma rays, follow	ed by an effectiveness	check;]			
		(²) and/or	[change in pH to pH	5 for tw	o hours, followed	d by an effectiveness c	heck;]			
		(²) and/or [heat treatment of at least 80 °C throughout their substance, followed by an effectivene check.]]								

Treated blood products, excluding those of equidae, for the manufacture of derived products for purposes outside the feed chain for farmed animals

			the re	ed chain for farmed animals			
II.	Health informati	ion	II.a. Certificate reference No	II.b.			
(²) [II.6.	undergone one of and-mouth disea	of the following treatmase, vesicular stoma	om Suidae, Tayassuidae, poultry and other avian species, the products have ents guaranteeing the absence of pathogens of the following diseases: footies, swine vesicular disease, classical swine fever, African swine fever, ic avian influenza, as appropriate to the species:				
	(²) either	[heat treatment at a check;]	a temperature of 65 °C for at least three hours	, followed by an effectiveness			
	(²) and/or	[irradiation at 25 kG	y by gamma rays, followed by an effectiveness	check;]			
	(²) and/or	[heat treatment of at least 80 °C for Suidae/Tayassuidae (²) and at least 70 °C for poultry a other avian species (²) throughout the substance of the product, followed by an effectivene check]].					
(²) [II.7.		case of blood products derived from species other than those listed in point II.5 or II.6, the products have gone of the following treatment (please specify):]					
II.8.	The products wer	re:					
	(²) either	[packed in new or s	terilised bags or bottles,]				
	(²) or	[transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectant approved by the competent authority before use;] and					
	the outer packagi	ing or containers bear	labels indicating 'NOT FOR HUMAN OR ANIM	IAL CONSUMPTION';			
II.9.	the products were	e stored in enclosed s	etorage;				
II.10.	all precautions w	ere taken to avoid the	contamination of the products with pathogenic	agents after treatment;			
(²) [II.11.	The treated blood	d products described	above				
	(²) either	[is derived from other	er ruminants than bovine, ovine or caprine anim	nals.]]			
	(²) or	[is derived from bov	ine, ovine or caprine animals and does not con	tain and is not derived from:			
		conf	rine, ovine and caprine materials other than tho cinuously reared and slaughtered in a country o ligible BSE risk in accordance with Decision 200	or region classified as posing a			
		(²) or [(a)	specified risk material as defined in point 1 No 999/2001 of the European Parliament ar				
		(b) mechanically separated meat obtained from bones of bovine, ovine caprine animals, except from those animals that were born, continuous reared and slaughtered in a country or region classified as posing negligible BSE risk in accordance with Commission Decisi 2007/453/EC (4), in which there has been no indigenous BSE case,					
		(c)	animal by-product or derived product obt caprine animals which have been killed, a the central nervous tissue by means construment introduced into the cranial cavity into the cranial cavity, except for those continuously reared and slaughtered in a coposing a negligible BSE risk in accordance of	fter stunning, by laceration of of an elongated rod-shaped of, or by means of gas injected se animals that were born, country or region classified as			

COUNTRY

Treated blood products, excluding those of equidae, for the manufacture of derived products for purposes outside the feed chain for farmed animals

				outside the it	eeu chani for farmeu ammais
II.	Health information	II.a.	Certificate reference No)	II.b.
Note	es				
Part	:I:				
	Box reference I.6: Person responsible for the it is a certificate for a commodity to be transcommodity to be imported into the European	ted th	rough the European Uni		
_	Box reference I.11 and I.12: Approval numb issued by the competent authority.	er: the	e registration number of	the establishn	nent or plant, which has been
_	Box reference I.12: Place of destination: this in transit may only be stored in free zones, free				a transit commodity. Products
_	Box reference I.15: Registration number (rails to be provided. In the case of unloading a entry into the European Union.				
_	Box I.19: use the appropriate Harmonized Sys	stem (HS) code under the follow	wing headings:	05.11, 30.02, 35.02 or 35.04.
_	Box reference I.23: for bulk containers, the co	ntaine	r number and the seal nu	umber (if applic	able) must be included.
_	Box reference I.25: technical use: any use production or manufacturing of pet food.	othe	r than feeding of farme	d animals, oth	ner than fur animals, and the
_	Box reference I.26 and I.27: fill in according to	whet	her it is a transit or an im	port certificate.	
_	Box reference I.28 in case of Species: sel Ruminantia or Suidae, Pesca, Reptilian.	ect fr	om the following: Aves,	Ruminantia, S	Suidae, Mammalia other than
Part	II:				
(^{1a})	OJ L 300, 14.11.2009, p. 1.				
(1b)	OJ L 54, 26.2.2011, p. 1.				
(²)	Delete as appropriate.				
(^{2a})	OJ L 125, 23.5.1996, p. 3.				
(^{2b})	OJ L 125, 23.5.1996, p. 10.				
(³)	OJ L 147, 31.5.2001, p. 1.				
(4)	OJ L 172, 30.6.2007, p. 84.				
_	The signature and the stamp must be in a diff	erent (colour to that of the printi	ng.	
_	Note for the person responsible for the consignment in the European Union: this certificate is only for veterinary purposes and must accompany the consignment until it reaches the border inspection post of the European Union.				
Offic	cial veterinarian/Official inspector				
	Name (in capital letters):			Qualification	and title:
	Date:			Signature:	
	Stamp:				

(3) Chapter 6(B) is replaced by the following:

'CHAPTER 6(B)

Health certificate

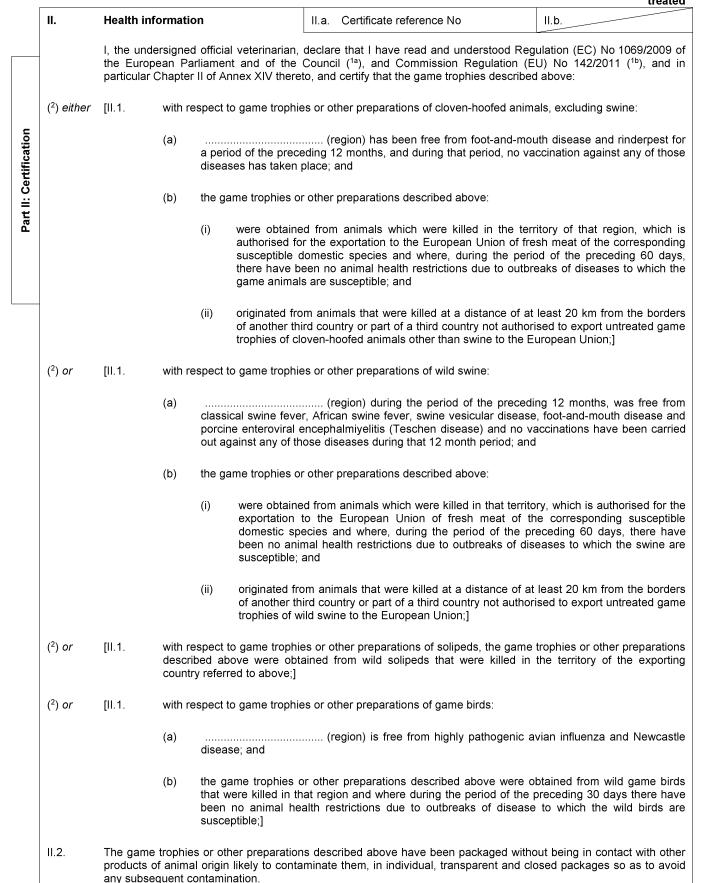
For game trophies or other preparations of birds and ungulates consisting of entire parts which have not been treated, intended for dispatch to or for transit through (2) the European Union

			n.o. Contrair compotent additionty					
		Address	1.4.	Local competen	t authority			
		Tel.						
	1.5.	Consignee	I.6. Person responsible for the load in EU					
in		Name		Name				
signme		Address		Address				
l con		Postcode		Postcode				
chec		Tel.		Tel.				
Part I : Details of dispatched consignment	I.7.	Country ISO code I.8. Region of Code of origin	1.9.	Country of destination	ISO code	I.10. Region of destination	Code	
s of								
Detail	l.11.	Place of origin	I.12.	Place of destina	tion			
art I:		Name Approval number				Custom warehouse		
۵		Address		Name		Approval number		
		Name Approval number		Address				
		Address						
		Name Approval number		Postcode				
		Address						
	I.13.	Place of loading	1.14.	Date of departur	re			
	I.15.	Means of transport	I.16.	Entry BIP in EU				
		Aeroplane ☐ Ship ☐ Railway wagon ☐						
		Road vehicle Other O	117	Number(s) of CI	TES			
		Identification	1.17.	rvamber(5) or or	120			
		Documentation references						
	I.18.	Description of commodity			I.19. Comm	nodity code (HS code)		
						,		
						I.20. Quantity		
	I.21.					I.22. Number of page	ckages	
	1.23.	Seal/Container No				I.24. Type of packaging		



1.25.	Commodities certified for:			
	Technical use □			
1.26.	For transit through EU to third	country	I.27. For import or admission into EU	
	Third country	ISO code		
1.28.	Identification of the commodition	es		
	Species (Scientific name)		Number of packages	
	Species (Scientific Harrie)		Number of packages	

Game trophies or other preparations of birds and ungulates consisting of entire parts which have not been treated



COUNTRY

Game trophies or other preparations of birds and ungulates consisting of entire parts which have not been treated

						tieateu			
II.	Health inf	formation		II.a.	Certificate reference No	II.b.			
(²) [II.3.	The game	ne trophies or other preparations described above							
	(²) either	[are derived	[are derived from other ruminants than bovine, ovine or caprine animals.]]						
	(²) or	[are derived	from bovine	e, ovine or	caprine animals and does not contain a	nd is not derived from:			
		(²) either	continuo	usly reare	d caprine materials other than those d and slaughtered in a country or r in accordance with Decision 2007/453/l	region classified as posing a			
		(²) or			ified risk material as defined in point 1 of Annex V to Regulation $99/2001$ of the European Parliament and of the Council (3);				
			° í a s a	animals, ex slaughtered accordance	ly separated meat obtained from bone ccept from those animals that were b in a country or region classified as p with Commission Decision 2007/453/E us BSE case,	orn, continuously reared and osing a negligible BSE risk in			
			r tl tl	animals who hervous tissended the cranial hose animal	product or derived product obtained finish have been killed, after stunning, sue by means of an elongated rod-shal cavity, or by means of gas injected intended in the cavity of the continuously reared assified as posing a negligible BSE ris [C.]]]	by laceration of the central ped instrument introduced into the cranial cavity, except for and slaughtered in a country			

Notes

Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is required to be filled in only if
 it is a certificate for a commodity to be transited through the European Union; it may be filled in if the certificate is for a
 commodity to be imported into the European Union.
- Box reference I.11 and I.12: Approval number: the registration number of the establishment or plant, which has been issued by the competent authority.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. Products in transit may only be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading in the European Union.
- Box reference I.19: use the appropriate HS code: 05.05; 05.06, 05.07, 05.11; 96.01 or 97.05.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) must be included.
- Box reference I.25: technical use: any use other than for animal consumption.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: Species: select from the following: Aves, Equidae, Tapiridae, Rhinoceritidae, Antilocaparidae, Bovidae, Camelidae, Cervidae, Giraffidae, Hippopotamindae, Moschidae Suidae, Tayassuidae, Tragulidae and Elephantidae.

COUNTRY

Game trophies or other preparations of birds and ungulates consisting of entire parts which have not been treated

II.	Health information	II.a. Certificate reference No		II.b.						
Part	Part II:									
(^{1a})	P) OJ L 300, 14.11.2009, p. 1.									
(1b)	OJ L 54, 26.2.2011, p. 1.									
(2)	Delete as appropriate.									
(3)	OJ L 147, 31.5.2001, p. 1.									
(4)	OJ L 172, 30.6.2007, p. 84.									
_	The signature and the stamp must be in a dif	fferent colour to that of the printin	g.							
_	Note for the person responsible for the consignment in the European Union: this certificate is only for veterinary purposes and must accompany the consignment until it reaches the border inspection post of the point of entry into the European Union.									
Offic	cial veterinarian/Official inspector									
	Name (in capital letters):		Qualification a	nd title:						
	Date:		Signature: '							
	Stamp:									

Type of packaging

I.23. Seal/Container No

(4) Chapter 8 is replaced by the following:

'CHAPTER 8

Health certificate

For animal by-products to be used for purposes outside the feed chain or for trade samples (2), intended for dispatch to or for transit through (2) the European Union

COUNTRY: Veterinary certificate to EU Consignor 1.2. Certificate reference No I.2.a. Name 1.3. Central competent authority Address 1.4. Local competent authority Tel. 1.5. Consignee 1.6. Person responsible for the load in EU Part I: Details of dispatched consignment Name Name Address Address Postcode Postcode Tel. Tel. 1.7. Country ISO code Region of Code 1.9. Country of ISO I.10. Region of Code destination destination of origin origin code I.11. Place of origin I.12. Place of destination Approval number Name Custom warehouse Address Name Approval number Name Approval number Address Address Name Approval number Postcode Address I.13. Place of loading I.14. Date of departure I.15. Means of transport I.16. Entry BIP in EU Aeroplane \square Ship 🔲 Railway wagon \square Road vehicle Other \square 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 \square Chilled Frozen

1.25.	Commodities certified for:										
	Technical use □	l									
1.26.	For transit throug	gh EU to third country	, 🗆	I.27. For import or admission into EU							
	Third country	ISO cod	de								
1.28.	Identification of t	he commodities									
			Approval number	of establishments							
(Sci	Species entific name)	Nature of commodity	Manufacturing plant	Number of packages	Net weight	Batch number					

Part II: Certification

COUNTRY

Animal by-products to be used for purposes outside the feed chain or for trade samples (2)

II. Health information Certificate reference No II a II b I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council (1a), and Commission Regulation (EU) No 142/2011 (1b), and in particular Chapter II of Annex XIV thereto, and certify that the animal by-products described above (2) either [are trade samples which consist of animal by-products intended for particular studies or analyses as referred to in the definition of trade samples in point 39 of Annex I to Regulation (EU) No 142/2011, that bear the label 'TRADE SAMPLE NOT FOR HUMAN CONSUMPTION'.] [satisfy the animal health requirements set out in point II.1.]; (2) or II.1. The animal by products described above II.1.1. have been imported (2) either [(a) obtained from materials from а third country. territory thereof:(3) authorised to export fresh meat to the European Union;] (2) and/or animals that either: (i) have remained in that third country, territory or part thereof eligible to export fresh meat to the European Union since birth or for a period of at least the preceding three months before the date of slaughter; and/or (ii) were killed in the wild in that third country, territory or part thereof (4);] [(c) derived from eggs, milk, rodents, lagomorphs, or aquatic animals or terrestrial or aquatic (2) and/or invertebrates;] (2) [II.1.2. in the case of materials other than materials derived from eggs, milk, rodents, lagomorphs, wool grease, aquatic animals, terrestrial or aquatic invertebrates and unprocessed furs, have been obtained from animals: (2) either [(a) coming from holdings: (i) where, for the following diseases for which the animals are susceptible, there has not been any case/outbreak of rinderpest, swine vesicular disease, Newcastle disease or highly pathogenic avian influenza during the period of the preceding 30 days, nor of classical or African swine fever during the period of the preceding 40 days; nor in the holdings situated in their vicinity within a 10 km radius, during the period of the preceding 30 days; and where there has not been any case/outbreak of foot-and-mouth disease during the (ii) period of the preceding 60 days, nor in the holdings situated in their vicinity within a 25 km radius, during the period of the preceding 30 days; and (b) which: (i) were not killed to eradicate any epizootic disease; remained on their holdings of origin for a period of at least 40 days before the date (ii) of departure and which were transported directly to the slaughterhouse without contact with other animals which did not comply with the same health conditions; (iii) at the slaughterhouse, passed the ante-mortem health inspection during the period of 24 hours before the time of slaughter and showed no evidence of the diseases referred to above for which the animals are susceptible; and were handled in the slaughterhouse before and at the time of slaughter or killing in (iv) accordance with the relevant provisions of Union legislation and complied with requirements at least equivalent to those laid down in Chapters II and III of Council Regulation (EC) No 1099/2009 (5)]

Animal by-products to be used for purposes outside the feed chain or for trade samples (2)

					the feed cha	ain or for trade samples (²)			
II.	Health inf	orm	ation		II.a. Certificate reference No	II.b.			
	(²) or	[(a) captured and k	killed in th	ne wild in an area:				
			follo rinc per	lowing dis iderpest, eriod of the	n a 25 km radius there has been no caseases for which the animals are susceptible. Newcastle disease or highly pathogenic e preceding 30 days nor of classical or Afee preceding 40 days; and	ole: foot-and-mouth disease, avian influenza during the			
			and	other terr	ated at a distance that exceeds 20 km fritory of a third country or part thereof, whice exportation of such material to the Europe	ch is not authorised at these			
		(b)		immediate	transported within a period of 12 hours for ely afterwards to a game establishmer				
(²) [II.1.3.	obtained i diseases i 30 days o exportation	n an refer r, in n to	establishment a red to in point I the event of a the European U	around w II.1.2 for case/out Jnion was	erials derived from fish or invertebrates can which, within a radius of 10 km, there has which the animals are susceptible during threak of one of those diseases, the prepayant of all representations of all representations.	s been no case/outbreak of g a period of the preceding paration of raw material for			
II.1.4.	have been	n ob requ	tained and prepired above, and	epared wi	ithout contact with other material which een handled so as to avoid contamination w	does not comply with the vith pathogenic agents;			
II.1.5.	disinfected sealed un PRODUC	nave been packed in new packaging which prevents any leakage or in packaging which has been cleaned and disinfected before use and, in the case of consignments shipped other than via parcel post, in containers sealed under the responsibility of the competent authority, bearing the label indicating 'ANIMAL BY-PRODUCTS ONLY FOR THE MANUFACTURE OF DERIVED PRODUCTS FOR USES OUTSIDE THE FEED CHAIN' and the name and address of the establishment of destination in the European Union;							
II.1.6.	consist on	ly of	the following ani	nimal by-p	roducts:				
	(²) either	[-	killed which we	ere deem	animals slaughtered or, in the case of game led fit for human consumption in accordanc animal by-products for commercial reason	e with Union legislation until			
	(²) and/or	[-	slaughterhouse ante-mortem i	se and ware inspection	wing parts originating either from animals ere considered fit for slaughter for humal n or bodies and the following parts of ar accordance with Union legislation:	n consumption following an			
			cor	nsumptio	bodies and parts of animals which were n in accordance with Union legislation, bease communicable to humans or animals;				
			(ii) hea	ads of po	ultry;				
			the		kins, including trimmings and splitting there ges and the carpus and metacarpus bor				
			(iv) pig	g bristles;					
			(v) fea	athers;]					
	(²) and/or	[-	Article 1(3)(d)) of Reg	m poultry and lagomorphs slaughtered or ulation (EC) No 853/2004 of the Europe not show any signs of disease communicab	ean Parliament and of the			
	(²) and/or	[-	humans or ani after having b	nimals, ob been con	n did not show any signs of disease com otained from animals that have been slaug isidered fit for slaughter for human cons ocordance with Union legislation;]	ghtered in a slaughterhouse			

Animal by-products to be used for purposes outside the feed chain or for trade samples (2)

				the feed cha	in or for trade samples (2)					
II.	Health information			II.a. Certificate reference No	II.b.					
	(²) and/or	[-		ising from the production of products intend one, greaves and centrifuge or separator slu						
	(²) and/or	[-	longer intended for h	origin, or foodstuffs containing products of animal origin, which are no human consumption for commercial reasons or due to problems of ckaging defects or other defects from which no risk to public or animal						
	(²) and/or	[-	derived products, which	tuffs of animal origin, or feedingstuffs containing animal by-products or ch are no longer intended for feeding for commercial reasons or due to curing or packaging defects or other defects from which no risk to public s;]						
	(²) and/or	[-		ol, feathers, hair, horns, hoof cuts and raw milk originating from live how signs of any disease communicable through that product to humans						
	(²) and/or	[-		parts of such animals, except sea mamma municable to humans or animals;]	ls, which did not show any					
	(²) and/or	[-		animal by-products from aquatic animals originating from establishments or products for human consumption;]						
	(²) and/or	[-		originating from animals which did not so that material to humans or animals:	how any signs of disease					
			(i) shells from	shellfish with soft tissue or flesh;						
			(ii) the following	ng originating from terrestrial animals:						
			— hatche	ery by-products;						
			— eggs;							
			— egg by	y-products, including egg shells;						
			(iii) day-old chi	cks killed for commercial reasons;]						
	(²) and/or	[-	animal by-products fro humans or animals;]	om aquatic or terrestrial invertebrates, other	than species pathogenic to					
	(²) and/or	[-	Category 1 material	ereof of the zoological orders of Rodentia as referred to in Article 8(a)(iii), (iv) an tegory 2 material as referred to in Article 9(a	d (v) of Regulation (EC)					
	(²) and/or	[-		dead animals that did not show clinion that product to humans or animals;]	cal signs of any disease					
II.1.7.		in s	uch a way that they will	of origin or have been preserved in accord not spoil between the time of dispatch and						
(²) (⁶) [II.1.8.										
(²) (⁷)										
either [II.1.8.1.		•	•	gnment come from animals that have bee						

territory or part thereof referred to in point II.1.1, where vaccination programmes against foot-and-mouth

disease are regularly carried out and officially controlled in domestic bovine animals.]]

Animal by-products to be used for purposes outside the feed chain or for trade samples (2)

				the feed cl	hain or for trade samples (²)					
II.	Health inf	ormation		II.a. Certificate reference No	II.b.					
(²) (⁸)										
and/or [II.1.8.2.	The anima	al by-produ	cts in this consi	gnment consist of animal by-products de	erived from offal or deboned					
(²) [II.1.9.	the animal	by-product	y-products described above							
	(²) either	[are derive	ed from other rum	ninants than bovine, ovine or caprine anima	als.]]					
	(²) or	[are derive	ed from bovine, o	vine or caprine animals and does not cont	ain and is not derived from:					
		(²) either	continuously	e and caprine materials other than those reared and slaughtered in a country or r E risk in accordance with Decision 2007/45	region classified as posing a					
		(²) or		d risk material as defined in point 1 of . 2001 of the European Parliament and of th						
			animals slaughte accorda	ically separated meat obtained from bone, except from those animals that were beered in a country or region classified as poince with Commission Decision 2007/453 indigenous BSE case,	orn, continuously reared and osing a negligible BSE risk in					
			animals nervous into the for thos country	by-product or derived product obtained fr which have been killed, after stunning, tissue by means of an elongated rod-s cranial cavity, or by means of gas injected e animals that were born, continuously or region classified as posing a negligible of 2007/453/EC.]]]	by laceration of the central chaped instrument introduced into the cranial cavity, except reared and slaughtered in a					
II.1.10	the animal	by-product	s described abov	e:						
	(²) either		ntain milk or milk imals, other than	products of ovine or caprine animal origin fur animals.]	or is not intended for feed for					
	(²) or			cts of ovine or caprine animal origin and is intended for feed for fanals, and the milk or milk products:						
				e and caprine animals which have been ke llowing conditions are fulfilled:	ept continuously since birth in					
		(i)	classical so	crapie is compulsorily notifiable;						
		(ii)	an awarene	ess, surveillance and monitoring system is	in place for classical scrapie;					
		(iii)		rictions apply to holdings of ovine or cap of TSE or the confirmation of classical scra						
		(iv)	ovine and o	caprine animals affected with classical scra	apie are killed and destroyed;					
		(v)	defined in t Health (OI	g to ovine and caprine animals of meat-arthe Terrestrial Animal Health Code of the NE), of ruminant origin has been banned antry for a period of at least the preceding se	World Organisation for Animal nd effectively enforced in the					
		(b) origin	ate from holdings	s where no official restrictions are imposed	due to a suspicion of TSE;					
			d of the precedi	is where no case of classical scrapie has ng seven years or, following the confirm						

Animal by-products to be used for purposes outside the feed chain or for trade samples (2)

II. Health info	rmation		II.a. Certificate reference No II.b.				
	(²) either	slaughtered carrying at	and caprine animals on the holding have been killed and destroyed or d, except for breeding rams of the ARR/ARR genotype, breeding ewes least one ARR allele and no VRQ allele and other ovine animals least one ARR allele;]				
	(²) or	destroyed, a since the da monitoring, accordance Annex X to	Is in which classical scrapie was confirmed have been killed and and the holding has been subjected for a period of at least two years date of confirmation of the last classical scrapie case to intensified TSE, including testing with negative results for the presence of TSE in e with the laboratory methods set out in point 3.2 of Chapter C of a Regulation (EC) No 999/2001, of all of the following animals which are ge of 18 months, except ovine animals of the ARR/ARR genotype:				
		— animal	ls which have been slaughtered for human consumption; and				
			ls which have died or been killed on the holding but which were not n the framework of a disease eradication campaign.]].				

Notes

Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is required to be filled in only if
 it is a certificate for a commodity to be transited through the European Union; it may be filled in if the certificate is for a
 commodity to be imported into the European Union.
- Box reference I.11: In the case of consignments for trade samples or analyses: indicate the name and address of the
 establishment only.
- Box reference I.11 and I.12: Approval number: the registration number of the establishment or plant, which has been issued by the competent authority.
- Box reference I.12: Place of destination: this box is to be filled in:
 - products for the manufacture of derived products for uses outside the feed chain: only if it is a certificate for a transit commodity. Products in transit may only be stored in free zones, free warehouses and custom warehouses.
 - products for trade samples or analyses: the plant in the European Union indicated in the authorisation of the competent authority where appropriate.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to be provided. In the case of unloading and reloading in the European Union, the consignor must inform the border inspection point of the point of entry into the European Union.
- Box reference I.19: use the appropriate Harmonized System (HS) code under the following headings: 04.01; 04.02; 04.03; 04.04; 04.08; 05.05; 05.06, 05.07; 05.11.91; 05.11.99, 23.01 or 30.01.
- Box reference I.23; for bulk containers, the container number and the seal number (if applicable) must be included.
- Box reference I.25: technical use: any use other than feeding of farmed animals, other than fur animals, and the
 production or manufacturing of pet food.
- Box reference I.25: for the purposes of the certificate, 'technical use' includes use as a trade sample.
- Box reference I.26 and I.27: except for trade samples, which are not sent in transit, fill in according to whether it is a transit or an import certificate.
- Box reference I.28:
 - products for the manufacture of derived products for uses outside the feed chain: Manufacturing plant: provide the veterinary control number of the approved establishment.
 - products for the particular technological studies or analyses: the plant in the European Union indicated in the authorisation of the competent authority where appropriate.
 - Species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or Suidae, Pesca, Mollusca, Crustacea, invertebrates other than Mollusca and Crustacea.

COUNTRY

Animal by-products to be used for purposes outside the feed chain or for trade samples (2)

		the feed cr	nain or for trade samples (2)					
II.	Health information	II.a. Certificate reference No	II.b.					
Part	II:							
(^{1a})	OJ L 300, 14.11.2009, p. 1.							
(1b)	OJ L 54, 26.2.2011, p. 1.							
(²)	Delete as appropriate.							
(^{2a})	OJ L 139, 30.4.2004, p. 55.							
(³)	The name and ISO code number of the exporting country as laid down in:							
_	Part 1 of Annex II to Commission Regulation (El	J) No 206/2010 (OJ L 73, 20.3.2010, p. 1);						
_	Annex I to Commission Regulation (EC) No 798	/2008 (OJ L 226, 23.8.2008, p. 1), and						
_	Annex I to Commission Regulation (EC) No 119	/2009 (OJ L 39, 10.2.2009, p. 12).						
	In addition the ISO code of territories and parts thereof referred to in the Annexes to Regulations (EU) No 206/2010, (EC) No 798/2008 and (EC) No 119/2009 referred to in this note (where applicable for the susceptible species concerned) must be included where applicable.							
(4)	Only for countries from where the game meat intended for human consumption of the same animal species is authorised for importation into the European Union.							
(⁵)	OJ L 303, 18.11.2009, p. 1.							
(⁶)	Supplementary guarantees to be provided where the material of domestic ruminants originated in the territory of a South American or South African country or part thereof from where only maturated and deboned fresh meat of domestic ruminants for human consumption is authorised for exportation to the European Union. The whole masseter muscles of bovine animals, incised in accordance with the requirements of Part B.1 of Chapter I of Section IV of Annex Ito Regulation (EC) No 854/2004 of the European Parliament and of the Council (OJ L 139 30.4.2004, p. 206), are also permitted.							
(⁷)	Only for certain South American countries.							
(8)	Only for certain South American and South Africa	can countries.						
(9)	OJ L 147, 31.5.2001, p. 1.							
(10)	OJ L 172, 30.6.2007, p. 84.							
_	The signature and the stamp must be in a different	ent colour to that of the printing.						
_	Note for the person responsible for the consigniand must accompany the consignment until it r Union.							
Offic	cial veterinarian/Official inspector							
	Name (in capital letters):	Qualification ar	nd title:					
	Date:	Signature:						
	Stamp:							

(5) Chapter 10(A), 10(B), 11 and 12 are replaced by the following:

'CHAPTER 10(A)

Health certificate

For rendered fats not intended for human consumption to be used as feed material, intended for dispatch to or for transit through (²) the European Union

COUNTRY: I.1. Consignor Name I.2. Certificate reference No I.2.a. I.3. Central competent authority

		Name					1.3.	Central compete	ent authority				
		Address					1.4.	Local competent	t authority				
		T-1											
	1.5.	Tel.					1.6.	.6. Person responsible for the load in EU					
+	1.5.	Consignee Name					1.6.	Name	ble for the loa	au III EU			
men		Address						Address					
signı		Address						Address					
con		Postcode						Postcode					
ped		Tel.						Tel.					
Part I : Details of dispatched consignment	1.7.	Country of origin	ISO code	1.8.	Region of origin	Code	1.9.	Country of destination	ISO code		Region of destination	Code	
ails c	l 11	Place of original	gin				112	Place of destination	tion				
: Det			5					Trade of decima					
art I		Name	Name Approval number							Custo	m warehouse		
ď		Address						Name		Appro	val number		
		Name					Address						
		Address											
		Name		Approv	val number		Postcode						
		Address											
	I.13.	Place of loa	ıding				I.14.	Date of departur	re				
	I.15.	Means of tra	ansport				I.16.	Entry BIP in EU					
			•					·					
		Aeroplane [☐ Ship		Railway wa	agon 🗖							
		Road vehic	le 🔲 Othe	r 🗖			l.17.						
		Identificatio	n					_					
		Documentation references											
	I.18.	Description of commodity							I.19. Comn	nodity co	de (HS code)		
								L		1.20.	Quantity		
	I.21.	Temperatur	e of product							1.22.	Number of pac	kages	
		Ambient \square			Chilled 🗆]		Frozen C	3				
	123	Seal/Contai	iner No							124	Type of packa	aina	



1.25.	Commodities certified for:										
	Animal feedingsto	uff 🗆	Manufactu	Manufacture of petfood \square Technical use \square							
1.26.	For transit throug	h EU to third country	у 🗆	I.27. For import or admission into EU							
	Third country	ISO co	ode								
1.28.	Identification of th	ne commodities	Approval number	of establishments							
	Species entific name)	Nature of commodity	Manufacturing plant	Number of packages	Net weight	Batch number					

	II.	Health information	on		II.a. Certificate reference No	II.b.							
		the European Pa	rliame	nt and of the C	declare that I have read and understood Regouncil (1a), and in particular Article 10 thereof Chapter II of Annex XIV thereto, and certify the	, and Commission Regulation							
_	II.1.												
icatio	II.2.												
II.2. consist of rendered fats not intended for human consumption; II.3. have been prepared and stored in a plant approved and supervised by the competent authority in accordance Article 24 of Regulation (EC) No 1069/2009 or in accordance with Article 4(2) of Regulation (EC) No 853/2 the European Parliament and of the Council (3), in order to kill pathogenic agents;													
g.	II.4.	have been prepared exclusively with the following animal by-products:											
		(²) either	[-	animals killed	carcases and parts of animals slaughtered or, in the case of game, bodies or parts of animals killed, and which are fit for human consumption in accordance with Union legislation, but are not intended for human consumption for commercial reasons;]								
		(²) and/or	[-	slaughtered in consumption f	carcases and the following parts originating either from animals that have been slaughtered in a slaughterhouse and were considered fit for slaughter for human consumption following an ante-mortem inspection or bodies and the following parts of animals from game killed for human consumption in accordance with Union legislation:								
				cor	 carcases or bodies and parts of animals which are rejected as unfit for human consumption in accordance with Union legislation, but which did not show any signs of disease communicable to humans or animals; 								
				(ii) heads of poultry;									
				inc	es and skins, including trimmings and split luding the phalanges and the carpus and m tatarsus bones;								
				(iv) pig	bristles;								
				(v) fea	thers;]								
		(²) and/or	[-	humans or ani after having b	als which did not show any signs of disease on mals, obtained from animals that have been sl een considered fit for slaughter for human co tion in accordance with Union legislation;]	aughtered in a slaughterhouse							
		(²) and/or	[-		oducts arising from the production of pr including degreased bone, greaves and centri g;]								
		(²) and/or	[-	products of animal origin, or foodstuffs containing products of animal origin, which are longer intended for human consumption for commercial reasons or due to problems manufacturing or packaging defects or other defects from which no risk to public or animhealth arise;]									
		(²) and/or	[-	or derived pro due to problen	eedingstuffs of animal origin, or feedingstuffs ducts, which are no longer intended for feedi ns of manufacturing or packaging defects or of mal health arises;]	ng for commercial reasons or							
		(²) and/or	[-		a, wool, feathers, hair, horns, hoof cuts and lid not show signs of any disease communi mals;]								

				used as feed material
II.	Health informa	ation	II.a. Certificate reference No	II.b.
	(²) and/or	[-	aquatic animals, and parts of such animals, except sea ma signs of diseases communicable to humans or animals;]	mmals, which did not show any
	(²) and/or	[-	animal by-products from aquatic animals originating from anufacturing products for human consumption;]	rom plants or establishments
	(²) and/or	[-	the following material originating from animals which did communicable through that material to humans or animals:	not show any signs of disease
			(i) shells from shellfish with soft tissue or flesh;	
			(ii) the following originating from terrestrial animals:	
			 hatchery by-products, 	
			— eggs,	
			 egg by-products, including egg shells; 	
			(iii) day-old chicks killed for commercial reasons;]	
II.5.	(²) either	[-	in the case of material of porcine origin, come from a cou country free from foot-and-mouth disease for the period of free from classical swine fever and African swine fever f 12 months;]	f the preceding 24 months and
	(²) and/or	[-	in the case of material of poultry origin, come from a concountry free from Newcastle disease and avian influenza 6 months;]	
	(²) and/or	[-	in the case of material of ruminant origin, come from a co- country free from foot-and-mouth disease for the period of free from rinderpest for the period of the preceding 12 month	f the preceding 24 months and
	(²) and/or	[-	where there has been an outbreak of one of the diseases the relevant period referred to in point II.5, and where the susceptible species, have been subjected to a heat tre 30 minutes or at least 90 °C for at least 15 minutes, and	e rendered fats derived from a
			details of the critical control points are recorded and reoperator or their representative and, as necessary, the control of the plant; the information must include temperature and, as appropriate, the absolute time, pressurand fat recycling rate.]	ompetent authority can monitor ude the particle size, critical
II.6.			t animals, were purified in such way that the maximum leved 0,15 % in weight;	els of remaining total insoluble
II.7.	the rendered fa	ıts:		
		(a)	have been subjected to processing in accordance with the Chapter II of Annex X to Regulation (EU) No 142/2011, or Section XII of Annex III to Regulation (EC) No 853/2004, in and	a treatment in accordance with
	(²) either	[(b)	are packaged in new containers or in containers that have necessary for the prevention of contamination, and all prevent their contamination;]	
	(²) or	[(b)	where bulk transport is intended, the pipe, pumps and be container or bulk road tanker used in the transportal manufacturing plant either directly on to the ship or into shave been checked under the responsibility of the compectean before use;]	ion of the product from the hore tanks or directly to plants
	and which bear	· labels ii	ndicating 'NOT FOR HUMAN CONSUMPTION';	

										used as feed material		
II.	Health info	rmation		II.a	i. C	Certificate r	eference N	0	ا	I.b.		
(²) [II.8.	the rendered	ed fats described above										
	(²) either	[is derived	is derived from other ruminants than bovine, ovine or caprine animals.]]									
	(²) or	[is derived	is derived from bovine, ovine or caprine animals and does not contain and is not derived from:									
		(²) either	er [bovine, ovine and caprine materials other than those derived from animals be continuously reared and slaughtered in a country or region classified as posing a negli BSE risk in accordance with Decision 2007/453/EC.]]									
		(²) or	[(a)					I in point 1 i		nnex V to Regulation (EC) e Council (4);		
			(b)	animals, slaughter accordar	exc red nce	cept from in a count	those anim ry or region mission De	als that were classified as	e bo s po:	s of bovine, ovine or caprine rn, continuously reared and sing a negligible BSE risk in /EC (⁵), in which there has		
			(c)	animals nervous into the of for those country of	whitiss cran e ar or re	ich have b sue by me nial cavity, on nimals that	een killed, ans of an or by means were borr ified as pos	after stunning after	ng, od-sh eted i	om bovine, ovine or caprine by laceration of the central laped instrument introduced nto the cranial cavity, except eared and slaughtered in a BSE risk in accordance with		
II.9.	the rendered	d fats descri	ibed above:									
	(²) either	-	contain milk imals, other	•			ne or caprir	ie animal oriç	gin o	or is not intended for feed for		
	(²) or		milk or milk ther than fu						d is i	ntended for feed for farmed		
		(a)					e animals v onditions are		een l	kept continuously since birth		
			(i)	classical	scr	apie is con	npulsorily no	otifiable;				
			(ii)	an awar scrapie;	ene	ess, surveil	lance and	monitoring s	syste	em is in place for classical		
			(iii)					s of ovine or on of classica		rine animals in the case of a rapie;		
			(iv)	ovine ar		caprine ai	nimals affe	cted with cl	lassi	cal scrapie are killed and		
			(v)	defined i Animal I	in th Hea	he Terrest alth (OIE),	rial Animal of rumina	Health Code nt origin has	e of s be	nd-bone meal or greaves, as the World Organisation for een banned and effectively least the preceding seven		
		(b)	originate f TSE;	rom holdi	ings	where no	official res	strictions are	imp	osed due to a suspicion of		
		(c)								s been diagnosed during the of classical scrapie:		

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

II. Health information	II.a. Certificate reference No	II.b.			
(²) either	[all ovine and caprine animals on the holding have slaughtered, except for breeding rams of the pewes carrying at least one ARR allele and not animals carrying at least one ARR allele;]	ARR/ARR genotype, breeding			
(²) or	[all animals in which classical scrapie was confirmed have been killed an destroyed, and the holding has been subjected for period of at least two year since the date of confirmation of the last classical scrapie case to intensifie TSE monitoring, including testing with negative results for the presence of TS in accordance with the laboratory methods set out in point 3.2 of Chapter C of Annex X to Regulation (EC) No 999/2001, of all of the following animals which are over the age of 18 months, except ovine animals of the ARR/AR genotype:				
	animals which have been slaughtered for hi	uman consumption; and			
	 animals which have died or been killed on skilled in the framework of a disease eradical 	· ·			

Notes

Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is required to be filled in only if
 it is a certificate for a commodity to be transited through the European Union; it may be filled in if the certificate is for a
 commodity to be imported into the European Union.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for a transit commodity. Products
 in transit may only be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in the case of unloading and reloading in the European Union.
- Box reference I.19: use the appropriate HS code: 04.05; 15.01; 15.02; 15.03; 15.04; 15.05; 15.06; 15.16.10 or 15.18.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) must be included.
- Box reference I.25: technical use: any use other than feeding of farmed animals, other than fur animals or pet animals, and the production or manufacturing of pet food.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28:
- Species: select from the following: Ruminantia, other than Ruminantia
- Manufacturing plant: provide the registration number of the treatment/processing establishment.

Part II:

- (1a) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.
- (3) OJ L 139, 30.4.2004, p. 55.

COUNTRY

II.	Health information	II.a. Certificate reference No		II.b.						
(4)	OJ L 147, 31.5.2001, p. 1.									
(5)	OJ L 172, 30.6.2007, p. 84.									
_	The signature and the stamp must be in a different colour to that of the printing.									
_	 Note for the person responsible for the consignment in the European Union: this certificate is only for veterinary purposes and must accompany the consignment until it reaches the border inspection post of the European Union. 									
Offic	cial veterinarian/Official inspector									
	Name (in capital letters):		Qualification a	and title:						
	Date:		Signature:							
	Stamp:									

Veterinary certificate to EU

CHAPTER 10(B)

Health certificate

For rendered fats not intended for human consumption to be used for certain purposes outside the feed chain, intended for dispatch to or for transit through (2) the European Union

	1.1.	Consignor	1.2.	Certificate reference No	I.2.a.				
		Name	I.3. Central competent authority						
		Address	1.4.	I.4. Local competent authority					
		Tel.							
	1.5.	Consignee	1.6.	Person responsible for the	e load in EU				
ent		Name		Name					
gnm		Address		Address					
Part I : Details of dispatched consignment		Destands		Dantanda					
o pe		Postcode		Postcode					
tch		Tel.		Tel.					
lispa	1.7.	Country ISO code I.8. Region of Code of origin	1.9.	Country of ISO destination code	I.10. Region of Code destination				
of c									
tails	1.11.	Place of origin	1.12.	Place of destination					
: De									
art		Name Approval number			Custom warehouse				
۵		Address		Name	Approval number				
		Name Approval number		Address					
		Address							
		Name Approval number		Postcode					
		Address							
	1.13.	Place of loading	1.14.	Date of departure					
	I.15.	Means of transport	I.16.	Entry BIP in EU					
		,		•					
		Aeroplane 🗆 Ship 🗖 Railway wagon 🗖							
		Road vehicle Other	1.17.						
		Identification							
		Documentation references							
	I.18.	Description of commodity		I.19. Co	ommodity code (HS code)				
					I.20. Quantity				
	I.21.	Temperature of product			I.22. Number of packages				
		Ambient Chilled Chilled		Frozen					
	1.23.	Seal/Container No			I.24. Type of packaging				



1.25.	Commodities certifi	ed for:				
	Technical use					
1.26.	For transit through	EU to third country		I.27. For imp	ort or admission into EU	
	Third country	ISO code				
1.28.	Identification of the		roval number	of establishme	nts	
(\$	Species scientific name)	Manufacturing plant	Number of	packages	Net weight	Batch number

Rendered fats not intended for human consumption for certain purposes outside the feed chain

								certaiı	n purp	oses ou	tside the fee	ed chain	
	II.	Health information	on		II.a.	Certificate	reference N	lo		II.b.			
		I, the undersigned European Parliar Regulation (EU) N described above:	ment a No 142	and of the Cou	uncil (1	^{la}), and in _l	particular A	articles 8, 9	and	10 there	of, and Con	nmission	
_	II.1.	consist of rendere	ed fats	not intended fo	r huma	an consumpt	ion that sat	isfy the heal	th requ	uirements	s below;		
icatio	II.2.	have been prepar	ed ex	clusively with th	e follov	wing animal	by-products	3:					
Part II: Certification	(²) [II.2.1.	in the case of materials destined for the production of renewable fuels referred to in point L of Section 2 of Chapter IV of Annex IV to Regulation (EU) No 142/2011, biodiesel or oleochemical products, animal by-products referred to in Articles 8, 9 and 10 of Regulation (EC) No 1069/2009;]											
Ä	(²) [II.2.2.	in the case of materials destined for the production of renewable fuels referred to in point J of Section 2 of Chapter IV of Annex IV to Regulation (EU) No 142/2011, the materials have been prepared exclusively from animal by-products referred to in Articles 9 and 10 of Regulation (EC) No 1069/2009;]											
	(²) [II.2.3.		in the case of materials destined for purposes other than cosmetics, pharmaceuticals or medical devices, the materials have been prepared exclusively from:									ces, the	
		(²) either	[-	animal by-preexceeding the									
		(²) and/or	[-	products of animal origin which have been declared unfit for human consumption due to the presence of foreign bodies in those products;]								ue to the	
		(²) and/or	[-	animals and parts of animals, other than those referred to in Articles 8 and 10 of Regulation (EC) No 1069/2009, that died other than being slaughtered or killed for human consumption including animals killed for disease control purposes;									
		(²) and/or	[-	carcasses and animals killed legislation, but	d, and	which are	fit for hu	man consu	mption	in acc	ordance wit		
		(²) and/or	[-	carcasses and in a slaughter an ante-morte human consul	house a	and were co	onsidered fi dies and th	t for slaught e following p	er for l	human c	onsumption t	following	
				consun	nption i		ce with Uni	on legislatio			l as unfit for d not show a		
				(ii) heads	of poult	try;							
				(iii) hides a the pha	and skii alanges	ns, including and the car	g trimmings rpus and m	and splittin etacarpus bo	ng there	eof, horn arsus and	is and feet, i d metatarsus	including bones;	
				(iv) pig bris	stles;								
				(v) feather	rs;]								
		(²) and/or	[-	blood of anim humans or an after having b mortem inspec	imals o been co	obtained fror onsidered fi	n animals t for slaugh	that have be nter for hum	een sla nan coi	ughtered	l in a slaugh	terhouse	
		(²) and/or	[-	animal by-proconsumption, milk processing	includi								

Rendered fats not intended for human consumption for certain purposes outside the feed chain

						certain pui	poses outside the feed chain
II.	Health informati	on		II.a.	Certificate reference No		II.b.
	(²) and/or	[-	longer intended	d for	human consumption for co	mmercial re	of animal origin, which are no easons or due to problems of hich no risk to public or animal
	or derived pro-				which are no longer intended ufacturing or packaging defe	d for feeding	containing animal by-products for commercial reasons or due defects from which no risk to
	(²) and/or	[-		id no	t show signs of any diseas		raw milk originating from live icable through that product to
	(²) and/or	[-	•		d parts of such animals, exce mmunicable to humans or an	•	nmals, which did not show any
	(²) and/or	[-			from aquatic animals or cts for human consumption;]		om plants or establishments
	(²) and/or	[-			al originating from animals of that material to humans of		ot show any signs of disease
			(i) shells from	om sl	nellfish with soft tissue or fles	h;	
			(ii) the follow	wing	originating from terrestrial an	mals:	
			— ha	tcher	y by-products,		
			— eg	gs,			
			— eg	g by-	products, including egg shells	S,	
			(iii) day-old	chick	s killed for commercial reaso	ns;]	
	(²) and/or	[-	aquatic and ter	restri	al invertebrates other than sp	ecies patho	genic to humans or animals;]
	(²) and/or	[-	Category 1 m	ateria	I as referred to in Article	8(a)(iii), (iv	entia and Lagomorpha, except) and (v) of Regulation (EC) e 9(a) to (g) of that Regulation;]
	(²) and/or	[-					r originating from dead animals ugh that product to humans or
	(²) and/or	[-	that material to were consider	hum ed fi	ans or animals, which were s	slaughtered	disease communicable through in a slaughterhouse and which on following an ante-mortem
(²) [II.2.4.	in the case of m cosmetics, pharm					on of orgar	ic fertilisers or soil improvers,
	(²) either	[-			al as defined in Article 3(1)(t and of the Council (^{2b});]	(g) of Regul	ation (EC) No 999/2001 of the
	(²) and/or	[-			rts of dead animals contain gulation (EC) No 999/2001 at		ed risk material as defined in disposal;]
	(²) and/or	[-		t as	lefined in Article 1(2)(d) of Co		which have been submitted to tive 96/22/EC (2c) or Article 2(b)

Rendered fats not intended for human consumption for certain purposes outside the feed chain

						certain pui	poses outside the feed chain
II.	Health i	nformation		II.a. Certificate re	eference No		II.b.
	(²) and/o	r [-	contaminants the permitted	listed in Group B(3)	of Annex I to Di Union legislation	irective 96/	bstances and environmental 23/EC, if such residues exceed absence thereof, by legislation
II.3.	the rend	ered fats:					
	'n						(indicate the processing 011, in order to kill pathogenic
				ipment to the Euro tration of at least 250			riheptanoate (GTH), so that a fat is achieved,
	` '	n the case of r emoved,	endered fats of	ruminant origin, ins	oluble impurities	s in excess	of 0,15% in weight have been
	(d) h	nave been trans	sported under c	onditions which prev	ent their contam	ination, and	d
	(e) b	ear labels on t	the packaging o	r container indicating	, "NOT FOR HUI	MAN OR A	NIMAL CONSUMPTION";
(²) [II.4.		se of materials ered fats desci		ganic fertilisers, cos	metics, pharmac	euticals, m	edical devices or soil improvers
	(²) either	are derive	ed from other ru	minants than bovine	, ovine or caprine	e animals.]	
	(²) or	[are derive	ed from bovine,	ovine or caprine ani	mals and does n	ot contain a	and is not derived from:
		(²) either	continuously	•	red in a country	or region of	derived from animals born, classified as posing a negligible
		(²) or		ed risk material as 9/2001 of the Europe			Annex V to Regulation (EC) ouncil (³);
			animal slaugh accord	s, except from the tered in a country	ose animals that or region class	at were b ified as po	s of bovine, ovine or caprine orn, continuously reared and osing a negligible BSE risk in (4), in which there has been no
			which means by me born, c	have been killed, aft of an elongated ro ans of gas injected	ter stunning, by I d-shaped instrur into the cranial o and slaughtered	laceration of ment introd cavity, exce in a countr	ovine, ovine or caprine animals of the central nervous tissue by luced into the cranial cavity, or ept for those animals that were y or region classified as posing 7/453/EC.]]]

Notes

Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is required to be filled in only if it
 is a certificate for a commodity to be transited through the European Union; it may be filled in if the certificate is for a
 commodity to be imported into the European Union.
- Box reference I.11 and I.12: Approval number: the registration number of the establishment or plant, which has been issued by the competent authority.

COUNTRY

Rendered fats not intended for human consumption for certain purposes outside the feed chain

				certain pu	rposes outside the feed chain		
II.	Health information	II.a.	Certificate reference No		II.b.		
_	Box reference I.12: Place of destination: this transit may only be stored in free zones, free				a transit commodity. Products in		
_	Box reference I.15: Registration number (rail- to be provided. In the case of unloading a inspection post of the point of entry into the E	nd rele	pading in the European U				
_	Box I.19: use the appropriate Harmonized S 15.04; 15.05; 15.06; 15.16 or 15.18.	System	n (HS) code under the foll	lowing heading	gs: 04.05; 15.01, 15.02; 15.03;		
_	Box reference I.23: for bulk containers, the container number and the seal number (if applicable) must be included.						
_	Box reference I.25: technical use: any use of the production or manufacturing of pet food.	her tha	an feeding of farmed anima	als, other than	fur animals or pet animals, and		
_	Box reference I.26 and I.27: fill in according to	o whet	her it is a transit or an imp	ort certificate.			
-	Box reference I.28:						
	Species: select from the following: Ruminanti	a, othe	er than Ruminantia				
	Manufacturing plant: provide the registration i	numbe	er of the treatment/process	ing establishm	ent.		
Part	II:						
(^{1a})	OJ L 300, 14.11.2009, p. 1.						
(1b)	OJ L 54, 26.2.2011, p. 1.						
(2)	Delete as appropriate.						
(^{2a})	OJ L 125, 23.5.1996, p. 10.						
(^{2b})	OJ L 147, 31.5.2001, p. 1.						
(^{2c})	OJ L 125, 23.5.1996, p. 3.						
(3)	OJ L 147, 31.5.2001, p. 1.						
(4)	OJ L 172, 30.6.2007, p. 84.						
_	The signature and the stamp must be in a diff	erent	colour to that of the printing	g.			
_	Note for the person responsible for the consignment in the European Union: this certificate is only for veterinary purpose and must accompany the consignment until it reaches the border inspection post of the point of entry into the Europea Union.						
Offic	ial veterinarian/Official inspector						
	Name (in capital letters):			Qualification a	and title:		
	Date:			Signature:			
	Stamp:						

Veterinary certificate to EU

CHAPTER 11

Health certificate

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

	l.1.	Consignor	1.2.	Certificate reference No		I.2.a.		
		Name	1.3.	Central competent author	ity			
		Address	1.4.	Local competent authority	•			
		T-1						
		Tel.						
	1.5.	Consignee	1.6.	Person responsible for the	e load in	EU		
ent		Name		Name				
ignr		Address		Address				
Part I : Details of dispatched consignment		Postcode		Postcode				
Dec		Tel.		Tel.				
atcl	1.7.	Country ISO code I.8. Region of Code	1.9.	Country of ISO	I 1	0. Region of	Code	
disp		of origin origin	1.0.	destination code		destination		
s of								
etail	l.11.	Place of origin	1.12.	Place of destination				
.: O								
art		Name Approval number			С	ustom warehouse		
<u> </u>		Address		Name	А	pproval number		
		Name Approval number		Address				
		Address						
		Name Approval number		Postcode				
		Address						
	I.13.	Place of loading	I.14.	Date of departure				
	1.15.	Means of transport	I.16.	Entry BIP in EU				
		,		•				
		Aeroplane ☐ Ship ☐ Railway wagon ☐						
		Road vehicle Other	1.17.					
		Identification						
		Documentation references						
	I.18.	Description of commodity		I.19. Co	ommodit	ty code (HS code)		
					1.3	20. Quantity		
	I.21.	Temperature of product			1.3	22. Number of pac	kages	
		Ambient ☐ Chilled ☐		Frozen 🗖				
	1.23.	Seal/Container No			1.	24. Type of packa	ging	



1.25.	Commodities certific	ed for:				
	Animal feedingstuff		Manufactu	re of petfood \square	Technical ι	se 🗖
1.26.	For transit through I	EU to third country		I.27. For import or a	admission into EU	
	Third country	ISO code				
1.28.	Identification of the		oval number	of establishments		
(5	Species Scientific name)	Manufacturing plant	Number of	packages N	let weight	Batch number

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

,							chain
	II.	Health information	n		II.a.	Certificate reference No	II.b.
		the European Pa	arliame 11 (^{1b}),	nt and of the	e Council	e that I have read and understood Reg (^{1a}), and in particular Article 10 thereof, oter I of Annex XIV thereto, and certify	and Commission Regulation
	II.1.	consists of gelati	ine/colla	agen (²) that	satisfy the	health requirements below;	
ation	II.2.	consist exclusive	ely of ge	elatine/collag	jen (²) not	intended for human consumption;	
Part II: Certification	II.3.					roved and supervised by the competen order to kill pathogenic agents;	t authority in accordance with
art II:	II.4.	has been prepar	ed excl	usively with	the followi	ng animal by-products:	
a		(²) either	[-	animals ki	lled, and	of animals slaughtered or, in the case which are fit for human consumption t intended for human consumption for co	n in accordance with Union
		(²) and/or	[-	slaughtered consumption	d in a sl on followin	following parts originating either from aughterhouse and were considered g an ante-mortem inspection or bodie Iled for human consumption in accordar	fit for slaughter for human es and the following parts of
				con	sumption	podies and parts of animals which are in accordance with Union legislation, se communicable to humans or animals	but which did not show any
(ii) heads of poultry;					try;		
	the					ns, including trimmings and splitting the s and the carpus and metacarpus bo	
				(iv) pig	bristles;		
				(v) feat	thers;]		
		(²) and/or	[-		n, includir	arising from the production of prong degreased bone, greaves and centrif	
		(²) and/or	[-	ionger inte	nded for h ing or pac	rigin, or foodstuffs containing products on numan consumption for commercial re- kaging defects or other defects from wh	asons or due to problems of
		(²) and/or	[-	or derived due to prob	products, plems of m	tuffs of animal origin, or feedingstuffs of which are no longer intended for feeding anufacturing or packaging defects or oth alth arises;]	ng for commercial reasons or
		(²) and/or	[-			parts of such animals, except sea mam nmunicable to humans or animals;]	mals, which did not show any
		(²) and/or	[-			from aquatic animals originating fro cts for human consumption;]	m plants or establishments
II.5. the gelatine/collagen (²):							
			(a)	and in par	ticular wr	ged, stored and transported under sa apping and packaging took place in ed under Union legislation were used.	

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

					chain
II.	Health info	rmation	II.a. Certifica	ate reference No	II.b.
			Wrappings and packages GELATINE/COLLAGEN(2) SU	containing gelatine/collag	
	(²) either	[(b)	Category 3 material was subje more rinses, involving pH ad	ected to a treatment with acid djustment, extraction by hea	at ensured that unprocessed d or alkali, followed by one or ting one or several times in nd sterilisation, in order to kill
	(²) or	[(b)	Category 3 material was subje	cted to a treatment involving	at ensured that unprocessed washing, pH adjustment using and extrusion, in order to kill
(²) [II.6.	in the case	e of gelatine/c	llagen (²) from materials other	than hides and skins	
	(²) either	[is derived fr	m other ruminants than bovine	e, ovine or caprine animals.]]	
	(²) or	[is derived fr	m bovine, ovine or caprine ani	mals and does not contain and	d is not derived from:
		(²) either		ntered in a country or region c	derived from animals born, lassified as posing a negligible
		(²) or		as defined in point 1 of a copean Parliament and of the 0	Annex V to Regulation (EC) Council (3);
			animals, except from slaughtered in a coun	those animals that were bo try or region classified as po nission Decision 2007/453/EC	s of bovine, ovine or caprine orn, continuously reared and osing a negligible BSE risk in C (4), in which there has been
			animals which have be tissue by means of an cavity, or by means of that were born, conti	en killed, after stunning, by lar elongated rod-shaped instrum gas injected into the cranial ca nuously reared and slaught	om bovine, ovine or caprine ceration of the central nervous tent introduced into the cranial avity, except for those animals ered in a country or region accordance with Decision
II.7.	in the case	e of gelatine/c	llagen (²) from materials other	than hides and skins described	d above:
	(²) either		ntain milk or milk products of cals, other than fur animals.]	ovine or caprine animal origin	or is not intended for feed for
	(²) or		or milk products of ovine or rthan fur animals, and the milk		intended for feed for farmed
			rived from ovine and caprine a the following conditions are ful		nuously since birth in a country
		(i)	classical scrapie is com	npulsorily notifiable;	
		(ii)	an awareness, surveilla	ance and monitoring system is	in place for classical scrapie;
		(iii)		ly to holdings of ovine or cap confirmation of classical scra	orine animals in the case of a pie;
					"

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

					chain
II.	Health information	า	II.a.	Certificate reference No	II.b.
		(iv) ovi	ne and ca	aprine animals affected with classical scr	apie are killed and destroyed;
		def He	ined in thalth (OIE	to ovine and caprine animals of meat- e Terrestrial Animal Health Code of the), of ruminant origin has been banned ry for a period of at least the preceding s	World Organisation for Animal and effectively enforced in the
	(b)	originate from ho	ldings wh	nere no official restrictions are imposed d	ue to a suspicion of TSE;
	(c)			nere no case of classical scrapie has bee ars or, following the confirmation of a ca	
		sla car	ughtered, rying at	nd caprine animals on the holding have except for breeding rams of the ARR/, least one ARR allele and no VRQ a east one ARR allele;]	ARR genotype, breeding ewes
		des sin mo acc Ani	etroyed, a ce the da nitoring, cordance nex X to I	in which classical scrapie was con and the holding has been subjected for the of confirmation of the last classical s including testing with negative results with the laboratory methods set out Regulation (EC) No 999/2001, of all of t of 18 months, except ovine animals of t	a period of at least two years crapie case to intensified TSE for the presence of TSE in in point 3.2 of Chapter C of the following animals which are
		_	animal	s which have been slaughtered for huma	n consumption; and
		_		s which have died or been killed on the tramework of a disease eradication	•

Notes

Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a
 certificate for a commodity to be transited through the European Union; it may be filled in if the certificate is for a
 commodity to be imported into the European Union.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. Products in transit may only be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to be provided. In the case of unloading and reloading in the European Union, the consignor must inform the border inspection post of the point of entry into the European Union.
- Box I.19: use the appropriate Harmonized System (HS) code under the following headings: 35.03 or 35.04.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) must be included.
- Box reference I.25: technical use: any use other than feeding of farmed animals, other than fur animals, and the
 production or manufacturing of pet food.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: Species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or Suidae, Pesca.

COUNTRY

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

II.	Health information	II.a.	Certificate reference N	lo	II.b.					
Part	Part II:									
(^{1a})	OJ L 300, 14.11.2009, p. 1.									
(1b)	OJ L 54, 26.2.2011, p. 1.									
(2)	Delete as appropriate.									
(3)	OJ L 147, 31.5.2001, p. 1.									
(4)	OJ L 172, 30.6.2007, p. 84.									
-	The signature and the stamp must be in a d	lifferent c	olour to that of the printir	ng.						
_	Note for the person responsible for the con and must accompany the consignment until				s only for veterinary purposes					
Offic	cial veterinarian/Official inspector									
	Name (in capital letters):			Qualification a	nd title:					
	Date:			Signature:						
	Stamp:									
L										

Veterinary certificate to EU

CHAPTER 12

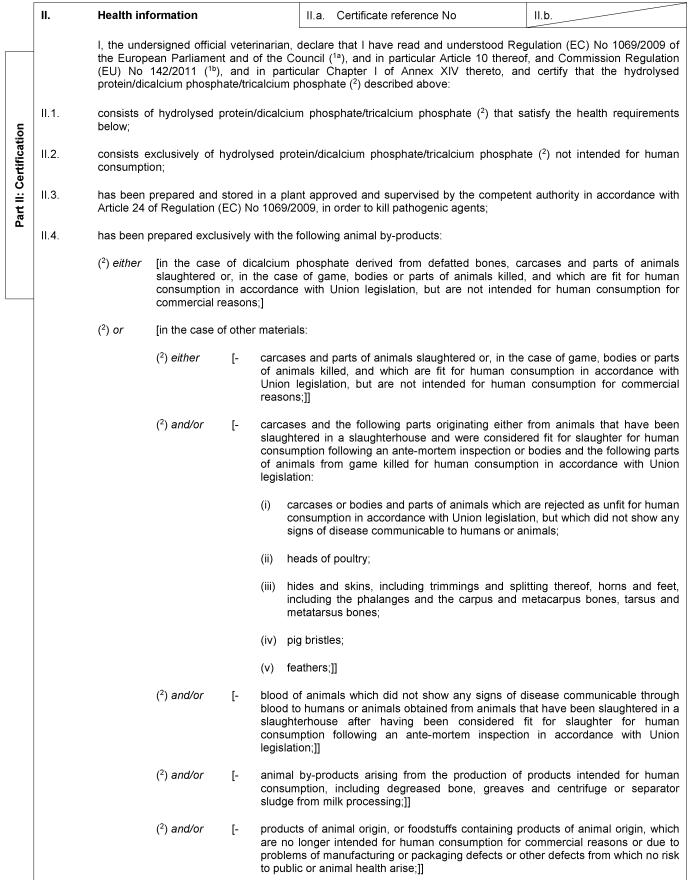
Health certificate

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

	l.1.	Consignor	1.2.	Certificate reference No		I.2.a.		
		Name	1.3.	Central competent author	ity			
		Address	1.4.	Local competent authority	•			
		T-1						
		Tel.						
	1.5.	Consignee	1.6.	Person responsible for the	e load in	EU		
ent		Name		Name				
ignr		Address		Address				
Part I : Details of dispatched consignment		Postcode		Postcode				
pet		Tel.		Tel.				
atcl	1.7.	Country ISO code I.8. Region of Code	1.9.	Country of ISO	I 1	0. Region of	Code	
disp		of origin origin	1.0.	destination code		destination		
s of								
etail	l.11.	Place of origin	1.12.	Place of destination				
.: O								
art		Name Approval number			С	ustom warehouse		
<u> </u>		Address		Name	А	pproval number		
		Name Approval number		Address				
		Address						
		Name Approval number		Postcode				
		Address						
	I.13.	Place of loading	I.14.	Date of departure				
	1.15.	Means of transport	I.16.	Entry BIP in EU				
		,		•				
		Aeroplane ☐ Ship ☐ Railway wagon ☐						
		Road vehicle Other	1.17.					
		Identification						
		Documentation references						
	I.18.	Description of commodity		I.19. Co	ommodit	ty code (HS code)		
					1.3	20. Quantity		
	I.21.	Temperature of product			1.3	22. Number of pac	kages	
		Ambient ☐ Chilled ☐		Frozen 🗖				
	1.23.	Seal/Container No			1.	24. Type of packa	ging	



1.25.	Commodities cer	tified for:				
	Animal feedingst	uff 🗖	Manufactu	re of petfood \square	Technical us	е 🗆
1.26.	For transit throug	h EU to third count	try 🗖	I.27. For import or	admission into EU	
	Third country	ISO c	code			
1.28.	Identification of the	he commodities	Approval number	of establishments		
(Sci	Species entific name)	Nature of commodity	Manufacturing plant	Number of packages	Net weight	Batch number



COUNTRY

					used as feed material or for uses outside the feed chain
II.	Health inf	formati	on		II.a. Certificate reference No II.b.
		(²) and	d/or	[-	petfood and feedingstuffs of animal origin, or feedingstuffs containing animal by- products or derived products, which are no longer intended for feeding for commercial reasons or due to problems of manufacturing or packaging defects or other defects from which no risk to public or animal health arises;]]
		(²) and	d/or	[-	blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk originating from live animals that did not show signs of any disease communicable through that product to humans or animals;]]
		(²) and	d/or	[-	aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of diseases communicable to humans or animals;]]
		(²) and	d/or	[-	animal by-products from aquatic animals originating from plants or establishments manufacturing products for human consumption;]]
		(²) and	d/or	[-	the following material originating from animals which did not show any signs of disease communicable through that material to humans or animals:
					(i) shells from shellfish with soft tissue or flesh;
					(ii) the following originating from terrestrial animals:
					 hatchery by-products,
					— eggs,
					egg by-products, including egg shells;
					(iii) day-old chicks killed for commercial reasons;]]
II.5.	the hydrol	ysed pr	otein/dic	alciun	m phosphate/tricalcium phosphate (²):
		(a)	CONSI particul	JMPT ar the	ed and packaged in packaging which bear labels indicating 'NOT FOR HUMAN TION' and was stored and transported under satisfactory hygiene conditions, and in the wrapping and packaging took place in a dedicated room, and only preservatives ander Union legislation were used; and
	(²) either	[(b)			of hydrolysed protein, was produced by a process involving appropriate measures to ntamination of raw Category 3 material.
			produc	ed in a	of hydrolysed proteins entirely or partly derived from ruminants hides and skins, was a processing plant dedicated only to hydrolysed proteins production, using a process e preparation of the raw Category 3 material by brining, liming and intensive washing
			(i)	temp	exposure of the material to a pH of more than 11 for more than 3 hours at a perature of more than 80 °C and subsequently by heat treatment at a temperature of e than 140 °C for 30 minutes at more than 3,6 bar; or
			(ii)		exposure of the material to a pH of 1 to 2, followed by a pH of more than 11, followed a heat treatment at a temperature of more than 140 °C for 30 minutes at 3 bar.]
	(²) or	[(b)	in the c	ase o	of dicalcium phosphate, was produced by a process that:
			(i)	and t	ures that all Category 3 bone-material is finely crushed and degreased with hot water treated with dilute hydrochloric acid (at a minimum concentration of 4 % and a pH of than 1,5) over a period of at least two days,
			(ii)		owed by a treatment of the obtained phosphoric liquor with lime, resulting in a cipitate of dicalcium phosphate at pH 4 to 7, and

					used as feed material or fo	r uses outside the feed chain			
II.	Health inf	formati	on		II.a. Certificate reference No	II.b.			
			(iii)		es this precipitate, with an inlet temperature of between 30 °C and 65 °C.]	of 65 °C to 325 °C and an end			
	(²) or	[(b)	in the	case of tricalci	um phosphate, was produced by a process e	nsuring:			
			(i)		gory 3 bone-material is finely crushed and dec chips less than 14 mm),	greased in counter-flow with hot			
			(ii)	the continuo	us cooking with steam at 145 °C during 30 mi	nutes at 4 bars,			
			(iii)	the separation	on of the protein broth from the hydroxyapa n, and	atite (tricalcium phosphate) by			
			(iv)	the granulati 200 °C.]	on of the tricalcium phosphate after drying	in a fluidised bed with air at			
(²) [II.6.	the hydrol	ysed pr	protein/dicalcium phosphate/tricalcium phosphate (²) described above						
	(²) either	[is de	[is derived from other ruminants than bovine, ovine or caprine animals.]]						
	(²) or	[is de	rived fro	ed from bovine, ovine or caprine animals and does not contain and is not derived from:					
		(²) eiti	her	er [bovine, ovine and caprine materials other than those derived from continuously reared and slaughtered in a country or region classifie negligible BSE risk in accordance with Decision 2007/453/EC.]]					
		(²) or			ed risk material as defined in point 1 of d/2001 of the European Parliament and of the				
				animals slaught accord	nically separated meat obtained from bone s, except from those animals that were be ered in a country or region classified as p ance with Commission Decision 2007/453/Edgenous BSE case,	orn, continuously reared and osing a negligible BSE risk in			
				animals tissue t cavity, that w classifie	by-product or derived product obtained for some subject of the solution of the	aceration of the central nervous ment introduced into the cranial cavity, except for those animals tered in a country or region			
II.7.	the hydrol	ysed pr	otein/dic	calcium phosph	nate/tricalcium phosphate (²) described above	x:			
	(²) either			itain milk or m ls, other than f	ilk products of ovine or caprine animal origin rur animals.]	or is not intended for feed for			
	(²) or				ucts of ovine or caprine animal origin and is als, and the milk or milk products:	s intended for feed for farmed			
		(a)			ne and caprine animals which have been ke llowing conditions are fulfilled:	pt continuously since birth in a			
			(i)	classical scra	apie is compulsorily notifiable;				
			(ii)	an awarenes	s, surveillance and monitoring system is in pla	ace for classical scrapie;			
			(iii)		etions apply to holdings of ovine or caprine an econfirmation of classical scrapie;	imals in the case of a suspicion			

COUNTRY

Hydrolysed protein, dicalcium phosphate and tricalcium phosphate not intended for human consumption to be used as feed material or for uses outside the feed chain

		acca ac reca material er re-	acce caterae the reca chain
I	II. Health information	II.a. Certificate reference No	II.b.

- (iv) ovine and caprine animals affected with classical scrapie are killed and destroyed;
- (v) the feeding to ovine and caprine animals of meat-and-bone meal or greaves, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health (OIE), of ruminant origin has been banned and effectively enforced in the whole country for a period of at least the preceding seven years;
- (b) originate from holdings where no official restrictions are imposed due to a suspicion of TSE;
- (c) originate from holdings where no case of classical scrapie has been diagnosed during the period of the preceding seven years or, following the confirmation of a case of classical scrapie:
 - (2) either [all ovine and caprine animals on the holding have been killed and destroyed or slaughtered, except for breeding rams of the ARR/ARR genotype, breeding ewes carrying at least one ARR allele and no VRQ allele and other ovine animals carrying at least one ARR allele:]
 - (²) or [all animals in which classical scrapie was confirmed have been killed and destroyed, and the holding has been subjected for a period of at least two years since the date of confirmation of the last classical scrapie case to intensified TSE monitoring, including testing with negative results for the presence of TSE in accordance with the laboratory methods set out in point 3.2 of Chapter C of Annex X to Regulation (EC) No 999/2001, of all of the following animals which are over the age of 18 months, except ovine animals of the ARR/ARR genotype:
 - animals which have been slaughtered for human consumption; and
 - animals which have died or been killed on the holding but which were not killed in the framework of a disease eradication campaign.]]

Notes

Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is required to be filled in only if
 it is a certificate for a commodity to be transited through the European Union; it may be filled in if the certificate is for a
 commodity to be imported into the European Union.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for a transit commodity. Products
 in transit can only be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading.
- Box reference I.19: use the appropriate HS code: 05.08, 28.35.25; 28.35.26, 29.22; 35.02; 35.03 or 35.04.
- Box reference I.23; for bulk containers, the container number and the seal number (if applicable) must be included.
- Box reference I.25: technical use: any use other than feeding of farmed animals, other than fur animals, and the
 production or manufacturing of pet food.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28:
 - Species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or Suidae, Pesca,
 Mollusca, Crustacea, invertebrates other than Mollusca and Crustacea.

II.	Health information	II.a. Certificate reference No		II.b.							
	Nature of commodity: specify if hydroly:	sed protein, dicalcium phosphat	e or tricalcium	phosphate.							
	Manufacturing plant: provide the registr	ration number of treatment/proce	essing establis	nment.							
Part	Part II:										
(^{1a})	a) OJ L 300, 14.11.2009, p. 1.										
(1b)	OJ L 54, 26.2.2011, p. 1.										
(2)	Delete as appropriate.										
(3)	OJ L 147, 31.5.2001, p. 1.										
(4)	OJ L 94, 1.4.2006, p. 28.										
_	The signature and the stamp must be in a diff	ferent colour to that of the printin	ng.								
_	 Note for the person responsible for the consignment in the European Union: this certificate is only for veterinary purposes and must accompany the consignment until it reaches the border inspection post of the point of entry into the European Union. 										
Offic	cial veterinarian/Official inspector										
	Name (in capital letters):		Qualification a	and title:							
	Date:		Signature:								
	Stamp:										

Type of packaging

I.23. Seal/Container No

(6) Chapter 18 is replaced by the following:

'CHAPTER 18

Health certificate

For horns and horn products, excluding horn meal, and hooves and hoof products, excluding hoof meal, intended for the production of organic fertilisers or soil improvers intended for dispatch to or for transit through (2) the European Union

COUNTRY: Veterinary certificate to EU Consignor 1.2. Certificate reference No I.2.a. Name 1.3. Central competent authority Address 1.4. Local competent authority Tel. 1.5. Consignee 1.6. Person responsible for the load in EU Part I: Details of dispatched consignment Name Name Address Address Postcode Postcode Tel. Tel. 1.7. Country ISO code Region of Code 1.9. Country of ISO I.10. Region of Code destination of origin origin destination code I.11. Place of origin I.12. Place of destination Name Approval number Custom warehouse Address Name Approval number Name Approval number Address Address Name Approval number Postcode 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 Other \square Road vehicle I.17. Number(s) of CITES Identification Documentation references I.18. Description of commodity I.19. Commodity code (HS code) 05.07 I.20. Quantity I.21. Temperature of product I.22. Number of packages Chilled Frozen Ambient \square

1.25.	Commodities certified for:						
	Further process	Tech	Technical use □				
1.26.	For transit through EU to third	d country	ı	1.27. For import or admission into EU			
	Third country	ISO code					
1.28.	Identification of the commodi		umber of	establishments			
	Species (Scientific name)	Manufacturing plant	t	Net weight	Batch number		

Horns and horn products, excluding horn meal, and hooves and hoof products, excluding hoof meal, intended for the production of organic fertilisers or soil improvers

	II.	Health information			II.a. Certificate reference No	II.b.						
		the Europ particular	ean Parliame Chapter II of	ent and of the Co Annex XIV thereto	eclare that I have read and understood Regulation (EU) ouncil (1a), and Commission Regulation (EU) o, and certify that the horns and horn products f meal (2) described above) No 142/2011 (^{1b}), and in						
_	II.1.	originate from animals										
tificatio		(²) either	[that were slaughtered in a slaughterhouse, after undergoing ante-mortem inspection, and were fit, as a result of such inspection, for slaughter for human consumption;]									
Part II: Certification		(2) or [that did not show clinical signs of any disease communicable through that product to animals;]										
Pa	II.2.		horns, horn products, hooves and hoof products must have undergone a heat treatment for one hour at a core temperature of at least 80 °C;									
II.3. horns must have been removed without opening the cranial cavity;												
	 II.4.		at any stage of processing, storage or transport every precaution must have been taken to avoid cross-contamination.									
	II.5.	the horns packed:	and horn pr	oducts, excluding	horn meal, and hooves and hoof products,	excluding hoof meal, were						
		(²) either	[in new pack	kaging or containe	rs;]							
		(²) or	[in vehicles authority;]	or bulk containers	s disinfected prior to loading using a product	approved by the competent						
			'NOT FOR H		ked so as to indicate the type of the animal by MAL CONSUMPTION' and the name and addr							
	(²)[II.6.	The horns above	and horn pro	oducts, excluding	horn meal, and hooves and hoof products, exc	cluding hoof meal described						
		(²) either	[is derived fi	rom other ruminan	its than bovine, ovine or caprine animals.]]							
		(²) or	[is derived fi	rom bovine, ovine	or caprine animals and does not contain and is	not derived from:						
			(²) either	continuously rea	and caprine materials other than those de red and slaughtered in a country or region clas rdance with Decision 2007/453/EC.]							
			(²) or		risk material as defined in point 1 of An 2001 of the European Parliament and of the Co							
				animals, slaughter accordar	cally separated meat obtained from bones of except from those animals that were borned in a country or region classified as positive with Commission Decision 2007/453/EC (separate SEE case,	n, continuously reared and ng a negligible BSE risk in						
		(c) animal by-product or derived product obtained from bovine, ovine or capring animals which have been killed, after stunning, by laceration of the central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranic cavity, or by means of gas injected into the cranial cavity, except for those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]]										

Horns and horn products, excluding horn meal, and hooves and hoof products, excluding hoof meal, intended for the production of organic fertilisers or soil improvers

			for the produc	tion of organic t	fertilisers or soil improvers					
II.	Health information	II.a.	Certificate reference N	lo	II.b.					
Note	es									
Part	Part I:									
_	Box reference I.6: Person responsible for the consignment in the European Union: this box is required to be filled in only if it is a certificate for a commodity to be transited through the European Union; it may be filled in if the certificate is for a commodity to be imported into the European Union.									
_	Box reference I.11 and I.12: Approval number issued by the competent authority.	er: the	registration number of	the establishme	nt or plant, which has been					
_	Box reference I.12: Place of destination: this be in transit must only be stored in free zones, free				transit commodity. Products					
_	Box reference I.15: Registration number (railwinformation is to be provided in the event of un				nber (aircraft) or name (ship);					
_	Box reference I.23: for bulk containers, the cor	ntainer	number and the seal nu	umber (if applicab	ole) must be given.					
_	Box reference I.25: technical use: any use other	er than	for animal consumption	١.						
_	Box reference I.26 and I.27: fill in according to	wheth	er it is a transit or an im	port certificate.						
_	Box reference I.28: Nature of commodity.									
Part	: II:									
(^{1a})	OJ L 300, 14.11.2009, p. 1.									
(1b)	OJ L 54, 26.2.2011, p. 1.									
(2)	Delete as appropriate.									
(3)	Type of product: horns, horn products, hooves	, hoof _l	oroducts.							
(4)	OJ L 147, 31.5.2001, p. 1.									
(⁵)	OJ L 172, 30.6.2007, p. 84.									
_	The signature and the stamp must be in a diffe	erent co	plour to that of the printi	ng.						
_	 Note for the person responsible for the consignment in the European Union: this certificate is only for veterinary purposes and must accompany the consignment until it reaches the border inspection post of the point of entry into the European Union. 									
Offic	cial veterinarian/Official inspector									
	Name (in capital letters):			Qualification an	d title:					
	Date:			Signature: '						
	Stamp:									

(7) Chapter 20 is replaced by the following:

'CHAPTER 20

Model declaration

Declaration for the import from third countries and for the transit through (²) the European Union of intermediate products to be used for the manufacture of medicinal products, veterinary medicinal products, medical devices for medical and veterinary purposes, active implantable medical devices, in vitro diagnostics medical devices for medical and veterinary purposes, laboratory reagents and cosmetic products

COUNTRY:	Veterinary certificate to EU
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	l.1.	Consignor					1.2.	Certificate refere	ence No	1.2.a.			
	Name Address						1.3.	I.3. Central competent authority					
		Address					1.4.	I.4. Local competent authority					
	1.5	Tel.					1.6	Poreon recepons	ible for the le	ad in EU			
+	I.5.	Consignee Name					1.6.	Person responsi	ible for the lo	ad in EU			
men		Address						Address					
sign		Address						Addiess					
Part I : Details of dispatched consignment		Postcode						Postcode					
pəy		Tel.						Tel.					
spate	1.7.	Country	ISO code	1.8.	Region of	Code	1.9.	Country of	ISO	I.10. Region of	Code		
of dis		of origin		I	origin			destination	code 	destination	1		
ails (l 11	Place of origin					112	Place of destina	tion				
Det	1.11. Trace of origin							Tidoo of dootilla					
art		Name		Appro	val number					Custom warehouse			
ď		Address						Name		Approval number			
		Name		Appro	val number		Address						
		Address											
		Name		Appro	val number			Postcode					
		Address											
	I.13.	Place of loa	ading				1.14.	Date of departur	re				
	I.15.	Means of to	ransport				I.16.	Entry BIP in EU					
		Aeroplane	☐ Ship		Railway wa	agon 🗖							
		Road vehic	cle 🔲 Othe	er 🔲			1.17.						
		Identification	on										
			ation reference										
	l.18.	Description	n of commodi	ity					I.19. Comr	modity code (HS code)			
								l		I.20. Quantity			
	I.21.	Temperatu	re of product	t						I.22. Number of pa	ackages		
		Ambient 🗀	<u> </u>		Chilled D]		Frozen []				
	1.23.	Seal/Conta	iner No							I.24. Type of pack	aging		

1.25.	Commodities certified fo	or:						
	Technical use \square							
1.26.	For transit through EU to	o third country		I.27. For import or admission into EU				
	Third country	ISO code						
1.28.	8. Identification of the commodities Approval number of establishments							
	Species (Scientific name)	Manufact	uring plant	Net weight	Batch number			

Part II: Certification

COUNTRY

Intermediate products to be used for the manufacture of medicinal products, veterinary medicinal products, medical devices for medical and veterinary purposes, active implantable medical devices, in vitro diagnostics medical devices for medical and veterinary purposes, laboratory reagents, and cosmetic products

II. Health information	II.a. Certificate reference No	II.b.
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DECLARATION

I, the undersigned, declare that the intermediate product referred to above is intended to be imported by me into or to be transited through the European Union and satisfies the definition of an intermediate product provided for in point 35 of Annex I to Commission Regulation (EU) No 142/2011 (^{1a}), and in particular that:

- (1) it is intended for the manufacture of:
 - (2) either [- medicinal products,]
 - (2) and/or [- veterinary medicinal products,]
 - (2) and/or [- medical devices for medical and veterinary purposes,]
 - (2) and/or [- active implantable medical devices,]
 - (2) and/or [- in vitro diagnostic medical devices for medical and veterinary purposes,]
 - (2) and/or [- laboratory reagents,]
 - (2) and/or [- cosmetic products;]
- (2) its design, transformation and manufacturing stages have been sufficiently completed in order to qualify the material directly or as a component of a product intended for that purpose, except for the fact that it requires further manufacturing or transformation such as mixing, coating, assembling or packaging to make it suitable for placing on the market or putting into service as a medicinal product, veterinary medicinal product, medical device for medical and veterinary purposes, an active implantable medical devices, an in vitro diagnostic medical device for medical and veterinary purposes or a cosmetic product in accordance with the European Union legislation (1b) applicable to those products or as a laboratory reagent;
- (3) it has been derived from:
 - (2) either [- material which may have originated from animals submitted to an illegal treatment as defined in Article 1(2)(d) of Council Directive 96/22/EC (2a) or in Article 2(b) of Council Directive 96/23/EC (2b);]
 - (2) and/or [- carcases and parts of animals slaughtered or, in the case of game, bodies or parts of animals killed, and which are fit for human consumption in accordance with Union legislation, but are not intended for human consumption for commercial reasons;]
 - (2) and/or [- carcases and the following parts originating either from animals that have been slaughtered in a slaughterhouse and were considered fit for slaughter for human consumption following an antemortem inspection or bodies and the following parts of animals from game killed for human consumption in accordance with Union legislation:
 - carcases or bodies and parts of animals which are rejected as unfit for human consumption in accordance with Union legislation, but which did not show any signs of disease communicable to humans or animals:
 - (ii) heads of poultry;
 - (iii) hides and skins, including trimmings and splitting thereof, horns and feet, including the phalanges and the carpus and metacarpus bones, tarsus and metatarsus bones, of animals other than ruminants;
 - (iv) pig bristles;
 - (v) feathers;]

Intermediate products to be used for the manufacture of medicinal products, veterinary medicinal products, medical devices for medical and veterinary purposes, active implantable medical devices, in vitro diagnostics medical devices for medical and veterinary purposes, laboratory reagents, and cosmetic products

II.	Health	infor	mation	II.a. Certificate reference No	II.b.						
	(²) and/or	[-	animals obtained from anima	not show any signs of disease communicable through blood to humans or nals other than ruminants that have been slaughtered in a slaughterhouse ared fit for slaughter for human consumption following an ante-mortem of the Union legislation;							
	(²) and/or	[-	,	m the production of products intended for had centrifuge or separator sludge from milk p							
	(²) and/or	[-	intended for human consum	r foodstuffs containing products of anima ption for commercial reasons or due to p fects from which no risk to public or animal	problems of manufacturing or						
	(²) and/or	[-	products, which are no longe	animal origin, or feedingstuffs containing animal by-products or derived er intended for feeding for commercial reasons or due to problems of defects or other defects from which no risk to public or animal health							
	(²) and/or	[-	blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk originating from live animals that did not show signs of any disease communicable through that product to humans or animals;]								
	(²) and/or	[-		aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of diseases communicable to humans or animals;]							
	(²) and/or	[-	animal by-products from aqu products for human consump	uatic animals originating from plants or etion;]	establishments manufacturing						
	(²) and/or	[-	the following material originat through that material to huma	ing from animals which did not show any si ns or animals:	igns of disease communicable						
			(i) shells from shellfish with	n soft tissue or flesh;							
			(ii) the following originating	g from terrestrial animals:							
			— hatchery by-produ	ucts,							
			— eggs,								
			 egg by-products, ii 	s, including egg shells;							
			(iii) day-old chicks killed for	commercial reasons;]							
	(²) and/or	[-	animal by-products from aqua or animals;]	atic or terrestrial invertebrates other than s	species pathogenic to humans						
	(²) and/or	[-		the zoological orders of Rodentia and Lagricle 8(a)(iii), (iv) and (v) and Category n (EC) No 1069/2009;]							
	(²) and/or	[-	products derived from or gene	erated by:							
				parts of such animals, except sea mammals, which did not show any signs able to humans or animals,							
			 aquatic or terrestrial inventor 	vertebrates other than species pathogenic to humans or animals,							
			Category 1 material as	ereof of the zoological orders of Rodentia and Lagomorpha, except is referred to in Article 8(a)(iii), (iv) and (v) and Category 2 material as i) to (g) of Regulation (EC) No 1069/2009;]							

COUNTRY

Date:

Intermediate products to be used for the manufacture of medicinal products, veterinary medicinal products, medical devices for medical and veterinary purposes, active implantable medical devices, in vitro diagnostics medical devices for medical and veterinary purposes, laboratory reagents, and cosmetic products

Signature: '.

						lab	oratory reagents	s, and cosmetic products
II.	Health	infor	matic	on	II.a.	Certificate reference N	lo	II.b.
	(²) and/or	[-		nals and parts of animal 1069/2009,	s, othe	er than those referred to	in Article 8 or Ar	ticle 10 of Regulation (EC)
			(i)	that died other than b killed for disease contr			for human consu	umption, including animals
			(ii)	foetuses;				
			(iii)	oocytes, embryos and	semei	n which are not destined	I for breeding purp	poses; and
			(iv)	dead-in-shell poultry;]				
	(²) and/or	[-	anim	nal by-products other tha	ın Cat	egory 1 material or Cate	gory 3 material;]	
(4)	4) its outer packaging is labelled 'FOR MEDICINAL PRODUCTS / VETERINARY MEDICINAL PRODUCTS / MEDICAL DEVICES FOR MEDICAL AND VETERINARY PURPOSES / ACTIVE IMPLANTABLE MEDICAL DEVICES / IN VITRO DIAGNOSTIC MEDICAL DEVICES FOR MEDICAL AND VETERINARY PURPOSES / LABORATORY REAGENTS / COSMETIC PRODUCTS ONLY' and it is not intended to be diverted at any stage within the European Union for any other use;							
(5)	the consignm point I.12 of the				o the	place of destination in	n the European	Union as indicated under
	(2) either [an establishment or plant for the production of medicinal products, veterinary medicinal products, medical devices for medical and veterinary purposes, active implantable medical devices, in vitro diagnostic medical devices for medical and veterinary purposes, laboratory reagents or cosmetic products, which has been registered in accordance with Article 23 of Regulation (EC) No 1069/2009],							
	(²) or	No	1069					24(1)(i) of Regulation (EC) or plant referred to in the
Note	es							
_	2007/275/EC	of 17	' April		f anim	als and products to be	subject to controls	ith Commission Decision at border inspection posts
_	Box reference	1.25	tech	nical use: any use other	than f	or animal consumption.		
(^{1a})	OJ L 54, 26.2	.2011	I, p. 1					
(^{1b})	Directive 2001/82/EC of the European Parliament and of the Council of 6 November 2001 on the Community code relating to veterinary medicinal products (OJ L 311, 28.11.2001, p. 1), Directive 2001/83/EC of the European Parliament and of the Council of 6 November 2001 on the Community code relating to medicinal products for human use (OJ L 311, 28.11.2001, p. 67), Council Directive 93/42/EEC of 14 June 1993 concerning medical devices (OJ L 169, 12.7.1993, p. 1) and Directive 98/79/EC of the European Parliament and the Council of 27 October 1998 on in vitro diagnostic medical devices (OJ L 331, 7.12.1998, p. 1), Regulation (EC) No 1223/2009 of the European Parliament and of the Council of 30 November 2009 on cosmetic products (OJ L 342, 22.12.2009, p. 59), as appropriate.							
(2)	Delete as app	ropri	ate.					
(^{2a})	OJ L 125, 23.	5.199	96, p.	3.				
(^{2b})	OJ L 125, 23.	5.199	96, p.	10.				
The	importer							
	Name (in capi	ital le	tters)	:			Address:	