

Brussels, 23 October 2018 (OR. en)

13380/18 ADD 1

AGRILEG 167 VETER 71

COVER NOTE

From:	European Commission
date of receipt:	17 October 2018
To:	General Secretariat of the Council
No. Cion doc.:	D052851 ANNEXES 1 to 2
Subject:	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

Delegations will find attached document D052851 ANNEXES 1 to 2.

Encl.: D052851 ANNEXES 1 to 2

13380/18 ADD 1 OT/mb

LIFE.2.B EN



Brussels, XXX SANTE/7008/2017 ANNEX (POOL/G2/2017/7008/7008-EN ANNEX.doc) D052851/03 [...](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

ANNEX I

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

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

'SECTION B

Health certificate requirements

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

reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC;

or

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

or

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

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

- 3. In addition to the requirements of points 1 and 2 of this Section, imports of the animal by-products and derived products referred to in Section A, containing milk or milk products of ovine or caprine animal origin and intended for feed, shall be subject to the presentation of a health certificate which has been completed with the following attestation:
 - (a) the ovine and caprine animals from which those animal by-products or derived products have been derived have been kept continuously since birth in a country where the following conditions are fulfilled:
 - (i) classical scrapie is compulsorily notifiable;
 - (ii) an awareness, surveillance and monitoring system is in place;
 - (iii) official restrictions apply to holdings of ovine or caprine animals in the case of a suspicion of TSE or a confirmation of classical scrapie;
 - (iv) ovine and caprine animals affected with classical scrapie are killed and completely destroyed;

- (v) the feeding to ovine and caprine animals of meat-and-bone meal or greaves of ruminant origin, as defined in the OIE Terrestrial Animal Health Code, has been banned and effectively enforced in the whole country for a period of at least the preceding seven years;
- (b) the milk and milk products of ovine or caprine animals originate from holdings where no official restrictions are imposed due to a suspicion of TSE;
- (c) the milk and milk products of ovine or caprine animals originate from holdings where no case of classical scrapie has been diagnosed during a period of at least the preceding seven years or, following the confirmation of a case of classical scrapie:
 - (i) all ovine and caprine animals on the holding have been killed and destroyed or slaughtered, except for breeding rams of the ARR/ARR genotype, breeding ewes carrying at least one ARR allele and no VRQ allele and other ovine animals carrying at least one ARR allele;

or

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

ANNEX II

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

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

'CHAPTER 1

Health certificate

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

COUNTRY: Veterinary certificate to EU

	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 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 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 Ambient □ Chilled □	I.22. Number of packages					
	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										
Approval number of establishments											
	Species (Scientific name)	Nature of commodity	Manufacturing plant	Net weight	Batch number						

Health information

Part II: Certification

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

	11.	пеанн	шогшаноп		11.a.	Certificate	reference No	1	11.0.	
		I, the undersigned official veterinarian, declare that I have read and understood Regulati (EC) No 1069/2009 of the European Parliament and of the Council(^{1a}) and in particular Artic 10 thereof, and Commission Regulation (EU) No 142/2011(^{1b}), and in particular Section 1 Chapter II of Annex X, and Chapter I of Annex XIV thereto and certify that:								
	II.1.		cessed animal pronot intended for				above contain	ıs exclu	sively processed animal	
(a) has been prepared and stored in an establishment or plant approved and supervision the competent authority in accordance with Article 24 of Regulation No 1069/2009, and										
		(b)	has been prepa	ared excl	usively	with the fo	ollowing anim	al by-p	roducts:	
			(²)either [-	bodies consum	or par nption	rts of anii in accord	nals killed, a	and wh	or, in the case of game, nich are fit for human egislation, but are not cial reasons;]	
		(²)and/or [- carcases and the following parts originating either from anima have been slaughtered in a slaughterhouse and were considered slaughter for human consumption following an anterminspection or bodies and the following parts of animals from killed for human consumption in accordance with Union legisla							d were considered fit for wing an ante-mortem s of animals from game	
							•		Is which are rejected as ccordance with Union	

II a Certificate reference No.

- (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;

legislation, but which did not show any signs of disease

- (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;]

communicable to humans or animals;

- (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;]
- (²) and/or [- aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of diseases communicable to humans

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. Certificate reference No	II.b.			
		or an	mals;]				
	(²)and/or	estab	l by-products from aquatic anii ishments or plants manufacturing mption;]	mals originating from products for human			
	(²)and/or	any s	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 terrestri	al animals:			
			- hatchery by-products,				
			- eggs,				
			egg by-products, including egg s	hells;			
		(iii)	day-old chicks killed for commercial re	easons;]			
	(²)and/or		c and terrestrial invertebrates other that or animals and other than insects;]	an species pathogenic to			
	(²)and/or	Lago Artic	Is and parts thereof of the zoological morpha, except Category 1 mater e 8(a)(iii), (iv) and (v) and Category 2 e 9(a) to (g) of Regulation (EC) No 100	ial as referred to in material as referred to in			
	and						
	(c) has been	subjected to	the following processing standard:				
	(²)either	without int	a core temperature of more than 133°C erruption at a pressure (absolute) of at eam, with a particle size prior to procees;	least 3 bars produced by			
	(²)or	method 1-2	of non-mammalian protein other than 3-4-5-7(indicat Chapter III of Annex IV to Regulation	e the processing method)			
	(²)or	[in the 7Chapter III	case of fishmeal the processing(indicate the processing of Annex IV to Regulation (EU) No 14				
	(²)or	(indicate the Regulation	e of porcine blood, the processing me e processing method) as set out in Ch (EU) No 142/2011, where in case of m 0 °C has been applied throughout its su	apter III of Annex IV to nethod 7 a heat treatment			
II.2.	the competent aut to comply with the		ned a random sample immediately prior and ards(3):	to dispatch and found it			
	Salmonella:	Absen	e in 25 g: $n = 5$, $c = 0$, $m = 0$, $M = 0$				
	Enterobacteriacea	e: $n = 5$,	= 2, $m = 10$, $M = 300$ in 1g;				
II.3.	the product has u after treatment;	indergone al	precautions to avoid recontamination	with pathogenic agents			
II.4.	the end product:						
	(²)either [was	packed in no	w or sterilised bags,]				
	(2) <i>or</i> [was	transported	in bulk in containers or other mean	s of transport that were			

(2)either

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

			ption including mixtures a tfood containing such prote	-						
II.	Health information	II.	a. Certificate reference No	II.b.						
	thoroughly cleaned	d and	disinfected before use,]							
	which bear labels indicating 'NOT FOR HUMAN CONSUMPTION';									
II.5. the end product was stored in enclosed storage;										
(²)[II.6.	(²)[II.6. the processed animal protein or product described above contains or is derived from animal-b products of ruminant origin and:									
	negligible BSE	E risk	a country or region, which is in accordance with Decision 200 adigenous BSE case, and]]							
	in accordance indigenous BS derived from a of ruminants was defined in t	e with SE can with the C	country or region classified as posith Decision 2007/453/EC in what ase, and the animal by-product calls born after the date from which meat-and-bone meal and greaves DIE Terrestrial Animal Health Country or region, and]	ich there has been an or derived product were in the ban on the feeding derived from ruminants,						
	(2)either [is derived from	m otł	her ruminants than bovine, ovine o	r caprine animals.]						
	(²)or [is derived from not derived from		vine, ovine or caprine animals and	d does not contain and is						
	f	from coun	animals born, continuously rearctry or region classified as posing and ance with Decision 2007/453/EC	ed and slaughtered in a a negligible BSE risk in						
	(²)or [[(a)	specified risk material as defined to Regulation (EC) No 999/2 Parliament and of the Council(4);							
		(b)	mechanically separated meat of bovine, ovine or caprine animanimals that were born, conslaughtered in a country or region negligible BSE risk in accordance Decision 2007/453/EC(5), in whindigenous BSE case,	als, except from those ntinuously reared and on classified as posing a ance with Commission						
		(c)	animal by-product or derived bovine, ovine or caprine animals after stunning, by laceration of the by means of an elongated introduced into the cranial cavity, of that were born, continuously read country or region classified as prisk in accordance with Decision	which have been killed, ne central nervous tissue rod-shaped instrument ty, or by means of gas except for those animals red and slaughtered in a posing a negligible BSE						
II.7.	the processed animal protein or p	prodi	act described above:							

[contains milk or milk products of ovine or caprine animal origin and is intended $(^{2})or$ for feed for farmed animals, other than fur animals, which:

intended for feed for farmed animals, other than fur animals.]

are derived from ovine and caprine animals which have been kept

[does not contain milk or milk products of ovine or caprine animal origin or is not

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

II.	Health info	ormation	1		II.a.	. Ce	rtifica	te refe	erence N	lo	II.b.	
			continue		y sinc	ce bi	rth in	a cou	ntry wh	ere the	following condition	ons are
			(i)	cla	ıssical	l scra	apie is	comp	ulsorily	notifiab	e;	
			(ii)		aware assical			eillan	ce and	monitorii	ng system is in p	lace for
			(iii)	in		ase o					ovine or caprine confirmation of c	
			(iv)		ine ar led an				nals aff	ected wi	th classical scra	pie are
			(v)	or the ori	greave Worligin ha	ves, a rld C nas b	as defi Organis een b	ned ir sation anned	the Te for An and ef	errestrial iimal He fectively	of meat-and-box Animal Health (alth (OIE), of ru enforced in the ing seven years;	Code of uminant
		(b)	originat suspicio				gs whe	ere no	official	restriction	ons are imposed	due to a
		(c)	diagnos	ed	during	g a	period	d of a	at least		ssical scrapie ha ceding seven ye crapie:	
			(²)eithei	an AF all	d dest RR/AF	troye RR g nd n	ed or genoty o VRO	slaugh pe, br Qalle	ntered, or reeding	except for ewes can	olding have been breeding rams rying at least or ine animals carr	of the are ARR
			(²)or	kill per las inc acc Ch the	lled and riod of classic classic classic cordand apter follows and cordand apter follows.	nd dof at assicant as	estroyeleast to all screeting with the fanne g anim	ed, and wo ye apie with ne lab ex X to mals	ears since case to egative a coratory of Regula which a	olding he the date intense results for methods ation (EC	as confirmed has been subjected e of confirmation of the presence of the set out in point the age of 18 the age of 18 the enotype:	ed for a n of the nitoring, TSE in t 3.2 of of all of
				_			als w imptic			been si	aughtered for	human
				_	b	but w	which '	were 1			n killed on the framework of a	
II.8.											derived from an ne Consignor ref	
	(²)either	[not ir animal		for	the p	orodu	ection	of fe	ed for	farmed	animals, other t	han fur
	(²)(⁶)or	fur an Inspec	imals, a tion Post	nd of	the C	Cons will	ignor be pro	has ovided	undertal d with t	ken to o	med animals, othersure that the sof the analyses ex VI to Com	Border carried

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.

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

Notes

Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is required to be filled in only if it is a certificate for a commodity to be transited through the European Union; it may be filled in if the certificate is for a commodity that is to be imported into the European Union.
- 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 (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 feeding of farmed animals, other than fur animals, and the production or manufacturing of pet food.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: Species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or Suidae, Pesca, Mollusca, Crustacea, invertebrates other than Mollusca and Crustacea. In the case of farmed fish, specify the scientific name of the fish.

Part II:

- (la) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.
- (3) Where:
 - n = number of samples to be tested;
 - m = threshold value for the number of bacteria; the result is considered satisfactory if the number of bacteria in all samples does not exceed m;
 - M = maximum value for the number of bacteria; the result is considered unsatisfactory if the number of bacteria in one or more samples is M or more; and
 - c = number of samples the bacterial count of which may be between m and M, the sample still being considered acceptable if the bacterial count of the other samples is m or less.
- (4) OJ L 147, 31.5.2001, p. 1.
- (5) OJ L 172, 30.6.2007, p. 84.
- (6) The Person responsible for the load referred to in Box I.6 must ensure that, if the processed animal protein or product described in this health certificate is intended to be used for the production of feed for non-ruminant farmed animals, other than fur animals, the consignment must be analysed, in accordance with the methods set out in Annex VI to Regulation (EC) No 152/2009, in order to verify the absence of unauthorised constituents of animal origin. The information on the result of such analysis must be attached to this health certificate when presenting the consignment at an EU border inspection post.
- (⁷) OJ L 54, 26.2.2009, p. 1.

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.						
_	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.								
Offi	cial veterinarian/Official inspector								
	Name (in capital letters):	Qu	alification and title:						
	Date:	Sig	gnature:						
	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 through² 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				
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				
ils (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 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 Ambient □ Chilled □	I.22. Number of packages				
	I.23. Seal/Container No	I.24. Type of packaging				
	I.25. Commodities certified for:					
	Animal feedingstuff □ Tech	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	Approval 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(^{1a}) and 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:

- 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:
 - 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);]
 - (2) 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);]
 - (2) and/or [- Field Cricket (Gryllus assimilis).]

and

(c) has been processed by method [1]-[2]-[3]-[4]-[5]-[7](2) 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(3):

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

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.					
	after treatme	nt;	ı							
II.4.	the end prod	and product:								
	$(^2)$ either	[was packed in new	s packed in new or sterilised bags,]							
	$(^2)or$		was transported in bulk in containers or other means of transport that was acroughly cleaned and disinfected before use,]							
	PROTEIN -	labels indicating 'NOT FOR HUMAN CONSUMPTION/ PROCESSED INSECT – SHALL NOT BE USED IN FEED FOR FARMED ANIMALS EXCEPT TURE AND FUR ANIMALS';								
II.5.	the end prod	uct was stored in er	nclos	ed storage;						
(²)[II.6.	5. the processed animal protein or product described above contains or is derived from animal-by products of ruminant origin and:									
	(²)ei	BSE risk in a	ccor	country or region, which is classified dance with Decision 2007/453/EC, as BSE case, and]]						
	(2)01	(2) or [originates from a country or region classified as posing a negligible BSE risin accordance with Decision 2007/453/EC in which there has been a indigenous BSE case, and the animal by-product or derived product we 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 ruminant as defined in the OIE Terrestrial Animal Health Code, has been effective enforced in that country or region, and]]								
	(²)ei	ither [is derived fro	m ot	her ruminants than bovine, ovine or o	caprine animals.]]					
	$(^{2})oi$	r [is derived fro not derived fro		ovine, ovine or caprine animals and d	loes not contain and is					
			from cour	ine, ovine and caprine materials of animals born, continuously reared atry or region classified as posing a rdance with Decision 2007/453/EC.]	and slaughtered in a negligible BSE risk in					
		(²)or	[(a)	specified risk material as defined i to Regulation (EC) No 999/200 Parliament and of the Council(4);						
		•	(b)	mechanically separated meat obta bovine, ovine or caprine animals animals that were born, conti- slaughtered in a country or region negligible BSE risk in accordan Decision 2007/453/EC(5), in which indigenous BSE case,	s, except from those nuously reared and classified as posing a ce with Commission					
			(c)	animal by-product or derived pr bovine, ovine or caprine animals w after stunning, by laceration of the by means of an elongated re- introduced into the cranial cavity, injected into the cranial cavity, ex- that were born, continuously reared country or region classified as pos- risk in accordance with Decision 20	hich have been killed, central nervous tissue od-shaped instrument or by means of gas cept for those animals d and slaughtered in a sing a negligible BSE					

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

II.	Health inf	ormatio	n	II.a. Certificate reference No II.b.					
II.7.	the process	sed anim	al proteir	n or product described above:					
	(²)either	intended for feed for farmed animals, other than fur animals.]							
	$(^2)or$		[contains milk or milk products of ovine or caprine animal origin and is intende for feed for farmed animals, other than fur animals, which:						
		(a)		erived from ovine and caprine animals which have been kep uously since birth in a country where the following conditions are ed:					
			(i)	classical scrapie is compulsorily notifiable;					
			(ii)	an awareness, surveillance and monitoring system is in place for classical scrapie;					
			(iii)	official restrictions apply to holdings of ovine or caprine animals in the case of a suspicion of TSE or the confirmation of classica scrapie;					
			(iv)	ovine and caprine animals affected with classical scrapie are killed and destroyed;					
			(v)	the feeding to ovine and caprine animals of meat-and-bone mea or greaves, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health (OIE), of ruminan origin has been banned and effectively enforced in the whole country for a period of at least the preceding seven years;					
		(b)		ate from holdings where no official restrictions are imposed due to a ion of TSE;					
		(c)	diagno	ate from holdings where no case of classical scrapie has beer osed during a period of at least the preceding seven years or ring the confirmation of a case of classical scrapie:					
			(²)eithe	er [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 ARF allele and no VRQ allele and other ovine animals carrying a least one ARR allele;]					
			(²) <i>or</i>	[all animals in which classical scrapie was confirmed have beer killed and destroyed, and the holding has been subjected for a period of at least two years since the date of confirmation of the last classical scrapie case to intensified TSE monitoring including testing with negative results for the presence of TSE in accordance with the laboratory methods set out in point 3.2 of Chapter C of Annex X to Regulation (EC) No 999/2001, of all of the following animals which are over the age of 18 months except ovine animals of the ARR/ARR genotype:					
				 animals which have been slaughtered for human consumption; and 					
				 animals which have died or been killed on the holding but which were not killed in the framework of a disease eradication campaign.]] 					

products of non-ruminant origin and is, according to the statement of the Consignor referred to

Processed animal protein 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.
	in Box I.1,			
	(²)either	[not intended for thanimals.]	e production of feed for farmed an	imals, other than fur
	(²)(⁶)or	fur animals, and the inspection post of ent of the analyses carrie	duction of feed for non-ruminant farm the Consignor has undertaken to entry into the European Union will be producted out in accordance with the methods sion (EC) No 152/2009(7).]	sure that the border ovided with the results

Notes

Part I:

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

Part II:

- (1a) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.
- (3) Where:
 - n = number of samples to be tested;
 - m = threshold value for the number of bacteria; the result is considered satisfactory if the number of bacteria in all samples does not exceed m;
 - M = maximum value for the number of bacteria; the result is considered unsatisfactory if the number of bacteria in one or more samples is M or more; and
 - 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

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.						
	information on the result of such analysis must be attached to this health certificate when presenting the consignment at an EU Border Inspection Post.								
(7)	OJ L 54, 26.2.2009, p. 1.								
_	The signature and the stamp must be	in a different colour to that of the prir	iting.						
_	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.								
Offi	cial veterinarian/Official inspector								
	Name (in capital letters):	Qualifi	cation and title:						
	Date:	Signatu	ire:						
	Stamp:								

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

CU	OUNIKY:			veterin	ary certificate to	EU			
	I.1. Consignor Name		I.2. Certificate reference	No	I.2.a.				
	Address		I.3. Central competent authority						
änt	Tel.		I.4. Local competent auth	nority					
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.						
atch									
of disp	I.7. Country of ISO code I.8. I origin	Region of origin Code	I.9. Country of destination	ISO code	I.10. Region of destination	Code			
tails	I.11. Place of origin	-	I.12. Place of destination	I					
Part I : De	Name Address Name Address Name	Approval number 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		I.16. Entry BIP in EU						
	Aeroplane	Railway wagon 🗆	I.17. Number(s) of CITES						
	I.18. Description of commodity	<u>'</u>	I.19. Commodity code (HS code)						
			L		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 ☐ Technical use ☐	Further p	process		Production of petfoc	od 🗆			
	I.26. For transit through EU to third cou	untry	I.27. For import or ac	lmission into EU	U \square				
	Third country	ISO code							
ŀ	I.28. Identification of the commodities								
	Species (Scientific name)	Approval number of establi Manufacturing plant		weight	Batch nu	ımber			

Part II: Certification

II.	Health	informatio	on	II.a. Certificate reference No	II.b.				
	(EC) N Article Section milk(2),	I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council(^{1a}), and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011(^{1b}), and in particular Section 4 of Chapter II of Annex X, and Chapter I of Annex XIV thereto, and certify that the milk(²), the milk-based products(²) and milk-derived products(²) referred to in box I.28 comply with the following conditions:							
II.1.	Annex I and-mo	$O(^3)$, If to Commuth disease	nission Regulat e (FMD) and	in	h is listed in Part I of has been free from foot- is immediately prior to				
II.2.	show cl had bee subject	inical signs n kept for to official	s of any disease a period of at restrictions due	lk derived from animals which at the e transmissible through milk to human least 30 days prior to production on to foot-and-mouth disease or rinderpe	s or animals, and which holdings that were not				
II.3.	-		ilk products tha						
	(²)eithe	r[have un point II.4		of the treatments or combinations	s thereof described in				
	(²) <i>or</i>		whey was collec	d to animals of species susceptible to cted from milk subjected to one of the					
		(²)either	[the whey was below 6;]	vas collected at least 16 hours after clotting and has a pH					
		(²)(⁵)or		as been produced at least 21 days before the shipping and eriod no cases of FMD have been detected in the exporting					
		(²)(⁵)or	foreseen voya	as been produced on//, this date, in consideration of the age duration, being at least 21 days before the consignment is a border inspection post of the European Union;]]					
II.4.	they hav	e been sul	oject to one of t	he following treatments:					
	(²)eithe	equivalen		time pasteurisation at 72°C for at a achieving a negative reaction to a ph:h:					
		(²)either	at least 15 sec	second high temperature short time particles or an equivalent pasteurisation ion to a phosphatase test in bovine mi	which itself achieves a				
		(²) <i>or</i>		drying process that in the case of mill hadditional heating to 72°C or higher					
		(²) <i>or</i>	[a subsequent hour at a level	process by which the pH is reduced a below 6;]	and kept for at least one				
		$^{(2)(5)}or$	prior to the da	that the milk/milk product has been pate of shipping and during that period in the exporting country;]					
		(²)(⁵)or	date, in consi days prior to	deration of the foreseen voyage dure the date that the consignment is st of the European Union;]	ation, being at least 21				
		.0.	- •						

[sterilisation at a level of at least F_03 ;]]

[ultra high temperature treatment at 132°C for at least one second in combination

 $(^{2})or$

 $(^{2})or$

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

II.	Health info	rmatio	n	II.a. Certificate reference No II.b.				
	wit	h:						
	$(^2)\epsilon$	either		nt drying process that in the case of milk intended for feeding is rith additional heating to 72°C or higher;]				
	$(^2)\alpha$	or		nt process by which the pH is reduced and kept for at least one rel below 6;]				
	(2)((⁵)or	prior to the	on that the milk/milk product has been produced at least 21 days date of shipping and during that period no cases of FMD has ed in the exporting country;]				
	(2)((⁵)or	date, in condays prior	ilk product has been produced on/(insert the date), this insideration of the foreseen voyage duration, being at least 21 to the date that the consignment is presented to a border post of the European Union;]]				
II.5.	every preca derived prod			o avoid contamination of the milk/milk-based product/milk-g;				
II.6.	the milk/mil	k-base	d product/m	lk-derived product was packed:				
	(²)either	[in n	ew container	s;]				
	$(^2)or$	_		bulk containers disinfected prior to loading using a product competent authority;]				
	and	prod	uct/milk-der	e marked so as to indicate the nature of the milk/milk-based wed product and bear labels indicating that the product is ial and not intended for human consumption;				
II.7.	the milk, mi	lk-base	ed products a	nd milk-derived products described above:				
	(²)either	-		milk or milk products of ovine or caprine animal origin or is not for farmed animals, other than fur animals.]				
	$(^2)or$			milk products of ovine or caprine animal origin and is intended ed animals, other than fur animals, which:				
		(a)		yed from ovine and caprine animals which have been kept usly since birth in a country where the following conditions are				
			(i) c	lassical scrapie is compulsorily notifiable;				
				n awareness, surveillance and monitoring system is in place for lassical scrapie;				
			i	fficial restrictions apply to holdings of ovine or caprine animals a the case of a suspicion of TSE or the confirmation of classical crapie;				
				vine and caprine animals affected with classical scrapie are illed and destroyed;				
			ti o	ne feeding to ovine and caprine animals of meat-and-bone meal r greaves, as defined in the Terrestrial Animal Health Code of ne World Organisation for Animal Health (OIE), of ruminant rigin has been banned and effectively enforced in the whole ountry for a period of at least the preceding seven years;				
		(b)	originate suspicion	from holdings where no official restrictions are imposed due to a of TSE;				
		(c)	diagnosed	from holdings where no case of classical scrapie has been during a period of at least the preceding seven years or, the confirmation of a case of classical scrapie:				
			(²)either [all ovine and caprine animals on the holding have been killed				

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

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

Notes

Part I:

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

Part II:

- (1a) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.
- (3) For completion if the authorisation to import into or transit through the European Union is restricted to certain regions of the third country concerned.
- (4) OJ L 175, 10.7.2010, p. 1.
- (5) this condition applies only to third countries listed in column 'A' of Annex I to Regulation (EU) No 605/2010.

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

II.	Health information	II.a. Certificate reference No	О	II.b.				
(6)	OJ L 147, 31.5.2001, p. 1.							
(7)	OJ L 172, 30.6.2007, p. 84.							
_	The signature and the stamp must be in a 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.							
Offi	cial veterinarian/Official inspector							
	Name (in capital letters):		Qualifi	cation and title:				
	Date:		Signatu	ıre:				
	Stamp:							

CHAPTER 2(B)

Health certificate

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

CO	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.	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					
	L15 Manus of transport	I.16. Entry BIP in EU					
	I.15. Means of transport Aeroplane Road vehicle Other Other	I.17. Number(s) of CITES					
	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 Frozen □					
	I.23. Seal/Container No	I.24. Type of packaging					
	I.25. Commodities certified for: Animal feedingstuff □ Furth Technical use □	er process Production of petfood					
	126 For transit through EU to third country	1.27 For import or admirrion into FU					
	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 estr Manufacturing pla						

Notes

			animals not for human consumption								
		II.	Health info	ormation	II.a. Certificate reference No II.b.						
Ī	uo		(EC) No Article 10 Section 4	1069/2009 of the thereof, and Coo of Chapter II of A ²) or the colostrur	veterinarian, declare that I have read and understood Regulation European Parliament and of the Council(1a), and in particular mmission Regulation (EU) No 142/2011(1b), and in particular than X and Chapter I of Annex XIV thereto, and certify that the m products(2) referred to in box I.28 comply with the following						
	Part II: Certification	II.1.	country)(3) Commission disease (FM	they were produced and derived in							
	Part	II.2.	show clinic which had	cal signs of any di been kept for a per	ostrum derived from animals which at the time of milking did not isease transmissible through colostrum to humans or animals, and criod of at least 30 days prior to the date of production on holdings ial restrictions due to foot-and-mouth disease or rinderpest;						
		II.3.	temperatur	e short time past ion achieving a n	trum products of bovine animals that have been subject to high teurisation at 72°C for at least 15 seconds, or an equivalent negative reaction to a phosphatase test in bovine colostrum, in						
			(²)(⁵)either	a period at least 2	at the colostrum or colostrum products have been produced during 21 days before the date of shipping and during this period no cases en detected in the exporting country,]						
			(²)(⁵)or	date), this date, i	or colostrum products have been produced on/(insert the in consideration of the foreseen voyage duration, being at least 21 consignment is presented to a border inspection post of the ,]						
			and		ined from animals subject to regular veterinary inspections to come from holdings on which all bovine herds are:						
				$(^{2})(^{5})either$ [2]	recognised as officially tuberculosis and brucellosis free(6),]						
					not restricted under the national legislation of the third country of origin for the eradication of tuberculosis and brucellosis,]						
			and	$(^{2})(^{5})either$ [2]	recognised as official enzootic-bovine-leukosis-free(6),]						
				included in an official system for the control of enzootic bovine eukosis and there has been no evidence as result of clinical and aboratory testing of this disease in the herd during the period of he preceding two years,]]							
		II.4.	every preca		aken to avoid contamination of the colostrum/colostrum product						
		II.5.	the colostru	ım or colostrum pr	roduct was packed:						
			(²)either	er [in new containers,]							
			$(^2)or$		bulk containers disinfected prior to loading using a product competent authority,]						
			and	product and bear	re marked so as to indicate the nature of the colostrum/colostrum r labels indicating that the product is Category 3 material and not nan consumption;						
		II.6.	the colostru animal orig		product does not contain milk or milk products of ovine or caprine						

Colostrum and colostrum products from bovine animals not for human consumption

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

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 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 feeding of farmed animals, other than fur animals, and the production or manufacturing of pet food..
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: 'Manufacturing plant': provide the registration number of the treatment or processing establishment.

Part II:

- (1a) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.
- (3) For completion if the authorisation for introduction into the European Union is restricted to certain regions of the third country concerned.
- (4) OJ L 175, 10.7.2010, p. 1.
- (5) This condition applies only to third countries authorised in column 'A' of Annex I to Commission Regulation (EU) No 605/2010 (OJ L 175, 10.7.2010, p. 1).
- (6) 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-leukosis-free 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

U	OUNTRY:						Veterii	nary certificat	e to EU	
	I.1. Consignor				I.2.	Certificate reference	e No	I.2.a.		
	Name Address				I.3.	Central competent a	uthority			
int	Tel.				I.4.	Local competent au	thority			
me	I.5. Consignee				I.6.	Person responsible f	for the load in E	EU		
ısign	Name Address					Name Address				
cor	Postcode					Postcode				
ched	Tel.					Tel.				
ispa	I.7. Country of	ISO code	I.8. Region of origin	Code	I.9. C	Country of	ISO code	I.10. Region of	Code	
Part I: Details of dispatched consignment	origin	1				estination		destination	ĺ	
tails	I.11. Place of orig	in			I.12.	Place of destination			<u> </u>	
Ğ	Name		Approval number	r				om warehouse		
t I :	Address Name		Approval numbe			Name Address	Appro	oval number		
ar	Address		Approvai numoc	1		Address				
1	Name Address		Approval numbe	r	Postcode					
•	I.13. Place of load	ling			I.14. Date of departure					
	I.15. Means of tra	nsport			I.16.	Entry BIP in EU				
	Aeroplane	Shi	p Railway wagor	n 🗆						
	Road vehicle	e 🗆 Oth	ier 🗆		I.17.					
	Identification Documentati	ion references								
	I.18. Description	of commodity			I.19. Commodity code (HS code) 23.09					
								I.20. Quantity		
•	I.21. Temperature Ambient	of product	Chilled □			Frozen		I.22. Number of p	ackages	
	I.23. Seal/Contain	ier No	Cilinea 🗆		riozen 🗆			I.24. Type of pack	I.24. Type of packaging	
•	I.25. Commodities	s certified for:								
	Petfood			Techr	nical us	е				
	I.26. For transit th	rough EU to th	ird country			I.27. For import or a	admission into I	EU		
	Third countr	y	ISO code							
	I.28. Identification	n of the commo	dities							
		22 222 00111110								
	Species (Scientif	fic name)	Approval nur Manu	nber of es facturing			t weight	Ва	atch number	

COUNTRY Canned Petfood

II. Health information II.a. Certificate reference No II.b. I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC)

I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council(^{1a}), and in particular Articles 8 and 10 thereof, and Commission Regulation (EU) No 142/2011(^{1b}), and in particular Chapter II of Annex XIII and Chapter II of Annex XIV, thereto and certify that the petfood described above:

- II.1. has been prepared and stored in an establishment or plant approved and supervised by the competent authority in accordance with Article 24 of Regulation (EC) No 1069/2009;
- II.2. has been prepared exclusively with the following animal by-products:
 - (2) either [- carcases and parts of animals slaughtered or, in the case of game, bodies or parts of animals killed, and which are fit for human consumption in accordance with Union legislation, but are not intended for human consumption for commercial reasons;]
 - (²) and/or [- carcases and the following parts originating either from animals that have been slaughtered in a slaughterhouse and were considered fit for slaughter for human consumption following an 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 [- animal by-products from poultry and lagomorphs slaughtered on the farm as referred to in Article 1(3)(d) of Regulation (EC) No 853/2004 of the European Parliament and of the Council(2ª), which did not show any signs of disease communicable to humans or animals]
 - (²) and/or [- blood of animals which did not show any signs of disease communicable through blood to humans or animals, obtained from animals that have been slaughtered in a slaughterhouse after having been considered fit for slaughter for human consumption following an ante-mortem inspection in accordance with Union legislation;]
 - (²) and/or [- animal by-products arising from the production of products intended for human consumption, including degreased bone, greaves and centrifuge or separator sludge from milk processing;]
 - (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 [- 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;]
 - (2) and/or [- blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk originating from live animals that did not show signs of any disease communicable through that product to humans or animals;]
 - (²) and/or [- aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of diseases communicable to humans or animals;]

COUNTRY Canned Petfood

II.	Health inf	format	ion		II.a.	Certificate	reference No		II.b.		
	(²)and/or	[-			ts from aquatic animals originating from plants or ufacturing products for human consumption;]						
	(²)and/or	[-			rial originating from animals which did not show any signs cable through that material to humans or animals:						
						•	sue or flesh;				
			(ii) the f	Collowing o	origina	iting from te	rrestrial anim	als:			
			-	hatchery b	y-pro	ducts,					
			-	eggs,							
			-	egg by-pro	oducts	, including e	egg shells;				
			(iii) day-	old chicks	killed	for comme	rcial reasons;]]			
	(²)and/or	[-	animal by pathogenic				errestrial inve	ertebr	ates other	than spec	ies
	(²)and/or	[-	except Ca	tegory 1 i i (EC) No	materia 1069/2	al as referre 2009 and Ca	gical orders of ed to in Artic ategory 2 mate	ele 8	(a)(iii), (iv)	and (v)	of
	(²)and/or	[-	are prohib	oited by C mitted in	ouncil	Directive	en treated wi 96/22/EC(^{2b}), Article 35(a)	the	import of	the mater	rial
II.3.	has been containers		cted to hea	at treatme	nt to	a minimun	n Fc value o	of 3	in hermetic	cally seal	led
II.4.		/ diagn	ostic meth				samples from				
	has under	_	-		d conta	amination w	ith pathogenio	c age	nts after tre	atment.	
()[11	(²)either				inants	than boyine	, ovine or cap	rine	animals 1		
	$^{(2)}or$	[is de					animals and		-	n and is 1	not
			ther [bovir born, as po	continuou	sly rea	ired and sla	als other than ughtered in a risk in a	coun	try or regio	on classifi	ied
		(²)o.	r [(a)		on (EC		s defined in 001 of the Eur				
			(b)	or caprir continuou as posing	ne ani usly re g a ne 2007/	mals, exceptared and slaggligible BS	eat obtained for from those aughtered in a SE risk in ac in which the	e and cour corda	imals that ntry or region nce with	were bo on classifi Commissi	orn, ied ion
			(c)	or caprin laceration rod-shape means o	ne ani n of the ed ins f gas	mals which ne central natrument int injected in	ved product on have been ervous tissue roduced into to the crania ontinuously r	kille by the al ca	ed, after s means of a cranial ca vity, excep	tunning, n elongat vity, or ot for the	by ted by ose

COUNTRY Canned Petfood

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.23: for bulk containers, the container number and the seal number (if applicable) must be given.
- Box reference I.25: technical use: any use other than feeding of farmed animals, other than fur animals, and the production or manufacturing of pet food..
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: Species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or Suidae, Pesca, Mollusca, Crustacea, invertebrates other than Mollusca and Crustacea.

Part II:

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

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

CO	UNTRY:	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 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					
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 I.17.					
	Aeroplane						
	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	•					
	Approval number of Species (Scientific name) Approval number of Manufacturin						
	. , , , , , , , , , , , , , , , , , , ,						

		II.	Health informat	tion		II.a.	Certificate reference No		II.b.		
I, the undersigned official veterinaria No 1069/2009 of the European Parlia 10 thereof, and Commission Regula Annex XIII and Chapter II of Annex							d of the Council(1a), and J) No 142/2011(1b), and	in pai	rticular Articles 8 and rticular Chapter II of		
	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;								
	fica	II.2.	has been prepared exclusively with the following animal by-products:								
	Part II: Certification		(²) either [-	(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;]							
	Pari	(²)and/or [- carcases and the following parts originating either from animals that slaughtered in a slaughterhouse and were considered fit for slaughter consumption following an ante-mortem inspection or bodies and the parts of animals from game killed for human consumption in according to the carcases and the slaughtered in a slaughterhouse and were considered fit for slaughter consumption or bodies and the parts of animals from game killed for human consumption in according to the carcases and the following parts originating either from animals that slaughtered in a slaughterhouse and were considered fit for slaughter consumption or bodies and the parts of animals from game killed for human consumption in according to the carcases and the slaughterhouse and were considered fit for slaughter consumption or bodies and the parts of animals from game killed for human consumption in according to the carcases are carcases.									
					human consump	otion i	d parts of animals whic n accordance with Union disease communicable to	legis	slation, but which did		
	(ii) heads of poult				heads of poultry	.,					
				, ,		nalang	ding trimmings and splitt es and the carpus and me				
				(iv)	pig bristles;						
				(v)	feathers;]						
		referred to in Article					oultry and lagomorphs s of Regulation (EC) No ncil(^{2a}), which did not s animals]	853/2	2004 of the European		
			(²)and/or [-	throug slaugl for h	gh blood to hur htered in a slaug	nans of the state	id not show any signs or animals, obtained from ouse after having been collowing an ante-mortem	m an	imals that have been lered fit for slaughter		
							from the production of pr				

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;]

petfood and feedingstuffs of animal origin, or feedingstuffs containing animal $(^2)$ and/or [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;]

blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk originating $(^2)$ and/or [from live animals that did not show signs of any disease communicable through that product to humans or animals;]

 $(^2)$ and/or [aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of diseases communicable to humans or animals;]

Processed petfood other than canned petfood

						essea petroot					
II.	Health inf	orma	tion		II.a.	Certificate refe	erence No	II.b.			
	(2)and/or	[-		al by-products lishments manuf		•	-		s or		
	(²)and/or	[-		ollowing materia sease communica		signs					
			(i)	shells from shellfish with soft tissue or flesh;							
			(ii)	ŕ							
				- hatchery by-products,							
				- eggs,							
				- egg by-pro	oducts,	, including egg	shells,				
			(iii)	day-old chicks k	cilled f	or commercial	reasons;]				
	(2)and/or	[-		imal by-products from aquatic or terrestrial invertebrates other than spe thogenic 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;]								
1.2	(²)and/or	[-	are p	material from animals which have been treated with certain substances whare prohibited by Council Directive 96/22/EC(2b), the import of the mate being permitted in accordance with Article 35(a)(ii) of Regulation (EC) 1069/2009;]							
II.3.	(²)either	Luzac	cubie	cted to a heat trea	tment	of at least 90 °C	T throughout its	cubetance:1			
	()etiner (²)or	[was		aced as regards			_	-	ducts		
		(a)		e case of animates							
		(b)	in the	e case of milk and	d milk	based products,	,				
			(i)	column B of A	nnex I pasteu	to Commission	n Regulation	d countries liste (EU) No 605/20 o produce a neg	$10(^{3})$		
			(ii)	countries listed	in colu to a	mn C of Annex pasteurisation	x I to Regulation	ries or parts of n (EU) No 605/2 ficient to produ	2010,		
			(iii)		Regul able he	ation (EU) No eat treatment wh	605/2010, subm here each treatr	tries listed in col nitted to a sterilisment was sufficie	ation		
			(iv)	C of Annex I to outbreak of foor	Regut-and-r	lation (EU) No mouth disease is oot-and-mouth	605/2010, when the preceding	tries listed in colere there has been 12 months or we carried out in	en an where		
			either								
				 a sterilisat achieved 	ion pro	ocess whereby a	an Fc value equ	al or greater than	1 3 is		
				or							

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

either

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

or

- an acidification process such that the pH has been maintained at less than 6 for at least one hour;
- (c) in the case of gelatine, produced using a process that ensures that unprocessed Category 3 material is subjected to a treatment with acid or alkali, followed by one or more rinses with subsequent adjustment of the pH and subsequent, if necessary repeated, extraction by heat, followed by purification by means of filtration and sterilisation;
- (d) in the case of hydrolysed protein produced using a production process involving appropriate measures to minimise contamination of raw Category 3 material, 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

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

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;
- (l) in the case of dicalcium phosphate produced by a process that
 - (i) ensures that all Category 3 bone-material is finely crushed and degreased with hot water and treated with dilute hydrochloric acid (at a minimum concentration of 4 % and a pH of less than 1,5) over a period of at least two days;
 - (ii) following the procedure referred to in (i), applies a treatment of the obtained phosphoric liquor with lime, resulting in a precipitate of dicalcium phosphate at pH 4 to 7; and
 - (iii) finally, air dries the precipitate of dicalcium phosphate with inlet temperature of 65 °C to 325 °C and end temperature between 30 °C and 65 °C ;
- (m) in the case of tricalcium phosphate produced by a process that ensures
 - (i) that all Category 3 bone-material is finely crushed and degreased in counter-flow with hot water (bone chips less than 14 mm);
 - (ii) continuous cooking with steam at 145 °C during 30 minutes at 4 bar;
 - (iii) separation of the protein broth from the hydroxyapatite (tricalcium phosphate) by centrifugation; and
 - (iv) granulation of the tricalcium phosphate after drying in a fluid bed with air at 200 $^{\circ}\text{C}$;
- (n) in the case of flavouring innards, produced according to a treatment method and parameters, which ensure that the product complies with the microbiological standards referred to in point II.4.]
- (2) or [was subject to a treatment such as drying or fermentation, which has been authorised by the competent authority;]
- (2) or [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. was analysed by a random sampling of at least five samples from each processed batch taken during or after storage at the processing plant and complies with the following standards(4):

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

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;
- II.6. was packed in new packaging, which, if the petfood is not dispatched in ready-to-sell packages on which it is clearly indicated that the content is destined for feeding to pets only, bear labels indicating "NOT FOR HUMAN CONSUMPTION";

(2)[II.7. the petfood described above

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

(2) or [is derived from bovine, ovine or caprine animals and does not contain and is not

II.	Health infor	mation		II.a. Certificate reference No	II.b.			
		derived from	n:					
		(²) either	[bovine, ovine and caprine materials other than those derived from animals born, continuously reared and slaughtered in a country of region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]					
	(²) <i>or</i>		[(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 boving ovine or caprine animals, except from those animals that were born, continuously reared and slaughtered in a country of region classified as posing a negligible BSE risk is accordance with Commission Decision 2007/453/EC(5), is which there has been no indigenous BSE case,					
			ovi stu of cra cav rea pos	•				
Mate								

Notes

Part I:

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

Part II:

- ^(1a) OJ L 300, 14.11.2009, p. 1.
- ^(1b) OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.

Processed petfood other than canned petfood

II.	Healt	h information	II.a. Certificate reference No	o	II.b.				
(^{2a})	OJ L	139, 30.4.2004, p. 55.							
(2b)	OJ L	125, 23.5.1996, p. 3.							
(3)	OJ L 175, 10.7.2010, p. 1.								
(4)	Where	2:							
	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 being considered acceptable if the							
(5)	OJ L	147, 31.5.2001, p. 1.							
(6)	OJ L	172, 30.6.2007, p. 84.							
-	The si	gnature and the stamp must be in	a different colour to that of the	e printir	ng.				
-	Note for the person responsible for the consignment in the European Union: This certificate is only for veterinary purposes and must accompany the consignment until it reaches the border inspection post of entry into the European Union.								
Offi	cial vet	erinarian/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

CO	UNTRY:	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 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 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	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.
	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(1a), and in particular Article 10 of that Regulation, and Commission Regulation (EU) No 142/2011(1b), and in particular Chapter II of Annex XIII and Chapter II of Annex XIV thereto, and certify that the 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;]
 - (²) 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,

COUNTRY Dogchews

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

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;
- 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,
 - (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 feeding of farmed animals, other than fur animals, and the production or manufacturing of pet food.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: Species: select from the following: Aves, Ruminantia, Suidae, Mammalia
 Other Than Ruminantia or Suidae, Pesca, Mollusca, Crustacea, Invertebrates Other Than

COUNTRY Dogchews

II.	Health	information	II.a. Certificate reference	No	II.b.					
	Mollusca And Crustacea.									
Part	Part II:									
(1a)	OJ L 300, 14.11.2009, p. 1.									
(1b)	OJ L 54	1, 26.2.2011, p. 1.								
(2)	Delete a	as appropriate.								
(^{2a})	OJ L 12	25, 23.5.1996, p. 3. ⁽³⁾ Where:								
	n =	number of samples to be test	ed;							
	m =	threshold value for the num number of bacteria in all san		is conside	ered satisfactory if the					
	M =	maximum value for the num number of bacteria in one or			ed unsatisfactory if the					
	c =	number of samples the bactestill being considered accept								
(4)	OJ L 14	47, 31.5.2001, p. 1.								
(5)	OJ L 17	2, 30.6.2007, p. 84.								
_	The sign	nature and the stamp must be i	n a different colour to that o	f the prin	ting.					
_	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.									
Offic	Official veterinarian/Official inspector									
	Name (in capital letters): Qualification and title:									
	Date: Signature:									
	Stamp:									

CHAPTER 3(D)

Health certificate

For raw petfood for direct sale or animal by-products to be fed to fur animals, 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 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	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	I.22. Number of packages
	Ambient ☐ Chilled ☐ I.23. Seal/Container No	Frozen ☐ I.24. Type of packaging
	L25 Commodities certified for:	
		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	
		pproval number of establishments
	Species (Scientific name) Nature of commodity	Manufacturing plant Net weight Batch number

II. H	Iealth information	II.a.	Certificate reference No	II.b.
-------	--------------------	-------	--------------------------	-------

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

- II.1. consist of animal by-products that satisfy the health requirements below;
- II.2. consist of animal by-products:
 - (a) derived from meat which satisfies the relevant animal and public health requirements laid down in:
 - Commission Regulation (EU) No 206/2010(3) 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(4), and provided that the animals from which the meat is derived come from the third countries, territories or parts thereof..... (ISO code in the case of a country, or codes in the case of territories or parts thereof) as listed in that Regulation which has been free from Newcastle disease and avian influenza for the last 12 months;
 - and/or Commission Regulation (EC) No 119/2009(5), and provided that the animals from which the meat is derived come from the third countries, territories or parts thereof...... (ISO code in the case of a country, or codes in the case of territories or parts thereof) as listed in that Regulation which has been free from foot and mouth disease, rinderpest, classical swine fever, African swine fever, swine vesicular disease, Newcastle disease and avian influenza for the preceding 12 months and where no vaccination has taken place during that time (only where relevant for the susceptible species);
 - (b) derived from animals that, at the slaughterhouse, have passed the ante-mortem health inspection during the period of 24 hours before the time of slaughter and have shown no evidence of the diseases referred in the Regulations referred to in point (a) for which the animals are susceptible; and
 - (c) derived from animals that have been handled in the slaughterhouse before and at the time of slaughter or 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(6); or
 - (d) in the case of feed for fur animals, are derived from aquatic animals which satisfy the relevant animal and public health requirements laid down in Commission Decision 2006/766/EC(7), and come from countries or territories thereof (ISO code of the country) as listed in Annex II to that Decision;
- II.3.1. consist only of the following animal by-products:
 - (a) carcases and parts of animals slaughtered or, in the case of game, bodies or parts of animals killed which were deemed fit for human consumption in accordance with Union legislation until irreversibly declared as animal by-products for commercial reasons;
 - (b) parts of slaughtered animals, which are rejected as unfit for human consumption but are not affected by any signs of diseases communicable to humans or animals and derived from carcases that are fit for human consumption in accordance with Union legislation;
- II.3.2.in the case of feed for fur animals in addition to II.3.1. consist also of the following animal by-products:
 - (²)either [- animal by-products from poultry and lagomorphs slaughtered on the farm as referred to in Article 1(3)(d) of Regulation (EC) No 853/2004 of the European Parliament and of the Council(²a), which did not show any signs of

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

				· -						
II.	Health in	forma	ntion	II.a. Certificate reference No	II.b.					
			disease commun	nicable to humans or animals;]						
	(²)and/or	[-	through blood to slaughtered in a	Is which did not show any signs of the ohumans or animals, obtained from slaughterhouse after having been con tumption following an ante-mortem is slation;]	animals that have been asidered fit for slaughter					
	(²)and/or	[-	human consum	ucts arising from the production of otion, including degreased bone, great from milk processing;]						
	(²)and/or	[-	which are no lor or due to proble	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 arises;]						
	(²)and/or	[-	by-products or commercial rea	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;]						
	(²)and/or	[-	blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk originating from live animals that did not show signs of any disease communicable through that product to humans or animals;]							
	(²)and/or	[-	aquatic animals, and parts of such animals, except sea mammals, which dinot show any signs of diseases communicable to humans or animals;]							
	(²)and/or	[-	animal by-products from aquatic animals originating from plants establishments manufacturing products for human consumption;]							
	(²)and/or	[-		aterial originating from animals which nunicable through that material to hun						
			(i) shells from	n shellfish with soft tissue or flesh;						
			(ii) the follow	ing originating from terrestrial animal	s:					
			- hatc	hery by-products,						
			- eggs	3,						
			- egg	by-products, including egg shells,						
			(iii) day-old ch	nicks killed for commercial reasons;]						
	(2)and/or	[-	• •	icts from aquatic or terrestrial inverter imans or animals;]	brates other than species					
	(²)and/or	[-	Lagomorpha, ex (iv) and (v) of	arts thereof of the zoological or scept Category 1 material as referred Regulation (EC) No 1069/2009 and ticle 9(a) to (g) of that Regulation;]	d to in Article 8(a)(iii),					
II.4.	have been obtained and prepared without contact with other material which does not comply with the conditions laid down in the Regulation (EC) No 1069/2009, and it has been handled									
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;									

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.		
II.6.	in the	case of raw p	etfood:					
	(a) 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 and							
	(b)				oling of at least five samples and complies with the following			
		Salmonella:	abser	nce in	25 g: n=5, c=0, m=0, M=0			
		Enterobacter	iaceae: n=5,	c=2,	m=10, M=5000 in 1 gram;			
(²)[II					o be fed to fur animals describ uminant origin and:	ed above contains or is		
		(²)either	negligible BS	E ris	country or region, which is k in accordance with Decision n no indigenous BSE case, and]	n 2007/453/EC, and in		
	(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	-		er ruminants than bovine, ovine			
		(²)or	_	m bo	vine, ovine or caprine animals a			
				from count	ne, ovine and caprine materials animals born, continuously rearry or region classified as posin cordance with Decision 2007/45	red and slaughtered in a g a negligible BSE risk		
			(²)or	[(a)	specified risk material as defin V to Regulation (EC) No 999 Parliament and of the Council(9/2001 of the European		
				(b)	mechanically separated meat of bovine, ovine or caprine animal that were born, continuously real country or region classified BSE risk in accordance with 2007/453/EC(10), in which indigenous BSE case,	als, except from animals eared and slaughtered in as posing a negligible Commission Decision		
(c) animal by-product or derived product obtained from bovine, ovine or caprine animals which have bee killed, after stunning, by laceration of the central nervous tissue by means of an elongated rod-shape instrument introduced into the cranial cavity, or be means of gas injected into the cranial cavity, except for those animals that were born, continuously reare and slaughtered in a country or region classified a posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]]								
NT = 4								
Note								
Part	1:							

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

Part II:

- (1a) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.
- (^{2a}) OJ L 139, 30.4.2004, p. 55.
- (3) OJ L 73, 20.3.2010, p. 1.
- (4) OJ L 226, 23.8.2008, p. 1.
- (5) OJ L 39, 10.2.2009, p. 12.
- (6) OJ L 303, 18.11.2009, p. 1.
- (⁷) OJ L 320, 18.11.2006, p. 53.
- (8) Where:
 - n = number of samples to be tested;
 - m = threshold value for the number of bacteria; 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.
- (9) OJ L 147, 31.5.2001, p. 1.
- (10) OJ L 172, 30.6.2007, p. 84.

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.		
-	The signature and the stamp must be in a different colour to that of the printing.						
-	- Note for the person responsible for the consignment in the European Union: This certificate is only for veterinary purposes and must accompany the consignment until it reaches the border inspection post of the European Union.						
Offic	cial veterinarian/Official inspector						
	Name (in capital letters): Qualification and title:						
	Date: Signature:						
	Stamp:						

CHAPTER 3(E)

Health certificate

For flavouring innards for use in the manufacture of petfood, intended for dispatch to or for transit through $(^2)$ the European 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						
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	I.9. Country of 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	Custom warehouse Name Approval number Address Postcode						
	Address 1.13. 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 Ambient □ Chilled □	I.22. Number of packages						
	1.23. Seal/Container No	I.24. Type of packaging						
	I.25. Commodities certified for:	-						
	Petfood □ Tech	chnical 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) Nature of commodity	Approval number of establishments Manufacturing plant Net weight Batch number						

Health information

II.

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 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 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;] carcases and the following parts originating either from animals that have (2)and/or 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 $(^2)$ 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 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;]
- (²) and/or [- blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk originating from live animals that did not show signs of any disease communicable through that product to humans or animals;]
- (²) and/or [- aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of diseases communicable to humans or animals;]

Flavouring innards for use in the manufacture of petfood

				<u> </u>			
II.	Health in	nformati	on	II.a. Certificate reference No	II.b.		
	(²)and/or			cts from aquatic animals origina nufacturing products for human con			
	(2)and/or			terial originating from animals white municable through that material to			
		(i) shells from s	shellfish with soft tissue or flesh;			
		(ii) the followin	g originating from terrestrial animal	s:		
			- hatcher	y by-products,			
			- eggs,				
			- egg by-	products, including egg shells;			
		(iii) day-old chic	ks killed for commercial reasons;]			
	(2)and/or			ets from aquatic or terrestrial invecto humans or animals;]	ertebrates other than		
	(²)and/or	I (Lagomorpha, exceiv) and (v) of Re	est s thereof of the zoological order the category 1 material as referred gulation (EC) No 1069/2009 and Cole 9(a) to (g) of that Regulation;]	to in Article 8(a)(iii),		
	(²)and/or	v r	vhich are prohibi	imals which have been treated witted by Council Directive 96/22/EComitted in accordance with Article 3 [19;]	(2a), the import of the		
II.3.				ng in accordance with Chapter II rder to kill pathogenic agents;	I of Annex XIII to		
II.4.				g of at least five samples from each sing plant and complies with the fol	_		
	Salmone	ella:	absence in 2	5g: $n = 5$, $c = 0$, $m = 0$, $M = 0$,			
	Enteroba	acteriacea	ae: $n = 5, c = 2,$	m = 10, M = 300 in 1 gramme;			
II.5.	the end p	roduct w	as:				
	(²)either	[packe	d in new or sterili	sed bags,]			
	(²) <i>or</i>	thorou		n containers or other means of nd disinfected with a disinfectar fore use,]			
	and whic	h bear lal	bels indicating 'N	OT FOR HUMAN CONSUMPTIO	N';		
II.6.	the end p	roduct w	as stored in enclo	sed storage;			
II.7.	the produ		indergone all pre-	cautions to avoid contamination w	ith pathogenic agents		
(²)[II.8.	the flavo	uring inn	ards products des	cribed above			
	(²)either	is deriv	ed from other run	ninants than bovine, ovine or caprine	e animals.]]		
	(2) or [is derived from bovine, ovine or caprine animals and does not contain and is no derived from:						
		(²) eithe	animals bor region class	ine and caprine materials other than, continuously reared and slaught ified as posing a negligible BSE ris 07/453/EC.]]	ered in a country or		
		(²) <i>or</i>	Regi	ified risk material as defined in polalation (EC) No 999/2001 of the of the Council(4);			

Flavouring innards for use in the manufacture of petfood

II.	Health information		II.a. Certificate reference No II.b.
		ovine born regio acco	hanically separated meat obtained from bones of bovine, he or caprine animals, except from those animals that were a, continuously reared and slaughtered in a country or on classified as posing a negligible BSE risk in ordance with Commission Decision 2007/453/EC(5), in the there has been no indigenous BSE case,
		ovine stunr of ar crani cavit reare posir	nal by-product or derived product obtained from bovine, he or caprine animals which have been killed, after ning, by laceration of the central nervous tissue by means an elongated rod-shaped instrument introduced into the hial cavity, or by means of gas injected into the cranial ty, except for those animals that were born, continuously ed and slaughtered in a country or region classified as ng a negligible BSE risk in accordance with Decision 7/453/EC.]]]

Notes

Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is required to be filled in only if it is a certificate for a commodity to be transited through the European Union; it may be filled in if the certificate is for a commodity to be imported into the European Union.
- Box reference I.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 feeding of farmed animals, other than fur animals, and the production or manufacturing of pet food.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28:
 - species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or Suidae, Pesca, Mollusca, Crustacea, Invertebrates other than Mollusca and crustacea
 - define the innard product.

Part II:

- (1a) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.
- (^{2a}) OJ L 125, 23.5.1996, p. 3.
- (3) Where:
 - n = number of samples to be tested;
 - m = threshold value for the number of bacteria; the result is considered satisfactory if the

Flavouring innards for use in the manufacture of petfood

II.	Не	ealth information	II.a.	Certificate reference	e No	II.b.		
		number of bacteria in all sample	s does	not exceed m;				
	M =	maximum value for the number number of bacteria in one or mo				unsatisfactory if the		
	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 1	147, 31.5.2001, p. 1.						
(5)	OJ L 172, 30.6.2007, p. 84.							
_	The sign	nature and the stamp must be in a	differ	ent colour to that of	the printir	ıg.		
_	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.							
Offi	cial veter	inarian/Official inspector						
	Name (i	in capital letters):	Qualification and title:			ion 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	UNTRY:	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 Address	Custom warehouse Name Approval number Address						
Ь	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 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:							
	Manufacture of petfood \Box Furth	rther process 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 n	umber of establishments						
		nufacturing plant Number of packages Net weight Batch number						

	or periodu									
	II.	Health inform	matio	n	II.a. Certificate reference No	II.b.				
	I, the undersigned official veterinarian, declare that I have read and understood Re (EC) No 1069/2009 of the European Parliament and of the Council(la) and Con Regulation (EU) No 142/2011(lb), and in particular Chapter II of Annex XIV the certify that the animal by-products described above:									
n n	II.1.1.	consist of a	ınimal	by-pro	ducts that satisfy the animal health	requirements below;				
atic	II.1.2.	have been o	obtain	ed in th	e territory of:	(1c) from animals:				
Part II: Certification		(²)either	[(a)		we remained in this territory since nonths preceding the date of slaug	-				
ŭ		$(^2)or$	[(b)	killed	in the wild in this territory(1d);]					
ırt II:		$(^2)or$	[(c)		d from rodents, lagomorphs, aq c invertebrates;]	quatic animals or terrestrial or				
Pa	II.1.3.	have been o	have been obtained from or produced by animals:							
		(²)either	[(a)	coming	g from holdings:					
				s v i	where, for the following diseas susceptible, there has been no cast vesicular disease, Newcastle dise influenza during the period of the classical or African swine fever du 40 days; nor in the holdings situated from radius, during the period of the	se/outbreak of rinderpest, swine ase or highly pathogenic avian the preceding 30 days, nor of aring the period of the preceding ted in their vicinity within a 10				
				(oreak of foot-and-mouth disease ag 60 days, nor in the holdings 25 km radius, during the period					
			(b)	which:						
				(i) v	were not killed to eradicate any ep	izootic disease;				
				(nave remained in their holdings of days before the date of departure a directly to the slaughterhouse wanimals which did not comply with	and which have been transported vithout any contact with other				
				i	at the slaughterhouse, have particular the period of 2 slaughter and have shown no evidabove for which the animals are su	24 hours preceding the time of dence of the diseases referred to				
				s U t	nave been handled in the slaughter slaughter or killed in accordance Union legislation and have met rechose laid down in Chapters II and No 1099/2009(4)]	with the relevant provisions of quirements at least equivalent to				
		$(^{2})or$	[(a)	capture	ed and killed in the wild in an area	ı:				
				a s c	n which within a 25 km radius the any of the following diseases susceptible: foot-and-mouth disease or highly pathogenic avian influ- preceding 30 days, nor of classica the period of the preceding 40 days	for which the animals are se, rinderpest, Newcastle disease tenza during the period of the all or African swine fever during				
				t	situated at a distance of at least 20 the territory of a country not author Union of poultry material during	orised for export to the European				

II.	Health information	II.a. Certificate reference No II.b.							
		porcine material during the preceding 40 days; and							
(b) which after killing were transported within a period of 12 hours following the killing for chilling either to a collection centre and immediately afterwards to a game handling establishment, or directly to a game handling establishment;]									
II.1.4.	have been obtained in an establishment around which, within a radius of 10 km, there has been no case/outbreak of the diseases referred to in point II.1.3 for which the animals are susceptible during the period of the preceding 30 days or, in the event of a case of disease, the preparation of raw material for exportation to the European Union has been authorised only after the removal of all meat, and the total cleaning and disinfection of the establishment under the control of an official veterinarian;								
II.1.5.	comply with the	ed and prepared without contact with any other material that does not conditions required above, and it has been handled so as to avoid th pathogenic agents;							
II.1.6.	have been packed in new packaging preventing any leakage and in officially sealed containers bearing the label indicating 'RAW MATERIAL ONLY FOR THE MANUFACTURE OF PET FOOD' and the name and address of the establishment of destination in the European Union or during the transit through the European Union;								
II.1.7.	•	e following animal by-products:							
	(2)either [- carcases and parts of animals slaughtered or, in the case of game, bod or parts of animals killed which were deemed fit for hum consumption in accordance with Union legislation until irreversit declared as animal by-products 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;							
	2	(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;]							
	(2) and/or [- products of animal origin, or foodstuffs containing products of anim origin, which are no longer intended for human consumption commercial reasons or due to problems of manufacturing or packag defects or other defects from which no risk to public or animal hearise;]								
	(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							

					or petroou				
II.	Health i	nfor	matic	n	II.a. Certificate reference No	II.b.			
				establ	ishments manufacturing products f	for human consumption;]			
	(²)and	l/or	[-		llowing material originating from a of disease communicable throughs:				
				(i)	shells from shellfish with soft tissu	e or flesh;			
				(ii)	the following originating from terr	estrial animals:			
					- hatchery by-products,				
					- eggs,				
					 egg by-products, including egg 	g shells;			
				(iii)	day-old chicks killed for commercial	ial reasons;]			
	(²)and	l/or	[-		I by-products from aquatic or terms pathogenic to humans or animals				
	(²)and	l/or	[-	Lagor 8(a)(ii	Is and parts thereof of the zool morpha, except Category 1 mate (ii), (iv) and (v) of Regulation (EC) (ial as referred to in Article 9(a) to (erial as referred to in Article) No 1069/2009 and Category 2			
	(²)and	l/or	[-	which the m	ial from animals which have been are prohibited by Council Direct laterial being permitted in accordation (EC) No 1069/2009;]	tive 96/22/EC(4a), the import of			
II.1.8.	have been deep-frozen at the plant of origin or have been preserved in accordance we European Union legislation in such a way that they will not spoil between dispatch a delivery to the plant of destination in the European Union or during the transit through European Union;								
II.1.9.	substa	nces	proh	ibited b	terial derived from animals which have been treated with certain by Directive 96/22/EC for the manufacture of petfood, the import rdance with Article 35(a)(ii) of Regulation (EC) No 1069/2009:				
		(a) it has been marked in the third country before entry into the territory of the European Union by a cross of liquefied charcoal or activated carbon on each outer side of each frozen block, or, when the raw material is transported in pallets which are not divided into separate consignments during transport to the petfood plant of destination in the European Union or during the transit through the European Union on each outer side of each pallet, in a way that the marking covers at least 70 % of the diagonal length of the frozen block and be of at least 10 cm width;							
		third with	coun lique	try before	erial which is not frozen, the raw is ore entry into the territory of the arcoal or by applying charcoal provisible on the material; and	European Union by spraying it			
		refer	red to	above	by-products are made up of raw material, and other non-treated raw material, d to in point (a) and (b) above.				
(²)(⁵)[II.2.	Specif	fic re	quire	ments					
(²)(⁶)[II.2.1	territo	ry re	eferre	d to in	s consignment come from anima point (II.1.2), where vaccination gregularly carried out and officiall	programmes against foot-and-			
(²)(⁷)[II.2.2	trimm	ed o	ffal of	f domes	s consignment consist only of an tic ruminants, which have maturat period of at least three hours, or in	ed at an ambient temperature of			

Animal by-products for the manufacture of petfood

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

bovine animals and deboned meat of domestic animals, for a period of at least 24 hours.]]

(²)[II.3. the animal by-products for the manufacture of petfood 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, and]]

(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(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(9), 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.

Animal by-products for the manufacture of petfood

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

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

Part II:

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

II.	Health information	II.a. Certificate reference No	II.b.					
	for veterinary purposes and must accompany the consignment until it reaches the border inspection post of the European Union.							
Offi	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

I.1.	Consignor										
	Name				I.2.	Certificate reference	e No	I.2	2.a.		
	Address				I.3. Central competent authority						
	Tel.				I.4.						
I.5.	Consignee Name Address Postcode Tel.				I.6. Person responsible for the load in EU Name Address Postcode Tel.						
		ISO code	I.8. Region of origin	Code	I.9.	Country of destination	ISO code	I.10.	Region of destination	Code	
I.11.	Place of origin	n	-		I.12	. Place of destination					
	Name Address Name Address		Approval number	r		Name Address					
Address											
I.13.	Place of loadi	ing			I.14. Date of departure						
I.15. Means of transport						I.16. Entry BIP in EU					
	Road vehicle Identification	Oth			I.17.						
I.18.	Description o	f commodity					I.19. Commo	dity cod	de (HS code)		
						L		I.20.	Quantity		
I.21.	Temperature Ambient	of product	Chilled □		Frozen			I.22.	I.22. Number of packages		
I.23.	Seal/Containe	er No							I.24. Type of packaging		
I.25.				Monu	.fo.atuu	ra of matfood			Tachmical		
I.26. For transit through EU to third country Third country ISO code						I.27. For import or admission into EU					
I.28.	Identification	of the commo	dities								
Species (Scientific name) Nature of commodity									Batch number		
	I.7. (c I.11. I.13. I.15. I.18. I.21. I.23. I.25.	I.15. Consignee Name Address Postcode Tel. I.7. Country of origin I.11. Place of origin Name Address Name Address Name Address I.13. Place of loads I.15. Means of trar Aeroplane Road vehicle Identification Documentatic I.18. Description of I.21. Temperature Ambient I.23. Seal/Contained I.25. Commodities Animal feedin I.26. For transit the Third country I.28. Identification	Address Tel. 1.5. Consignee Name Address Postcode Tel. 1.7. Country of ISO code origin I.11. Place of origin Name Address Name Address Name Address Name Address I.13. Place of loading I.15. Means of transport Aeroplane	Address Tel. 1.5. Consignee Name Address Postcode Tel. 1.7. Country of ISO code I.8. Region of origin origin Name Address I.13. Place of loading 1.15. Means of transport Aeroplane Ship Railway wagon Road vehicle Other Identification Documentation references 1.18. Description of commodity 1.21. Temperature of product Ambient Chilled Chi	Address Tel. 1.5. Consignee Name Address Postcode Tel. 1.7. Country of ISO code I.8. Region of origin Code origin Name Approval number Address I.13. Place of loading 1.15. Means of transport Railway wagon Road vehicle Other Identification Documentation references 1.18. Description of commodity 1.21. Temperature of product Ambient Chilled Manual I.25. Commodities certified for: Animal feedingstuff Manual I.26. For transit through EU to third country Third country ISO code	Address Tel. I.5. Consignee Name Address Postcode Tel. I.7. Country of ISO code I.8. Region of origin Code origin Name Address Name Address Name Address Name Address Name Address Name Adproval number Address Name Address Name Approval number Address Name Actual Contains I.14 I.15. Means of transport Aeroplane Ship Railway wagon I.18. Description of commodity I.17 I.18. Description of commodity I.21. Temperature of product Ambient Chilled I.23. Seal/Container No I.25. Commodities certified for: Animal feedingstuff Manufactur I.26. For transit through EU to third country Third country ISO code I.28. Identification of the commodities	Address Tel. 1.4. Local competent au 1.5. Consignee Name Address Postcode Tel. 1.7. Country of ISO code I.8. Region of origin Code origin 1.8. Page of destination 1.19. Place of origin Name Address Name Approval number Address Name Actoplane Other Othe	Address Tel. 1.3. Central competent authority 1.4. Local competent authority 1.5. Consignee Name Address Postcode Tel. 1.7. Country of ISO code I.8. Region of origin Code origin 1.8. Postcode Tel. 1.9 Country of ISO code origin Name Address Name Address Name Approval number Address Name Address Name Approval number Address Name Address Name Approval number Name Address Name Address Name Address Name Address Name Address Name Approval number Name Name Address	Address Tel. 1.3. Central competent authority 1.4. Local competent authority 1.5. Consignee Name Address Postcode Tel. 1.7. Country of ISO code I.8. Region of origin Code origin 1.8. Region of origin Code of Tel. 1.9. Country of destination 1.10. Place of origin 1.11. Place of origin 1.12. Place of destination 1.13. Place of destination 1.14. Date of departure 1.15. Means of transport Address 1.16. Person responsible for the load in EU Name Address Name Approval number Postcode 1.15. Means of transport Acroplane Ship Railway wagon Documentation references 1.18. Description of commodity 1.19. Commodity country 1.10. Interperature of product Ambient Prozen 122. 1.20. Scall/Container No 122. 1.21. Temperature of product Ambient Manufacture of petfood 122. 1.22. Commodities certified for: Animal feedingstuff Manufacture of petfood 127. For import or admission into EU 1.24. Identification of the commodities Approval number of establishments	Address Tel. 1.3. Central competent authority 1.4. Local competent authority 1.5. Consignee Name Address Postcode Tel. 1.6. Person responsible for the load in EU Name Address Postcode Tel. 1.7. Country of origin ISO code I.8. Region of origin Code I.9. Country of destination 1.11. Place of origin Name Address Name Approval number of establishments	

		•	
			Į.

	II.	Health info	rmation	II.a. Certificate reference No	II.b.		
		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council(^{1a}) and Commission Regulation (EU) No 142/2011(^{1b}) and certify that the blood products described above:					
	II.1.	consist of blood products that satisfy the health requirements below;					
ion	II.2.	consist excl	usively of blood prod	ucts not intended for human consump	tion;		
ificat	II.3. have been prepared and stored in a plant, approved and supervised by the comauthority in accordance with Article 24 of Regulation (EC) No 1069/2009;						
ert	II.4.	have been p	repared exclusively v	with the following animal by-products:			
Part II: Certification		(²)either	accordance with	tered animals, which is fit for human consumption in Union legislation, but which is not intended for human ommercial reasons;]			
Pa		(²)and/or	[blood of slaughtered animals, which has been rejected as unfit for human consumption in accordance with Union legislation, but which did not show any signs of diseases communicable to humans or animals, which has been derived from carcases that have been slaughtered in a slaughterhouse and which were considered fit for human consumption following an ante-mortem inspection in accordance with Union legislation;]				
	II.5.	in order to inactivate pathogenic agents, have been submitted					
		(2) either [to processing in accordance with processing method(3) as set of Chapter III of Annex IV to Regulation (EU) No 142/2011;]					
	(2) or [to a method and parameters which ensure that the product complies microbiological standards set out in Chapter I of Annex X to Regulati No 142/2011;]						
(2)or [in the case of blood products, including spray dried blood and bloof porcine origin intended for the feeding of porcine animals treatment at a temperature of at least 80°C throughout the substated dry blood and blood plasma does not contain more than 8% w/w in 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	[packed in new or	sterilised bags;]			
	(2)or [transported in bulk in containers or other means of transport that thoroughly cleaned and disinfected with a disinfectant approved competent authority before use,]						
		and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION';					
	II.7.	the end product was stored in enclosed storage;					
	II.8.	the product has undergone all precautions to avoid contamination with pathogenic agents after treatment;					
		(²)and	of porcine origin is	od products, including spray dried ble ntended for the feeding of porcine and conditions under room temperature for	imals, has been stored		
	II.9.	have been examined prior to dispatch under the responsibility of the competent authority by taking a random sample during or on removal from storage which was found to comply with the following standards(⁴):					
		Salmonella: absence in 25g: $n = 5$, $c = 0$, $m = 0$, $M = 0$,					
		Enterobacteriaceae: $n = 5$, $c = 2$, $m = 10$, $M = 300$ in 1 gram;					

II.	Health in	ıformation		II.a. Certificate reference No	II.b.	
(²)[II.10. the blood products described above						
() [(²) either [is derived from other ruminants than bovine, ovine or caprine animals.]]					
	(²)or	-	from bovir	ne, ovine or caprine animals and does		
(²) either [bovine, ovine ar animals born, co region classified		ovine and caprine materials other that born, continuously reared and slaugh lassified as posing a negligible BSE ri 2007/453/EC.]]	tered in a country or			
		(²)or	R	pecified risk material as defined in pecegulation (EC) No 999/2001 of the nd of the Council(4);		
			o b re	nechanically separated meat obtained to vine or caprine animals, except from the orn, continuously reared and slaught egion classified as posing a negl occordance with Commission Decision which there has been no indigenous BSI	nose animals that were tered in a country or igible BSE risk in n 2007/453/EC(5), in	
			o si o cr cr re	nimal by-product or derived product of vine or caprine animals which have tunning, by laceration of the central need for an elongated rod-shaped instrument ranial cavity, or by means of gas injudy, except for those animals that we eared and slaughtered in a country of osing a negligible BSE risk in accos 007/453/EC.]]]	we been killed, after rvous tissue by means t introduced into the ected into the cranial ere born, continuously r region classified as	
II.11. the blood products or products described above:		lescribed above:				
	(²)either	-	[does not contain milk or milk products of ovine or caprine animal origin or is not intended for feed for farmed animals, other than fur animals.]			
	(2) <i>or</i>			r milk products of ovine or caprine for farmed animals, other than fur anim		
\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \		ed from ovine and caprine animals wasly since birth in a country where the ed:	-			
			(i) c	lassical scrapie is compulsorily notifial	ole;	
				n awareness, surveillance and monitor or classical scrapie;	ing system is in place	
			a	fficial restrictions apply to holdings nimals in the case of a suspicion of TS f classical scrapie;		
				vine and caprine animals affected wit illed and destroyed;	h classical scrapie are	
			m C ru	ne feeding to ovine and caprine animal or greaves, as defined in the Terrode of the World Organisation for Anuminant origin has been banned and ene whole country for a period of at least	estrial Animal Health simal Health (OIE), of effectively enforced in	

Health informa	tion	II.a. Certificate reference No	II.b.	
(t	o) originate fr	om holdings where no official rest	crictions are imposed due	
(0	diagnosed	during the period of at least the p	receding seven years or,	
	an Al Al	d destroyed or slaughtered, except RR/ARR genotype, breeding ewe RR allele and no VRQ allele a	for breeding rams of the es carrying at least one	
	be for co TS the me Re wl	ten killed and destroyed, and the hor a period of at least two yes infirmation of the last classical sets. Et monitoring, including testing we presence of TSE in accordance thods set out in point 3.2 of Chegulation (EC) No 999/2001, of all hich are over the age of 18 months	olding has been subjected ears since the date of trapie case to intensified with negative results for the with the laboratory papter C of Annex X to of the following animals	
	-	animals which have been s consumption; and	slaughtered for human	
	_	animals which have died or been which were not killed in the eradication campaign.]]		
the blood products described above contain or are derived from animal-by products of non-ruminant origin, and are, according to the statement of the Consignor referred to in Box I.1,				
		the production of feed for farmed	animals, other than fur	
th in	[intended for the production of feed for non-ruminant farmed animals, other than fur animals, and the Consignor has undertaken to ensure that the border inspection post of entry will be provided with the results of the analyses carried out in accordance with the methods set out in Annex VI to Commission Regulation (EC) No 152/2009(8).]			
	the blood produruminant origin, (2)either [ran (2)(7)or [ith	(b) originate from to a suspicific to a suspicific originate from the diagnosed of following to the following to the suspicific originate from the suspicific original following to the suspicific originate from the suspicific originate from the suspicific original following to the suspicific original follow	to a suspicion of TSE; (c) originate from holdings where no case of claignosed during the period of at least the p following the confirmation of a case of classic (2)either [all ovine and caprine animals on the and destroyed or slaughtered, except ARR/ARR genotype, breeding ewe ARR allele and no VRQ allele a carrying at least one ARR allele;] (2)or [all animals in which classical scrabeen killed and destroyed, and the hofor a period of at least two years confirmation of the last classical scrabeen killed and destroyed, and the hofor a period of at least two years confirmation of the last classical scrabeen killed and destroyed, and the hofor a period of at least two years confirmation of the last classical scrabeen killed and destroyed, and the hofor a period of at least two years confirmation of the last classical scrabeen killed and destroyed, and the hofor a period of at least two years confirmation of the last classical scrabeen killed and destroyed, and the hofor a period of at least two years confirmation of the last classical scrabeen killed and destroyed, and the hofor a period of at least two years confirmation of the last classical scrabeen killed and destroyed, and the hofor a period of at least two years confirmation of the last classical scrabeen killed and destroyed, and the hofor a period of at least two years confirmation of the last classical scrabeen killed and destroyed, and the hofor a period of at least two years confirmation of the last classical scrabeen killed and destroyed, and the hofor animals which have dead or been which are over the age of 18 months the ARR/ARR genotype: — animals which have died or been which were not killed in the eradication campaign.]] the blood products described above contain or are derived from an ruminant origin, and are, according to the statement of the Consigno (2)either [not intended for the production of feed for non-rumina than fur animals, and the Consignor has undertaken	

Notes

Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is required to be filled in only if it is a certificate for a commodity that is to be transited through the European Union; it may be filled in if the certificate is for a commodity that is to be imported into the European Union.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for a transit commodity. Products in transit may only be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in the case of unloading and reloading in the European Union.

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

- Box reference I.19: use the appropriate HS code: 05.11.91, 05.11.99, 35.02 or 35.04.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.
- Box reference I.25: technical use: any use other than feeding of farmed animals, other than fur animals, and the production or manufacturing of pet food.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
 - Box reference I.28: Species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or Suidae, Pesca, Reptilia.

Part II:

- (1a) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.
- (3) Insert method 1 to 5 or method 7 as applicable.
- (4) Where:
 - n = number of samples to be tested;
 - m = threshold value for the number of bacteria; the result is considered satisfactory if the number of bacteria in 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.
- (5) OJ L 147, 31.5.2001, p. 1.
- (6) OJ L 172, 30.6.2007, p. 84.
- (7) The person responsible for the load referred to in Box I.6 must ensure that, if the blood products or the 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 a border inspection post of the European Union.
- (8) OJ L 54, 26.2.2009, p. 1.
- The signature and the stamp must be in a different colour to that of the printing.
- Note for the person responsible for the consignment in the European Union: this certificate is only
 for veterinary purposes and must accompany the consignment until it reaches the border
 inspection post of the point of entry into the European Union.

Official veterinarian/Official inspector				
Name (in capital letters):	Qualification 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 ISO code I.8. Region of origin Code origin	I.9. Country of ISO code I.10. Region of destination Code			
ails c	I.11. Place of origin	I.12. Place of destination			
Part I : Det	Name Approval number Address Name Approval number Address	Custom warehouse Name Approval number Address			
7	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 □	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 ☐	1.27. For import or admission into EU			
	Third country ISO code	1.27. For import of dames on the 2e			
	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

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(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; they have been prepared and stored in a plant supervised by the competent authority or II.3. in the establishment of collection, 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 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;] blood and blood products originating from live animals that did not (2)and/or show signs of any disease communicable through that product to humans or animals;] (2)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^(2a) or Article 2(b) of Council Directive 96/23/EC^(2b); (2)and/or animal by-products containing residues of other substances and environmental 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;] II.4. the blood, that such products were manufactured from, was 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 $(^{2})[II.5.$ in the case of blood products obtained from animals belonging to the taxa Artiodactyla, Perissodactyla and Proboscidea, including crossbreds between species of those taxa, the blood was collected in a country or region where no case of rinderpest, peste des petits ruminants and Rift Valley fever has been recorded for a period of at least the preceding 12 months and in which vaccination has not been carried out against those diseases for a

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

II.	Health inform	ation	II.a. Certificate reference No II.	.b.
	period of a	t least the preceding	12 months, and;	
(²)either	[in third c	of a country, or cocase of foot-and-1 the preceding 12 i	or parts thereof (insert ISO country odes (3) in the case of territories or parts mouth disease has been recorded for a months and in which vaccination has no e for a period of at least the preceding 12	thereof) where no period of at least of been carried out
	(²) <i>or</i>	in the case of a como case of foot-and the preceding 12 foot-and-mouth d	territories or parts thereof (insert ountry or codes(3) for territories or part demouth disease has been recorded for a months and in which vaccination profisease are being officially carried out an animals for a period of at least the contract of the series of the contract of the	rts thereof) where a period of at least ogrammes against and controlled in
(²)[II.5.1.	in the case which:	of animals other tha	n Suidae and Tayassuidae, in third coun	ntries or regions in
	(²)either	of seropositive an preceding 12 mon	ular stomatitis and bluetongue(2) (including lands) has been recorded for a perion of the and in which vaccination has not asses for a period of at least the preceding	od of at least the t been carried out
	$(^{2})or$	[vesicular stomatic	tis and bluetongue(2) seropositive anima	als are present ⁽⁴⁾ ;]]
(²)[II.5.2.	swine vesion for a period	cular disease, classic d of at least the prec se diseases for a per	assuidae, in third countries or regions in the sal swine fever and African swine fever leding 12 months and vaccination has not riod of at least the preceding 12 months	has been recorded of been carried out
	(²)either	animals) has been and in which vacc	cular stomatitis (including the presence recorded for a period of at least the precination has not been carried out against the preceding 12 months;]]	eceding 12 months
	$(^{2})or$	[vesicular stomati	tis seropositive animals are present(4);]]]]
(²)[II.6.		•	derived from poultry or other avian spe territory of the country or region with	
			veastle disease and highly pathogenic a al Health Code of the OIE,	avian influenza as
		a period of at least an influenza,	the preceding 12 months has not carrie	ed out vaccination
	against Ne	ewcastle disease wi	ch the products are derived, have not th vaccines prepared from a Newcast enicity than lentogenic virus strains;]	
II.7.	the produc	ts were:		
	$(^2)$ either	[packed in new or	sterilised bags or bottles,]	
	(²)or		alk in containers or other means of traded and disinfected with a disinfectant ty before use,]	
the outer packaging or containers bear labels indic ANIMAL CONSUMPTION';			iners bear labels indicating 'NOT FO	OR HUMAN OR
II.8.	the produc	ts were stored in end	closed storage;	

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

II.	Health informa	ation		II.a. Certificate reference No	II.b.
II.9.	all precaut agents duri			o avoid contamination of the produ	ucts with pathogenic
(²)[II.10.	the untreate	ed blood	l products de	escribed above	
	(²)either	[is der	rived from o	ther ruminants than bovine, ovine or	caprine animals.]]
	$(^2)or$		rived from b derived from	ovine, ovine or caprine animals and m:	does not contain and
	(²) either [bovine, ov animals bor region class		ine and caprine materials other that rn, continuously reared and slaught sified as posing a negligible BSE ris 007/453/EC.]]	ered in a country or	
(²)or [(a) spec Regu		ecified risk material as defined in point 1 of Annex V to gulation (EC) No 999/2001 of the European Parliament of the Council(4);			
(b) mecl ovin- born regio acco		hanically separated meat obtained find or caprine animals, except from the continuously reared and slaughted on classified as posing a neglion ordance with Commission Decision of there has been no indigenous BSE	ose animals that were ered in a country or gible BSE risk in a 2007/453/EC(5), in		
			(c) animal by-product or derived product obtained from borovine or caprine animals which have been killed, stunning, by laceration of the central nervous tissue by m of an elongated rod-shaped instrument introduced into cranial cavity, or by means of gas injected into the cracavity, except for those animals that were born, continuor reared and slaughtered in a country or region classifie posing a negligible BSE risk in accordance with Deci 2007/453/EC.]]]		e been killed, after ryous tissue by means introduced into the ected into the cranial re born, continuously region classified as
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)

COUNTRY

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

II.	Health information	II.a. Certificate reference	No	II.b.		
	must be included.					
-	Box reference I.25: technical use: any u animals, and the production or manufactu		rmed an	imals, other than fur		
-	Box reference I.26 and I.27: fill in accord	ling to whether it is a transit	or an in	port certificate.		
-	Box reference I.28 Species: select from other than Ruminantia or Suidae, Pesca,		minantia	, Suidae, Mammalia		
Part	: П:					
(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)	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).					
(4)	In this case following the veterinary checks provided for in Directive 97/78/EC (OJ L 24, 30.1.1998, p. 9), and in accordance with the conditions laid down in Article 8(4) of that Directive, the products must be transported directly to the establishment at the place of destination.					
(5)	Code of the territory as it appears in I 798/2008 (OJ L 226, 23.8.2008, p. 1).	Part 1 of Annex I to Com	mission	Regulation (EC) No		
(6)	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 th	ne printir	ıg.		
-	Note for the person responsible for the co- for veterinary purposes and must acc- inspection post of the point of entry into	ompany the consignment				
Offic	Official veterinarian/Official inspector					
	Name (in capital letters):	Q	ualificat	ion and title:		
	Date:	S	ignature	:		
	Stamp:					

CHAPTER 4(D)

Health certificate

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

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.			
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 Code destination			
ails c	I.11. Place of origin	I.12. Place of destination			
Part I : Det	Name Approval number Address Name Approval number Address	Custom warehouse Name Approval number 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 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	·			
		number of establishments nufacturing plant Batch number			

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

II. 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 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 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, 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;] blood of slaughtered animals, which is rejected as unfit for human $(^2)$ and/or [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;] blood and blood products originating from live animals that did not show $(^2)$ and/or [clinical signs of any disease communicable through these products to humans or animals;] (2)and/or [blood and blood products derived from the production of products intended for human consumption;] animal by-products which have been derived from animals which have $(^2)$ and/or [been submitted to illegal treatment as defined in Article 1(2)(d) of Council Directive 96/22/EC^(2a) or Article 2(b) of Council Directive 96/23/EC^(2b);] animal by-products containing residues of other substances and $(^2)$ and/or [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 (2)either an effectiveness check;] (2)and/or [irradiation at 25 kGy by gamma rays, followed by an effectiveness check;]

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

II.	Health info	ormation		II.a. Certificate reference No	II.b.	
	(²)and/or [change in pH to pH :		in pH to pH	for two hours, followed by an effectiveness check;]		
	(2)and/or [heat treatment of at least 80 °C throughout their substance, followed by effectiveness check.]]					
(²)[II.6.	species, the absence of swine vesion	pathogens cular diseas	have undergof the followingse, classical s	ived from Suidae, Tayassuidae, por gone one of the following treatme ing diseases: foot-and-mouth disease swine fever, African swine fever, N as appropriate to the species:	nts guaranteeing the , vesicular stomatitis,	
	(²)either		eatment at a t	emperature of 65 °C for at least threak;]	e hours, followed by	
	(2)and/or	[irradiat	ion at 25 kGy	y by gamma rays, followed by an effe	ectiveness check;]	
	(²)and/or	for poul	try and other	least 80 °C for Suidae/Tayassuidae avian species(²) throughout the substiveness check]].		
(²)[II.7.				ed from species other than those liste following treatment (please specify)		
II.8.	The produc	cts were:				
	$(^2)$ either	[packed in	new or sterili	ised bags or bottles,]		
		[transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectant approved by the competent authority before use;] and				
		the outer packaging or containers bear labels indicating 'NOT FOR HUMAN OR ANIMAL CONSUMPTION';				
II.9.	the product	ts were stor	ed in enclose	d storage;		
II.10.	all precauti		iken to avoid	the contamination of the products w	ith pathogenic agents	
(²)[II.11.	The treated	d blood prod	ducts describe	ed above		
	(²)either [i	is derived f	rom other run	minants than bovine, ovine or caprine	animals.]]	
		is derived f lerived fron		ovine or caprine animals and does n	ot contain and is not	
	(2	(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.]]			ered in a country or	
	(2) or [(a) specified risk material as defined in point 1 of Annex V Regulation (EC) No 999/2001 of the European Parliame 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,			ose animals that were bred in a country or gible BSE risk in 2007/453/EC(5), in		
			ovin	nal by-product or derived product of e or caprine animals which have ning, by laceration of the central ner	e been killed, after	

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.
------------------------	--------------------------------	-------

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 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 feeding of farmed animals, other than fur animals, and the production or manufacturing of pet food.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28 in case of Species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or Suidae, Pesca, Reptilian.

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 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.

COUNTRY

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.	
Official veterinarian/Official ins	pector			
Name (in capital letters):		Qualification and title:		
Date:		Signature	·	
Stamp:				

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

'CHAPTER 6(B)

Health certificate

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

UIN	ΓRY:	Veterina	ary certificate to EU
	Consignor	I.2. Certificate reference No	I.2.a.
	Name Address	I.3. Central competent authority	
	Γel.	I.4. Local competent authority	
	Consignee Name Address Postcode Fel.	I.6. Person responsible for the load in EU Name Address Postcode Tel.	T
	ountry of ISO code I.8. Region of origin Code igin	I.9. Country of destination ISO code	I.10. Region of Code destination
I.11.	Place of origin	I.12. Place of destination	
	Name Approval number Address Name Approval number Address Name Approval number	Name Approv Address	n warehouse al 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 Cher Other Chentification Documentation references	I.17. Number(s) of CITES	
	Description of commodity	I.19. Commodi	ty 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:		
	Γechnical 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		
C	cies (Scientific name) Nu	mber of packages	
>ne			

_			
		•	
			l.

II.2

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. 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 (ii) originated from animals that were killed at a distance of at least 20 km from the borders of another third country or part of a third country not authorised to export untreated game trophies of wild swine to the European Union;] $(^2)or$ [II.1. 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 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;]

The game trophies or other preparations described above have been packaged without being in

 $(^2)or$

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

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

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

(2) or [are 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(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.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.

COUNTRY

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

II.	Health information	II.a. Certificate reference	e No	II.b.		
-	 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	и:					
(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.					
1 1	The signature and the stamp must be in a Note for the person responsible for the conformation of the person post of the point of entry into the person post of the point of entry into the person post of the point of entry into the person post of the point of entry into the person post of the point of entry into the person post of the perso	onsignment in the Europea ompany the consignmen	ın Union: 1	this certificate is only		
Offic	cial veterinarian/Official inspector					
	Name (in capital letters):		Qualificat	tion and title:		
	Date:		Signature	:		
	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 samples², 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 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	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	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				
-	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	har of artablishments				
		nber of establishments turing plant Number of packages Net weight Batch number				

Health information

II.

II.b.

Part II: Certification

I, the undersigned	official veterinarian, declare th	nat I have read and understood
Regulation (EC) No	1069/2009 of the European Parli	iament and of the Council(1a), and
Commission Regulat	tion (EU) No 142/2011(1b), and	in particular Chapter II of Annex
XIV thereto, and cert	ify that the animal by-products de	escribed above

(2)either [are trade samples which consist of animal by-products intended for particular studies or analyses as referred to in the definition of trade samples in point 39 of Annex I to Regulation (EU) No 142/2011, that bear the label 'TRADE SAMPLE NOT FOR HUMAN CONSUMPTION'.]

II.a. Certificate reference No

(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

- (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 either:
 - (i) have remained in that third country, territory or part thereof eligible to export fresh meat to the European Union since birth or for a period of at least the preceding three months before the date of slaughter; and/or
 - (ii) were killed in the wild in that third country, territory or part thereof(4);]
- (²) and/or [(c) derived from eggs, milk, rodents, lagomorphs, or aquatic animals or terrestrial or aquatic invertebrates;]
- (²)[II.1.2. in the case of materials other than materials derived from eggs, milk, rodents, lagomorphs, wool grease, aquatic animals, terrestrial or aquatic invertebrates and unprocessed furs, have been obtained from animals:
 - (2) either [(a) coming from holdings:
 - (i) where, for the following diseases for which the animals are susceptible, there has not been any case/outbreak of rinderpest, swine vesicular disease, Newcastle disease or highly pathogenic avian influenza during the period of the preceding 30 days, nor of classical or African swine fever during the period of the preceding 40 days; nor in the holdings situated in their vicinity within a 10 km radius, during the period of the preceding 30 days; and
 - (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

II. Hea	lth information		II.a. Certificate reference No	II.b.
		wh	ich did not comply with the same heal	th conditions;
		(iii) at du she	the slaughterhouse, passed the ante-mring the period of 24 hours before the owed no evidence of the diseases refer animals are susceptible; and	ortem health inspection e time of slaughter and
		sla Ur eq	re handled in the slaughterhouse bet ughter or killed in accordance with the tion legislation and complied with uivalent to those laid down in Chapte gulation (EC) No 1099/2009(5)]	e relevant provisions of requirements at least
	(²) <i>or</i> [(a)	captured	and killed in the wild in an area:	
		ang sus dis the	there within a 25 km radius there has being of the following diseases for we exceptible: foot-and-mouth disease, the ease or highly pathogenic avian influe to preceding 30 days nor of classical ring the period of the preceding 40 days	which the animals are rinderpest, Newcastle nza during the period of or African swine fever
		se _l wł	at is situated at a distance that exceeds corating another territory of a third contich is not authorised at these dates for terial to the European Union; and	country or part thereof,
	(b)	chilling	fter killing were transported within a either to a collection centre and imme ablishment, or directly to a game estab	ediately afterwards to a
(²)[II.1.3.	the wild, have be there has been animals are sus case/outbreak of the European U	een obtain no case/or ceptible du f one of the Union was	her than materials derived from fish of ed in an establishment around which, we atbreak of diseases referred to in polaring a period of the preceding 30 datus ose diseases, the preparation of raw material authorised only after the removal of the of the establishment under the	within a radius of 10 km, nt II.1.2 for which the ys or, in the event of a aterial for exportation to all meat, and the total
П.1.4.		e conditio	prepared without contact with other n ns required above, and it has been benic agents;	
II.1.5.	has been cleaned other than via p authority, beari MANUFACTU	ed and disi arcel post, ng the lab RE OF D	packaging which prevents any leakage infected before use and, in the case of in containers sealed under the responsed indicating 'ANIMAL BY-PRODUCTS FOR USES of daddress of the establishment of destablishment of destablishment.	f consignments shipped sibility of the competent CTS ONLY FOR THE OUTSIDE THE FEED
II.1.6.	consist only of t	he following	ng animal by-products:	
	(²)either [-	bodies o	and parts of animals slaughtered or r parts of animals killed which were tion in accordance with Union legis as animal by-products for commercial	deemed fit for human lation until irreversibly
	(²)and/or [-	were sla	and the following parts originating e aughtered in a slaughterhouse and v r for human consumption following an	vere considered fit for

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.
			and the following parts of anima assumption in accordance with Union	
		unfi legis	ases or bodies and parts of animals t for human consumption in a slation, but which did not show municable to humans or animals;	accordance with Union
		(ii) head	ls of poultry;	
		and	s and skins, including trimmings ar feet, including the phalanges and the es, tarsus and metatarsus bones;	
		(iv) pig l	oristles;	
		(v) featl	ners;]	
	(²)and/or [-	farm as re of the Eur	-products from poultry and lagome ferred to in Article 1(3)(d) of Regu opean Parliament and of the Counc of disease communicable to humans	lation (EC) No 853/2004 il ^(2a) , which did not show
	(²)and/or [-	communic animals the been cons	animals which did not show table through blood to humans or nat have been slaughtered in a slaudered fit for slaughter for human communication in accordance with University of the showing the same inspection in accordance with University of the same inspection in a same inspect	animals, obtained from ughterhouse after having onsumption following an
	(²)and/or [-	for huma	-products arising from the product n consumption, including degrea or separator sludge from milk proce	sed bone, greaves and
	(²)and/or [-	origin, w	of animal origin, or foodstuffs containing are no longer intended for all reasons or due to problems of matter defects from which no risk to	human consumption for nufacturing or packaging
	(²)and/or [-	animal by for feedi manufactu	nd feedingstuffs of animal origin, or products or derived products, which ing for commercial reasons or uring or packaging defects or other polic or animal health arises;]	th are no longer intended due to problems of
	(²)and/or [-	originating	centa, wool, feathers, hair, horns, g from live animals that did not shable through that product to humans	ow signs of any disease
	(²)and/or [-		nimals, and parts of such animals not show any signs of diseases con	
	(²)and/or [-	animal lestablishment	1	0 0
	(²)and/or [-		ring material originating from anin of disease communicable through the	
		(i) shel	ls from shellfish with soft tissue or f	lesh;
		(ii) the f	Collowing originating from terrestria	l animals:
			<u> </u>	

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

II. Hea	lth information		II.a. Certificate reference No	II.b.				
		-	hatchery by-products;					
		-	eggs;					
		-	egg by-products, including egg shells	5;				
(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 [-	Lagomory Article 8(Category	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 [-	-	nating from dead animals that did no se communicable through that produc	_				
II.1.7.	European Union	n legislation	ne plant of origin or have been present in such a way that they will not spivery to the plant of destination.					
(²)(⁶)[II.1.8. (²)(⁷)								
	in the country,	territory or ainst foot-a	this consignment come from animals part thereof referred to in point II and-mouth disease are regularly can ne animals.]]	.1.1, where vaccination				
$(^{2})(^{8})$								
and/or[II.1.8	.2.The animal by- offal or deboned		this consignment consist of animal b	y-products derived from				
(²)[II.1.9.	the animal by-pr	roducts desc	cribed above					
	(²)either [are de	rived from	other ruminants than bovine, ovine or	caprine animals.]]				
		rived from rived from:	bovine, ovine or caprine animals and	does not contain and is				
	(²) eith	anim regio	ine, ovine and caprine materials other als born, continuously reared and slam classified as posing a negligible I Decision 2007/453/EC.]]	ughtered in a country or				
	(²) <i>or</i>	[(a)	specified risk material as defined in Regulation (EC) No 999/2001 of thand of the Council(4);	-				
		(b)	mechanically separated meat ob bovine, ovine or caprine anima animals that were born, con slaughtered in a country or region negligible BSE risk in accorda Decision 2007/453/EC(5), in whi indigenous BSE case,	ls, except from those tinuously reared and a classified as posing a nce with Commission				
		(c)	animal by-product or derived p bovine, ovine or caprine animals after stunning, by laceration of the by means of an elongated	which have been killed,				

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

II.	Health informa	tion		II.a. Certificate reference No	II.b.
				introduced into the cranial cavity, e injected into the cranial cavity, e that were born, continuously rear country or region classified as posi in accordance with Decision 2007/	xcept for those animals ed and slaughtered in a ng a negligible BSE risk
II.1.10	the animal	by-prod	ducts des	cribed above:	
	(²)either	-		milk or milk products of ovine or car feed for farmed animals, other than	
	(²) <i>or</i>	-		or milk products of ovine or caprired for farmed animals, other than fur a	_
		(a)		ved from ovine and caprine animals ously since birth in a country where lled:	
			(i)	classical scrapie is compulsorily not	ifiable;
			(ii)	an awareness, surveillance and m place for classical scrapie;	nonitoring system is in
			(iii)	official restrictions apply to holding animals in the case of a susp confirmation of classical scrapie;	
			(iv)	ovine and caprine animals affected villed and destroyed;	with classical scrapie are
			(v)	the feeding to ovine and caprine ar meal or greaves, as defined in the T Code of the World Organisation for of ruminant origin has been banned in the whole country for a period of seven years;	errestrial Animal Health r Animal Health (OIE), and effectively enforced
		(b)		e from holdings where no official suspicion of TSE;	restrictions are imposed
		(c)	diagnos	e from holdings where no case of cl ed during the period of the prec ng the confirmation of a case of classi	eding seven years or,
			(²)eithe	fall ovine and caprine animals on killed and destroyed or slaughters rams of the ARR/ARR genotype, be least one ARR allele and no VRQ animals carrying at least one ARR a	ed, except for breeding reeding ewes carrying at allele and other ovine
			(²)or	[all animals in which classical scrabeen killed and destroyed, and subjected for a period of at least two confirmation of the last classical so TSE monitoring, including testing the presence of TSE in accordant methods set out in point 3.2 of Changulation (EC) No 999/2001, of animals which are over the age of animals of the ARR/ARR genotype:	the holding has been by years since the date of rapie case to intensified with negative results for ce with the laboratory lapter C of Annex X to f all of the following 18 months, except ovine
				 animals which have been s consumption; and 	laughtered for human

II.	Health information	II.a	. Certificate reference No	II.b.
		_	animals which have died or b but which were not killed in the eradication campaign.]].	Č

Notes

Part I:

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

Part II:

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

COUNTRY

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.		
(2)	Delete as appropriate.			'			
(2a)	OJ L 139, 30.4.2004, p. 55.						
(3)	The name and ISO code number of t	he exp	porting country as laid do	own 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	on (E0	C) No 798/2008 (OJ L 22	26, 23.8.	2008, p. 1), and		
	- Annex I to Commission Regulation	on (E0	C) No 119/2009 (OJ L 39	, 10.2.2	009, p. 12).		
	In addition the ISO code of territori (EU) No 206/2010, (EC) No 798/2 applicable for the susceptible species	2008 a	and (EC) No 119/2009	referred	to in this note (where		
(4)	Only for countries from where the game meat intended for human consumption of the same animal species is authorised for importation into the European Union.						
(5)	OJ L 303, 18.11.2009, p. 1.						
(6)	Supplementary guarantees to be provided where the material of domestic ruminants originated in the territory of a South American or South African country or part thereof from where only maturated and deboned fresh meat of domestic ruminants for human consumption is authorised for exportation to the European Union. The whole masseter muscles of bovine animals, incised in accordance with the requirements of Part B.1 of Chapter I of Section IV of Annex Ito Regulation (EC) No 854/2004 of the European Parliament and of the Council (OJ L 139 30.4.2004, p. 206), are also permitted.						
(7)	Only for certain South American cou	untries	3.				
(8)	Only for certain South American and	d Sout	h African countries.				
(9)	OJ L 147, 31.5.2001, p. 1.						
(10)	OJ L 172, 30.6.2007, p. 84.						
-	The signature and the stamp must be			-	•		
-	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.						
Offi	Official veterinarian/Official inspector						
	Name (in capital letters):			Qualific	cation and title:		
	Date:			Signatu	re:		
	Stamp:						

EN 94 EN

(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

	UNIKI:	veterinary certificate to EU				
	I.1. Consignor Name	I.2. Certificate reference No I.2.a.				
	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				
atch	10.	Tel.				
of dispa	I.7. Country of 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				
art I : De	Name Approval number Address Name Approval number Address	Custom warehouse Name Approval number Address				
F	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:					
	Animal feedingstuff Manu	nufacture 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	umber of establishments				
		nufacturing plant Number of packages Net weight Batch number				

	II.	Health informa	ntion	II.a. Certificate reference No	II.b.				
		(EC) No 1069/2 Article 10 there	2009 of the Europeof, and Commiss	arian, declare that I have read and u bean Parliament and of the Council tion Regulation (EU) No 142/2011(and certify that the rendered fats desc	(la), and in particular (lb), and in particular				
0 0	II.1.	consist of rendered fats that satisfy the health requirements below;							
ati	II.2.	consist of rendered fats not intended for human consumption;							
Part II: Certification	II.3.	have been prepared and stored in a plant approved and supervised by the competent authority in accordance with Article 24 of Regulation (EC) No 1069/2009 or in accordance with Article 4(2) of Regulation (EC) No 853/2004 of the European Parliament and of the Council(3), in order to kill pathogenic agents;							
rt I	II.4.	have been prepa	ared exclusively wi	th the following animal by-products:					
Pa		(²)either [-	carcases and parts of animals slaughtered or, in the case of game, bodies o 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 [-	been slaughtered for human consumathe following part	following parts originating either from a slaughterhouse and were considered to the following an ante-mortem insects of animals from game killed for his Union legislation:	lered fit for slaughter pection or bodies and				
			human con	bodies and parts of animals which are sumption in accordance with Union to ow any signs of disease communications.	legislation, but which				
			(ii) heads of po	oultry;					
			feet, includ	kins, including trimmings and splitti ling the phalanges and the carpus ar metatarsus bones;					
			(iv) pig bristles	•					
			(v) feathers;]						
		(²)and/or [-	through blood to slaughtered in a slaughter for hur	which did not show any signs of d humans or animals, obtained from an a slaughterhouse after having been nan consumption following an ante- Union legislation;]	nimals that have been n considered fit for				
		(²)and/or [-	human consumpt	cts arising from the production of ption, including degreased bone, greatfrom milk processing;]					
		(²)and/or [-	origin, which a commercial reas	nal origin, or foodstuffs containing are 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				
		(²)and/or [-	animal by-production feeding for com-	dingstuffs of animal origin, or feed tests or derived products, which are no mercial reasons or due to problems s or other defects from which no ris	o longer intended for of manufacturing or				
		(2) 1/ 5	11 1 1	1 0 1 1 1 1 0	. 1 91				

(2) and/or [- blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk

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

II.	Health infor	mation		II.a.	Certificate refere	ence No	II.b.
					animals that did that product to h		signs of any disease mals;]
	(²)and/or [-				rts of such animaliseases communi		a mammals, which did ans or animals;]
	(²)and/or [-				rom aquatic ani cturing products f	_	ating from plants or assumption;]
	(²)and/or [-		_				ich did not show any o humans or animals:
		(i)	shells from	shellf	ish with soft tissu	ie or flesh;	
		(ii)	the followin	g orig	ginating from terr	estrial animal	s:
			- hatche	ery by	-products,		
			- eggs,				
			- egg by	y-pro	ducts, including e	gg shells;	
		(iii)	day-old chic	eks ki	lled for commerc	ial reasons;]	
II.5.	(²)either	the of t	territory of a	cour g 24	itry free from foo	ot-and-mouth e from class	n a country or part of disease for the period ical swine fever and 2 months;]
	(²)and/or	terr		m Ne	ewcastle disease		a country or part of a luenza for a period of
	(²)and/or	a to	erritory free	fron nonths	n foot-and-mout s and free from	h disease fo	m a country or part of or the period of the for the period of the
	(²)and/or	poir the to a	nt II.5. during rendered fats	g the s deri	relevant period relevant period relevant	referred to in ptible species	liseases referred to in point II.5, and where s, have been subjected or at least 90 °C for at
		the com info app	owner, open npetent auth ormation must	erator ority st inc absol	or their repre can monitor t lude the particle	sentative and the operation size, critica	nd maintained so that d, as necessary, the n of the plant; the l temperature and, as material feed rate and
II.6.					e purified in such a not exceed 0,15		e maximum levels of
II.7.	the rendered	fats:	_			_	
		t	of Section 3 142/2011, or	of (Chapter II of A atment in accorda	nnex X to ance with Sec	with the requirements Regulation (EU) No etion XII of Annex III ill pathogenic agents;
	(²)either	а	and disinfect	ed if		e prevention	hat have been cleaned of contamination, and ontamination;]
	$(^{2})or$	[(b) v	where bulk to	ranspo	ort is intended, th	ne pipe, pum	os and bulk tanks and

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

II.	Health i	nformation		II.a. Certificate reference No	II.b.
			of the proc ship or into	ulk container or bulk road tanker used luct from the manufacturing plant eit o shore tanks or directly to plants hav sibility of the competent authority and	her directly on to the e been checked under
	and whic	ch bear labels	s indicating	'NOT FOR HUMAN CONSUMPTIO	N';
$(^{2})[II.8.$	the rende	ered fats desc	cribed above		
	(²)either	[is derived	from other i	ruminants than bovine, ovine or caprin	e animals.]]
	(²) <i>or</i>	[is derived derived from		e, ovine or caprine animals and does	not contain and is not
		(²) either	animals b	ovine and caprine materials other that born, continuously reared and slaugh assified as posing a negligible BSE ris 2007/453/EC.]]	tered in a country or
		(²) <i>or</i>	Re	ecified risk material as defined in pogulation (EC) No 999/2001 of the d of the Council(4);	
			ov bo re ac	echanically separated meat obtained frine or caprine animals, except from thorn, continuously reared and slaught gion classified as posing a negl cordance with Commission Decision hich there has been no indigenous BSE	ose animals that were ered in a country or igible BSE risk in n 2007/453/EC(5), in
			ov str of er ca re	imal by-product or derived product of time or caprine animals which have unning, by laceration of the central new an elongated rod-shaped instrument anial cavity, or by means of gas injuvity, except for those animals that we have and slaughtered in a country of sing a negligible BSE risk in accord 107/453/EC.]]]	re been killed, after rvous tissue by means t introduced into the ected into the cranial ere born, continuously r region classified as
II.9. the	rendered	fats describe	d above:		
	(²)either	_		ilk or milk products of ovine or capri d for farmed animals, other than fur an	•
	(2) <i>or</i>			milk products of ovine or caprine or farmed animals, other than fur animals	
		(from ovine and caprine animals we since birth in a country where the following	
		(i) clas	sical scrapie is compulsorily notifiable	e;
		(awareness, surveillance and monitoring classical scrapie;	ng system is in place
		(anii	cial restrictions apply to holdings nals in the case of a suspicion of TS lassical scrapie;	
		(ne and caprine animals affected with ed and destroyed;	classical scrapie are
		(v) the	feeding to ovine and caprine anima	als of meat-and-bone

(b)

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

II.	Health information	II.a. Certificate re	eference No	II.b.	
	me	meal or greaves, as defined in the Terrestrial Animal Health			
	Co	Code of the World Organisation for Animal Health (OIE), of			
	rur	ninant origin has bee	en banned and e	ffectively enforced in	

a suspicion of TSE;

originate from holdings where no official restrictions are imposed due to

the whole country for a period of at least the preceding seven

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

Notes

Part I:

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

COUNTRY

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

II.	Health information	II.a. Certificate reference No	II.b.					
_	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	i II:							
(1a)	OJ L 300, 14.11.2009, p. 1.							
(1b)	OJ L 54, 26.2.2011, p. 1.							
$(^{2})$	Delete as appropriate.							
$(^{3})$	OJ L 139, 30.4.2004, p. 55.							
(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 print	ing.					
-	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):	Qualifica	ation and title:					
	Date:	Signatur	e:					
	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

CC	UNTRY:	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 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 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 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 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					
		1.24. 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					

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

II. **Health information** II.a. Certificate reference No II.b. I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council (^{1a}), and in particular Articles 8, 9 and 10 thereof, and Commission Regulation (EU) No 142/2011(1b), and in particular Chapter II of Annex XIV thereto, and certify that the rendered fats described 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: (2)[II.2.1. in the case of materials destined for the production of renewable fuels referred to in point L of Section 2 of Chapter IV of Annex IV to Regulation (EU) No 142/2011, biodiesel or oleochemical products, animal by-products referred to in Articles 8, 9 and 10 of Regulation (EC) No 1069/2009:1 (2)[II.2.2. in the case of materials destined for the production of renewable fuels referred to in point J of Section 2 of Chapter IV of Annex IV to Regulation (EU) No 142/2011, the materials have been prepared exclusively from animal by-products referred to in Articles 9 and 10 of Regulation (EC) No 1069/2009;] (2)[II.2.3. in the case of materials destined for purposes other than cosmetics, pharmaceuticals or medical devices, the materials have been prepared exclusively from: animal by-products containing residues of authorised substances or (2)either contaminants exceeding the permitted levels referred to in Article 15(3) of Council Directive 96/23/EC(^{2a});] (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;] animals and parts of animals, other than those referred to in Articles 8 and (2)and/or 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; feathers;] (v) (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

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.		
been slaughtered in a slaughterhouse after having been considered slaughter for human consumption following an ante-mortem inspec 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 arises;]		
	(²)and/or	[-	petfood and feeding stuffs of animal origin, or feeding stuffs 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	[-	blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk originating from live animals that did not show signs of any disease communicable through that product to humans or animals;]		
	(²)and/or	[-	aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of diseases communicable to humans or animals;]		
	(²)and/or	[-	animal by-products from aquatic animals originating from plants or establishments manufacturing products for human consumption;]		
	(²)and/or	[-	the following material originating from animals which did not show any signs of disease communicable through that material to humans or animals:		
			(i) shells from shellfish with soft tissue or flesh;		
			(ii) the following originating from terrestrial animals:		
			- hatchery by-products,		
			- eggs,		
			- egg by-products, including egg shells,		
	2	_	(iii) day-old chicks killed for commercial reasons;]		
	(²)and/or	[-	aquatic and 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 1069/2009and Category 2 material as referred to in Article 9(a) to (g) of that Regulation;]		
	(²)and/or	[-	hides and skins, hooves, feathers, wool, horns, hair and fur originating from dead animals that did not show any signs of disease communicable through that product to humans or animals;]		
	(²)and/or	[-	adipose tissue from animals which did not show any signs of disease communicable through that material to humans or animals, which were slaughtered in a slaughterhouse and which were considered fit for slaughter for human consumption following an ante-mortem inspection in accordance with Union legislation;]]		
(²)[II.2	2.4. in the case	of ma	terials destined for purposes other than the production of organic fertilisers		

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

II.	Healt	h info	ormati	on	II.a.	Certificate r	eference No	II.b.
	or soi	l impi	rovers,	cosmetics, pl	narmaceuti	cal or medical	devices:	
						erial as defined in Article 3(1)(g) of Regulation (EC) e European Union and of the Council ^(2b) ;]		
	(²)an	d/or	[-	entire bodies or parts of dead animals containing specified risk material a defined in Article 3(1)(g) of Regulation (EC) No 999/2001 at the time o disposal;]				
	(²)an	d/or	[-	animal by-products which have been derived from animals which have been submitted to illegal treatment as defined in Article 1(2)(d) of Council Directive 96/22/EC ^(2c) or Article 2(b) of Council Directive 96/23/EC;]				
	(²)an	d/or	[-	environment 96/23/EC, it	al contaming such restantion or, in	inants listed in sidues exceed	Group B(3) of the permitted	her substances and Annex I to Directive levels laid down by lation of the Member
II.3.	the re	ndere	d fats:					
 (a) have been subjected to processing in accordance with method processing method) as set out in Chapter III of Annex IV to No 142/2011, in order to kill pathogenic agents, (b) have been marked before shipment to the European Union with g (GTH), so that a homogenous minimum concentration of at leas kilogramme fat is achieved, 					n Chapter III			
	(c) in the case of rendered fats of ruminant in weight have been removed,					nant origin, in	soluble impuriti	es in excess of 0,15%
	(d)	have	been transported under conditions which prevent their con-				amination, and	
	(e)			s on the pac		container in	dicating "NOT	FOR HUMAN OR
(²)[II.4. in the case of materials destined for organic fertilisers, condevices or soil improvers the rendered fats described above					rmaceuticals, medical			
	(²)eit	her [a	are der	ived from oth	er ruminar	nts than bovine	e, ovine or caprii	ne animals.]
	(2) <i>or</i>		are der erived	_	vine, ovine	or caprine an	imals and does	not contain and is not
		(2	²) eithe	animals region o	born, cor	ntinuously rea as posing a ne	red and slaught	n those derived from tered in a country or sk in accordance with
		(2)	or	Re		(EC) No 999/2		int 1 of Annex V to opean Parliament and
				ov bo re w	vine or cap orn, contingion class ith Comm	orine animals, nuously reared ified as posing	except from the d and slaughter g a negligible BS n 2007/453/EC(om bones of bovine, ose animals that were red in a country or SE risk in accordance 5), in which there has
				OV	ine or c	aprine anima	als which have	btained from bovine, e been killed, after yous tissue by means

II.	Health information	II.a. Certificate reference No	II.b.
	of an	elongated rod-shaped instrument	introduced into the
	cranial	cavity, or by means of gas inject	eted into the cranial
	cavity,	except for those animals that were	e born, continuously
	reared	and slaughtered in a country or	region classified as
	posing	a negligible BSE risk in accord	ance with Decision

2007/453/EC.]]]

II a Cartificata reference No

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 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 feeding of farmed animals, other than fur animals or pet animals, and the production or manufacturing of pet food.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28:
 - Species: select from the following: Ruminantia, other than Ruminantia
 - Manufacturing plant: provide the registration number of the treatment/processing establishment.

Part II:

- (1a) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.
- (2a) OJ L 125, 23.5.1996, p. 10.
- (2b) OJ L 147, 31.5.2001, p. 1.
- (2c) OJ L 125, 23.5.1996, p. 3.
- (3) OJ L 147, 31.5.2001, p. 1.
- (4) OJ L 172, 30.6.2007, p. 84.
- The signature and the stamp must be in a 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

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

II. Health informat	ion II	.a.	Certificate reference N	0	II.b.				
inspection post of the	inspection post of the point of entry into the European Union.								
Official veterinarian/Officia	l inspector								
Name (in capital letter	s):		Qua	lificati	on and title:				
Date:			Sign	nature:					
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
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 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: Animal feedingstuff □ Manu	afacture of petfood □ 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 (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

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 in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011(^{1b}), and in particular Chapter I of Annex XIV thereto, and certify that the gelatine/collagen(2) described above: Part II: Certification II.1. consists of gelatine/collagen(2) that satisfy the health requirements below; 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: has been prepared exclusively with the following animal by-products: II.4. carcases and parts of animals slaughtered or, in the case of game, bodies (2)either 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;] carcases and the following parts originating either from animals that (2)and/or 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;] animal by-products arising from the production of products intended for (2)and/or human consumption, including degreased bone, greaves and centrifuge or separator sludge from milk processing;] products of animal origin, or foodstuffs containing products of animal (2)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 arises;] (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;]

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

II.	Health in	nformatio	n	II.a. Certificate reference No II.b.					
II.5.	the gelat	ne/collagen(²):							
		(a)	hygiene cond in a dedicat legislation w	_					
				nd packages containing gelatine/collagen(²) bear the words /COLLAGEN(2) SUITABLE FOR ANIMAL TON'; and					
	(²)either	(b)	in the case of gelatine, was produced by a process that ensured that unprocessed Category 3 material was subjected to a treatment with acid or alkali, followed by one or more rinses, involving pH adjustment, extraction by heating one or several times in succession, followed by purification by means of filtration and sterilisation, in order to kill pathogenic agents;]						
	(²)or	[(b)	in the case of collagen, was produced by a process that ensured that unprocessed Category 3 material was subjected to a treatment involving washing, pH adjustment using acid or alkali followed by one or more rinses, filtration and extrusion, in order to kill pathogenic agents;]						
(²)[II.6.	in the cas	he case of gelatine/collagen(2) from materials other than hides and skins							
	(²)either	(2) either [is derived from other ruminants than bovine, ovine or caprine animals.]]							
	(²) <i>or</i>		[is derived from bovine, ovine or caprine animals and does not contain and is a derived from:						
		(²) either	animals b	ovine and caprine materials other than those derived from orn, continuously reared and slaughtered in a country or ssified as posing a negligible BSE risk in accordance with 2007/453/EC.]]					
		(²) <i>or</i>	Reg	eified risk material as defined in point 1 of Annex V to ulation (EC) No 999/2001 of the European Parliament and ne Council(4);					
			ovir borr regi with	chanically separated meat obtained from bones of bovine, the or caprine animals, except from those animals that were an animals reared and slaughtered in a country or on classified as posing a negligible BSE risk in accordance a Commission Decision 2007/453/EC(5), in which there has a no indigenous BSE case,					
			ovir stun of a crar cavi rear posi	nal by-product or derived product obtained from bovine, ne or caprine animals which have been killed, after ning, by laceration of the central nervous tissue by means an elongated rod-shaped instrument introduced into the tial cavity, or by means of gas injected into the cranial ty, except for those animals that were born, continuously ed and slaughtered in a country or region classified as ng a negligible BSE risk in accordance with Decision 7/453/EC.]]]					
II.7.	in the cas	se of gelati	ne/collagen(2)	from materials other than hides and skins described above:					
	(²)eithei			or milk products of ovine or caprine animal origin or is not armed animals, other than fur animals.]					
	(2) <i>or</i>	[contain	s milk or milk	products of ovine or caprine animal origin and is intended					

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

for feed for farmed animals, other than fur animals, which:

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

Notes

Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is 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

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.

transit commodity. Products in transit may only be stored in free zones, free warehouses and custom warehouses.

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

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 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.

Offici	Official veterinarian/Official inspector									
1	Name (in capital letters):	Qualification and title:								
1	Date:	Signature:								
5	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

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 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 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 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	
		number of establishments nufacturing plant Number of packages Net weight Batch number

	II.	Health i	nformation	_	II.a. Certificate reference No	II.b.				
g.		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council(^{1a}), and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011(^{1b}), and in particular Chapter I of Annex XIV thereto, and certify that the hydrolysed protein/dicalcium phosphate/tricalcium phosphate(2) described above:								
icatio	II.1.		onsists of hydrolysed protein/dicalcium phosphate/tricalcium phosphate(2) that satisfy the ealth requirements below;							
ertif	II.2.		consists exclusively of hydrolysed protein/dicalcium phosphate/tricalcium phosphate(2) not intended for human consumption;							
Part II: Certification	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;							
Ь	II.4.	has been	has been prepared exclusively with the following animal by-products:							
		(²)either	[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;]							
		(2)or	[in the case of other materials:							
			(²)either [-	r [- 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	for [-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	intend	nal by-products arising from the led for human consumption, includi entrifuge or separator sludge from m	ng degreased bone, greaves				
			(²)and/or		ducts of animal origin, or foodstu al origin, which are no longer intend					

II.	Health information	ı	II.a. Certificate reference No	II.b.					
		packa	for commercial reasons or due to problems of manufacturing packaging defects or other defects from which no risk to public animal health arise;]] [-petfood and feedingstuffs of animal origin, or feedingst containing animal by-products or derived products, which are longer intended for feeding for commercial reasons or due problems of manufacturing or packaging defects or other def from which no risk to public or animal health arises;]]						
	(²)and/or	conta longe probl							
	(²)and/or	origin	[-blood, placenta, wool, feathers, hair, horns, hoof cuts and raw moriginating from live animals that did not show signs of any disease communicable through that product to humans or animals;]] [-aquatic animals, and parts of such animals, except sea mamma which did not show any signs of diseases communicable to human or animals;]]						
	(²)and/or	which							
	(2)and/or		[-animal by-products from aquatic animals originating from plant establishments manufacturing products for human consumption;]]						
	(²)and/or		following material originating from a igns of disease communicable throumals:						
		(i)	shells from shellfish with soft tissue	or flesh;					
		(ii)	the following originating from terres	strial animals:					
		-	hatchery by-products,						
		-	eggs,						
		-	egg by-products, including egg she	ells;					
		(iii)	day-old chicks killed for commercia	ıl reasons;]]					
II.5.	the hydrolysed prote	ein/dicalc	ium phosphate/tricalcium phosphate	(²):					
		'NOT FO under sar packagin	pped and packaged in packaging work HUMAN CONSUMPTION' and itsfactory hygiene conditions, and in g took place in a dedicated room of under Union legislation were used;	I was stored and transported particular the wrapping and m, and only preservatives					
			se of hydrolysed protein, was produ ate measures to minimise contami						
		ruminant only to l preparati	ase of hydrolysed proteins entire is hides and skins, was produced in a hydrolysed proteins production, usi on of the raw Category 3 materia washing followed by:	a processing plant dedicated ng a process involving the					
		3 h	exposure of the material to a pH of cours at a temperature of more than at treatment at a temperature of nutes at more than 3,6 bar; or	80 °C and subsequently by					
		mo	exposure of the material to a pH of ore than 11, followed by a heat tre ore than 140 °C for 30 minutes at 3 b	atment at a temperature of					

II.	Health i	nformatio	n	II.a.	Certificate reference No	II.b.
	(2) <i>or</i>	[(b)	in the ca	se of di	calcium phosphate, was pro	oduced by a process that:
			de (at	greased a min	l with hot water and treated	aterial is finely crushed and with dilute hydrochloric acid 6 and a pH of less than 1,5)
				ie, resu		ained phosphoric liquor with alcium phosphate at pH 4 to 7,
					r-dries this precipitate, with and an end temperature of	an inlet temperature of 65 °C between 30 °C and 65 °C.]
	$(^{2})or$	[(b)	in the ca	se of tr	icalcium phosphate, was pro	oduced by a process ensuring:
					ategory 3 bone-material is r-flow with hot water (bone	finely crushed and degreased chips less than 14 mm),
				contin	nuous cooking with steam a	t 145 °C during 30 minutes at
					ration of the protein bro m phosphate) by centrifugat	th from the hydroxyapatite ion, and
			` /	_	ulation of the tricalcium bed with air at 200 °C.]	phosphate after drying in a
(²)[II.6.	the hydr	olysed prot	tein/dicalc	ium ph	nosphate/tricalcium phospha	te(2) described above
	(²)eithe	r [is derive	d from otl	ner rum	ninants than bovine, ovine o	r caprine animals.]]
	(²) <i>or</i>	[is derived from bovine, ovine or caprine animals and does not contain and is no derived from:				
		anim regio			n, continuously reared and	ther than those derived from slaughtered in a country or BSE risk in accordance with
		(²) <i>or</i>	[(a)	Regula		d in point 1 of Annex V to the European Parliament and
			(b)	ovine born, region with C	or caprine animals, except continuously reared and classified as posing a negl	ained from bones of bovine, from those animals that were slaughtered in a country or igible BSE risk in accordance 453/EC(5), in which there has
			(c)	ovine stunning of an cranial cavity, reared posing	or caprine animals whing, by laceration of the cerelongated rod-shaped instruction cavity, or by means of except for those animals and slaughtered in a course.	oduct obtained from bovine, ch have been killed, after atral nervous tissue by means trument introduced into the gas injected into the cranial that were born, continuously antry or region classified as an accordance with Decision
II.7.	the hydr	olysed prot	tein/dicalc	ium ph	nosphate/tricalcium phospha	te(2) described above:
	(²)either	[does:	not contai	n milk	or milk products of ovine	or caprine animal origin or is

II.	Health ir	nformation	1	II.a. Certificate reference No II.b.
		not inte	ended for	r feed for farmed animals, other than fur animals.]
	(²) <i>or</i>			or milk products of ovine or caprine animal origin and is ed for farmed animals, other than fur animals, which:
		(a)		erived from ovine and caprine animals which have been kep amously since birth in a country where the following conditions lfilled:
			(i) (classical scrapie is compulsorily notifiable;
			` /	an awareness, surveillance and monitoring system is in place fo classical scrapie;
			i	official restrictions apply to holdings of ovine or caprine animal in the case of a suspicion of TSE or the confirmation of classica scrapie;
				ovine and caprine animals affected with classical scrapie ar killed and destroyed;
			t	the feeding to ovine and caprine animals of meat-and-bone meator greaves, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health (OIE), of ruminar origin has been banned and effectively enforced in the whol country for a period of at least the preceding seven years;
		(b)	_	ate from holdings where no official restrictions are imposed duspicion of TSE;
		(c)	diagno	ate from holdings where no case of classical scrapie has bee osed during the period of the preceding seven years or, following infirmation of a case of classical scrapie:
			(²)eith	[all ovine and caprine animals on the holding have bee killed and destroyed or slaughtered, except for breedin rams of the ARR/ARR genotype, breeding ewes carryin at least one ARR allele and no VRQ allele and othe ovine animals carrying at least one ARR allele;]
			(²)or	[all animals in which classical scrapie was confirme have been killed and destroyed, and the holding has bee subjected for a period of at least two years since the dat of confirmation of the last classical scrapie case t intensified TSE monitoring, including testing wit 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/ARI genotype:
				 animals which have been slaughtered for huma 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.]]

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

Part I:

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

Part II:

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

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 $(^2)$ the European Union

	JUNIKI:	veterinary certificate to EU			
	I.1. Consignor Name	I.2. Certificate reference No I.2.a.			
	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	I.6. Person responsible for the load in EU Name Address			
patched	Tel.	Postcode Tel.			
s of dis	I.7. Country of 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			
art I : De	Name Approval number Address Name Approval number Address	Custom warehouse Name Approval number Address			
P.	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. Number(s) of CITES			
	I.18. Description of commodity	I.19. Commodity code (HS code) 05.07			
		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:	·			
	Further process 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 estable Manufacturing plants				

Part II: Certification

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.

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 horns and horn products, excluding horn meal, and hooves and hoof products, excluding hoof meal(²) described above originate from animals

- (2) either [that were slaughtered in a slaughterhouse, after undergoing ante-mortem inspection, and were fit, as a result of such inspection, for slaughter for human consumption;]
- (2) or [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:

(2)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;]

and 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.

- (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(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

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.
-----	--------------------	--------------------------------	-------

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 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.

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

(7) Chapter 20 is replaced by the following:

'CHAPTER 20

Model declaration

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

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 dispa	I.7. Country of ISO code I.8. Region of origin Code origin	I.9. Country of 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	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 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			
	Approval number of establis Species (Scientific name) Manufacturing plant	shments Net weight Batch number		

•		•	

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^(1a), and in particular that:

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

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;]

(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

[- 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

[- blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk originating from live animals that did not show signs of any disease communicable through that product to humans or animals;]

(2)and/or

[- aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of diseases communicable to humans or animals;]

(2)and/or

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

(2)and/or

- [- the following material originating from animals which did not show any signs of disease communicable through that material to humans or animals:
 - (i) shells from shellfish with soft tissue or flesh;
 - (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) 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:
 - aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of disease communicable to humans or animals,
 - 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) 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;

- (ii) foetuses;
- (iii) oocytes, embryos and semen which are not destined for breeding purposes; and
- (iv) dead-in-shell poultry;]

(2) and/or [- animal by-products other than Category 1 material or Category 3 material;]

- (4) its outer packaging is labelled 'FOR MEDICINAL PRODUCTS / VETERINARY MEDICINAL PRODUCTS / MEDICAL DEVICES FOR MEDICAL AND VETERINARY PURPOSES / ACTIVE IMPLANTABLE MEDICAL DEVICES / IN VITRO DIAGNOSTIC MEDICAL DEVICES FOR MEDICAL AND VETERINARY PURPOSES / LABORATORY REAGENTS / COSMETIC PRODUCTS ONLY' and it is not intended to be diverted at any stage within the European Union for any other use;
- 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:

²⁾either [an establishment or plant for the production of medicinal products, veterinary medicinal products, medical devices for medical and veterinary purposes, active implantable medical devices, in vitro diagnostic medical devices for medical and veterinary purposes, laboratory reagents or cosmetic products, which has been registered in accordance with Article 23 of Regulation (EC) No 1069/2009],

[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.
- ^(1a) 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.
- ^(2a) OJ L 125, 23.5.1996, p. 3.
- ^(2b) OJ L 125, 23.5.1996, p. 10.

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

EN 131 EN