

Brussels, XXX SANTE/7008/2017 ANNEX CIS Rev. 4 (POOL/G2/2017/7008/7008R4-EN ANNEX CIS.doc) [...](2018) XXX draft

ANNEXES 1 to 2

#### **ANNEXES**

to the

# COMMISSION REGULATION (EU) .../...

amanding 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

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#### **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(<sup>2</sup>) the European Union

CC	OUNTRY:	Veterinary certificate to EU				
	I.1. Consignor Name	I.2. Certificate reference No I.2.a.				
	Address	I.3. Central competent authority				
ent	Tel.	I.4. Local competent authority				
Part I : Details of dispatched consignment	I.5. Consignee Name Address  Postcode Tel.	I.6. Person responsible for the load in EU Name Address  Postcode Tel.				
of disp	I.7. Country of ISO code I.8. Region of origin Code origin	I.9. Country of ISO code I.10. Region of Code destination destination				
tails	I.11. Place of origin	I.12. Place of destination				
Part I : De	Name Approval number Address Name Approval number Address Name Approval number	Custom warehouse  Name Approval number Address  Postcode				
	Address  I.13. Place of loading	I.14. Date of departure				
	I.15. Means of transport  Aeroplane  Ship  Railway wagon  Road vehicle  Other  Identification	I.16. Entry BIP in EU  I.17.				
	Documentation references  I.18. Description of commodity	I.19. Commodity code (HS code)				
	1.10. Description of commonly					
		I.20. Quantity				
	I.21. Temperature of product Ambient □ Chilled □	I.22. Number of packages Frozen □				
	I.23. Seal/Container No	I.24. Type of packaging				
	I.25. Commodities certified for:	·				
	Animal feedingstuff ☐ Tech	nical use $\square$ Manufacture of petfood $\square$				
	I.26. For transit through EU to third country	I.27. For import or admission into EU				
	Third country ISO code					
	I.28. Identification of the commodities  A Species (Scientific name) Nature of commodity	approval number of establishments  Manufacturing plant  Net weight  Batch number				

**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

# Part II: Certification

II.1.

II.	Health information	II.a. Certificate reference No	II.b.
		narian, declare that I have read and an Parliament and of the Council(1a) a	
			ind in particular Article

- 10 thereof, and Commission Regulation (EU) No 142/2011(1b), and in particular Section 1 of Chapter II of Annex X, and Chapter I of Annex XIV thereto and certify that: the processed animal protein or product described above contains exclusively processed animal
- protein not intended for human consumption that: has been prepared and stored in an establishment or plant approved and supervised by (a) the competent authority in accordance with Article 24 of Regulation (EC) No 1069/2009, and
  - (b) 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 ante-mortem 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;
      - heads of poultry; (ii)
      - 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 [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;]
    - $(^{2})$ and/or [products of animal origin, or foodstuffs containing products of animal origin, which are no longer 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;]
    - $(^{2})$ 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;]
    - $\binom{2}{and/or}$  [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 aquatic animals originating from establishments or plants manufacturing products for human

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 information		II.a. Certificate reference No	II.b.			
		consump	tion;]				
	( <sup>2</sup> )and/or [-		wing material originating from anims of disease communicable through				
			ells from shellfish with soft tissue or	flesh:			
		` '	following originating from terrestria				
		-	hatchery by-products,				
		-	eggs,				
		- egg by-products, including egg shells;					
	2	•	y-old chicks killed for commercial re				
	(²)and/or [-	humans o	and terrestrial invertebrates other that or animals and other than insects;]				
	(²)and/or [-	animals and parts thereof of the zoological orders of Rodentia Lagomorpha, except Category 1 material as referred to Article 8(a)(iii), (iv) and (v) and Category 2 material as referred Article 9(a) to (g) of Regulation (EC) No 1069/2009;]					
	and						
		jected to the	following processing standard:				
	wit sat	heating to a core temperature of more than 133°C for at least 20 minu vithout interruption at a pressure (absolute) of at least 3 bars produced attracted steam, with a particle size prior to processing of not more than 133°C.					
	(²)or [in me	50 millimetres;] [in the case of non-mammalian protein other than fishmeal, the proce method 1-2-3-4-5-7(indicate the processing me as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011;					
	(²) <i>or</i> [in 7	the case	e of fishmeal the processing(indicate the processing Annex IV to Regulation (EU) No 142	method 1-2-3-4-5-6- method) as set out in			
	(in Re	dicate the p gulation (EU	f porcine blood, the processing me rocessing method) as set out in Cha J) No 142/2011, where in case of m C has been applied throughout its sub	apter III of Annex IV to ethod 7 a heat treatment			
II.2.		y examined	a random sample immediately prior				
	Salmonella:	_	n 25 g: $n = 5$ , $c = 0$ , $m = 0$ , $M = 0$				
	Enterobacteriaceae:		2, m = 10, M = 300  in  1g;				
II.3.	the product has under after treatment;	gone all pr	ecautions to avoid recontamination	with pathogenic agents			
II.4.	the end product:						
(²)either [was packed in new or sterilised bags,]							
			bulk in containers or other means nd disinfected before use,]	of transport that were			
	<del>-</del>	-	FOR HUMAN CONSUMPTION';				
II.5.	the end product was st	ored in encl	osed storage;				
(²)II.6.	[the processed animal products of ruminant of		roduct described above contains or is	derived from animal-by			
	(²)either [orig	inates from	n a country or region, which is is is accordance with Decision 200				

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	rmation		II.a. Certificate reference No	II.b.			
	( <sup>2</sup> ) <i>a</i>	or [originate in according in digenous derived from the control of rumina as defined in the control of rumina and rumina as defined in the control of rumina and rumina as defined in the control of rumina as def	s from lance values BSE rom an ants with d in the	o indigenous BSE case.] a country or region classified as posin with Decision 2007/453/EC in which case, and the animal by-product or imals born after the date from which the meat-and-bone meal and greaves due OIE Terrestrial Animal Health Cod country or region.]	ch there has been an derived product were the ban on the feeding erived from ruminants,			
	and							
	$(^2)\epsilon$	either [is derive	d from	other ruminants than bovine, ovine or	caprine animals.]			
	$(^2)\epsilon$	or [is derived not derived		bovine, ovine or caprine animals and ::	does not contain and is			
		( <sup>2</sup> ) either	fro co	ovine, ovine and caprine materials of om animals born, continuously reared untry or region classified as posing a cordance with Decision 2007/453/EC.	l and slaughtered in a negligible BSE risk in			
		( <sup>2</sup> ) <i>or</i>						
			bo tha co ac	) mechanically separated meat obtavine, ovine or caprine animals, except were born, continuously reared untry or region classified as posing a cordance with Commission Decisionich there has been no indigenous BSE	pt from those animals and slaughtered in a negligible BSE risk in n 2007/453/EC(5), in			
			bo stu of cra ca rea po	o animal by-product or derived privine, ovine or caprine animals which inning, by laceration of the central near elongated rod-shaped instrumer anial cavity, or by means of gas in vity, except for those animals that wared and slaughtered in a country casing a negligible BSE risk in accountry of the control of the country of the countr	have been killed, after ervous tissue by means at introduced into the ected into the cranial ere born, continuously or region classified as			
II.7.	-	ed animal protei	n or pr	oduct described above:				
	( <sup>2</sup> )either			lk or milk products of ovine or caprine farmed animals, other than fur animal	_			
	$(^2)$ or			lk products of ovine or caprine anima nimals, other than fur animals, which:	origin and is intended			
		(a) are de contin						
		(i)	clas	sical scrapie is compulsorily notifiable	· ,			
		(ii)		wareness, surveillance and monitoring sical scrapie;	g system is in place for			
		(iii)	offic	cial restrictions apply to holdings of one case of a suspicion of TSE or the co				
		(iv)	ovir	ne and caprine animals affected with	n classical scrapie are			

killed and destroyed;

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 informati	on	II.a. Certi	ficate reference No	II.b.	
		or g the orig	reaves, as de World Orga in has been	vine and caprine animals of efined in the Terrestrial Anisation for Animal Heal banned and effectively of iod of at least the preceding	Animal Health Code of lth (OIE), of ruminant enforced in the whole	
	(b)	originate from holdings where no official restrictions are imposed due to a suspicion of TSE;				
	(c)	diagnosed d	originate from holdings where no case of classical scrapie has beer diagnosed during a period of at least 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;]				
		kille peri last incl acco Cha the	d and destrood of at least classical standing testing rdance with oter C of An following a pt ovine animals	which classical scrapie way oyed, and the holding has to two years since the date scrapie case to intensify with negative results for the laboratory methods anex X to Regulation (EC) nimals which are over the mals of the ARR/ARR general which have been sla	s been subjected for a of confirmation of the ied TSE monitoring, the presence of TSE in set out in point 3.2 of No 999/2001, of all of he age of 18 months, notype:	
		_	but whic	tion; and which have died or been h were not killed in the fi on campaign.]]		
II.8.			oduct descri	bed above contains or is d ling to the statement of the		
	(²)either [not anim		e productio	on of feed for farmed ar	nimals, other than fur	
	fur Inspe	animals, and the ection Post of ex	ne Consigno ntry will be	eed for non-ruminant farm or has undertaken to en provided with the results tethods set out in Anne	sure that the Border of the analyses carried	

## 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).]

- Box reference I.12: Place of destination: this box is required to be filled in only if it is a certificate
  for a commodity to be transited through the European Union. 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

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 information	II.a. Certificate reference No	II.b.
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(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: 05.05; 05.06; 05.07; 05.11; 23.01 or 23.09.
- 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, Ruminantia, Mammalia other than Ruminantia, Pesca, Mollusca, Crustacea, invertebrates other than Mollusca and Crustacea. In the case of farmed fish, specify the scientific 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.
- (<sup>3</sup>) 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
  - 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.
- (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.
- (<sup>7</sup>) 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.

Official veterinarian/Official inspector								
Name (in capital letters):	Qualification and title:							
Date:	Signature:							
Stamp:								

#### 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<sup>2</sup> the European Union

**COUNTRY:** 

CO	OUNTRY:	Veterinary certificate to EU			
	I.1. Consignor	I.2. Certificate reference No I.2.a.			
	Name Address	I.3. Central competent authority			
ınt	Tel.	I.4. Local competent authority			
Part I: Details of dispatched consignment	I.5. Consignee Name Address  Postcode Tel.	I.6. Person responsible for the load in EU Name Address  Postcode Tel.			
f dispa	I.7. Country of ISO code I.8. Region of origin Code origin	I.9. Country of ISO code I.10. Region of Code destination			
s of					
ail	I.11. Place of origin	I.12. Place of destination			
art I : Det	Name Approval number Address Name Approval number Address	Custom warehouse Name Approval number Address			
F	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			
	A avanlana Chin Dailway wasan C				
	Aeroplane	I.17.			
	Identification	1.17.			
	Documentation references  I.18. Description of commodity	I.19. Commodity code (HS code)			
		I.20. Quantity			
	I.21. Temperature of product Ambient □ Chilled □	I.22. Number of packages			
	I.23. Seal/Container No	I.24. Type of packaging			
	I.25. Commodities certified for:				
	Animal feedingstuff □ Techn	nical use   Manufacture of petfood			
	I.26. For transit through EU to third country	I.27. For import or admission into EU			
	Third country ISO code				
	I.28. Identification of the commodities				
	Species (Scientific name)  Nature of commodity	pproval number of establishments  Manufacturing plant  Net weight  Batch number			

Part II: Certification

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.

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(<sup>1a</sup>) and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011(<sup>1b</sup>), and in particular Section 1 of Chapter II of Annex X, and Chapter I of Annex XIV thereto and certify that:

- II.1. the processed animal protein derived from farmed insects or product described above contains exclusively processed animal protein not intended for human consumption that:
  - (a) has been prepared and stored in an establishment or plant approved and supervised by the competent authority in accordance with Article 24 of Regulation (EC) No 1069/2009, and
  - (b) has been prepared exclusively from farmed insects of the following species:
    - (2) either [- Black Soldier Fly (Hermetia illucens);]
    - (<sup>2</sup>)and/or [- Common Housefly (Musca domestica);]
    - (2) and/or [- Yellow Mealworm (Tenebrio molitor);]
    - (2) and/or [- Lesser Mealworm (Alphitobius diaperinus);]
    - (2) and/or [- House cricket (Acheta domesticus);]
    - (2) and/or [- Banded cricket (Gryllodes sigillatus);]
    - (<sup>2</sup>)and/or [- Field Cricket (*Gryllus assimilis*).]

and

(c) has been processed by method [1]-[2]-[3]-[4]-[5]-[7](<sup>2</sup>) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011;

and

- (d) the substrate for the feeding of farmed insects may only contain products of non-animal origin or the following products of animal origin of Category 3 material:
  - fishmeal;
  - blood products from non-ruminants;
  - di and tricalcium phosphate of animal origin;
  - hydrolysed proteins from non-ruminants;
  - 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;

and

- (e) the substrate for the feeding of insects and the insects or their larvae have not been in contact with any other materials of animal origin than those referred to in point (d) and the substrate did not contain manure, catering waste or other waste.
- II.2. the competent authority examined a random sample immediately prior to dispatch and found it to comply with the following standards(<sup>3</sup>):

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

Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1g;

- II.3. the product has undergone all precautions to avoid recontamination with pathogenic agents after treatment;
- II.4. the end product:

(2) either [was packed in new or sterilised bags,]

(²) or [was transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected before use,]

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

			contai	ning such protein	
II.	Health info	ormation		II.a. Certificate reference No	II.b.
	PROTEIN		T BE	F FOR HUMAN CONSUMPTION USED IN FEED FOR FARME IMALS';	
II.5.	the end pro	duct was stored	in enclo	osed storage;	
( <sup>2</sup> )II.6.	- 1	sed animal prote ruminant origin		roduct described above contains or i	is derived from animal-b
	( <sup>2</sup> ).	BSE risk	in acco	a country or region, which is classiful ordance with Decision 2007/453/EG bus BSE case.]	
	(2)	or [originate in accord indigenou derived fi of rumina as defined	s from a lance was BSE com animals with d in the	a country or region classified as pos- with Decision 2007/453/EC in w case, and the animal by-product mals born after the date from which h meat-and-bone meal and greaves to OIE Terrestrial Animal Health Country or region.]	thich there has been and or derived product were the ban on the feedings derived from ruminants
	and	cinoreca	in that c	ountry of region.	
				other ruminants than bovine, ovine	or caprine animals.]
	$\binom{2}{2}$		d from b	bovine, ovine or caprine animals ar	
		(²) either	[bo froi cou	ovine, ovine and caprine materials om animals born, continuously reauntry or region classified as posing cordance with Decision 2007/453/E	red and slaughtered in g a negligible BSE risk i
	( <sup>2</sup> )		specifie n (EC)	ed risk material as defined in p No 999/2001 of the European	point 1 of Annex V to
		(b) m ca co as D	nechanic aprine ontinuou s posing	cally separated meat obtained from animals, except from those are usly reared and slaughtered in a cog a negligible BSE risk in accord 2007/453/EC(5), in which there expended.	nimals that were born ountry or region classified dance with Commission
		ca of in in bo	aprine and the center in the c	y-product or derived product obtain unimals which have been killed, afte entral nervous tissue by means of nt introduced into the cranial cav- into the cranial cavity, except for ntinuously reared and slaughtered as posing a negligible BSE of 2007/453/EC.]]	er stunning, by laceration an elongated rod-shaped rity, or by means of gathose animals that were l in a country or region
II.7.	-	-	-	oduct described above:	
	( <sup>2</sup> )either			k or milk products of ovine or capri farmed animals, other than fur anim	
	( <sup>2</sup> ) <i>or</i>			lk products of ovine or caprine anir nimals, other than fur animals, whic	
			uously	from ovine and caprine animals since birth in a country where the	-

classical scrapie is compulsorily notifiable;

fulfilled:

(i)

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

II.	Health informat	ion	II.a. C	ertificate reference No	II.b.	
		` '	wareness sical scra	, surveillance and monitorin	g system is in place for	
		(iii) offi in t	cial restri	ctions apply to holdings of of a suspicion of TSE or the c	-	
			ne and ca	aprine animals affected wit stroyed;	h classical scrapie are	
		(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 fro suspicion of	-	s where no official restriction	ons are imposed due to a	
	(c)	diagnosed d	uring a	ngs where no case of clas period of at least the pre- ation of a case of classical sc	ceding seven years or,	
	(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 AR alleled and no VRQ alleled and other ovine animals carrying a least one ARR allele;]				r breeding rams of the rying at least one ARR	
		(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,				
		_	anima	animals of the ARR/ARR ge als which have been slamption; and	· =	
		-	but w	als which have died or been hich were not killed in the cation campaign.]]		
II.8.				scribed above contains or is ording to the statement of the		
		intended for t nals.]	ne produ	ction of feed for farmed a	nimals, other than fur	
	fur insp of th	animals, and t ection post of er ne analyses carri	he Consi try into the ed out in	of feed for non-ruminant farming gnor has undertaken to enter the European Union will be paccordance with the methods No 152/2009(7).]	ensure that the border provided with the results	
Notes						
Part I:		Person responsi	de for th	e consignment in the Europ	ean Union: this box is	
_ E	ook reference 1.0: I	erson responsit	ne for the	e consignment in the Europ	can Union; this dox is	

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.

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 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: insects, specify its scientific name.

#### Part II:

- (<sup>1a</sup>) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.
- (<sup>3</sup>) 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
  - 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.
- (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.
- (<sup>7</sup>) 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.

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

II.a. Certificate reference No	II.b.	
Qualification and title:		
Signature:		
	Qualific	

# 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

CO	OUNTRY:			Veterin	ary certificate to EU	
	I.1. Consignor Name	I	.2. Certificate reference l	No	I.2.a.	
	Address	I	I.3. Central competent authority			
ent	Tel.	I	.4. Local competent auth	ority		
Part I : Details of dispatched consignment	I.5. Consignee Name Address  Postcode Tel.	I	I.6. Person responsible for the load in EU Name Address Postcode Tel.			
of dispa	I.7. Country of ISO code I.8. origin	Region of origin Code I	.9. Country of destination	ISO code	I.10. Region of destination Code	
ails	I.11. Place of origin	I	.12. Place of destination			
Part I : Det	Name Address Name Address Name Address	Name Address Postcode	ess			
	I.13. Place of loading	I	.14. Date of departure			
	I.15. Means of transport  Aeroplane  Road vehicle  Identification  Ship  Other  Other	Railway wagon	.16. Entry BIP in EU .17. Number(s) of CITES			
	Documentation references  I.18. Description of commodity		1	I.19. Commod	lity code (HS code)	
					I.20. Quantity	
	I.21. Temperature of product Ambient □	Chilled □	Frozen $\square$		I.22. Number of packages	
	I.23. Seal/Container No	Cliffied 🗆	Piozeii 🗆	I.24. Type of packaging		
	I.25. Commodities certified for:					
	Animal feedingstuff □ Technical use □	Further pr	er process  Production of petfood  Production			
	I.26. For transit through EU to third co	untry	I.27. For import or admission into EU			
	Third country	ISO code				
	I.28. Identification of the commodities					
	Species (Scientific name)	Approval number of establish Manufacturing plant		weight	Batch number	

# Part II: Certification

II.	Health information	II.a. Certificate reference No	II.b.		
		inarian, declare that I have read and			
		C) 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 particula				
	Section 4 of Chapter II of Annex	X, and Chapter I of Annex XIV the	reto, and certify that the		

II a Cortificata reference No

milk(²), the milk-based products(²) and milk-derived products(²) referred to in box I.28 comply

- with the following conditions: they were produced and derived in ...... (insert name of exporting II.1.  $(insert\ name\ of\ region)(^3)$ , which is listed in Part I of Annex II to Commission Regulation (EU) No 605/2010(4), and which has been free from footand-mouth disease (FMD) and rinderpest for a period of 12 months immediately prior to
- 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;

export and has not practised vaccination against rinderpest during that period;

- II.3. they are milk or milk products that:
  - (2)either[have undergone one of the treatments or combinations thereof described in point II.4;]
  - $(^2)$ or [comprise whey to be fed to animals of species susceptible to foot-and-mouth disease, and that whey was collected from milk subjected to one of the treatments described in point II.4 and:
    - (2) either [the whey was collected at least 16 hours after clotting and has a pH below 6;]
    - $(^{2})(^{5})$ or [the whey has been produced at least 21 days before the shipping and during that period no cases of FMD have been detected in the exporting country;]
    - $(^{2})(^{5})$ or [the whey has been produced on ../../.., this date, in consideration of the foreseen voyage duration, being at least 21 days before the consignment is presented to a border inspection post of the European Union;]]
- II.4. they have been subject to one of the following treatments:
  - (2)either[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 milk, in combination with:
    - (2) either [a subsequent second high temperature short time pasteurisation at 72°C for at least 15 seconds or an equivalent pasteurisation which itself achieves a negative reaction to a phosphatase test in bovine milk;]
    - $(^2)$ or [a subsequent drying process that in the case of milk intended for feeding is combined with additional heating to 72°C or higher;]
    - $(^2)$ or [a subsequent process by which the pH is reduced and kept for at least one hour at a level below 6;]
    - $^{(2)(5)}or$ [the condition that the milk/milk product has been produced at least 21 days prior to the date of shipping and during that period no cases of FMD have been detected in the exporting country;]
    - $(^{2})(^{5})or$ [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;]
    - $(^2)$ or [sterilisation at a level of at least  $F_03$ ;]]
  - [ultra high temperature treatment at 132°C for at least one second in combination  $(^2)or$ 
    - (2) either [a subsequent drying process that in the case of milk intended for feeding is combined with additional heating to 72°C or higher;]

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

II.	Health info	ormation	1	II.a.	Certificate reference No		II.b.
			nt process by which the pH is reduced and kept for at least one				
	( <sup>2</sup> )(	-	nour at a lev		/ 6;] ne milk/milk product has be	en n	roduced at least 21 days
	prior to the da				shipping and during that p	-	•
	25				exporting country;]	, ,	
	(-)(				uct has been produced on . on of the foreseen voyage		
	days prior to the date that the consignment is presented to a bo inspection post of the European Union;]]						
II.5.	every precaution was taken to avoid contamination of the milk/milk-based product/milk derived product after processing;				ilk-based product/milk-		
II.6.	_		•		ed product was packed:		
	(²)either	-	w containers				
	$(^2)or$				ntainers disinfected prior to nt authority;]	o lo	pading using a product
	and				d so as to indicate the na		
					duct and bear labels ind ot intended for human cons		
II.7.	the milk, mi	_	•		derived products described		
	(²)either	_			milk products of ovine or ca		C
	$(^2)$ or	[contai	ins milk or i	nilk pro	ducts of ovine or caprine a s, other than fur animals, w	nima	al origin and is intended
		(a)			ovine and caprine anim		
				birth in a country where t		_	
			(i) cl	assical s	scrapie is compulsorily noti	fiabl	e;
				awarer assical s	ness, surveillance and moni scrapie;	torin	ig system is in place for
			in		strictions apply to holdings e of a suspicion of TSE or		
					l caprine animals affected destroyed;	l wi	th classical scrapie are
	(v) the or the original that t		greave e World gin has	g to ovine and caprine anii s, as defined in the Terrest d Organisation for Animal s been banned and effection or a period of at least the pro-	rial Hea vely	Animal Health Code of alth (OIE), of ruminant enforced in the whole	
		(b)		om holo	lings where no official restr		•
	(c) originate fro diagnosed d		rom ho during	ldings where no case of a period of at least the irmation of a case of classic	pred	ceding seven years or,	
	( <sup>2</sup> )either [all and AR alle		d destro RR/ARI lele and	and caprine animals on the oyed or slaughtered, except genotype, breeding ewes no VRQ allele and other ARR allele;]	ot fo	r breeding rams of the rying at least one ARR	
			(²)or [a ki	ll anima lled and	als in which classical scrap d destroyed, and the holdin at least two years since the	ng h	as been subjected for a

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

II.	Health information	II.a. Certificate reference No II.b.
		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
Notes		but which were not killed in the framework of a disease eradication campaign.]]

#### 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 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: 'Manufacturing plant': provide the registration number of treatment or processing establishment.

#### Part II:

- (<sup>1a</sup>) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (<sup>2</sup>) Delete as appropriate.
- For completion if the authorisation to import into or transit through the European Union is restricted to certain regions of the third country concerned.
- OJ L 175, 10.7.2010, p. 1.
- this condition applies only to third countries listed in column 'A' of Annex I to Regulation (EU) No 605/2010.
- 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 is only for veterinary purposes and must accompany the consignment until it reaches the border inspection post.

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

	II.b.			
Official veterinarian/Official inspector				
Qualification and title:				
Signature:				
	_			

# 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

CO	OUNTRY:			Veter	inary certificate to	EU
	I.1. Consignor	I.2.	. Certificate reference N		I.2.a.	
	Name Address	I.3.	. Central competent aut	hority		
ent	Tel.	I.4.	. Local competent author	ority		
Part I: Details of dispatched consignment	I.5. Consignee Name Address  Postcode Tel.	1.6.	Person responsible for Name Address Postcode Tel.	the load in EU	J	
of dispa	I.7. Country of ISO code I.8. Regio origin	n of origin Code I.9.	Country of destination	ISO code	I.10. Region of destination	Code
tails	I.11. Place of origin	I.12	2. Place of destination	<u>l</u>		
Part I : De	Address Name App Address	roval number roval number roval number	Name Address Postcode		n warehouse   wal number	
	I.13. Place of loading	I.14	4. Date of departure			
	L15 Manageftunger	11	C. Fatas DID in EU			
	I.15. Means of transport		6. Entry BIP in EU			
	Aeroplane	ilway wagon  I.1'	7. Number(s) of CITES			
	I.18. Description of commodity	<u> </u>	I	.19. Commod	ity code (HS code)	
			L		I.20. Quantity	
	I.21. Temperature of product Ambient □	Chilled	Frozen		I.22. Number of package	es
	I.23. Seal/Container No				I.24. Type of packaging	
	I.25. Commodities certified for:					
	Animal feedingstuff □ Technical use □	Further proc	ess 🗆		Production of petfoo	d 🗆
	I.26. For transit through EU to third country		I.27. For import or add	mission into E	U $\square$	
	Third country I	ISO code				
	I.28. Identification of the commodities		I.			
	App. Species (Scientific name)	roval number of establishn Manufacturing plant	nents Net w	veight	Batch nu	ımber

# Part II: Certification

# 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

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(<sup>1a</sup>), and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011(<sup>1b</sup>), and in particular Section 4 of Chapter II of Annex X and Chapter I of Annex XIV thereto, and certify that the colostrum(<sup>2</sup>) or the colostrum products(<sup>2</sup>) referred to in box I.28 comply with the following conditions:

- II.2. they were produced from colostrum derived from animals which at the time of milking did not show clinical signs of any disease transmissible through colostrum to humans or animals, and which had been kept for a period of at least 30 days prior to the date of production on holdings that were not subject to official restrictions due to foot-and-mouth disease or rinderpest;
- 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 colostrum or colostrum products have been produced on .../../....(insert the date), this date, in consideration of the foreseen voyage duration, being at least 21 days before the consignment is presented to a border inspection post of the European Union,]
  - and have been obtained from animals subject to regular veterinary inspections to ensure that they come from holdings on which all bovine herds are:
    - (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,] [recognised as official enzootic-bovine-leukosis-free(<sup>6</sup>),]
    - (²)(⁵)either [recognised as official enzootic-bovine-leukosis-free(⁶),]
      (²)(⁵)or [included in an official system for the control of enzootic bovine leukosis and there has been no evidence as result of clinical and laboratory testing of this disease in the herd during the period of the preceding two years,]]
- II.4. every precaution has been taken to avoid contamination of the colostrum/colostrum product after processing;
- II.5. the colostrum or colostrum 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 the containers are marked so as to indicate the nature of the colostrum/colostrum product and bear labels indicating that the product is Category 3 material and not

intended for human consumption; the colostrum or colostrum product does not contain milk or milk products of ovine or caprine

II.6. the colostrum or colostrum product does not contain milk or milk products of ovine or caprine animal origin.

#### Notes

and

#### 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

## Colostrum and colostrum products from bovine animals not for human consumption

II. Health information II.a. Certificate reference No II.b.	
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- 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 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 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: 'Manufacturing plant': provide the registration number of the treatment or processing establishment.

#### Part II:

- (<sup>1a</sup>) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- Delete as appropriate.
- For completion if the authorisation for introduction into the European Union is restricted to certain regions of the third country concerned.
- OJ L 175, 10.7.2010, p. 1.
- This condition applies only to third countries authorised in column 'A' of Annex I to Commission Regulation (EU) No 605/2010 (OJ L 175, 10.7.2010, p. 1).
- Officially tuberculosis-free and brucellosis-free herd as laid down in Annex A to Council Directive 64/432/EEC (OJ 121, 29.7.1964, p. 1977/64) and officially enzootic-bovine-leukosisfree herd as laid down in Chapter I of Annex D to that Directive.
- The signature and the seal must be in a different colour from that of the printing.

Note for the importer: this certificate is only for veterinary purposes and must accompany the consignment until it reaches the border inspection post of the European Union.				
Official veterinarian/Official inspector				
Name (in capital letters):	Qualification and title:			
Date:	Signature:			
Stamp:				

# CHAPTER 3(A)

# Health certificate

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

	UNIKI:	veterinary certificate to EU		
	I.1. Consignor Name	I.2. Certificate reference No I.2.a.		
	Address	I.3. Central competent authority		
ınt	Tel.	I.4. Local competent authority		
nsignme	I.5. Consignee Name Address	I.6. Person responsible for the load in EU Name Address		
itched co	Postcode Tel.	Postcode Tel.		
of dispa	I.7. Country of origin ISO code I.8. Region of origin Code origin	I.9. Country of destination ISO code I.10. Region of destination Code		
tails	I.11. Place of origin	I.12. Place of destination		
Part I: Details of dispatched consignment	Name Approval number Address Name Approval number Address Name Approval number	Custom warehouse  Name Approval number Address  Postcode		
	Address I.13. Place of loading	I.14. Date of departure		
	1.13. Take of folding	1.14. Date of departure		
	I.15. Means of transport  Aeroplane  Ship  Railway wagon  Identification  Road vehicle  Identification	I.16. Entry BIP in EU  I.17.		
	Documentation references  I.18. Description of commodity	I.19. Commodity code (HS code)		
		23.09   I.20. Quantity		
	I.21. Temperature of product Ambient □ Chilled □	I.22. Number of packages		
	I.23. Seal/Container No	I.24. Type of packaging		
	I.25. Commodities certified for:			
	Petfood  Techn	nical use		
	I.26. For transit through EU to third country	I.27. For import or admission into EU		
	Third country ISO code			
	I.28. Identification of the commodities			
	Species (Scientific name)  Approval number of expectation of the second			

COUNTRY Canned Petfood

	II.	Health informa	ntion	II.a. Certificate reference No	II.b.		
				, declare that I have read and under			
ication	II.1. II.2.	10 thereof, and Annex XIII and has been prep competent auth	I Commission Regulation I Chapter II of Annex X ared and stored in an ority in accordance with	nent and of the Council( <sup>1a</sup> ), and in pa on (EU) No 142/2011( <sup>1b</sup> ), and in pa (IV), thereto and certify that the petfor establishment or plant approved a the Article 24 of Regulation (EC) No 1a efollowing animal by-products:	articular Chapter II of od described above: and supervised by the		
Part II: Certification		( <sup>2</sup> )either [-	carcases and parts of animals slaughtered or, in the case of game, bod parts of animals killed, and which are fit for human consumption in according with Union legislation, but are not intended for human consumption commercial reasons;]				
Part 1		( <sup>2</sup> )and/or [-	carcases and the following parts originating either from animals that have I slaughtered in a slaughterhouse and were considered fit for slaughter for hu consumption following an ante-mortem inspection or bodies and the follow parts of animals from game killed for human consumption in accordance Union legislation:				
	•		human consum	dies and parts of animals which are ption in accordance with Union leg- gns of disease communicable to hungs:	islation, but which did		
			(iii) hides and skins including the p metatarsus bon	, including trimmings and splitting t halanges and the carpus and metaca			
			(iv) pig bristles;				
		( <sup>2</sup> )and/or [-	referred to in Article	from poultry and lagomorphs slaug 1(3)(d) of Regulation (EC) No 853 the Council(2 <sup>a</sup> ), which did not show mans or animals	/2004 of the European		
		( <sup>2</sup> )and/or [-	blood of animals w through blood to hu slaughtered in a slau	hich did not show any signs of or mans or animals, obtained from an ghterhouse after having been consi- tion following an ante-mortem ins-	nimals that have been dered fit for slaughter		
	(2)and/or [- animal by-products arising fro			rising from the production of producing degreased bone, greaves and c			
		(²) and/or [- products of animal origin, or foodstuffs containing pr which are no longer intended for human consumption for due to problems of manufacturing or packaging defect which no risk to public or animal health arise;]			commercial reasons or		
	(2) and/or [- petfood and feedingstuffs of animal origin, or feedingstuffs of by-products or derived products, which are no longer intended commercial reasons or due to problems of manufacturing or products from which no risk to public or animal health a				tended for feeding for g or packaging defects		
		( <sup>2</sup> )and/or [-	blood, placenta, woo	l, feathers, hair, horns, hoof cuts and t did not show signs of any disease of	d raw milk originating		
		( <sup>2</sup> )and/or [-		parts of such animals, except sea ma eases communicable to humans or a			
		( <sup>2</sup> )and/or [-	animal by-products	from aquatic animals originat facturing products for human consun	ing from plants or		
		( <sup>2</sup> )and/or [-		al originating from animals which d	=		

COUNTRY Canned Petfood

II. Health information	II.a. Certificate reference No	II.b.
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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;]
- (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 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. has been subjected to heat treatment to a minimum Fc value of 3 in hermetically sealed containers;
- II.4. has undergone all precautions to avoid contamination with pathogenic agents after treatment.
- (2)[II.5. the petfood described above

 $(^2)$ or

- (<sup>2</sup>)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.]
    - [(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(<sup>4</sup>);
      - (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,
      - (c) animal by-product or derived product obtained from bovine, ovine or caprine 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 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 Canned Petfood** 

II. Health information II.a. Certificate reference No II.b.	II.	Health information	II.a. Certificate reference No	II.b.	
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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 in the European Union.
- 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 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, Ruminantia, Mammalia other than Ruminantia, Pesca, Mollusca, Crustacea, invertebrates other than Mollusca and Crustacea.

#### Part II:

- (<sup>1a</sup>) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.(2) Delete as appropriate.
- (<sup>2a</sup>) OJ L 139, 30.4.2004, p. 55.
- (2b) OJ L 125, 23.5.1996, p. 3.
- (<sup>3</sup>) 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 different colour to that of the printing.

for veterinary purposes and must accompany the opost.	1
Official veterinarian/Official inspector	
Name (in capital letters):	Qualification and title:
Date:	Signature:
Stamp:	

# (CHAPTER 3(B)

# Health certificate

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

	UNIKI:	veterinary certificate to EU
Part I : Details of dispatched consignment	I.1. Consignor Name	I.2. Certificate reference No I.2.a.
	Address	I.3. Central competent authority
	Tel.	I.4. Local competent authority
	I.5. Consignee Name Address  Postcode Tel.	I.6. Person responsible for the load in EU Name Address  Postcode Tel.
isp	I.7. Country of ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10. Region of Code
of d	origin	destination destination
ails	I.11. Place of origin	I.12. Place of destination
Part I : Deta	Name Approval number Address Name Approval number Address Name Approval number Address	Custom warehouse  Name Approval number Address  Postcode
	I.13. Place of loading	I.14. Date of departure
	I.15. Means of transport  Aeroplane  Road vehicle  Other  Other	I.16. Entry BIP in EU  I.17.
	Identification  Documentation references	
	I.18. Description of commodity	I.19. Commodity code (HS code)
		I.20. Quantity
	I.21. Temperature of product Ambient □ Chilled □	I.22. Number of packages
	I.23. Seal/Container No	I.24. Type of packaging
	I.25. Commodities certified for:	
	Petfood  Techn	nical use $\square$
	I.26. For transit through EU to third country	I.27. For import or admission into EU
	Third country ISO code	
	I.28. Identification of the commodities	
	Approval number of Species (Scientific name) Manufacturin	
	1	

	II.	Health informa	tion	II.a. Certificate reference No	II.b.			
	-	No 1069/2009 of 10 thereof, and	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( <sup>1a</sup> ), and in particular Articles 8 and 10 thereof, and Commission Regulation (EU) No 142/2011( <sup>1b</sup> ), and in particular Chapter II of Annex XIII and Chapter II of Annex XIV thereto, and certify that the petfood described above:					
tion	II.1.	has 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;						
<u> </u>	II.2.							
Part II: Certification		(²) either [- carcases and parts of animals slaughtered or, in the case of game, b of animals killed, and which are fit for human consumption in acc Union legislation, but are not intended for human consumption fo reasons;]						
Part ]		( <sup>2</sup> )and/or [-	slaughtered in a slaug consumption following	wing parts originating either from a hterhouse and were considered fit for a gan ante-mortem inspection or bod a game killed for human consumption	or slaughter for human lies and the following			
	(i) carcases or bodies and parts human consumption in accord			lies and parts of animals which are otion in accordance with Union legi- gns of disease communicable to hum	slation, but which did			
(iii)		(iii) hides and skins, including the pl	• •					
			<ul><li>(iv) pig bristles;</li><li>(v) feathers;</li></ul>					
	(²) and/or [- animal by-products from poultry and lagomorp referred to in Article 1(3)(d) of Regulation (EC Parliament and of the Council(²a), which did communicable to humans or animals]		1(3)(d) of Regulation (EC) No 853/2 e Council( <sup>2a</sup> ), which did not show	2004 of the European				
		( <sup>2</sup> )and/or [-	or [- blood of animals which did not show any signs of disease commun through blood to humans or animals, obtained from animals that have slaughtered in a slaughterhouse after having been considered fit for slau for human consumption following an ante-mortem inspection in according with Union legislation;]		imals that have been lered fit for slaughter			
		( <sup>2</sup> )and/or [-		rising from the production of producing degreased bone, greaves and ceressing;]				
(²)and/or [- products of animal origin, or foodstuf which are no longer intended for human due to problems of manufacturing or which no risk to public or animal health (²)and/or [- petfood and feedingstuffs of animal or by-products or derived products, which commercial reasons or due to problem or other defects from which no risk to products or derived products.		ntended for human consumption for chanufacturing or packaging defects	commercial reasons or					
		d products, which are no longer into the due to problems of manufacturing	ended for feeding for or packaging defects					
	(²)and/or [- blood, placenta, wool		, feathers, hair, horns, hoof cuts and t did not show signs of any disease c s or animals;]					
		( <sup>2</sup> )and/or [-		parts of such animals, except sea ma eases communicable to humans or an				
	(²)and/or [- animal by-products establishments manuf			from aquatic animals origination acturing products for human consum				
			l originating from animals which di	d not show any signs				

# Processed petfood other than canned petfood

COUNTRY				Trocessed pe	enood other than	
II. Health information			tion_	II.a. Certificat	te reference No	II.b.
			of di (i) (ii)	ease communicable through that shells from shellfish with soft tis the following originating from to hatchery by-products, eggs,	ssue or flesh;	or animals:
				- egg by-products, including		
	( <sup>2</sup> )and/or	[-	anim	day-old chicks killed for comme al by-products from aquatic or t genic to humans or animals;]		tes other than species
	( <sup>2</sup> )and/or	[-	animals and parts thereof of the zoological orders of Rodentia and Lagomorpha, except 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		(iii), (iv) and (v) of	
W 0	( <sup>2</sup> )and/or	[-	9(a) to (g) of that Regulation;] material from animals which have been treated with certain substances which are prohibited by Council Directive 96/22/EC( <sup>2b</sup> ), the import of the material being permitted in accordance with Article 35(a)(ii) of Regulation (EC) No 1069/2009;]			
II.3.	( <sup>2</sup> )either	[was	suhie	ted to a heat treatment of at least	t 90 °C throughout its	s substance:]
	$\binom{2}{0}$ or	[was	-	ced as regards ingredients of a	_	
		(a)		e case of animal by-products of a heat treatment	-	
		(b)	in th	case of milk and milk based pro	ducts,	
			(i) if they are from third countries or parts of third countries listed is column B of Annex I to Commission Regulation (EU) No 605/2010( submitted to a pasteurisation treatment sufficient to produce a negative		(EU) No 605/2010( <sup>3</sup> )	
			phosphatase test;  (ii) with a pH reduced to less than 6 from third countries or parts of thi countries listed in column C of Annex I to Regulation (EU) No 605/201 first submitted to a pasteurisation treatment sufficient to produce negative phosphatase test;		on (EU) No 605/2010,	
			(iii) if they are from third countries or parts of third countries listed in colu C of Annex I to Regulation (EU) No 605/2010, submitted to a sterilisat process or a double heat treatment where each treatment was sufficien produce a negative phosphatase test on its own;		nitted to a sterilisation	
			(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 achieved		al or greater than 3 is	
				or - an initial heat treatment vachieved by a pasteurisat 15 seconds and sufficient phosphatase test, followed	tion process of at le	ast 72 °C for at least
				either - a second heat treatment v achieved by the initial hea		

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

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;
- (i) in the case of non-mammalian processed protein with the exclusion of fishmeal submitted to any of the processing methods 1 to 5 or 7 as referred to in 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;

# Processed petfood other than canned petfood

II.	Health inf	orma	tion		II.a. Certificate reference No	II.b.
	(l) in the case of dicalcium phosphate produced by a process that			hat		
	with hot water			ith hot water oncentration o	Category 3 bone-material is finely c and treated with dilute hydrochloric of 4 % and a pH of less than 1,5) over	acid (at a minimum
			ob	tained phosp	procedure referred to in (i), applie phoric liquor with lime, resulting phate at pH 4 to 7; and	
			te		ies the precipitate of dicalcium p 65 °C to 325 °C and end temperatur	
		(m)	in the ca	ase of tricalciu	am phosphate produced by a process	that ensures
					ory 3 bone-material is finely crush ith hot water (bone chips less than 14	
			(ii) co	ntinuous cool	king with steam at 145 °C during 30 i	minutes at 4 bar;
				•	the protein broth from the hydro centrifugation; and	xyapatite (tricalcium
				anulation of to 200 °C;	he tricalcium phosphate after drying	in a fluid bed with air
		(n)	paramet		ing innards, produced according to a usure that the product complies with in point II.4.1	
	$(^2)or$		subject t	o a treatment	such as drying or fermentation, which	h has been authorised
	2	•	-	ent authority;		
	( <sup>2</sup> ) <i>or</i>	[in the case of aquatic and terrestrial invertebrates other than species pathogenic to humans 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;]				
II.4.	•		4			
Salmonella: absence in 25g: $n = 5$ , $c = 0$ , in		n = 5, c = 0, m = 0, M = 0,				
Enterobacteriaceae: $n = 5$ , $c = 2$ , $m = 10$ , $M = 300$ in 1 gramme;						
II.5. has undergone all precautions to avoid contamination with pathogenic agents after treatment;		s 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						
	( <sup>2</sup> )eit	her [i	s derived	from other ru	minants than bovine, ovine or caprine	e animals.]
(²) or [is derived from bovine, ovine or capaderived from:		, ovine or caprine animals and does	not contain and is not			
		( <sup>2</sup>	either	animals boregion class	vine and caprine materials other that orn, continuously reared and slaught sified as posing a negligible BSE ris 007/453/EC.]	tered in a country or
		( <sup>2</sup>	e)or		ied risk material as defined in poi (EC) No 999/2001 of the European	
					nically separated meat obtained from	
	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				in a country or region k in accordance with	

II.	Health information	II.a. Certificate reference No	II.b.
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no indigenous BSE case,

(c) animal by-product or derived product obtained from bovine, ovine or caprine 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 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) 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 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, ruminantia, mammalia other than ruminantia, pesca, mollusca, crustacea, invertebrates other than mollusca and crustacea.

#### Part II:

- <sup>(1a)</sup> OJ L 300, 14.11.2009, p. 1.
- <sup>(1b)</sup> OJ L 54, 26.2.2011, p. 1.
- Delete as appropriate.
- (<sup>2a</sup>) OJ L 139, 30.4.2004, p. 55.(<sup>2b</sup>) OJ L 125, 23.5.1996, p. 3.
- <sup>(3)</sup> OJ L 175, 10.7.2010, p. 1.
- (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 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.
- (5) OJ L 147, 31.5.2001, p. 1.
- <sup>(6)</sup> 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 entry into the European Union.

# Processed petfood other than canned petfood

II. Health information	II.a. Certificate reference No II.b.				
Official veterinarian/Official inspector	Official veterinarian/Official inspector				
Name (in capital letters):	Qualification and title:				
Date:	Signature:				
Stamp:					

# CHAPTER 3(C)

# Health certificate

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

CC	OUNTRY:	Veterinary certificate to EU		
	I.1. Consignor	I.2. Certificate reference No I.2.a.		
Part I: Details of dispatched consignment	Name Address	I.3. Central competent authority		
	Tel.	I.4. Local competent authority		
	I.5. Consignee Name Address  Postcode Tel.	I.6. Person responsible for the load in EU Name Address  Postcode Tel.		
f dispa	I.7. Country of origin ISO code I.8. Region of origin Code origin	I.9. Country of ISO code I.10. Region of Code destination		
ils o	III Non-ef-vicin	L12 Phos of destination		
Part I : Detai	I.11. Place of origin  Name Approval number Address Name Approval number Address Name Approval number	I.12. Place of destination  Custom warehouse  Name Approval number Address  Postcode		
	Address			
	I.13. Place of loading	I.14. Date of departure		
	I.15. Means of transport	I.16. Entry BIP in EU		
	Aeroplane	I.17.		
	Documentation references I.18. Description of commodity	I.19. Commodity code (HS code)		
		I.20. Quantity		
	I.21. Temperature of product Ambient □ Chilled □	I.22. Number of packages		
	I.23. Seal/Container No	I.24. Type of packaging		
	I.25. Commodities certified for:			
	Petfood  Tech	nical use		
	I.26. For transit through EU to third country	I.27. For import or admission into EU		
	Third country ISO code			
	I.28. Identification of the commodities			
	Species (Scientific name)  Approval number of e Manufacturing			

COUNTRY Dogchews

#### 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(<sup>1a</sup>), and in particular Article 10 of that Regulation, and Commission Regulation (EU) No 142/2011(<sup>1b</sup>), and in particular Chapter II of Annex XIII and Chapter II of Annex XIV thereto, and certify that the dogchews described above:

- II.1. have 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 ante-mortem 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 [- 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;]
  - (<sup>2</sup>)and/or [- aquatic animals, and parts of such animals, expect sea mammals, which did not show any signs of disease communicable to humans or animals;]
  - (²) and/or [- animal by-products from aquatic animals originating from plants or establishments manufacturing products for human consumption;]
  - (²) and/or [- material from animals which have been treated with certain substances which are prohibited by Council Directive 96/22/EC(²a), the import of the material being permitted in accordance with Article 35(a)(ii) of Regulation (EC) No 1069/2009;]
- II.2. have been subjected
  - (²)either [in the case of dogchews made from hides and skins of ungulates or from fish, to a treatment sufficient to destroy pathogenic organisms (including salmonella); and the dogchews are dry;]
  - (²) and/or [in the case of dogchews made from animal by-products other than hides and skins of ungulates or from fish, to a heat treatment of at least 90°C throughout their substance;]
- II.3. were examined by random sampling of at least five samples from each processed batch taken during or after storage at the processing plant and complies with the following standards(3):

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

Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gramme;

II.4. have undergone all precautions to avoid contamination with pathogenic agents after treatment;

COUNTRY Dogchews

II. H	ealth information	II.a.	Certificate reference No		II.b.
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### II.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.]

 $(^{2})or$ 

- [(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(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,
- (c) animal by-product or derived product obtained from bovine, ovine or caprine 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 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); 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 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, Ruminantia, Mammalia Other Than Ruminantia, Pesca, Mollusca, Crustacea, Invertebrates Other Than Mollusca And Crustacea.

### Part II:

- <sup>(1a)</sup> OJ L 300, 14.11.2009, p. 1.
- <sup>(1b)</sup> OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.
- (<sup>2a</sup>) 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

**COUNTRY** Dogchews

II.	I. Health information		II.a.	Certificate reference	No	II.b.
		number of bacteria in all san	nples	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					
		number of samples the bactostill being considered accept		•		
(4)	OJ L 147	, 31.5.2001, p. 1.				
(5)	OJ L 172,	30.6.2007, p. 84.				
_	<ul> <li>The signature and the stamp must be in a different colour to that of the printing.</li> <li>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 entry into the European Union.</li> </ul>					
Offi	Official veterinarian/Official inspector					
	Name (in capital letters):				Qualific	ation and title:
Date: Signature:			re:			
	Stamp:					

# 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

CC	OUNTRY:	Veterinary certificate to EU		
	I.1. Consignor	I.2. Certificate reference No I.2.a.		
	Name Address	I.3. Central competent authority		
ent	Tel.	I.4. Local competent authority		
Part I: Details of dispatched consignment	I.5. Consignee Name Address  Postcode Tel.	I.6. Person responsible for the load in EU Name Address Postcode Tel.		
	I.7. Country of ISO code I.8. Region of origin Code origin	I.9. Country of ISO code I.10. Region of Code destination destination		
ails	I.11. Place of origin	I.12. Place of destination		
Part I: Det	Name Approval number Address Name Approval number Address Name Approval number Address	Custom warehouse  Name Approval number  Address  Postcode		
	I.13. Place of loading	I.14. Date of departure		
	I.15. Means of transport	I.16. Entry BIP in EU		
	Aeroplane Ship Railway wagon Road vehicle Other Identification Documentation references	I.17.		
	I.18. Description of commodity	I.19. Commodity code (HS code)		
		I.20. Quantity		
	I.21. Temperature of product	I.22. Number of packages		
	Ambient ☐ Chilled ☐  I.23. Seal/Container No	Frozen   I.24. Type of packaging		
	I.25. Commodities certified for:			
	Petfood □ Tech	nical use		
	I.26. For transit through EU to third country	I.27. For import or admission into EU		
	Third country ISO code			
	I.28. Identification of the commodities	<u> </u>		
	Δ	pproval number of establishments		
	Species (Scientific name) Nature of commodity	Manufacturing plant Net weight Batch number		

# 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(<sup>1a</sup>) and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011(<sup>1b</sup>), 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:
    - Commission Regulation (EU) No 206/2010(<sup>3</sup>) 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);
    - and/or Commission Regulation (EC) No 798/2008(<sup>4</sup>), 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(<sup>5</sup>), 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 killed 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(<sup>6</sup>); 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(<sup>7</sup>), 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

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

II.	Health inform	ation	II.a. Certificate reference No	II.b.
		slaughtered in a s	humans or animals, obtained from a claughterhouse after having been consemption following an ante-mortem instation;]	idered fit for slaughter
	( <sup>2</sup> )and/or [-	human consumpt	cts arising from the production of gion, including degreased bone, greafrom milk processing;]	-
	( <sup>2</sup> )and/or [-	which are no long or due to problem	al origin, or foodstuffs containing proger intended for human consumption for sof manufacturing or packaging deak to public or animal health arises;]	or commercial reasons
	( <sup>2</sup> )and/or [-	petfood and feedi by-products or de commercial reas	ngstuffs of animal origin, or feedingst crived products, which are no longer ir ons or due to problems of manufa efects from which no risk to public or	ntended for feeding for acturing or packaging
	( <sup>2</sup> )and/or [-	blood, placenta, originating from	wool, feathers, hair, horns, hoof live animals that did not show arough that product to humans or animals.	cuts and raw milk signs of any disease
	( <sup>2</sup> )and/or [-		and parts of such animals, except seans of diseases communicable to human	
	( <sup>2</sup> )and/or [-		acts from aquatic animals origina anufacturing products for human cons	
	( <sup>2</sup> )and/or [-	the following mat	terial originating from animals which on the control of the contro	did not show any signs
		` '	shellfish with soft tissue or flesh;	
		(ii) the following	ng originating from terrestrial animals:	
		- hatch	ery by-products,	
		- eggs,		
			y-products, including egg shells,	
	.2	•	cks killed for commercial reasons;]	
	( <sup>2</sup> )and/or [-	animal by-produce pathogenic to hur	ets from aquatic or terrestrial invertebre nans or animals;]	ates other than species
	( <sup>2</sup> )and/or [-	Lagomorpha, exc (iv) and (v) of R	rts thereof of the zoological orderept Category 1 material as referred degulation (EC) No 1069/2009 and Cocle 9(a) to (g) of that Regulation;]	to in Article 8(a)(iii),
II.4.	with the condi-	tions laid down in t	without contact with other material whe Regulation (EC) No 1069/2009, an outhogenic agents:	
II.5.	so as to avoid contamination with pathogenic agents; have been packed in final packaging which bear labels indicating 'RAW PET FOOD — NOT FOR HUMAN CONSUMPTION' or 'ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS — NOT FOR HUMAN CONSUMPTION' and then placed in leak-proof and officially sealed boxes/containers or in new packaging preventing any leakage and officially sealed boxes/containers which bear labels indicating 'RAW PET FOOD — NOT FOR HUMAN CONSUMPTION' or 'ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS — NOT FOR HUMAN CONSUMPTION', and the name and the address of the establishment of destination;			
II.6.	in the case of ra	aw petfood:		
	(a) has been	n prepared and sto	red in a plant approved and supervi a Article 24 of Regulation (EC) No 100	
	(b) was exa	mined by random	sampling of at least five samples fr	rom each batch taken

II.	Health information	II.a.	Certificate reference No	II.b.
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during storage (before dispatch) and complies with the following standards(8):

Salmonella: absence in 25 g: n=5, c=0, m=0, M=0 Enterobacteriaceae: n=5, c=2, m=10, M=5000 in 1 gram;

(²)II.7. [the processed animal protein or product described above contains or is derived from animal-by products of ruminant origin and:

(²)either [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 indigenous BSE case.]

(²) or [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 an indigenous BSE case, and the animal by-product or derived product were 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.]

and

(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.]

 $(^2)or$ 

- [(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(9);
- (b) mechanically separated meat obtained from bones of bovine, ovine or caprine animals, except from 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(10), in which there has been no indigenous BSE case,
- (c) animal by-product or derived product obtained from bovine, ovine or caprine 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 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

# 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.	
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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 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:

Nature of commodity: select raw petfood or animal by-product.

In the case of raw material for the manufacture of raw pet food indicate the scientific name of the species.

In case of raw material for manufacture of feed for fur animals select from the following: Aves, Ruminantia, Mammalia Other Than Ruminantia, Pesca, Mollusca, Crustacea, Invertebrates other than Mollusca And Crustacea.

#### Part II:

- (<sup>1a</sup>) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p.1
- (2) Delete as appropriate.
- (<sup>2a</sup>) OJ L 139, 30.4.2004, p. 55.
- (<sup>3</sup>) OJ L 73, 20.3.2010, p. 1.
- (4) OJ L 226, 23.8.2008, p. 1.
- <sup>(5)</sup> OJ L 39, 10.2.2009, p. 12.
- (<sup>6</sup>) OJ L 303, 18.11.2009, p. 1.
- (<sup>7</sup>) 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; 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.
- (<sup>9</sup>) OJ L 147, 31.5.2001, p. 1.
- (<sup>10</sup>) 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.

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

II.a. Certificate reference No	II.b.		
Official veterinarian/Official inspector			
Qual	ification and title:		
Sign	ature:		
	Qual		

### 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

CO	UNTRY:	Veterinary certificate to EU		
	I.1. Consignor	I.2. Certificate reference No I.2.a.		
	Name Address	I.3. Central competent authority		
ınt	Tel.	I.4. Local competent authority		
Part I: Details of dispatched consignment	I.5. Consignee Name Address  Postcode Tel.	I.6. Person responsible for the load in EU Name Address  Postcode Tel.		
of dispa	I.7. Country of origin ISO code I.8. Region of origin Code origin	I.9. Country of ISO code I.10. Region of destination destination		
ails	I.11. Place of origin	I.12. Place of destination		
Part I: Det	Name Approval number Address Name Approval number Address Name Approval number Address	Custom warehouse  Name Approval number Address  Postcode		
	I.13. Place of loading	I.14. Date of departure		
	I.15. Means of transport	I.16. Entry BIP in EU		
	Aeroplane	I.17.		
	I.18. Description of commodity	I.19. Commodity code (HS code)		
		I.20. Quantity		
	I.21. Temperature of product Ambient □ Chilled □	I.22. Number of packages		
	I.23. Seal/Container No	I.24. Type of packaging		
	I.25. Commodities certified for:	•		
	Petfood □ Tech	nical use		
	I.26. For transit through EU to third country	I.27. For import or admission into EU		
	Third country ISO code			
	I.28. Identification of the commodities	<u>I</u>		
	Species (Scientific name)  Nature of commodity	Approval number of establishments  Manufacturing plant  Net weight  Batch number		

#### 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 8 and 10 thereof, and Commission Regulation (EU) No 142/2011(1b), and in particular Chapter III of Annex XIII and Chapter II of Annex XIV thereto, and certify that the flavouring innards products described above: Part II: Certification II.1. consist of animal by-products that satisfy the animal health requirements below; II.2. have been prepared and include the following animal by-products which are exclusively: carcases and parts of animals slaughtered or, in the case of game, bodies or (<sup>2</sup>)either 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 ante-mortem 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; (iv) pig bristles; (v) feathers;] $(^{2})$ 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; ] $(^2)$ and/or [products of animal origin, or foodstuffs containing products of animal origin, which are no longer 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:1 $(^2)$ and/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 arise;] (<sup>2</sup>)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;] (<sup>2</sup>)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;] (<sup>2</sup>)and/or animal by-products from aquatic animals originating from plants or establishments manufacturing products for human consumption;] (<sup>2</sup>)and/or the following material originating from animals which did not show any

# Flavouring innards for use in the manufacture of petfood

II.	Health inforn	nation	II.a. Certificate reference No	II.b.	
		<ul><li>(i) shells from s</li><li>(ii) the following</li></ul>	mmunicable through that material thellfish with soft tissue or flesh; goriginating from terrestrial animaly by-products,		
		<ul> <li>eggs,</li> <li>egg by-products, including egg shells;</li> <li>(iii) day-old chicks killed for commercial reasons;]</li> </ul>			
	( <sup>2</sup> )and/or [-	• •	animal by-products from aquatic or terrestrial invertebrates other than species pathogenic to humans or animals;] animals and parts thereof of the zoological orders of Rodentia and Lagomorpha, except 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 [-	Lagomorpha, exce (iv) and (v) of Reg			
	( <sup>2</sup> )and/or [-	which are prohibit	material from animals which have been treated with certain substances which are prohibited by Council Directive 96/22/EC( <sup>2a</sup> ), the import of the material being permitted in accordance with Article 35(a)(ii) of Regulation		
II.3.			g in accordance with Chapter I der to kill pathogenic agents;	II of Annex XIII to	
II.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( $^3$ ):  Salmonella: absence in 25g: n = 5, c = 0, m = 0, M = 0,  Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gramme;				
II.5.	the end produc		iii = 10, 141 = 300 iii 1 grainiie,		
	(²)either [pa				
	(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 bea	ar labels indicating 'No	OT FOR HUMAN CONSUMPTIO	N';	
II.6.		ct was stored in enclos	•		
II.7.	the product har		cautions to avoid contamination w	ith pathogenic agents	
$(^{2})II.8.$	the flavouring	innards products desc	cribed above		
			ninants than bovine, ovine or capring ovine or caprine animals and does		
		ved from:	ovine of captine animals and does	not contain and is not	
		ither [bovine, ovi animals born	ne and caprine materials other than, continuously reared and slaugh fied as posing a negligible BSE ri 07/453/EC.]	tered in a country or	
	( <sup>2</sup> )01	- · · · -	d risk material as defined in poi EC) No 999/2001 of the European		
		(b) mechan ovine of born, co classifie Commis	ically separated meat obtained from the caprine animals, except from the ontinuously reared and slaughtered dot as posing a negligible BSE rissesion Decision 2007/453/EC(5), in tenous BSE case,	ose animals that were in a country or region k in accordance with	
			by-product or derived product of caprine animals which have been		

# Flavouring innards for use in the manufacture of petfood

II. Hea	alth information	II.a. Certificate reference No	II.b.
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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 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); information is to be provided in the event of unloading and reloading in the European Union.
- Box reference I.19: use the appropriate HS code: 05.04; 05.06 or 05.11.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be given.
- 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, ruminantia, mammalia other than ruminantia, pesca, mollusca, crustacea, invertebrates other than mollusca and crustacea
  - define the innard product.

### Part II:

- (<sup>1a</sup>) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.
- (<sup>2a</sup>) OJ L 125, 23.5.1996, p. 3.
- (<sup>3</sup>) 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.
- (<sup>5</sup>) 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.

# Flavouring innards for use in the manufacture of petfood

II. Health information	II.a. Certificate reference No II.b.		
Official veterinarian/Official inspector			
Name (in capital letters):	Qualification and title:		
Date:	Signature:		
Stamp:			

# CHAPTER 3(F)

# Health certificate

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

CO	OUNTRY:	Veterinary certificate to EU
	I.1. Consignor	I.2. Certificate reference No I.2.a.
	Name Address	I.3. Central competent authority
nt	Tel.	I.4. Local competent authority
Part I: Details of dispatched consignment	I.5. Consignee Name Address  Postcode Tel.	I.6. Person responsible for the load in EU Name Address  Postcode Tel.
of dispa	I.7. Country of ISO code I.8. Region of origin Code origin	I.9. Country of ISO code I.10. Region of Code destination
ails	I.11. Place of origin	I.12. Place of destination
art I : Det	Name Approval number Address Name Approval number	Custom warehouse  Name Approval number Address
P	Address Name Approval number Address	Postcode
	I.13. Place of loading	I.14. Date of departure
	I.15. Means of transport	I.16. Entry BIP in EU
	Aeroplane ☐ Ship ☐ Railway wagon ☐	
	Road vehicle Other Identification	I.17.
	Documentation references  I.18. Description of commodity	I.19. Commodity code (HS code)
		I.20. Quantity
	I.21. Temperature of product Ambient □ Chilled □	I.22. Number of packages
	I.23. Seal/Container No	I.24. Type of packaging
	I.25. Commodities certified for:	4
	Manufacture of petfood $\square$ Furth	er process $\square$ Technical use $\square$
	I.26. For transit through EU to third country	I.27. For import or admission into EU
	Third country ISO code	
	I.28. Identification of the commodities	
	Approval n	number of establishments
	**	nufacturing plant Number of packages Net weight Batch number

# Animal by-products for the manufacture of petfood

				T					
	II. Health information		II.a. Certificate reference No	II.b.					
Part II: Certification	II.1.1. II.1.2.	(EC) No 106 Regulation (I certify that the consist of ani have been ob (2)either [ (2)or [	59/2009 of EU) No 14. The animal by imal by-proportion of the three in	al veterinarian, declare that I have the European Parliament and of the 2/2011(1b), and in particular Chaptey-products described above: ducts that satisfy the animal health the territory of:  ave remained in this territory since months preceding the date of slauging the wild in this territory(1d);]	the Council( <sup>1a</sup> ) and Commission ter II of Annex XIV thereto, and a requirements below;( <sup>1c</sup> ) from animals: the birth or for a period of at least there or production;				
II:		$(^2)or$ [		aquatic invertebrates;]					
art	II.1.3.	have been ob	tained fron	or produced by animals:					
P		(²)either [	(a) comin	g from holdings:					
			(i)	where, for the following diseas susceptible, there has been no case vesicular disease, Newcastle dise influenza during the period of classical or African swine fever du 40 days; nor in the holdings situated km radius, during the period of the	se/outbreak of rinderpest, swine case or highly pathogenic avian the preceding 30 days, nor of uring the period of the preceding sted in their vicinity within a 10				
			(ii) where there has been no case/outbreak of foot-and during the period of the preceding 60 days, nor situated in their vicinity within a 25 km radius, du of the preceding 30 days; and		ng 60 days, nor in the holdings				
		(	b) which	:					
			(i)	were not killed to eradicate any ep	izootic disease;				
			(iii)	have remained in their holdings of days before the date of departure a directly to the slaughterhouse we animals which did not comply with at the slaughterhouse, have particularly dispersion during the period of a slaughter and have shown no evice	and which have been transported without any contact with other h the same health conditions; assed the ante-mortem health 24 hours preceding the time of				
			(iv)	have been handled in the slaughte slaughter or killed in accordance Union legislation and have met rethose laid down in Chapters II and	rhouse before and at the time of with the relevant provisions of quirements at least equivalent to				
		(2) or [		No 1099/2009( <sup>4</sup> )] ed and killed in the wild in an area	3.				
		( <sup>2</sup> )or [	(i)	ed and killed in the wild in an area in which within a 25 km radius the any of the following diseases susceptible: foot-and-mouth disease or highly pathogenic avian influenceding 30 days, nor of classical the state of the second state of the seco	ere has been no case/outbreak of for which the animals are se, rinderpest, Newcastle disease tenza during the period of the all or African swine fever during				
		(	(ii)	the period of the preceding 40 day situated at a distance of at least 20 the territory of a country not author Union of poultry material during porcine material during the preced after killing were transported	0 km from any country or part of orised for export to the European g the preceding 30 days or of ing 40 days; and				
				ring the killing for chilling eith					

# Animal by-products for the manufacture of petfood

II.	Health inform	nation	II.a. Certificate reference No II.b.				
			mmediately afterwards to a game handling establishment, or directly to				
П.1.4.	a game handling establishment;]  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.	comply with	h the	d and prepared without contact with any other material that does no conditions required above, and it has been handled so as to avoid a pathogenic agents;				
II.1.6.	containers MANUFAC	bearin TURE	I in new packaging preventing any leakage and in officially sealed g the label indicating 'RAW MATERIAL ONLY FOR THE OF PET FOOD' and the name and address of the establishment of European Union;				
II.1.7.	consist only	of 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 which were deemed fit for human consumption in accordance with Union legislation until irreversibly declared as animal by-products for commercial reasons;]						
	( <sup>2</sup> )and/or	(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 ante-mortem inspection or bodies and the following parts of animals from game killed for human consumption in accordance with Union legislation:					
		(	<ol> <li>carcases or bodies and parts of animals which are rejected as unfi for human consumption in accordance with Union legislation, bu which did not show any signs of disease communicable to humans or animals;</li> </ol>				
		(	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;				
		(	pig bristles;				
	2		(v) feathers;]				
	( <sup>2</sup> )and/or	ŀ	animal by-products arising from the production of products intended for numan consumption, including degreased bone, greaves and centrifuge or separator sludge from milk processing;]				
	( <sup>2</sup> )and/or	(²) and/or [- products of animal origin, or foodstuffs containing products of animal origin, which are no longer 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;]					
	( <sup>2</sup> )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 aquatic animals originating from plants o establishments manufacturing products for human consumption;]				
	( <sup>2</sup> )and/or		the following material originating from animals which did not show any signs of disease communicable through that material to humans of animals:				
		(	i) shells from shellfish with soft tissue or flesh;				

# Animal by-products for the manufacture of petfood

II. I	Health in	formatio	on	II.a. Certificate	reference No	II.b.
			(ii) t	he following orig	inating from ter	restrial animals:
			-	hatchery by-p	oroducts,	
			-	eggs,		
<ul> <li>egg by-products, including egg shells;</li> </ul>						
			(iii) (	day-old chicks kil	led for commerc	cial reasons;]
	(2) and/or [- animal by-products from aquatic or terrestrial invertebrates, other than					
	2		_	s pathogenic to hu		
	( <sup>2</sup> )and/o	or [-	Lagon 8(a)(iii	norpha, except (i), (iv) and (v) of	Category 1 mat Regulation (EC	logical orders of Rodentia and erial as referred to in Article (2) No 1069/2009 and Category 2 (g) of that Regulation;]
	( <sup>2</sup> )and/o	or [-	which the m	are prohibited by	y Council Direc mitted in accor	n treated with certain substances tive 96/22/EC( <sup>4a</sup> ), the import of dance with Article 35(a)(ii) of
II.1.8.	Europea	an Unior	n legislat		y that they will	en preserved in accordance with not spoil between dispatch and
II.1.9.	substan	ces proh	ibited by	yDirective 96/22/	EC for the mar	have been treated with certain nufacture of petfood, the import gulation (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, 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;					r activated carbon on each outer al is transported in pallets which transport to the petfood plant of ide of each pallet, in a way that
	th w	ird coun	ntry befo	re entry into the	territory of the lying charcoal	material has been marked in the European Union by spraying it powder in such a way that the
	re	eferred to	above a		ted raw material	aterial which has been treated as I, all the raw materials have been
$(^{2})(^{5})[II.2.$	Specific	e require	ments			
( <sup>2</sup> )( <sup>6</sup> )II.2.1.	(2)(6)II.2.1. The by-products in this consignment come from animals that have been kept in the territory referred to in point (II.1.2), where vaccination programmes against foot-and-mouth disease are being regularly carried out and officially controlled in domestic bovine animals.					
$(^{2})(^{7})II.2.2.$						nimal by-products derived from
	trimmed offal of domestic ruminants, which have maturated at an ambient temperature of more than + 2 °C for a period of at least three hours, or in the case of masseter muscles of bovine animals and deboned meat of domestic animals, for a period of at least 24 hours.]					the case of masseter muscles of
( <sup>2</sup> )II.3. [the						tains or is derived from animal-
	products		-	-		
	( <sup>2</sup> )e	neg	gligible I	BSE risk in accor	dance with Deci	hich is classified as posing a sion 2007/453/EC, and in which
	there has been no indigenous BSE case.]  (2) or [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 an indigenous BSE case, and the animal by-product or derived product were					

# Animal by-products for the manufacture of petfood

II.	Health information	II.a. Certificate reference No II.b.			
	of ruminan as defined	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.]			
	and				
	_	from other ruminants than bovine, ovine or caprine animals.]			
	(²)or [is derived not derived	from bovine, ovine or caprine animals and does not contain and is I from:			
	( <sup>2</sup> ) 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.]			
	( <sup>2</sup> )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(8);			
		(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( <sup>9</sup> ), in which there has been no indigenous BSE case,			
		(c) animal by-product or derived product obtained from bovine, ovine or caprine 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 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: 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 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, Ruminantia, Mammalia other than Ruminantia,

# Animal by-products for the manufacture of petfood

II.	Health information	II.a. Certificate reference No	II.b.
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Pesca, Mollusca, Crustacea, invertebrates other than Mollusca and Crustacea;

- Manufacturing plant: provide the veterinary control number of the approved establishment.

#### Part II:

- (<sup>1a</sup>) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (1c) The name and ISO code number of the exporting country as laid down in:
  - Part 1 of Annex II to Regulation (EU) No 206/2010;
  - Part 1 of Annex I to Regulation (EC) No 798/2008, and
  - Part 1 of Annex I to Regulation (EC) No 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 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 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.

Official veterinarian/Official inspector					
Name (in capital	letters):	Qualification and title:			
Date:		Signature:			
Stamp:					

"

(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

CC	DUNTRY:	Veterinary certificate to EU
	I.1. Consignor	I.2. Certificate reference No I.2.a.
	Name Address	I.3. Central competent authority
ınt	Tel.	I.4. Local competent authority
Part I: Details of dispatched consignment	I.5. Consignee Name Address  Postcode Tel.	I.6. Person responsible for the load in EU Name Address  Postcode Tel.
of dispa	I.7. Country of origin ISO code I.8. Region of origin Code origin	I.9. Country of destination ISO code I.10. Region of destination Code
ails	I.11. Place of origin	I.12. Place of destination
Part I : Det	Name Approval number Address Name Approval number Address Name Approval number	Custom warehouse  Name Approval number Address  Postcode
	Address	
	I.13. Place of loading	I.14. Date of departure
	I.15. Means of transport  Aeroplane □ Ship □ Railway wagon □	I.16. Entry BIP in EU
	Road vehicle ☐ Other ☐ Identification  Documentation references	I.17.
	I.18. Description of commodity	I.19. Commodity code (HS code)
		I.20. Quantity
	I.21. Temperature of product Ambient □ Chilled □	I.22. Number of packages Frozen □
	I.23. Seal/Container No	I.24. Type of packaging
	I.25. Commodities certified for:	
	Animal feedingstuff □ Man	ufacture of petfood $\square$ Technical use $\square$
	I.26. For transit through EU to third country	I.27. For import or admission into EU
	Third country ISO code	
	I.28. Identification of the commodities	·
	Species (Scientific name) Nature of commodity	Approval number of establishments  Manufacturing plant  Batch number

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

	II.	Health informa	tion	II.a.	Certificate refere	ence No	II.b.	
		(EC) No 1069/2	2009 of the Eur	opean		of the Counci	inderstood Regulation l( <sup>1a</sup> ) and Commission	
	II.1.				e health requirem	•	escribed above.	
u	II.2.		-	•	ot intended for hu		tion;	
icatio	II.3.	have been prep	ared and stored	l in a		and supervis	ed by the competent	
tifi.	II.4.	=			e following anima			
Part II: Certification		ac	ther [blood of slaughtered animals, which is fit for human consumption in accordance with Union legislation, but which is not intended for human consumption for commercial reasons;]					
Part		co ar de w	•					
	II.5.	in order to inactivate pathogenic agents, have been submitted						
		(2) either [to processing in accordance with processing method(3) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011;]						
		(2) or [to a method and parameters which ensure that the product complies with microbiological standards set out in Chapter I of Annex X to Regulation (F No 142/2011;]						
		(²) or [in the case of blood products, including spray dried blood and blood plass of porcine origin intended for the feeding of porcine animals, to a lateratment at a temperature of at least 80°C throughout the substance and dry blood and blood plasma does not contain more than 8% w/w moisture was a water activity (Aw) of less than 0,60.]				e animals, to a heat the substance and the		
	II.6.	the end product	was:					
		(²)either [p	acked in new or	sterilis	sed bags;]			
		th		ed and	l disinfected with		f transport that were ant approved by the	
			_		FOR HUMAN C	CONSUMPTIO	N';	
	II.7.	•	was stored in end		•			
	II.8.	after treatment;					rith pathogenic agents	
		(²) and [in the case of blood products, including spray dried blood and blood plast of porcine origin intended for the feeding of porcine animals, has been stor in dry warehouse conditions under room temperature for a period of at least weeks.]				imals, has been stored		
	II.9.	have been examined prior to dispatch under the responsibility of the competent authority be taking a random sample during or on removal from storage which was found to comply with the following standards(4):						
		Salmonella:		_	n = 5, c = 0, m =			
		Enterobacteriace	eae: $n = 5, c =$	2, m =	= 10, M = 300 in	1 gram;		
	( <sup>2</sup> )II.10.	the blood produc	cts described abo	ve				
	/11.10.				ints than bovine,	ovine or caprin	e animals.]	
		•					not contain and is not	

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

II.	Health info	ormation		II.a.	Certificate reference No	II.b.
	d	erived fror	n:	•		
	(2	<sup>2</sup> ) 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	<sup>2</sup> )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(4);			
			ovine born, class Com	e or ca , continified a missio	ly separated meat obtained finding animals, except from the nuously reared and slaughtered as posing a negligible BSE rin Decision 2007/453/EC(5), in bus BSE case,	ose animals that were in a country or region sk in accordance with
			ovine by la elong cavit for t slaug	e or ca aceration gated y, or b those thtered	product or derived product of prine animals which have been on of the central nervous timed-shaped instrument intrody means of gas injected into the animals that were born, continuated in a country or region of the accordance with Desire the product of the prod	a killed, after stunning, assue by means of an uced into the cranial e cranial cavity, except attinuously reared and assified as posing a
II.11.	the blood p	roducts or	_	-		
	(²)either				milk products of ovine or cap	_
	.2.				farmed animals, other than fur	-
	$(^2)or$	intende	d for feed f	or farn	products of ovine or caprine ned animals, other than fur anim	nals, which:
				ly sinc	ovine and caprine animals to birth in a country where the	
			(i) cla	assical	scrapie is compulsorily notifia	ble;
					eness, surveillance and monito ical scrapie;	ring system is in place
			an	nimals	restrictions apply to holdings in the case of a suspicion of T cal scrapie;	_
					nd caprine animals affected wind destroyed;	th classical scrapie are
			m Co ru th	eal or ode of minan	ing to ovine and caprine anir greaves, as defined in the Ter the World Organisation for A t origin has been banned and le country for a period of at lea	restrial Animal Health nimal Health (OIE), of effectively enforced in
			=	rom ho	oldings where no official restriction	ctions are imposed due
		(c)	originate fi diagnosed	rom he	oldings where no case of class the period of at least the pre- firmation of a case of classical	ceding seven years or,
			(²)either [a	ll ovin	e and caprine animals on the horoyed or slaughtered, except for	olding have been killed

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

II.	Health information		II.a. Certificate reference No	II.b.
		A ca	RR/ARR genotype, breeding ewes RR allele and no VRQ allele and arrying at least one ARR allele;]	other ovine animals
	( <sup>2</sup> )or	bo fo CC T th m R	all animals in which classical scrapic een killed and destroyed, and the hold or a period of at least two year confirmation of the last classical scrap SE monitoring, including testing with the presence of TSE in accordance tethods set out in point 3.2 of Chap egulation (EC) No 999/2001, of all of which are over the age of 18 months, extra ARR/ARR genotype:  animals which have been slat consumption; and animals which have died or been kill which were not killed in the frat eradication campaign.]]	ing has been subjected is since the date of pie case to intensified the negative results for with the laboratory of the following animals accept ovine animals of the laboratory of the following animals accept ovine animals of the laboratory of the following animals of the laboratory of the following animals of the laboratory of the la
II.12.			ove contain or are derived from animaing to the statement of the Consignor r	
	( <sup>2</sup> )either [not intende animals.]	d for	the production of feed for farmed as	nimals, other than fur
	than fur anii inspection p carried out	nals, ost o in a	production of feed for non-ruminant and the Consignor has undertaken to f entry will be provided with the recordance with the methods set calation (EC) No 152/2009(8).]	ensure that the border esults of the analyses
Notes Port I				

### 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 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, Ruminantia, Mammalia other than Ruminantia, Pesca, Reptilia.

### Part II:

- (<sup>1a</sup>) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.

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

II.	Health information	II.a. Certificate reference No	II.b.			
( <sup>2</sup> )	Delete as appropriate.					
( <sup>3</sup> )	Insert method 1 to 5 or method 7 as ap	plicable.				
( <sup>4</sup> )	Where:					
	n = number of samples to be tested;					
	m = threshold value for the numb number of bacteria in all sample	er of bacteria; the result is consider les does not exceed m;	red satisfactory if the			
	M = maximum value for the numb number of bacteria in one or m	er of bacteria; the result is considered ore samples is M or more; and	ed unsatisfactory if the			
		al count of which may be between m the bacterial count of the other sample				
( <sup>5</sup> )	OJ L 147, 31.5.2001, p. 1.					
( <sup>6</sup> )	OJ L 172, 30.6.2007, p. 84.					
( <sup>7</sup> )	The person responsible for the load re					
	or the product described in this health					
	for non-ruminant farmed animals, oth accordance with the methods set out					
	verify the absence of unauthorised con					
	such analysis must be attached to the		the consignment at a			
	border inspection post of the European Union.					
_	The signature and the stamp must be in	-	-			
_	Note for the person responsible for the					
	for veterinary purposes and must accompany the consignment until it reaches the border inspection post of the point of entry into the European Union.					
O.CC.	· · · · · · · · · · · · · · · · · · ·					
Offi	cial veterinarian/Official inspector					
	Name (in capital letters):	Qualific	eation and 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

CC	OUNTRY:	Veterinary certificate to EU					
	I.1. Consignor	I.2. Certificate reference No I.2.a.					
	Name Address	I.3. Central competent authority					
ınt	Tel.	I.4. Local competent authority					
Part I: Details of dispatched consignment	I.5. Consignee Name Address  Postcode Tel.	I.6. Person responsible for the load in EU Name Address  Postcode Tel.					
of dispa	I.7. Country of origin ISO code I.8. Region of origin Code origin	I.9. Country of destination ISO code I.10. Region of destination Code					
tails	I.11. Place of origin	I.12. Place of destination					
Part I : De	Name Approval number Address Name Approval number Address Name Approval number Address	Custom warehouse  Name Approval number Address  Postcode					
	I.13. Place of loading	I.14. Date of departure					
	I.15. Means of transport  Aeroplane  Ship  Railway wagon	I.16. Entry BIP in EU					
	Road vehicle Other Identification Documentation references	I.17.					
	I.18. Description of commodity	I.19. Commodity code (HS code)					
		I.20. Quantity					
	I.21. Temperature of product Ambient □ Chilled □	I.22. Number of packages					
	I.23. Seal/Container No	I.24. Type of packaging					
	I.25. Commodities certified for:  Technical use □						
	I.26. For transit through EU to third country	I.27. For import or admission into EU					
	Third country ISO code						
	I.28. Identification of the commodities	1					
		nufacturing 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

#### Health information II.a. Certificate reference No II. 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 8(c) and Article 8(d) and Article 10 thereof, and Commission Regulation (EU) No 142/2011(1b), and in particular Chapter II of Annex XIV thereto, and certify that: Part II: Certification II.1. the blood products described above consist of blood products that satisfy the health requirements below; II.2. they consist exclusively of blood products not intended for human or animal consumption; II.3. they have been prepared and stored in a plant supervised by the competent authority or in the establishment of collection, exclusively with the following animal by-products: blood of slaughtered animals, which is fit for human consumption in (2) either accordance with Union legislation, but is not intended for human consumption for commercial reasons;] (<sup>2</sup>)and/or blood of slaughtered animals, which is 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, derived from carcases that have been slaughtered in a slaughterhouse and were considered fit for human consumption following an antemortem inspection in accordance with 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 blood and blood products derived from the production of products intended for human consumption;] (2)and/or blood and blood products originating from live animals that did not show signs of any disease communicable through that product to humans or animals;] (<sup>2</sup>)and/or animal by-products derived from animals which have been submitted to illegal treatment as defined in Article 1(2)(d) of Council Directive 96/22/EC<sup>(2a)</sup> or Article 2(b) of Council Directive 96/23/EC<sup>(2b)</sup>; (<sup>2</sup>)and/or animal by-products containing residues of other substances and environmental contaminants listed in Group B(3) of Annex I to Directive 96/23/EC, if such residues exceed the permitted level laid down in Union legislation or, in the absence thereof, in national legislation;] the blood that such products was manufactured from was collected in slaughterhouses II.4. approved in accordance with Union legislation, in slaughterhouses approved and supervised by the competent authority of the country of collection or from live animals in facilities approved and supervised by the competent authority of the country of $(^{2})[II.5.$ in the case of blood products derived from animals belonging to the taxa Artiodactyla, Perissodactyla and Proboscidea, including crossbreds between species of those taxa, the products come: II.5.1. from a country 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;

II. H	ealth inform	ation	II.a. Certificate reference No	II.b.			
(²)II.5.2.	either	country code in the parts thereof) when for a period of at 1	countries, territories or parts there case of a country, or codes ( <sup>3</sup> )in the re no case of foot-and-mouth disease the preceding 12 months and ed out against this disease for a plans;]	e case of territories or use has been recorded in which vaccination			
	or	code in the case of where no case of fat least the precedagainst foot-and-r	es, territories or parts thereof of a country or codes(3) for territo cot-and-mouth disease has been rec ding 12 months and in which vac mouth disease are being officia nestic ruminant animals for a pe hs(4);]]	ries or parts thereof) corded for a period of cination programmes lly carried out and			
$(^{2})[II.5.3.$	In addition	, in the case of anima	als other than Suidae and Tayassuid	ae:			
	( <sup>2</sup> )either	bluetongue( <sup>2</sup> ) (included recorded for a per	region of origin no case of ves luding the presence of seropositive riod of at least the preceding 12 in the been carried out against those dise 12 months;]	re animals) has been months and in which			
	( <sup>2</sup> ) <i>or</i>	[in the country or seropositive animal	region of origin vesicular stomatils are present(4);]]	tis and bluetongue( <sup>2</sup> )			
$(^{2})[II.5.4.$	In addition	In addition, in the case of Suidae and Tayassuidae:					
II.5.4.1.	in the country or region of origin no case of swine vesicular disease, classical swine fever and African swine fever has been recorded for a period of at least the preceding 12 months and vaccination has not been carried out against those diseases for a period of at least the preceding 12 months in the susceptible species and:						
( <sup>2</sup> )II.5.4.2.	either	the presence of ser least the precedin	region of origin no case of vesicular ropositive animals) has been record g 12 months and in which vacci t this disease for a period of at le	ded for a period of at ination has not been			
(²)II.5.4.2.	or	[in the country or are present(4);]	region of origin vesicular stomatitis	s seropositive animals			
( <sup>2</sup> )[II.6.			derived from poultry or other avian territory of the country or region w				
			castle disease and highly pathoger al Health Code of the OIE,	nic avian influenza as			
		a period of at least t an influenza,	he preceding 12 months has not ca	arried out vaccination			
	against Ne	ewcastle disease with	h the products are derived, have h vaccines prepared from a New enicity than lentogenic virus strains;	castle disease master			
II.7.	the produc	ts were:					
	(²)either	[packed in new or	sterilised bags or bottles,]				
	( <sup>2</sup> ) <i>or</i>		lk in containers or other means of and disinfected with a disinfect y before use,]				
	the outer packaging or containers bear labels indicating 'NOT FOR HUMA ANIMAL CONSUMPTION';						
II.8.	the produc	ts were stored in encl	losed storage;				

II.	Health information			II.a.	Certificate 1	reference No	II.b.	
II.9.	all precautions were taken to agents during transport;			avoid	contaminati	on of the produ	cts with pathogenic	
$(^{2})II.10.$	_	C	d products des	cribec	l above			
	(²)eithei	r [is derived fi	rom other rum	inants	than bovine,	ovine or caprine	animals.]	
	$(^2)or$	[is derived f derived from		vine c	or caprine and	imals and does n	ot contain and is not	
		animals bor region classi			rine and caprine materials other than those derived from rn, continuously reared and slaughtered in a country or sified as posing a negligible BSE risk in accordance with 007/453/EC.]			
		( <sup>2</sup> ) <i>or</i>	r [(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(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,					
			ovine or by lacer elongated cavity, o for thos slaughter	caprination d rod r by m e anime	of the cent -shaped inst neans of gas mals that we a country	rhich have been k ral nervous tissurument introductinjected into the covere born, conti- or region class	tained from bovine, cilled, after stunning, ue by means of an ed into the cranial cranial cavity, except nuously reared and sified as posing a sion 2007/453/EC.]]	
Notes								

### 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 for animal consumption.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.

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

Health information	II.a. Certificate reference No	II.b.			
Box reference I.28 Species: select from the following: Aves, Ruminantia, Suidae, Mammalia					
•					
** *					
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).					
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					
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).					
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 is only					
for veterinary purposes and must accompany the consignment until it reaches the border					
inspection post of the point of entry into the	he European Union.				
Official veterinarian/Official inspector					
Name (in capital letters):	Qualifica	ation and title:			
Date:	Signature	e:			
Stamp:					
	Box reference I.28 Species: select from other than Ruminantia, Pesca, Reptilia.  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 120.3.2010, p. 1).  In this case following the veterinary of 30.1.1998, p. 9), and in accordance with the products must be transported directly to Code of the territory as it appears in P798/2008 (OJ L 226, 23.8.2008, p. 1).  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 conformal of the product of the person responsible for the conformal of the point of entry into the conformal of the point of entry into the conformal of the point of the point of entry into the conformal of the point of the point of entry into the conformal of the point of entry into the conformal of the point of the point of entry into the conformal of the point of the point of entry into the conformal of the point of the point of the point of entry into the conformal of the point of the point of the point of the point of entry into the conformal of the point of	Box reference I.28 Species: select from the following: Aves, Ruminantia other than Ruminantia, Pesca, Reptilia.  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) N 20.3.2010, p. 1).  In this case following the veterinary checks provided for in Directive 30.1.1998, p. 9), and in accordance with the conditions laid down in Article the products must be transported directly to the establishment at the place of Code of the territory as it appears in Part 1 of Annex I to Commission 798/2008 (OJ L 226, 23.8.2008, p. 1).  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 printi Note for the person responsible for the consignment in the European Union: for veterinary purposes and must accompany the consignment until it inspection post of the point of entry into the European Union.  ial veterinarian/Official inspector  Name (in capital letters):  Qualification of the point of entry into the European Union.			

### 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(<sup>2</sup>) the European Union

CC	OUNTRY:	Veterin	ary certificate to EU				
I.1. Consignor Name		I.2. Certificate reference No	I.2.a.				
	Address	I.3. Central competent authority					
ent	Tel.	I.4. Local competent authority					
Part I: Details of dispatched consignment	I.5. Consignee Name Address  Postcode Tel.	I.6. Person responsible for the load in EU Name Address  Postcode Tel.					
of dispa	I.7. Country of origin ISO code I.8. Region of origin Code origin	I.9. Country of destination	I.10. Region of destination Code				
tails	I.11. Place of origin	I.12. Place of destination					
Part I: De	Name Approval number Address Name Approval number Address Name Approval number		n warehouse   val number				
	Address I.13. Place of loading	I.14. Date of departure					
		•					
	I.15. Means of transport  Aeroplane  Ship  Railway wagon  Identification  Railway wagon  Identification	I.16. Entry BIP in EU  I.17.					
	Documentation references  I.18. Description of commodity	I 19 Commod	ity code (HS code)				
	1.16. Description of commounty	1.17. Commod					
			I.20. Quantity				
	I.21. Temperature of product Ambient □ Chilled □	Frozen	I.22. Number of packages				
	I.23. Seal/Container No		I.24. Type of packaging				
	I.25. Commodities certified for:  Technical use □						
	I.26. For transit through EU to third country	I.27. For import or admission into EU	J $\square$				
	Third country ISO code						
	I.28. Identification of the commodities						
		number of establishments nufacturing plant	Batch number				

Health information

II.

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

II.b.

II.a. Certificate reference No

#### 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 8(c) and Article 8(d) and Article 10 thereof, and Commission Regulation (EU) No 142/2011(1b), and in particular Chapter II of Annex XIV thereto, and certify that: the blood products described above consist of blood products that satisfy the requirements II.1. Part II: Certification II.2. they consist exclusively of blood products not intended for human or animal consumption; II.3. they have been prepared and stored in a plant supervised by the competent authority, exclusively with the following animal by-products: $(^2)$ 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 accordance with Union legislation, but which did not show any signs of diseases communicable to humans or animals, derived from carcases that have been slaughtered in a slaughterhouse and were considered fit for human consumption following an ante-mortem inspection in accordance with 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 [blood and blood products originating from live animals that did not show clinical signs of any disease communicable through these products to humans or animals: animal by-products which have been derived from animals which have $\binom{2}{and/or}$ [been submitted to illegal treatment as defined in Article 1(2)(d) of Council Directive 96/22/EC<sup>(2a)</sup> or Article 2(b) of Council Directive 96/23/EC<sup>(2b)</sup>;] $(^{2})$ and/or [animal by-products containing residues of other substances and environmental contaminants listed in Group B(3) of Annex I to Directive 96/23/EC, if such residues exceed the permitted levels laid down by Union legislation or, in the absence thereof, in national legislation;] II.4. the blood that these products were manufactured from was been collected in slaughterhouses approved in accordance with Union legislation, in slaughterhouses approved and supervised by the competent authority of the country of collection or from live animals in facilities approved and supervised by the competent authority of the country of collection. $(^{2})[II.5.$ In the case of blood products derived from Artiodactyla, Perissodactyla and Proboscidea including their crossbreeds, other than Suidae and Tayassuidae, the products have undergone one of the following treatments, guaranteeing the absence of pathogens of foot-and-mouth disease, vesicular stomatitis, rinderpest, peste des petits ruminants, Rift Valley fever and bluetongue: [heat treatment at a temperature of 65 °C for at least three hours, followed by (<sup>2</sup>)either an effectiveness check;] (<sup>2</sup>)and/or [irradiation at 25 kGy by gamma rays, followed by an effectiveness check;] (<sup>2</sup>)and/or [change in pH to pH 5 for two hours, followed by an effectiveness check;] (2)and/or [heat treatment of at least 80 °C throughout their substance, followed by an effectiveness check.]] (<sup>2</sup>)[II.6. In the case of blood products derived from Suidae, Tayassuidae, poultry and other avian species, the products have undergone one of the following treatments guaranteeing the

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

II.	Health in	formation		II.a.	Certifica	te reference No		II.b.
	swine ves	of pathogens of the following diseases: foot-and-mouth disease, vesicular stomatitis, resicular disease, classical swine fever, African swine fever, Newcastle disease and pathogenic avian influenza, as appropriate to the species:						
	(2)either	[heat tr	[heat treatment at a temperature of 65 °C for at least three hours, followed an effectiveness check;]					
	(2)and/or (2)and/or	for pou	eatment of at 1	least 8 avian s	80 °C for Species(2)	Suidae/Tayassui	idae(	ctiveness check;] <sup>(2)</sup> and at least 70°C tance of the product,
( <sup>2</sup> )[II.7.		e of blood p	roducts derived	d from	species of			d in point II.5 or II.6,
II.8.	The produ	icts were:						
	(²)either	[packed in	new or sterilis	ed bag	gs or bottle	es,]		
	( <sup>2</sup> ) <i>or</i>	thoroughly		lisinfec				transport that were yed by the competent
		packaging of MPTION';	or containers be	ear lab	els indica	ting 'NOT FOR	HU	MAN OR ANIMAL
II.9.	the produc	cts were sto	red in enclosed	storag	ge;			
II.10.	-	precautions were taken to avoid the contamination of the products with pathogenic agents or treatment;						
$(^{2})II.11.$	The treate	ed blood pro	ducts described	d abov	e			
	(2)either [is derived from other ruminants than bovine, ovine or caprine animals.]					animals.]		
		[is derived from bovine, ovine or caprine animals and does not contain a derived from:				ot contain and is not		
		( <sup>2</sup> ) either	animals born	n, cont fied as	inuously sposing a	reared and slau	ighte	those derived from ered in a country or a in accordance with
		( <sup>2</sup> ) <i>or</i>						t 1 of Annex V to Parliament and of the
			ovine or born, con classified Commis	caprintinuo d as p sion D	ne animal usly reare oosing a n	s, except from d and slaughtere egligible BSE 007/453/EC(5),	thos ed ir risk	m bones of bovine, the animals that were a a country or region in accordance with which there has been
			(c) animal be ovine or by lacer elongate cavity, o for thos slaughter	by-pro caprir cation d rod- or by m se anii red in	duct or done animals of the co-shaped ineans of gamals that	lerived product which have be entral nervous instrument intro as injected into a were born, c try or region	en k tissu oduce the c contin	tained from bovine, cilled, after stunning, are by means of an ed into the cranial cranial cavity, except nuously reared and sified as posing a sion 2007/453/EC.]]
Notes Part I:								

Stamp:

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 reference No	II.b.			
-	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 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 BIP of entry into the European Union.					
-	Box I.19: use the appropriate Harmonized System (HS) code under the following headings: 05.11, 30.02, 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 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 in case of Species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia, Pesca, Reptilia.					
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 co	-	•			
-	Note for the person responsible for the confor veterinary purposes and must accommospection post of the European Union.					
Offic	cial veterinarian/Official inspector					
	Name (in capital letters):	Qualificat	ion and title:			
	Date:	Signature:				

"

(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

CO	OUNTRY:	Veterinary certificate to EU				
	I.1. Consignor	I.2. Certificate reference No I.2.a.				
	Name Address	I.3. Central competent authority				
nt	Tel.	I.4. Local competent authority				
Part I : Details of dispatched consignment	I.5. Consignee Name Address  Postcode Tel.	I.6. Person responsible for the load in EU Name Address  Postcode Tel.				
s of dispat	I.7. Country of origin ISO code I.8. Region of origin Code origin	I.9. Country of destination ISO code I.10. Region of destination Code				
tail	I.11. Place of origin	I.12. Place of destination				
art I : De	Name Approval number Address Name Approval number Address	Custom warehouse Name Approval number Address				
P	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	I.17. Number(s) of CITES				
	Documentation references  I.18. Description of commodity	I.19. Commodity code (HS code)				
		I.20. Quantity				
		·				
	I.21.	I.22. Number of packages				
	I.23. Seal/Container No	I.24. Type of packaging				
	I.25. Commodities certified for:	<u>'</u>				
	Technical use □					
	I.26. For transit through EU to third country	I.27. For import or admission into EU				
	Third country ISO code					
	I.28. Identification of the commodities					
	Species (Scientific name) Nu	umber of packages				

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

### II. II.a. Certificate reference No **Health information** 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 game trophies described above: $(^{2})$ either [II.1. with respect to game trophies or other preparations of cloven-hoofed animals, Part II: Certification excluding swine: ..... (region) has been free from foot-and-mouth disease and rinderpest for a period of the preceding 12 months, and during that period, no vaccination against any of those diseases has taken place; and the game trophies or other preparations described above: were obtained from animals which were killed in the territory of that region, which is authorised for the exportation to the European Union of fresh meat of the corresponding susceptible domestic species and where, during the period of the preceding 60 days, there have been no animal health restrictions due to outbreaks of diseases to which the game animals are susceptible; and originated from animals that were killed at a distance of at least 20 km (ii) from the borders of another third country or part of a third country not authorised to export untreated game trophies of cloven-hoofed animals other than swine to the European Union;] $(^2)$ or [II.1. with respect to game trophies or other preparations of wild swine: ..... (region) during the period of the preceding 12 months, was free from classical swine fever, African swine fever, swine vesicular disease, foot-and-mouth disease and porcine enteroviral encephalmiyelitis (Teschen disease) and no vaccinations have been carried out against any of those diseases during that 12 month period; and the game trophies or other preparations described above: were obtained from animals which were killed in that territory, which is authorised for the exportation to the European Union of fresh meat of the corresponding susceptible domestic species and where, during the period of the preceding 60 days, there have been no animal health restrictions due to outbreaks of diseases to which the swine are susceptible; and originated from animals that were killed at a distance of at least 20 km (ii) from the borders of another third country or part of a third country not authorised to export untreated game trophies of wild swine to the European Union: $(^2)$ or with respect to game trophies or other preparations of solipeds, the game trophies or other preparations described above were obtained from wild solipeds that were killed in the territory of the exporting country referred to above;] $(^{2})or$ [II.1. with respect to game trophies or other preparations of game birds: ..... (region) is free from highly pathogenic avian influenza and Newcastle disease; and the game trophies or other preparations described above were obtained from wild game birds that were killed in that region and where during the period of the preceding 30 days there have been no animal health restrictions due to outbreaks of disease to which the wild birds are susceptible;] II.2. The game trophies or other preparations described above have been packaged without being in contact with other products of animal origin likely to contaminate them, in individual,

transparent and closed packages so as to avoid any subsequent contamination.

(2)II.3. The game trophies or other preparations described above

### 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.
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(2) either [is derived from other ruminants than bovine, ovine or caprine animals.]

<sup>2</sup>)or [is derived from bovine, ovine or caprine animals and does not contain and is not derived from:

(<sup>2</sup>) 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.]

 $(^2)or$ 

- [(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(<sup>4</sup>);
- (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.
- (c) animal by-product or derived product obtained from bovine, ovine or caprine 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 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.

### Part II:

- (<sup>1a</sup>) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.

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.						
( <sup>3</sup> )	OJ L 147, 31.5.2001, p. 1.						
( <sup>4</sup> )	OJ L 172, 30.6.2007, p. 84.						
_	<ul> <li>The signature and the stamp must be in a different colour to that of the printing.</li> <li>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.</li> </ul>						
Offic	cial veterinarian/Official inspector						
	Name (in capital letters):	Qualifica	ation and title:				
	Date:	Signatur	e:				
	Stamp:						

### (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<sup>2</sup>, intended for dispatch to or for transit through(<sup>2</sup>) the European Union

I.1. Consignor Name Address  Tel.  I.5. Consignee Name Address  Postcode Tel.  I.7. Country of ISO code origin  I.8. Region of origin  I.9. Country of destination  I.10. Country of destination  I.11. Place of origin  I.12. Certificate reference No  I.2.a.  I.3. Central competent authority  I.4. Local competent authority  I.5. Person responsible for the load in EU Name Address  Postcode Tel.  I.7. Country of ISO code I.8. Region of origin Code origin  I.11. Place of origin  I.12. Place of destination  Custom warehouse Address  Name Address  Name Address  Address  Address  Address  Address  Address  Address  Address  Address	
Address I.3. Central competent authority	
I.5. Consignee I.6. Person responsible for the load in EU Name Name	
Name Name	
Postcode Postcode	
Tel. Tel.	
I.7. Country of ISO code I.8. Region of origin Code I.9. Country of destination ISO code I.10. Region of destination	Code
origin destination destination	
I.11. Place of origin I.12. Place of destination	
Name Approval number Custom warehouse	
Address 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 ☐ I.17.	
Documentation references	
I.18. Description of commodity I.19. Commodity code (HS code)	
I.20. Quantity	
I.21. Temperature of product Ambient □ Chilled □ Frozen □	kages
Ambient ☐ Chilled ☐ Frozen ☐  I.23. Seal/Container No  I.24. Type of packag	ging
I.25. Commodities certified for:	
Technical use □	
I.26. For transit through EU to third country   I.27. For import or admission into EU	
Third country ISO code	
I.28. Identification of the commodities	
Approval number of establishments	tala muli
Approval number of establishments	tch number
Approval number of establishments	tch number

II.

II.b.

# Part II: Certification

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

II.a. Certificate reference No

(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'.]

(2) or [satisfy the animal health requirements set out in point II.1.];

- II.1. The animal by products described above
- II.1.1. have been

**Health information** 

- (²)either [(a) obtained from materials imported from a third country, territory or part thereof:......(³) authorised to export fresh meat to the European Union;]
- (²) and/or [(b) obtained in the exporting third country, territory or part thereof:......(³) from animals that
  - (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(<sup>4</sup>);]
- (²) and/or [(c) derived from eggs, milk, rodents, lagomorphs, or aquatic animals or terrestrial or aquatic invertebrates;]
- II.1.2. (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 fur, 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
    - (ii) where there has not been any 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) remained on their holdings of origin for a period of at least 40 days before the date 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

II.	Health information		II.a.	Ce	rtificat	e refere	ence No		II.b.
		the (iv) we sla Uı eq	e anima ere han aughter nion le uivaler	als andled or legisle	re susce d in the killed in lation those	eptible; e slaug n accor and co	; and ghterhound rdance womplied own in (	ise before with the with	ore and at the time of e relevant provisions of requirements at least as II and III of Council
	$(^{2})or$ [(a)	captured	_	,	,		` '	:	
		an su dis the	y of sceptib sease of prece	the le: r hig eding	follow foot-a ghly pa g 30 d	ving di ind-mou thogeni ays nor	iseases 1th dis ic avian	for wease, influer	een no case/outbreak of hich the animals are rinderpest, Newcastle iza during the period of or African swine fever s; and
		se <sub>j</sub>	parating	g ar not	nother authori	territor	y of a	third c	20 km from the borders ountry or part thereof, the exportation of such
	(b)	which a chilling	fter kil either	lling to a	were	transpo tion ce	orted wi	thin a d imme	period of 12 hours for diately afterwards to a ishment;]
II.1.3.	II.1.3. ( <sup>2</sup> )in the case of materials other than materials derived from fish or invertebrates caught in the wild, have been obtained in an establishment around which, within a radius of 10 km, there has been no case/outbreak of diseases referred to in point II.2.2 for which the animals are susceptible during a period of the preceding 30 days or, in the event of a case/outbreak of one of those diseases, the preparation of raw material for exportation to the European Union was authorised only after the removal of all meat, and the total cleaning and disinfection of the establishment under the control of an official veterinarian;						h, within a radius of 10 int II.2.2 for which the vs or, in the event of a terial for exportation to all meat, and the total		
II.1.4.		conditio	ns req	uire	d abov				aterial which does not andled so as to avoid
II.1.5.	have been packe has been cleaned other than via pa authority, bearin MANUFACTUR	in new and distant post, the lab	packag infected in con el indi ERIVI	ging d be itain catii	which fore users sea ng 'Al' PROD	se and, lled und NIMAL UCTS	in the ler the r BY-PF FOR U	case of espons RODUO JSES O	or in packaging which consignments shipped ibility of the competent CTS ONLY FOR THE DUTSIDE THE FEED nation in the European
II.1.6.	consist only of the ( <sup>2</sup> )either [-	carcases oodies o consum	and por parts	oarts s of n ac	of an anima cordan	imals s ls kille ice with	d which h Unior	n were n legisl	in the case of game, deemed fit for human ation until irreversibly reasons;]
	(²)and/or [-	carcases were sla slaughte or bodie numan c (i) ca	and the aughter for hues and consum	ne for ed uman the ption or b	ollowir in a s n consu follow n in ac oodies	ng parts laughte umptior ving pa cordanc and par	original crhouse of followerts of a creater	ating end wing and animals Union lanimals	ther from animals that vere considered fit for ante-mortem inspection from game killed for egislation: which were rejected as cordance with Union

II.	Health information		II.a. Certificate reference No	II.b.			
		cor	nmunicable to humans or animals;				
			ds of poultry;				
		and	es and skins, including trimmings and lefeet, including the phalanges and the less, tarsus and metatarsus bones;				
			bristles;				
	2	` '	thers;]				
	( <sup>2</sup> )and/or [-	farm as r of the Eu	y-products from poultry and lagomore eferred to in Article 1(3)(d) of Regula ropean Parliament and of the Council of disease communicable to humans	ntion (EC) No 853/2004 ( <sup>2a)</sup> , which did not show			
	(²)and/or [-	communi animals to been con	f animals which did not show cable through blood to humans or that have been slaughtered in a slaughtered fit for slaughter for human co- tem inspection in accordance with Uni	animals, obtained from ghterhouse after having nsumption following an			
	( <sup>2</sup> )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;]					
	( <sup>2</sup> )and/or [-	origin, w	of animal origin, or foodstuffs contain which are no longer intended for haid reasons or due to problems of man r other defects from which no risk to	uman consumption for ufacturing or packaging			
	( <sup>2</sup> )and/or [-	animal by for feed manufact	and feedingstuffs of animal origin, or a y-products or derived products, which ing for commercial reasons or uring or packaging defects or other blic or animal health arises;]	are no longer intended due to problems of			
	( <sup>2</sup> )and/or [-	originatir	acenta, wool, feathers, hair, horns, hag from live animals that did not sho cable through that product to humans	w signs of any disease			
	( <sup>2</sup> )and/or [-	_	animals, and parts of such animals, d not show any signs of diseases com	_			
	(²)and/or [-	animal establishi consump	nents or plants manufacturing	als originating from products for human			
	(²)and/or [-	any signs animals:	wing material originating from anima of disease communicable through that	at material to humans or			
		` '	lls from shellfish with soft tissue or fle	*			
		(ii) the	following originating from terrestrial	animals:			
		-	hatchery by-products; eggs;				
		-	egg by-products, including egg shells	S:			
		(iii) day	y-old chicks killed for commercial reas				
	( <sup>2</sup> )and/or [-	animal by	y-products from aquatic or terrestrial is athogenic to humans or animals;]				
	(²)and/or [-	animals	and parts thereof of the zoological	orders of Rodentia and			

II. Heal	th information	II.a. Certificate reference No	II.b.				
11. IIcai							
	Article	orpha, except Category 1 materia 8(a)(iii), (iv) and (v) of Regulation (I ry 2 material as referred to in Artiction;]	EC) No 1069/2009 and				
	(²)and/or [- fur orig	ginating from dead animals that did not ease communicable through that product					
II.1.7.	7. 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 the time of dispatch and the time of delivery to the plant of destination.						
$(^{2})(^{6})[II.1.8.$	Specific requirements						
( <sup>2</sup> )( <sup>7</sup> )II.1.8.1.	country, territory or p	consignment come from animals that hat art thereof referred to in point II.2 ot-and-mouth disease are regularly carryine animals.	2.1, where vaccination				
( <sup>2</sup> )( <sup>8</sup> )II.1.8.2.	The by-products in this c deboned meat.]	onsignment consist of animal by-produc	ets derived from offal or				
$(^{2})II.1.9.$	the animal by-products de						
	•	other ruminants than bovine, ovine or c	=				
	(²) or [is derived from not derived from	n bovine, ovine or caprine animals and n:	does not contain and is				
	an reg	ovine, ovine and caprine materials other imals born, continuously reared and slaugion classified as posing a negligible I th Decision 2007/453/EC.]	aghtered in a country or				
	(²)or [(a	specified risk material as defined in egulation (EC) No 999/2001 of the Europe Council(4);					
	(b)	ovine or caprine animals, except from born, continuously reared and slaug region classified as posing a ne accordance with Commission Decis which there has been no indigenous B	those animals that were htered in a country or gligible BSE risk in ion 2007/453/EC( <sup>5</sup> ), in				
	(c)	animal by-product or derived product ovine or caprine animals which h stunning, by laceration of the central rof an elongated rod-shaped instrume cranial cavity, or by means of gas i cavity, except for those animals that reared and slaughtered in a country posing a negligible BSE risk in acc 2007/453/EC.]]	ave been killed, after nervous tissue by means ent introduced into the njected into the cranial were born, continuously or region classified as				
II.1.10	the animal by-products or	r product described above:					
		in milk or milk products of ovine or cap for feed for farmed animals, other than f					
		or milk products of ovine or caprin feed for farmed animals, other than fur a					
	contin	erived from ovine and caprine animals nuously since birth in a country where t lfilled:					
	(i)	classical scrapie is compulsorily noti	fiable;				

## Animal by-products to be used for purposes outside the feed chain or for trade samples<sup>(2)</sup>

II.	Health information		II.a. Certificate reference No II.b.	
		(ii)	an awareness, surveillance and me place for classical scrapie;	onitoring system is in
		(iii)	official restrictions apply to holdin animals in the case of a suspi confirmation of classical scrapie;	
		(iv)	ovine and caprine animals affected w killed and destroyed;	vith classical scrapie are
		(v)	the feeding to ovine and caprine and meal or greaves, as defined in the Te Code of the World Organisation for of ruminant origin has been banned a in the whole country for a period of seven years;	errestrial Animal Health Animal Health (OIE), and effectively enforced
	(b)		e from holdings where no official resuspicion of TSE;	estrictions are imposed
	(c)	diagnos	e from holdings where no case of classed during the period of the precong the confirmation of a case of classic	eding seven years or,
		( <sup>2</sup> )eithe	r [all ovine and caprine animals on killed and destroyed or slaughtere rams of the ARR/ARR genotype, br least one ARR allele and no VRQ animals carrying at least one ARR al	d, except for breeding eeding ewes carrying at allele and other ovine
		( <sup>2</sup> )or	[all animals in which classical scrapbeen killed and destroyed, and subjected for a period of at least two confirmation of the last classical scraps of the presence of TSE in accordance methods set out in point 3.2 of Characteristics (EC) No 999/2001, of animals which are over the age of 1 animals of the ARR/ARR genotype:	the holding has been by years since the date of rapie case to intensified with negative results for the with the laboratory apter C of Annex X to all of the following 8 months, except ovine
			<ul> <li>animals which have been slaconsumption; and</li> <li>animals which have died or bee but which were not killed in the eradication campaign.]].</li> </ul>	n killed on the holding

### 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

## Animal by-products to be used for purposes outside the feed chain or for trade samples<sup>(2)</sup>

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

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 for animal consumption.
- 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, Mammalia other than Ruminantia, Pesca, Mollusca, Crustacea, invertebrates other than Mollusca and Crustacea.

### Part II:

- <sup>(1a)</sup> OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.
- <sup>(2a)</sup> 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 (EU) 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.

- 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.
- <sup>(5)</sup> OJ L 303, 18.11.2009, p. 1.
- 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.
- Only for certain South American countries.
- (8) Only for certain South American and South African countries.

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

II.	Health information	II.a. Certificate reference No	II.b.				
(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 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.						
Offic	cial veterinarian/Official inspector						
	Name (in capital letters):	Quali	fication and 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

OUNTRY:					Veterir	nary certificate to	) EU
I.1. Consignor Name			I.2.	Certificate reference	No No	I.2.a.	
Address			I.3.	Central competent a	uthority		
Tel.			I.4.	Local competent aut	hority		
I.5. Consignee			I.6.	Person responsible f	or the load in E	EU	
Name Address				Name Address			
Postcode				Postcode			
Tel.				Tel.			
I.7. Country of I origin	I.8. Region of origin	Code		Country of destination	ISO code	I.10. Region of destination	Code
I.11. Place of origin	I		I.12.	Place of destination			
Name	Approval numb	er			Custo	om warehouse	
Address Name	Approval numb	er		Name Address	Appro	oval number	
Address Name							
Address	Approval numb	er		Postcode			
I.13. Place of loading	9		I.14.	Date of departure			
I.15. Means of transp	port		I.16.	Entry BIP in EU			
Aeroplane	Ship  Railway wago	on $\square$					
Road vehicle  Identification	Other		I.17.				
Documentation							
I.18. Description of c	commodity				I.19. Commo	dity code (HS code)	
						I.20. Quantity	
I.21. Temperature of Ambient □	product Chilled			Frozen		I.22. Number of package	ges
I.23. Seal/Container				110zen 🗆		I.24. Type of packagin	g
I.25. Commodities co	ertified for:						
Animal feeding	stuff	Manu	ıfactur	e of petfood $\square$		Technical	use 🗆
I.26. For transit throu	ugh EU to third country			I.27. For import or a	dmission into E	EU	
Third country	ISO code						
I.28. Identification of							
Species (Scientific		1.1		of establishments uring plant Numb	per of packages	Net weight Batch	number
Species (Scientific	name, radic of commodity	iviai	iiuiaell	arms plant 14um	or packages	Thet weight Battill I	14111001

	II.	Health inform	ation	II.a. Certificate reference No	II.b.				
Part II: Certification	II.1. II.2. II.3.	(EC) No 1069. Article 10 ther Chapter II of A consist of render consist of render have been pre authority in acc with Article 4( Council( <sup>3</sup> ), in o	(2009 of the Europeof, and Commissionnex XIV thereto, ered fats that satisfyered fats not intended and stored cordance with Artical (2) of Regulation (1) order to kill pathoge	•	ed by the competent 2009 or in accordance Parliament and of the				
Part I	11.4.	( <sup>2</sup> )either [-	ave been prepared exclusively with the following animal by-products:  2) either [- carcases and parts of animals slaughtered or, in the case of game, bodies parts of animals killed, and which are fit for human consumption accordance with Union legislation, but are not intended for hum consumption for commercial reasons;]						
		( <sup>2</sup> )and/or [-	carcases and the been slaughtered for human consu the following par	following parts originating either from a slaughterhouse and were consimption following an ante-mortem in the soft animals from game killed for human legislation:	dered fit for slaughter spection or bodies and				
		<ul> <li>carcases or bodies and parts of animals which are rejected as unfinding human consumption in accordance with Union legislation, but with did not show any signs of disease communicable to human animals;</li> </ul>							
			feet, include tarsus and	skins, including trimmings and splitt ling the phalanges and the carpus a metatarsus bones;					
			<ul><li>(iv) pig bristles</li><li>(v) feathers;]</li></ul>	,					
		( <sup>2</sup> )and/or [-	blood of animals through blood to slaughtered in slaughter for hur	which did not show any signs of a humans or animals, obtained from a a slaughterhouse after having been nan consumption following an ante- Union legislation;]	animals that have been en considered fit for				
		( <sup>2</sup> )and/or [-	animal by-produ human consump	cts arising from the production of tion, including degreased bone, great from milk processing;]					
		( <sup>2</sup> )and/or [-	origin, which a	mal origin, or foodstuffs containing are no longer intended for hum ons or due to problems of manufa defects from which no risk to pu	nan consumption for acturing or packaging				
		( <sup>2</sup> )and/or [-	animal by-produ feeding for com	dingstuffs of animal origin, or feet cts or derived products, which are remercial reasons or due to problems s or other defects from which no ris	no longer intended for s of manufacturing or				
		( <sup>2</sup> )and/or [-	blood, placenta, originating from	wool, feathers, hair, horns, hoof live animals that did not show rough that product to humans or anim	signs of any disease				
		( <sup>2</sup> )and/or [-	aquatic animals,	and parts of such animals, except sea	a mammals, which did				

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

II.	Health info	rmation	II.a. Certificate reference No	II.b.
		not show any si	gns of diseases communicable to hum	ans or animals;]
	(²)and/or [		ducts from aquatic animals origina manufacturing products for human cor	
	( <sup>2</sup> )and/or [	signs of disease	naterial originating from animals wh communicable through that material t	•
		· /	m shellfish with soft tissue or flesh;	
			ving originating from terrestrial animal chery by-products,	ls:
		- nate	• • •	
			by-products, including egg shells;	
		= =	hicks killed for commercial reasons;]	
II.5.	( <sup>2</sup> )either	the territory of of the preced	material of porcine origin, come from f a country free from foot-and-mouth ing 24 months and free from class fever for the period of the preceding 1	disease for the period sical swine fever and
	(²)and/or		material of poultry origin, come from from Newcastle disease and avian inf 6 months;]	
	(²)and/or	a territory fr	material of ruminant origin, come fro ee from foot-and-mouth disease for months and free from rinderpest f months;]	or the period of the
	( <sup>2</sup> )and/or	point II.5. dur the rendered f	as been an outbreak of one of the cing the relevant period referred to in ats derived from a susceptible species ment for at least 70 °C for 30 minutes es, and	point II.5, and where s, have been subjected
		the owner, of competent au information m	critical control points are recorded a operator or their representative and athority can monitor the operation oust include the particle size, critical absolute time, pressure profile, rawate.]	d, as necessary, the n of the plant; the l temperature and, as
II.6.	remaining to	otal insoluble impurit	ls, were purified in such way that the ies does not exceed 0,15 % in weight;	e maximum levels of
II.7.	the rendered			
		of Section 142/2011, of	subjected to processing in accordance 3 of Chapter II of Annex X to or a treatment in accordance with Secon (EC) No 853/2004, in order to k	Regulation (EU) No etion XII of Annex III
	( <sup>2</sup> )either	and disinfe	ed in new containers or in containers to cted if necessary for the prevention ons have been taken to prevent their co	of contamination, and
	( <sup>2</sup> )or	any other b	transport is intended, the pipe, pumpulk container or bulk road tanker use luct from the manufacturing plant eit	d in the transportation
		ship or into	shore tanks or directly to plants have sibility of the competent authority a	e been checked under
	and which b		'NOT FOR HUMAN CONSUMPTIO	on';

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

II.	Health in	nformation	l	II.a. Certificate reference No	II.b.			
( <sup>2</sup> )II.8.	the rende	ered fats des	scribed above					
	(²)either (²)or	[is derived	l from bovine	rom other ruminants than bovine, ovine or caprine animals.] rom bovine, ovine or caprine animals and does not contain and is not				
		derived fro	[bovine, or animals boregion class	vine and caprine materials other that orn, continuously reared and slaught saffied as posing a negligible BSE ris	tered in a country or			
		( <sup>2</sup> )or	[(a) specif	007/453/EC.] ied risk material as defined in poi (EC) No 999/2001 of the European				
			(b) mecha ovine born, o classif Comm	onically separated meat obtained from the continuously reared and slaughtered fied as posing a negligible BSE rish assion Decision 2007/453/EC(5), in igenous BSE case,	ose animals that were in a country or region k in accordance with			
				(c) animal by-product or derived product obtained from bovine, ovine or caprine 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				
II.9. the	rendered t	fats describ		ible BSE risk in accordance with Dec	ISIOII 2007/433/LC.]]			
11.5. the	( <sup>2</sup> )either	· [does n	ot contain mil	k or milk products of ovine or capril for farmed animals, other than fur ar				
	( <sup>2</sup> )or			ailk products of ovine or caprine a farmed animals, other than fur animals				
				From ovine and caprine animals wisince birth in a country where the fol				
			(i) class	ical scrapie is compulsorily notifiable	·,			
				wareness, surveillance and monitoring lassical scrapie;	ng system is in place			
			anim	ial restrictions apply to holdings als in the case of a suspicion of TS assical scrapie;				
				e and caprine animals affected with d and destroyed;	classical scrapie are			
			meal Code rumi	feeding to ovine and caprine animal or greaves, as defined in the Terre of the World Organisation for Animant origin has been banned and ef whole country for a period of at leas s;	estrial Animal Health mal Health (OIE), of fectively enforced in			
			~	holdings where no official restriction	ns are imposed due to			
		(c)	originate from	n holdings where no case of classi uring the preceding seven years of a case of classical scrapie:				

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

animals which have died or been killed on the holding but which were not killed in the framework of a disease

II.	Health information	II.a. Certificate reference No	II.b.
	and ARR allele	ovine and caprine animals on the holdestroyed or slaughtered, except for L/ARR genotype, breeding ewes carry and no VRQ allele and other ovin one ARR allele;]	breeding rams of the ying at least one ARR
	been for p the l inclu in ac of C all c mon	animals in which classical scrapie killed and destroyed, and the holdir eriod of at least two years since the d ast classical scrapie case to intensit ding testing with negative results for ecordance with the laboratory method hapter C of Annex X to Regulation (of the following animals which are the, except ovine animals of the ARR nimals which have been slaugonsumption; and	ng has been subjected ate of confirmation of fied TSE monitoring, rethe presence of TSE ds set out in point 3.2 (EC) No 999/2001, of over the age of 18 /ARR genotype:

### 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.

eradication campaign.]]

- 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 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: Ruminantia, other than Ruminantia
  - Manufacturing plant: provide the registration number of the treatment/processing establishment.

### Part II:

- (<sup>1a</sup>) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.
- (<sup>3</sup>) OJ L 139, 30.4.2004, p. 55.
- (4) OJ L 147, 31.5.2001, p. 1.
- (<sup>5</sup>) OJ L 172, 30.6.2007, p. 84.
  - The signature and the stamp must be in a different colour to that of the printing.

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

II.	Health information	II.a. Certificate reference No	II.b.					
_	<ul> <li>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.</li> </ul>							
Offi	Official veterinarian/Official inspector							
	Name (in capital letters): Qualification and title:							
	Date:	Signature	2:					
	Stamp:							

### 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

CO	OUNTRY:	Veterinary certificate to EU
	I.1. Consignor Name	I.2. Certificate reference No I.2.a.
	Address	I.3. Central competent authority
ent	Tel.	I.4. Local competent authority
Part I: Details of dispatched consignment	I.5. Consignee Name Address  Postal code Tel.	I.6. Person responsible for the load in EU Name Address  Postal code Tel.
of dispa	I.7. Country of ISO code I.8. Region of origin Code origin	I.9. Country of ISO code I.10. Region of Code destination
ails	I.11. Place of origin	I.12. Place of destination
Part I: Det	Name Approval number Address Name Approval number Address Name Approval number	Custom warehouse  Name Approval number  Address  Postal code
	Address I.13. Place of loading	I.14. Date of departure
		_
	I.15. Means of transport	I.16. Entry BIP in EU
	Aeroplane	I.17.
	I.18. Description of commodity	I.19. Commodity code (HS code)
		I.20. Quantity
	I.21. Temperature of product Ambient □ Chilled □	Frozen □ I.22. Number of packages
	I.23. Seal/Container No	I.24. Type of packaging
	I.25. Commodities certified for:  Technical use □	
	I.26. For transit through EU to third country	I.27. For import or admission into EU
	Third country ISO code	
	I.28. Identification of the commodities	
	Approval number of establishments Species Manufacturing plant (scientific name)	Number of packages Net weight Batch number

**Health information** 

II.

II.b.

# Part II: Certification

I, the undersigned official veterinari	an, declare that I have read and understood Regulation
	in Parliament and of the Council (1a), and in particular
Articles 8, 9 and 10 thereof, and 0	Commission Regulation (EU) No 142/2011(1b), and in
particular Chapter II of Annex XIV	V thereto, and certify that the rendered fats described
above.	

II.a. Certificate reference No

- II.1. consist of rendered fats not intended for human consumption that satisfy the health requirements below;
- II.2. have been prepared exclusively with the following animal by-products:
- 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;
- 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;
- 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:
  - (2) either [- animal by-products containing residues of authorised substances or contaminants exceeding the permitted levels referred to in Article 15(3) of Council Directive 96/23/EC(<sup>2a</sup>);]
  - (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;]
  - (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 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 [- carcasses 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:
    - carcasses 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 [- blood of animals which did not show any signs of disease communicable through blood to humans or animals obtained from animals other than ruminants 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;]

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

II.	Health info	ormat	ion	II.a. Certificate reference No	II.b.
	( <sup>2</sup> )and/or	[-	human consumpti	tts arising from the production of pon, including degreased bone, greatrom milk processing;]	
	( <sup>2</sup> )and/or	[-	origin, which as	nal origin, or foodstuffs containing re no longer intended for huma ons or due to problems of manufa defects from which no risk to pub	an consumption for cturing or packaging
	( <sup>2</sup> )and/or	[-	animal by-produc feeding for comn	ing stuffs of animal origin, or feed ts or derived products, which are n hercial reasons or due to problems s or other defects from which no ris	o longer intended for of manufacturing or
	( <sup>2</sup> )and/or	[-	originating from	wool, feathers, hair, horns, hoof live animals that did not show sough that product to humans or ani	signs of any disease
	( <sup>2</sup> )and/or	[-		and parts of such animals, except signs of diseases communicable to	
	(2)and/or	[-		cts from aquatic animals origina anufacturing products for human co	
	( <sup>2</sup> )and/or	[-	the following ma	terial originating from animals whi communicable through that man	ch did not show any
			` '	shellfish with soft tissue or flesh;	1
				ng originating from terrestrial anima ry by-products,	IS:
			- eggs,	• • •	
				products, including egg shells, cks killed for commercial reasons;]	
	( <sup>2</sup> )and/or ( <sup>2</sup> )and/or	[-	•	estrial invertebrates other than sp	pecies pathogenic to
	( <sup>2</sup> )and/or	[-	Lagomorpha, exc (iv) and (v) of Re	ts thereof of the zoological order ept Category 1 material as referred egulation (EC) No 1069/2009and C cle 9(a) to (g) of that Regulation;]	to in Article 8(a)(iii),
	( <sup>2</sup> )and/or	[-	from dead animal	hooves, feathers, wool, horns, haids that did not show any signs of duct to humans or animals;]	
	( <sup>2</sup> )and/or	[-	communicable the slaughtered in a slaughter for hum	om animals which did not show a rough that material to humans or slaughterhouse and which were an consumption following an ante- Jnion legislation;]	animals, which were considered fit for
II.2.4.				purposes other than the production accutical or medical devices :	of organic fertilisers
	( <sup>2</sup> )either	[-	specified risk ma	terial as defined in Article 3(1)(g) ne European Union and of the Coun	
	( <sup>2</sup> )and/or	[-	entire bodies or p	arts of dead animals containing spect 3(1)(g) of Regulation (EC) No 99	cified risk material as

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

II.	Healt	h info	ormati	on	II.a. Certificate reference No	II.b.
	(²)an	d/or	[-	been submitted t	ts which have been derived from to illegal treatment as defined in the 96/22/EC <sup>(2c)</sup> or Article 2(b) o	n Article 1(2)(d) of
	(²)an	d/or	[-	environmental con 96/23/EC, if such	acts containing residues of oth intaminants listed in Group B(3) of the residues exceed the permitted by or, in the absence thereof, by legistern;]	Annex I to Directive levels laid down by
II.3.	the re	ndere	d fats:			
	(a)	proc	essing		essing in accordance with method out in Chapter III of Annex IV pathogenic agents,	
	(b)	have (GTI	been i H), so	narked before ship	oment to the European Union with s minimum concentration of at lea	
	(c)			of rendered fats of ave been removed,	ruminant origin, insoluble impuritie	es in excess of 0,15%
	(d)	have	been t	ransported under c	onditions which prevent their conta	mination, and
	(e)			on the packagin	g or container indicating "NOT;	FOR HUMAN OR
( <sup>2</sup> )II.4.				f materials destin r soil improvers	ed for organic fertilisers, cosmeti the rendered fats described al	-
	( <sup>2</sup> )eit	her [i	s deriv	ed from other rum	inants than bovine, ovine or caprine	animals.]
	( <sup>2</sup> ) <i>or</i>	_	s deriv erived		vine or caprine animals and does n	ot contain and is not
		( <sup>2</sup>	eithe)	animals born	ne and caprine materials other than a, continuously reared and slaughter fied as posing a negligible BSE rist 7/453/EC.]	ered in a country or
		( <sup>2</sup>	e)or	- ' '	d risk material as defined in poir EC) No 999/2001 of the European I	
				ovine or born, con classified Commiss	cally separated meat obtained fro caprine animals, except from those ntinuously reared and slaughtered in a posing a negligible BSE risk sion Decision 2007/453/EC(5), in venous BSE case,	se animals that were n a country or region in accordance with
				ovine or by lacer elongated cavity, o for thos slaughter	by-product or derived product ob caprine animals which have been lation of the central nervous tissed rod-shaped instrument introduct by means of gas injected into the eanimals that were born, continued in a country or region classed BSE risk in accordance with Deci	cilled, after stunning, ue by means of an ed into the cranial cranial cavity, except nuously reared and ssified as posing a
Notes Part I:						
- Box	x refere	nce I.	.6: Per	son responsible fo	or the consignment in the Europea	n Union: this box is

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

II. Health information	II.a. Certificate reference No	II.b.
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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 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: 04.05; 15.01, 15.02; 15.03; 15.04; 15.05; 15.06; 15.16 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 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: Ruminantia, other than Ruminantia

Manufacturing plant: provide the registration number of the treatment/processing establishment.

### Part II:

- <sup>(1a)</sup> OJ L 300, 14.11.2009, p. 1.
- <sup>(1b)</sup> OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.
- <sup>(2a)</sup> OJ L 125, 23.5.1996, p. 10.
- <sup>(2b)</sup> OJ L 147, 31.5.2001, p. 1.
- <sup>(2c)</sup> OJ L 125, 23.5.1996, p. 3.
- <sup>(3)</sup> 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 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):	Qualification and title:							
Date:	Signature:							
Stamp:								

### 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

CC	OUNTRY:	Veterinary certificate to EU
	I.1. Consignor	I.2. Certificate reference No I.2.a.
	Name Address	I.3. Central competent authority
ınt	Tel.	I.4. Local competent authority
Part I: Details of dispatched consignment	I.5. Consignee Name Address  Postcode Tel.	I.6. Person responsible for the load in EU Name Address  Postcode Tel.
of dispatcl	I.7. Country of ISO code I.8. Region of origin Code origin	I.9. Country of ISO code I.10. Region of Code destination
ails	I.11. Place of origin	I.12. Place of destination
Part I : Det	Name Approval number Address Name Approval number Address Name Approval number	Custom warehouse  Name Approval number Address  Postcode
	Address	114 D ( C)
	I.13. Place of loading	I.14. Date of departure
	I.15. Means of transport  Aeroplane  Ship  Railway wagon  Other  Other	I.16. Entry BIP in EU  I.17.
	Identification  Documentation references	
	I.18. Description of commodity	I.19. Commodity code (HS code)
		I.20. Quantity
	I.21. Temperature of product Ambient □ Chilled □	I.22. Number of packages
	I.23. Seal/Container No	I.24. Type of packaging
	I.25. Commodities certified for:	
	Animal feedingstuff   Manu	ifacture of petfood $\square$ Technical use $\square$
	I.26. For transit through EU to third country	I.27. For import or admission into EU
	Third country ISO code	
	I.28. Identification of the commodities	
	Approval number of establishments Species (Scientific name) Manufacturing plant	Number of packages Net weight Batch number

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

### Health information II.a. Certificate reference No II. 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(<sup>1a</sup>), and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011(<sup>1b</sup>), and in particular Chapter I of Annex XIV thereto, and certify that the gelatine/collagen(<sup>2</sup>) described above: consists of gelatine/collagen(2) that satisfy the health requirements below; II.1. Part II: Certification II.2. consist exclusively of gelatine/collagen(2) not intended for human consumption; II.3. has 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, in order to kill pathogenic agents; II.4. has been prepared exclusively with the following animal by-products: (<sup>2</sup>)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 ante-mortem 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; heads of poultry; (ii) 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 arising from the production of products intended for human consumption, including degreased bone, greaves and centrifuge or separator sludge from milk processing;] (<sup>2</sup>)and/or products of animal origin, or foodstuffs containing products of animal origin, which are no longer 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 (2)and/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/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 aquatic animals originating from plants or establishments manufacturing products for human consumption;] II.5. the gelatine/collagen(<sup>2</sup>): was wrapped, packaged, stored and transported under satisfactory

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 No	II.b.
			in a dedicated legislation wer Wrappings and	d packages containing gelatine/colla	rmitted under Union gen(2) bear the words
			GELATINE/C CONSUMPTI	COLLAGEN(2) SUITABLE ON'; and	FOR ANIMAL
	( <sup>2</sup> )either	[(b)	unprocessed C or alkali, follo extraction by	E gelatine, was produced by a proc Category 3 material was subjected to owed by one or more rinses, invol- heating one or several times in suc y means of filtration and sterilisatents;]	a treatment with acid lying pH adjustment, ccession, followed by
	( <sup>2</sup> ) <i>or</i>	[(b)	unprocessed C washing, pH a rinses, filtratio	collagen, was produced by a procategory 3 material was subjected to adjustment using acid or alkali follow and extrusion, in order to kill pathon	a treatment involving owed by one or more ogenic agents;]
$(^{2})II.6.$	in the case	e of gelati	ine/collagen(2) f	from materials other than hides and sl	kins
			•	minants than bovine, ovine or caprin	_
		[is derived f		ovine or caprine animals and does	not contain and is not
		( <sup>2</sup> ) either	animals bo region clas	rine and caprine materials other tha rn, continuously reared and slaught sified as posing a negligible BSE ris 007/453/EC.]	tered in a country or
		( <sup>2</sup> )or		ed risk material as defined in poi (EC) No 999/2001 of the European	
			ovine born, c classifi Comm	nically separated meat obtained from caprine animals, except from the continuously reared and slaughtered and as posing a negligible BSE risk ission Decision 2007/453/EC(5), in Igenous BSE case,	ose animals that were in a country or region k in accordance with
			ovine of by lact elonga cavity, for the slaught	by-product or derived product of or caprine animals which have been eration of the central nervous tiss ted rod-shaped instrument introdu or by means of gas injected into the ose animals that were born, cont tered in a country or region cla ble BSE risk in accordance with Dec	killed, after stunning, sue by means of an ced into the cranial cranial cavity, except inuously reared and assified as posing a cision 2007/453/EC.]]
II.7.	_	U	<b>O</b> , ,	m materials other than hides and skin	
	(²)either			or milk products of ovine or caprine armed animals, other than fur animals.	
	$(^2)or$			products of ovine or caprine animal mals, other than fur animals, which:	origin and is intended
		(a)	are derived fron	n ovine and caprine animals which we country where the following condition	-
				al scrapie is compulsorily notifiable;	
			(ii) an awa	reness, surveillance and monitoring	system is in place for

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 No	II.b.		
classical			al scrapie;		
	(iii)		restrictions apply to holdings of over case of a suspicion of TSE or the con- c;		
	(iv)		and caprine animals affected with and destroyed;	classical scrapie are	
	(v)	v) the feeding to ovine and caprine animals of meat-and-bone me or greaves, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health (OIE), of ruminal origin has been banned and effectively enforced in the who country for a period of at least the preceding seven years;			
(b)		te from h	oldings where no official restriction E;	s are imposed due to a	
(c)	diagnos	sed durir	holdings where no case of classing the period of the preceding seven of a case of classical scrapie:		
	( <sup>2</sup> )eithe	and de ARR/A	ine and caprine animals on the hole estroyed or slaughtered, except for ARR genotype, breeding ewes carry and no VRQ allele and other ovin the ARR allele;]	breeding rams of the ing at least one ARR	
	( <sup>2</sup> )or	killed a period last clincludi accorda Chapte the fol except — ar co	amals in which classical scrapie was and destroyed, and the holding has of at least two years since the date classical scrapie case to intensifing testing with negative results for tance with the laboratory methods set C of Annex X to Regulation (EC) lowing animals which are over the ovine animals of the ARR/ARR gentimals which have been slaughnsumption; and mimals which have died or been kill hich were not killed in the frantadication campaign.]]	been subjected for a of confirmation of the ed TSE monitoring, he presence of TSE in et out in point 3.2 of No 999/2001, of all of e age of 18 months, otype: ghtered for human ed on the holding but	

### 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.

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 No	II.b.			
-	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 us	se other than for animal consumpt	on.			
-	Box reference I.26 and I.27: fill in accor	ding to whether it is a transit or ar	import certificate.			
-	Box reference I.28: Species: select from Ruminantia, Pesca.	n the following: Aves, Ruminanti	a, 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.					
(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 different colour to that of the pri	nting.			
-	Note for the person responsible for the consignment in the European Union: this certificate is only					
	for veterinary purposes and must ac inspection post.	company the consignment until	it reaches the border			
Offi	Official veterinarian/Official inspector					
	Name (in capital letters):	Qualif	ication and title:			
	Date: Signature:					
	Stamp:					

### 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

CO	UNTRY:	Veteri	nary certificate to EU				
	I.1. Consignor Name	I.2. Certificate reference No	I.2.a.				
	Address	I.3. Central competent authority					
ınt	Tel.	I.4. Local competent authority					
Part I: Details of dispatched consignment	I.5. Consignee Name Address  Postcode Tel.	I.6. Person responsible for the load in F Name Address  Postcode Tel.	EU				
s of dispa	I.7. Country of origin ISO code I.8. Region of origin Code origin	I.9. Country of ISO code destination	I.10. Region of Code destination				
tails	I.11. Place of origin	I.12. Place of destination	1				
Part I : De	Name Approval number Address Name Approval number Address Name Approval number Address		om warehouse   oval number				
ŀ	I.13. Place of loading	I.14. Date of departure					
	I.15. Means of transport  Aeroplane  Road vehicle  Other  Identification  Other	I.16. Entry BIP in EU  I.17.					
ŀ	Documentation references  I.18. Description of commodity	I 19 Commo	odity code (HS code)				
	1.10. Description of commounty	1.19. Commo	-				
			I.20. Quantity				
	I.21. Temperature of product Ambient □ Chilled □	Frozen	I.22. Number of packages				
	I.23. Seal/Container No		I.24. Type of packaging				
	I.25. Commodities certified for:						
	Animal feedingstuff □ Manufacture of petfood □ Technical use □						
Ī	I.26. For transit through EU to third country						
	Third country ISO code						
•	I.28. Identification of the commodities						
	Approval number of establishments  Species (Scientific name) Nature of commodity Manufacturing plant Number of packages Net weight Batch number						

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

### II.a. Certificate reference No II. Health information 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(<sup>1a</sup>), and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011(<sup>1b</sup>), and in particular Chapter I of Annex XIV thereto, and certify that the hydrolysed protein/dicalcium phosphate/tricalcium phosphate(2) described above: Part II: Certification consists of hydrolysed protein/dicalcium phosphate/tricalcium phosphate(2) that satisfy the II.1. health requirements below; II.2. consists exclusively of hydrolysed protein/dicalcium phosphate/tricalcium phosphate(<sup>2</sup>) not intended for human consumption; II.3. has 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, in order to kill pathogenic II.4. has been prepared exclusively with the following animal by-products: II.4.1. in the case of dicalcium phosphate derived from defatted bones: 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; II.4.2. in the case of other materials: (<sup>2</sup>)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 ante-mortem 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; heads of poultry; (ii) (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 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 (<sup>2</sup>)and/or origin, which are no longer 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

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

II.	I. Health information			II.a.	Certificate referen	nce No	II.b.	
	( <sup>2</sup> )and/or	[-	animal b	y-prod ng for ging d	lucts or derived procommercial reason efects or other de	roducts, wh	or feedingstuffs containing nich are no longer intended problems of manufacturing which no risk to public or	
	(²)and/or	[-	blood, placenta, wool, feathers, hair, horns, hoof cuts and raw originating from live animals that did not show signs of any di communicable through that product to humans or animals;]			show signs of any disease		
	(²)and/or	[-	aquatic a	nimals show	, and parts of such	such animals, except sea mammals, which of diseases communicable to humans or		
	(²)and/or	[-					originating from plants or uman consumption;]	
	(²)and/or	[-	signs of animals:	diseas	se communicable	through th	als which did not show any nat material to humans or	
			* *	follov	m shellfish with so wing originating fro nery by-products,		· ·	
			- (iii) day		by-products, includ hicks killed for co			
II.5.	the hydrolys	sed pro	tein/dicalc	ium pł	osphate/tricalcium	n phosphate	( <sup>2</sup> ):	
		(a)	'NOT FO under sat packagin	OR HU isfacto g tool	MAN CONSUMF ory hygiene conditi	PTION' and in licated room	which bear labels indicating I was stored and transported particular the wrapping and m, and only preservatives and	
	(²)either	[(b)	in the cas	se of h	ydrolysed protein	, was produ	nced by a process involving nation of raw Category 3	
			ruminant only to l preparati	s hides nydroly on of	s and skins, was provided proteins process.	roduced in a luction, usi	ly or partly derived from a processing plant dedicated ng a process involving the all by brining, liming and	
			3 h	ours a t trea	t a temperature of	more than erature of	more than 11 for more than 80 °C and subsequently by more than 140 °C for 30	
			mo	re tha		a heat tre	1 to 2, followed by a pH of attment at a temperature of ar.]	
	(²)or	[(b)	in the cas (i) ens deg (at	e of di ures t greased a min	icalcium phosphate hat all Category 3 I with hot water ar	e, was produ 3 bone-mated was treated won of 4 %	uced by a process that: erial is finely crushed and with dilute hydrochloric acid and a pH of less than 1,5)	

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

II.	Health information	II.a. Certificate reference No II.b.
	(ii)	followed by a treatment of the obtained phosphoric liquor with lime, resulting in a precipitate of dicalcium phosphate at pH 4 to 7, and
	(iii	finally air-dries this precipitate, with an inlet temperature of 65 °C to 325 °C and an end temperature of between 30 °C and 65 °C.]
	$(^2)or$ [(b) in	he case of tricalcium phosphate, was produced by a process ensuring:
	(i)	that all Category 3 bone-material is finely crushed and degreased in counter-flow with hot water (bone chips less than 14 mm),
	(ii)	4 bars,
	(iii	(tricalcium phosphate) by centrifugation, and
	(iv	fluidised bed with air at 200 °C.]
( <sup>2</sup> )II.6.		dicalcium phosphate/tricalcium phosphate(2) described above
	•	om other ruminants than bovine, ovine or caprine animals.]
	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.]
		[(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( <sup>4</sup> );
		(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,
		(c) animal by-product or derived product obtained from bovine, ovine or caprine 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.]]
II.7.	the hydrolysed protein/	dicalcium phosphate/tricalcium phosphate(2) described above:
		contain milk or milk products of ovine or caprine animal origin or is ed for feed for farmed animals, other than fur animals.]
		milk or milk products of ovine or caprine animal origin and is or feed for farmed animals, other than fur animals, which:
	c	re derived from ovine and caprine animals which have been kept ontinuously since birth in a country where the following conditions re fulfilled:
	(i	) classical scrapie is compulsorily notifiable;
	(i	i) an awareness, surveillance and monitoring system is in place 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

			1			
II.	Health information		II.a	i. (	Certificate reference No	II.b.
		(iii) (i	fficia	ıl re cas		gs of ovine or caprine animals the confirmation of classical
					d caprine animals affected destroyed;	ed with classical scrapie are
		t (	or gre he W origin	ave orlo ha y fo	s, as defined in the Terred Organisation for Anims been banned and effector a period of at least the part of the p	- · · · · · · · · · · · · · · · · · · ·
	(b)	origina to a su			_	l restrictions are imposed due
	(c)	diagno	sed d	urir		of classical scrapie has been ling seven years or, following apie:
		( <sup>2</sup> )eith		[al kil rar at	l ovine and caprine anim led and destroyed or slan ns of the ARR/ARR geno	als on the holding have been aghtered, except for breeding type, breeding ewes carrying and no VRQ allele and other
		(²) <i>or</i>		har sul of int neg wir Ch of 18	ve been killed and destro- pjected for a period of at confirmation of the la ensified TSE monitori- gative results for the pre- th the laboratory metho- lapter C of Annex X to R all of the following anim	sical scrapie was confirmed yed, and the holding has been least two years since the date st classical scrapie case to ng, including testing with sence of TSE in accordance ds set out in point 3.2 of egulation (EC) No 999/2001, als which are over the age of animals of the ARR/ARR
				_	consumption; and	been slaughtered for human
				_		died or been killed on the e not killed in the framework 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

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

II. Health information	II.a. Certificate reference No	II.b.
25.04		

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 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, Ruminantia, Mammalia other than Ruminantia, Pesca, Mollusca, Crustacea, invertebrates other than Mollusca and Crustacea.
  - Nature of commodity: specify if hydrolysed protein, dicalcium phosphate or tricalcium phosphate.
  - Manufacturing plant: provide the registration number of treatment/processing establishment.

#### Part II:

- (<sup>1a</sup>) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.
- (<sup>3</sup>) 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 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.

post of the point of entry into the European Union	1.				
Official veterinarian/Official inspector					
Name (in capital letters):	Qualification and title:				
Date: Signature:					
Stamp:					

"

(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  $\binom{2}{2}$  the European Union

CC	OUNTRY:		Veterin	ary certificate to EU			
	I.1. Consignor	I.2. Certificate reference N	Ю	I.2.a.			
	Name Address	I.3. Central competent auth	hority				
ınt	Tel.	I.4. Local competent author	ority				
ıme	I.5. Consignee	I.6. Person responsible for	the load in E	U			
sign	Name Address	Name Address					
con							
peu	Postcode Tel.	Postcode Tel.					
atcl							
disp	I.7. Country of ISO code I.8. Region of origin Code	I.9. Country of	ISO code	I.10. Region of Code			
of (	origin	destination	1	destination			
Part I : Details of dispatched consignment	I.11. Place of origin	I.12. Place of destination	<u> </u>				
Del	Name Approval number		Custo	m warehouse			
t I :	Address Name Approval number	Name Address	Appro	val number			
Par	Address						
	Name Approval number Address	Postcode					
	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 $\square$ Other $\square$ Identification	I.17. Number(s) of CITES					
	Documentation references	1,	10 0	1, (10, 1)			
	I.18. Description of commodity		.19. Commod 5.07	lity code (HS code)			
				I.20. Quantity			
	I.21. Temperature of product Ambient ☐ Chilled ☐	Frozen		I.22. Number of packages			
	I.23. Seal/Container No	110Zell 🗆		I.24. Type of packaging			
	I.25. Commodities certified for:						
	Further process   Technical use						
	I.26. For transit through EU to third country						
	Third country ISO code						
	I.28. Identification of the commodities						
	Approval number of estab Species (Scientific name) Manufacturing pla		ght	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

# Part II: Certification

### 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(<sup>1a</sup>), and Commission Regulation (EU) No 142/2011(<sup>1b</sup>), and in particular Chapter II of Annex XIV thereto, and certify that the horns and horn products, excluding horn meal, and hooves and hoof products, excluding hoof meal(<sup>2</sup>) described above:

- II.1. (²)either [originate from animals that were slaughtered in a slaughterhouse, after undergoing ante-mortem inspection, and were fit, as a result of such inspection, for slaughter for human consumption;]
  - (<sup>2</sup>)or [originate from animals that did not show clinical signs of any disease communicable through that product to humans or animals;]
- 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 and horn products, excluding horn meal, and hooves and hoof products, excluding hoof meal, were packed:
  - (<sup>2</sup>)either [in new packaging or containers;]
  - (²)or [in vehicles or bulk containers disinfected prior to loading using a product approved by the competent authority;]
  - [the packaging or containers are marked so as to indicate the type of the animal by-product(3) and bear labels indicating 'NOT FOR HUMAN AND ANIMAL CONSUMPTION' and the name and address of the establishment of destination in the European Union.]
- (²)II.6. The horns and horn products, excluding horn meal, and hooves and hoof products, excluding hoof meal 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 [(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(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(<sup>5</sup>), in which there has been no indigenous BSE case,
      - (c) animal by-product or derived product obtained from bovine, ovine or caprine 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

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.
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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 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 a
  transit commodity. Products in transit must 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 in the European Union.
- 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 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: 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 products.
- (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 point of entry into the European Union.

"

### (7) Chapter 20 is replaced by the following:

### 'CHAPTER 20

### **Model declaration**

Declaration for the import from third countries and for the transit through 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

CO	OUNTRY:	Veterinary certificate to EU
	I.1. Consignor	I.2. Certificate reference No I.2.a.
	Name Address	I.3. Central competent authority
int	Tel.	I.4. Local competent authority
Part I: Details of dispatched consignment	I.5. Consignee Name Address  Postcode Tel.	I.6. Person responsible for the load in EU Name Address  Postcode Tel.
of dispa	I.7. Country of origin ISO code I.8. Region of origin Code origin	I.9. Country of destination ISO code I.10. Region of destination Code
ails	I.11. Place of origin	I.12. Place of destination
Part I: Det	Name Approval number Address Name Approval number Address Name Approval number	Custom warehouse  Name Approval number Address  Postcode
	Address I.13. Place of loading	I.14. Date of departure
	110. 1 100 01 10101119	The State of departure
	I.15. Means of transport  Aeroplane □ Ship □ Railway wagon □	I.16. Entry BIP in EU
	Road vehicle Other Identification Documentation references	I.17.
	I.18. Description of commodity	I.19. Commodity code (HS code)
		I.20. Quantity
	I.21. Temperature of product Ambient □ Chilled □	I.22. Number of packages Frozen □
	I.23. Seal/Container No	I.24. Type of packaging
	I.25. Commodities certified for:	
	Technical use □	
	I.26. For transit through EU to third country	I.27. For import or admission into EU
	Third country ISO code	
	I.28. Identification of the commodities	1
	Species (Scientific name)  Approval number of establic Manufacturing plant	shments  Net weight  Batch number

Part II: Certification

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.

### **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<sup>(1a)</sup>, 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<sup>(1b)</sup> 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<sup>(2a)</sup> or
    - in Article 2(b) of Council Directive 96/23/EC<sup>(2b)</sup>;]
  - (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 ante-mortem 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;]
  - (2) and/or [- blood of animals which did not show any signs of disease communicable through blood to humans or animals obtained from animals other than

		ruminants that have been slaughtered in a slaughterhouse after having been considered fit for slaughter for human consumption following an antemortem inspection in accordance with Union legislation;]
<sup>(2)</sup> 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;]
<sup>(2)</sup> and/or	[-	products of animal origin, or foodstuffs containing products of animal origin, which are no longer 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;]
<sup>(2)</sup> and/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;]
<sup>(2)</sup> 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;]
<sup>(2)</sup> 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;]
<sup>(2)</sup> and/or	[-	animal by-products from aquatic animals originating from plants or establishments manufacturing products for human consumption;]
<sup>(2)</sup> 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;
		(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-products from aquatic or terrestrial invertebrates other than
		species pathogenic to humans or animals;]
<sup>(2)</sup> and/or	[-	animals and parts thereof of the zoological orders of Rodentia and Lagomorpha, except Category 1 material as referred to in Article 8(a)(iii), (iv) and (v) and Category 2 material as referred to in Article 9(a) to (g) of Regulation (EC) No 1069/2009;]
(2)and/or	[-	products derived from or generated by:
		<ul> <li>aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of disease communicable to humans or animals,</li> </ul>
		<ul> <li>aquatic or terrestrial invertebrates other than species pathogenic to humans or animals,</li> </ul>
		- animals and parts thereof of the zoological orders of Rodentia and Lagomorpha, except Category 1 material as referred to in Article 8(a)(iii), (iv) and (v) and Category 2 material as referred to in Article 9(a) to (g) of Regulation (EC) No 1069/2009;]
(2)and/or	[-	animals and parts of animals, other than those referred to in Article 8 or Article 10 of Regulation (EC) No 1069/2009,
		(i) that died other than by being slaughtered or killed for human consumption, including animals killed for disease control purposes;
		<ul><li>(ii) foetuses;</li><li>(iii) oocytes, embryos and semen which are not destined for breeding</li></ul>
		purposes; and  (iv) dead in shell poultry:
(2)and/or	г	(iv) dead-in-shell poultry;] animal by-products other than Category 1 material or Category 3 material;]
		raging is labelled 'FOR MEDICINAL PRODUCTS / VETERINARY
(Ŧ) IIS OUICI	раск	aging is inscribed for MEDICHAL TRODUCTS / VETERINART

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;

- (2) (5) the consignment will be transported directly to the place of destination in the European Union as indicated under point I.12 of this declaration, that is:
  - 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,
  - an establishment or plant which has been approved in accordance with Article 24(1)(i) of Regulation (EC) No 1069/2009, from where they may only be dispatched to an establishment or plant referred to in the preceding indent of this point.

### **Notes**

- Box reference I.19: use appropriate Harmonised System (HS) code in accordance with Commission Decision 2007/275/EC of 17 April 2007 concerning lists of animals and products to be subject to controls at border inspection posts in accordance with Council Directives 91/496/EEC and 97/78/EC (OJ L 116, 4.5.2007, p.9)
- Box reference I.25: technical use: any use other than for animal consumption.
- <sup>(1a)</sup> OJ L 54, 26.2.2011, p. 1.
- 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 appropriate.
- <sup>(2a)</sup> OJ L 125, 23.5.1996, p. 3.
- (2b) OJ L 125, 23.5.1996, p. 10.

/ /1	
The importer	
Name (in capital letters):	Address:
Date:	Signature:

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