Commission Regulation (EU) No 142/2011 of 25 February 2011 implementing Regulation (EC) No 1069/2009 of the European Parliament and of the Council laying down health rules as regards animal by-products and derived products not intended for human consumption and implementing Council Directive 97/78/EC as regards certain samples and items exempt from veterinary checks at the border under that Directive (Text with EEA relevance)

### ANNEX XV

# MODEL HEALTH CERTIFICATES

The model health certificates in this Annex shall apply to the importation from third countries and to the transit through the European Union of the animal by-products and the derived products referred to in the respective model health certificates.

Notes

- (a) Veterinary certificates shall be produced by the exporting third country, based on the models set out in this Annex, according to the layout of the model that corresponds to the animal by-products or derived products concerned. They shall contain, in the numbered order that appears in the model, the attestations that are required for any third country and, as the case may be, those supplementary guarantees that are required for the exporting third country or part thereof.
- (b) Where the model certificate states that certain statements shall be kept as appropriate, statements which are not relevant may be crossed out and initialled and stamped by the certifying officer, or completely deleted from the certificate.
- (c) The original of each certificate shall consist of a single sheet of paper, both sides, or, where more text is required; it shall be in such a form that all sheets of paper needed are part of an integrated whole and indivisible.
- (d) It shall be drawn up in at least one of the official languages of the EU Member State in which the inspection at the border post shall be carried out and of the EU Member State of destination. However, these Member States may allow other languages, accompanied, if necessary, by an official translation.
- (e) If for reasons of identification of the items of the consignment, additional sheets of paper are attached to the certificate, these sheets of paper shall also be considered as forming part of the original of the certificate by the application of the signature and stamp of the certifying official veterinarian, in each of the sheets of paper.
- (f) When the certificate, including additional schedules referred to in e), comprises more than one page, each page shall be numbered (*page number*) of (*total number of pages*) at the bottom of the page and shall bear the code number of the certificate that has been designated by the competent authority at the top of the page.
- (g) The original of the certificate must be completed and signed by an official veterinarian. In doing so, the competent authorities of the exporting country shall ensure that the principles of certification equivalent to those laid down in Directive 96/93/EC are followed.
- (h) The colour of the signature shall be different to that of the printing. The same rule applies to stamps other than those embossed or watermark.
- (i) The original of the certificate must accompany the consignment at the EU border inspection post.
- (j) If health certificates are used for consignments in transit, box No I.5 ('Consignee') of the relevant health certificate shall be completed with the name and address of the border inspection post through which the consignment is intended to leave the European Union.

# CHAPTER 1

# **Health certificate**

For processed animal protein not intended for human consumption, including mixtures and products other than petfood containing such protein, for dispatch to or for transit through  $\binom{2}{1}$  the European Union

COUNTRY Veterinary certificate to							
	l.1.	Consignor	I.2. Certificate reference No I.2.a.				
		Name	I.O. Control comments at the city.				
		Address	I.3. Central competent authority				
		Tel.	I.4. Local competent authority				
Ħ	1.5.	Consignee	I.6. Person responsible for the load in EU				
i i		Name	Name				
sigr		Address	Address				
ő							
ped		Postcode Tel.	Postcode Tel.				
atch							
of dispatched consignment	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10. Region of Cod destination	ie			
8							
Part I: Details	l.11.	Place of origin	I.12. Place of destination				
Part I:		Name Approval number Address	Name Custom warehouse ☐ Address Approval number				
		Name Approval number Address					
		Name Approval number Address	Postcode				
	L13.	Place of loading	I.14. Date of departure				
		- Last of leading					
	l.15.	Means of transport	I.16. Entry BIP in EU				
		Aeroplane					
		Road vehicle Other O	1.17.				
		Identification  Documentation references					
	1.18.	Description of commodity	I.19. Commodity code (HS code)				
			I.20. Quantity				
	I.21.	Temperature of product	I.22. Number of packages				
			Frozen				
	1.23.	Seal/Container No	I.24. Type of packaging				
	1.25.	Commodities certified for:					
		Animal feedingstuff  Technical use					
	1.26.	For transit through EU to third country	I.27. For import or admission into EU				
		Third country ISO code					
	1.28.	Identification of the commodities					
	(		mber of establishments Net weight Batch numbe ufacturing plant	er			

# COUNTRY

# Processed animal protein not intended for human consumption including mixtures and products other than petfood containing such protein

COUNTRY			other than petfood containing such protein							
		II. Health information II.a. Certificate reference No II.b.								
			I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council ( <sup>1a</sup> ) and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011 ( <sup>1b</sup> ), and in particular Annex X, Chapter II, Section 1, and Annex XIV, Chapter I, thereof and certify that:							
	ntion	II.1.	the processed animal protein or product described above contains exclusively processed animal protein not intended for human consumption that:							
	Part II: Certification		(a) has been prepared and stored in an establishment or plant approved, validated and supervised by the competent authority in accordance with Article 24 of Regulation (EC) No 1069/2009, and							
	art II:		(b) has been prepared exclusively with the following animal by-products:							
	•		(²) either [- carcases and parts of animals slaughtered or, in the case of game, bodies or parts of animals killed, and which are fit for human consumption in accordance with Union legislation, but are not intended for human consumption for commercial reasons;]							
	(2) and/or [- carcases and the following parts originating either from animals that have been slaughtered in a slaughterhouse were considered fit for slaughter for human consumption following an ante-mortem inspection or bodies and the follow parts of animals from game killed for human consumption in accordance with Union legislation:									
			<ul><li>(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;</li></ul>							
			(ii) heads of poultry;							
			(iii) hides and skins, including trimmings and splitting thereof, horns and feet, including the phalanges and the carpus and metacarpus bones, tarsus and metatarsus bones, of animals, other than ruminants;							
			(iv) pig bristles;							
			(v) feathers;]							
			(²) and/or [- blood of animals which did not show any signs of disease communicable through blood to humans or animals, obtained from animals other than ruminants that have been slaughtered in a slaughterhouse after having been considered fit for slaughter for human consumption following an ante-mortem inspection in accordance with Union legislation;]							
			(²) and/or [- animal by-products arising from the production of products intended for human consumption, including degreased bone, greaves and centrifuge or separator sludge from milk processing;]							
			(²) and/or [- products of animal origin, or foodstuffs containing products of animal origin, which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects from which no risk to public or animal health arise;]							
			(2) and/or [- blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk originating from live animals that did not show signs of any disease communicable through that product to humans or animals;]							
			(2) and/or [- aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of diseases communicable to humans or animals;]							
			(2) and/or [- animal by-products from aquatic animals originating from establishments or plants manufacturing products for human 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,										
			<ul><li>eggs,</li><li>egg by-products, including egg shells;</li></ul>							
			(iii) day-old chicks killed for commercial reasons;]							

# COUNTRY

# Processed animal protein not intended for human consumption including mixtures and products other than petfood containing such protein

COUNT	COUNTRY other than petfood containing such protein							
II.	Health inform	nation	II.a. Certificate reference No	II.b.				
	(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) and Category 2 material as referred to in Article 9(a) to (g) of Regulation (EC No 1069/2009;]							
	and							
	(c) has been	subjected to the following processing standard:						
	(2) either	[heating to a core temperature of more than 133 $^{\circ}\text{C}$ least 3 bars produced by saturated steam, with						
	(²) or	[in the case of non-mammalian protein other than out in Annex IV, Chapter III, of Regulation (EU) N		-2-3-4-5-7 as set				
	(²) or	[in the case of fishmeal the processing method 1 Regulation (EU) No 142/2011;]	-2-3-4-5-6-7 as s	set out in Annex IV, Chapter III, of				
	(²) or	[in the case of porcine blood, the processing me Regulation (EU) No 142/2011, where in case of n its substance;]						
II.2.	the competent	authority examined a random sample immediately	prior to dispatch and found it to con	nply with the following standards (3):				
	Salmonella:	Absence in 25 g: $n = 5$ , $c = 0$ , $m = 0$ ,	M = 0					
	Enterobacteria	nceae: $n = 5$ , $c = 2$ , $m = 10$ , $M = 300$ in 1 g;						
II.3.	the end produ	ict:						
	(2) either	[was packed in new or sterilised bags,]						
	(2) or	[was transported in bulk in containers or other mea	ans of transport that were thoroughly	cleaned and disinfected before use,]				
	which bear lab	pels indicating 'NOT FOR HUMAN CONSUMPTION	٧';					
II.4.	the end produ	ct was stored in enclosed storage;						
II.5.	the product ha	as undergone all precautions to avoid recontaminat	ion with pathogenic agents after trea	atment;				
II.6.								
	(²) either	[the product does not contain and is not derived f 999/2001 of the European Parliament and of th bovine, ovine or caprine animals; and the anima stunning by means of gas injected into the crani central nervous tissue by means of an elongated	e Council (4) or mechanically separ als from which this product is derive ial cavity or killed by the same met	ated meat obtained from bones of ed have not been slaughtered after hod or slaughtered by laceration of				
	(²) or	[the product does not contain and is not derived animals born, continuously reared and slaughtere decision in accordance with Article 5(2) of Reguli	d in a country or region classified a					
Notes								
Part I:								
		Person responsible for the consignment in the Euro be filled in if the certificate is for import commodity		in only if it is a certificate for transit				
		Place of destination: this box is to be filled in only ones, free warehouses and custom warehouses.	if it is a certificate for transit commod	dity. The products in transit can only				
		Registration number (railway wagons or container ent of unloading and reloading.	and lorries), flight number (aircraft)	or name (ship); information is to be				
— Вох	reference I.19:	use the appropriate HS code: 05.05; 05.06; 05.07	or 23.01.					
— Вох	reference I.25:	technical use: any use other than for animal const	umption.					
— Вох	reference I.26	and I.27: fill in according to whether it is a transit	or an import certificate.					

cou	JNTRY	Processed animal protein not other than petfood containing	intended for human consumption such protein	including mixtures and products
II.	Health information		II.a. Certificate reference No	II.b.
Par	t II:			
(1a)	OJ L 300, 14.11.2009, p. 1.			
(1b)	OJ L 54, 26.2.2011, p. 1.			
( <sup>2</sup> )	Delete as appropriate.			
(3)	Where:			
	n = number of samples to be	tested;		
	m = threshold value for the nu m;	mber of bacteria; the result is consid	lered satisfactory if the number of bac	teria in all samples does not exceed
	M = maximum value for the nu or more; and	umber of bacteria; the result is consid	lered unsatisfactory if the number of b	pacteria in one or more samples is N
	c = number of samples the bi		een m and M, the sample still being o	considered acceptable if the bacteria
(4)	OJ L 147, 31.5.2001, p. 1.			
<u> </u> –	The signature and the stamp m	ust be in a different colour to that of	the printing.	
		le for the consignment in the Europ till it reaches the border inspection p	pean Union: This certificate is only ost.	for veterinary purposes and has to
Offi	icial veterinarian/Official inspecto	or		
	Name (in capital letters):		Qualification	and title:
	Date:		Signature:	
	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  $\binom{2}{2}$  the European Union

cou	COUNTRY						Veterinary certific	ate to EU
	l.1.	5		I.2. Certifi	icate reference No	)	I.2.a.	
		Name		10 0	-1			
		Addiese			al competent auth	ority		
					competent author	rity		
<u>+</u>	1.5.	Consignee		I.6. Perso	n responsible for	the load	in EU	
l e		Name		Name				
l iĝ		Address		Addre	ess			
Š								
D D		Postcode		Posto	ode			
tc.		Tel.		Tel.				
of dispatched consignment	1.7.	Country of origin ISO code I.8. Region of c	origin Code	I.9. Count		) code	I.10. Region of destination	Code
<u>s</u>								
Part I: Details	l.11.	Place of origin		I.12. Place	of destination			
불		Name Approval number	,	Name	•		Custom warehouse	
۳.		Address		Addre	ess		Approval number	
		Name Approval number Address			<b>-</b>			
		Name Approval number Address		Postcode				
	l.13.	Place of loading		I.14. Date of departure				
	l.15.	I.15. Means of transport			BIP in EU			
		Aeroplane Ship Railwa	y wagon 🗆	I.17. Number(s) of CITES				
		Road vehicle Other	,					
		Identification						
		Documentation references						
	I.18.	Description of commodity		I.19. Commodity code (HS code)				
						1.20. Qu	uantity	
	I.21.	Temperature of product				1.22. Nu	imber of packages	
		Ambient Chilled Chilled		Frozen				
	1.23.	Seal/Container No				1.24. Ty	rpe of packaging	
	1.25.	Commodities certified for:						
			her process		Technical			
	1.26.	For transit through EU to third country		I.27. For im	port or admission	into EU		
		Third country ISO code						
	1.28.	Identification of the commodities						
		Approvi	al number of estab Manufacturing pla		Ne	et weight	Batch r	number

## COUNTRY

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

	II.	Health information			II.a. Certificate reference No	II.b.				
		and of the Chapter II,	I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council ( <sup>1</sup> a), and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011 ( <sup>1</sup> b), and in particular Annex X, Chapter II, Section 4 and Annex XIV, Chapter I thereto, and certify that the milk ( <sup>2</sup> ), the milk-based products ( <sup>2</sup> ) and milk-derived products ( <sup>2</sup> ) referred to in box I.28 comply with the following conditions:							
Part II: Certification	II.1.	they were produced and derived in								
Part II: Ce	II.2.	they were produced from raw milk derived from animals which at the time of milking did not show clinical signs of any disease transmiss through milk to humans or animals, and which had been kept for at least 30 days prior to production on holdings that were not subject official restrictions due to foot-and-mouth disease or rinderpest;								
	II.3.	they are m	ey are milk or milk products that:							
		(2) either	[have unde	rgone one of the treatments or combination	ations thereof described in point II.4;]					
		(²) or	(2) or [comprise whey to be fed to animals of species susceptible to foot-and-mouth disease, and that whey was collected from milk subjected to one of the treatments described in point II.4 and:							
			(2) either	[the whey was collected at least 16 l	hours after clotting and has a pH belo	w 6;]				
			(²)(4) or	[the whey has been produced at lea have been detected in the exporting	st 21 days before the shipping and d country;]	uring that period no cases of FMD				
			(²)(4) or		/, this date, in consideration of the nt is presented to a border inspection					
	II.4.	they have t	oeen subject	to one of the following treatments:						
		(2) either		perature Short Time pasteurisation at 7: action to a phosphatase test in bovine		quivalent pasteurisation achieving a				
			(2) either		ature Short Time pasteurisation at 72 achieves a negative reaction to a pho					
			(²) or	[a subsequent drying process that in t 72 °C or higher;]	the case of milk intended for feeding is	combined with additional heating to				
			(²) or	[a subsequent process by which the	pH is reduced and kept for at least of	ne hour at a level below 6;]				
			(²)(⁴) or	[the condition that the milk/milk produ- period no cases of FMD have been	ct has been produced at least 21 days detected in the exporting country;]	before the shipping and during that				
			(²)(4) or		oduced on//,, this date, in corefore the consignment is presented t					
			(²) or	[sterilisation at a level of at least F <sub>0</sub> 3	5:]]					
		(²) or	[Ultra High	Temperature treatment at 132 °C for all	t least one second in combination with	n:				
			the case of milk intended for feeding is	combined with additional heating to						
			(²) or	[a subsequent process by which the	pH is reduced and kept for at least of	ne hour at a level below 6;]				
			(²)(4) or	[the condition that the milk/milk produ- period no cases of FMD has been d	ct has been produced at least 21 days letected in the exporting country;]	before the shipping and during that				
			(²)(4) or		oduced on//, this date, in corefore the consignment is presented t					

## COUNTRY

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

II.	Health inf	ormation	II.a. Certificate reference No	II.b.					
II.5.	every precaution was taken to avoid contamination of the milk/milk-based product/milk-derived product after processing;								
II.6.	the milk/mi	milk/milk-based product/milk-derived product was packed: either [in new containers:]							
	(²) or	[in vehicles or bulk containers disinfected prior to loading using a product approved by the competent authority:]							
	and	the containers are marked so as to indicate the nature of the milk/milk-based product/milk-derived product and bear labels indicating that the product is Category 3 material and not intended for human consumption;							
II.7.									
	(²) either	[the product does not contain and is not derived 999/2001 of the European Parliament and of the C ovine or caprine animals; and the animals from w means of gas injected into the cranial cavity or k tissue by means of an elongated rod-shaped inst	Council ( <sup>5</sup> ) or mechanically separated r rhich this product is derived have not illed by the same method or slaughte	neat obtained from bones of bovine, been slaughtered after stunning by red by laceration of central nervous					
	(²) or	[the product does not contain and is not deriver animals born, continuously reared and slaughtere decision in accordance with Article 5(2) of Regula	ed in a country or region classified as						
II.8.	in addition	as regards TSE:							
	(²) either	[in case of animal by-products intended for feeding the ovine and caprine animals from which these p three years on a holding where no official moveme the following requirements for the last three years	roducts are derived have been kept of int restriction is imposed due to a susp	ontinuously since birth or for the last					
		(i) it has been subject to regular official veterinal	ry checks;						
		(ii) no classical scrapie case, as defined in point following the confirmation of a classical scrap		999/2001, has been diagnosed or,					
		- all animals in which classical scrapie was	confirmed have been killed and destr	oyed, and					
		<ul> <li>all goats and sheep on the holding have genotype and breeding ewes carrying at le</li> </ul>							
		(iii) ovine and caprine animals, with the exception only if they come from a holding which comp							
	(²) or	[in case of animal by-products intended for feeding and destined to a Member State listed in the Ann animals from which these products are derived holding where no official movement restriction is requirements for the last seven years:	ex to Commission Regulation (EC) No nave been kept continuously since bir	546/2006 (6), the ovine and caprine th or for the last seven years on a					
		(i) it has been subject to regular official veterinal	ry checks;						
		(ii) no classical scrapie case, as defined in point following the confirmation of a classical scrap		999/2001, has been diagnosed or,					
		- all animals in which classical scrapie was	confirmed have been killed and destr	oyed, and					
		<ul> <li>all goats and sheep on the holding have genotype and breeding ewes carrying at le</li> </ul>							
		(iii) ovine and caprine animals, with the exception only if they come from a holding which comp							
Notes	;								

### Part I:

- Box reference I.6: Person responsible for the load in the European Union: this box is to be filled in only if it is a certificate for transit commodity.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity.

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

COUNTRY

Status: This is the original version (as it was originally adopted).

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) is to be provided. In 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) cod 35.04.	de of the World Customs Organisation:	23.09.10, 23.09.90, 35.01, 35.02 or				
-	Box reference I.23: for bulk containers, the container number and the	e seal number (if applicable) must be	included.				
-	Box reference I.25: technical use: any use other than for animal cor	sumption.					
_	Box reference I.26 and I.27: fill in according to whether it is a transi	t or an import certificate.					
_	Box reference I.28: 'Manufacturing plant': provide the registration nu	mber of treatment or processing estab	lishment.				
Par	t 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 the European Unio	on is restricted to certain regions of the	e third country concerned.				
( <sup>4</sup> )	this condition applies only to third countries listed in column 'A' of	Annex I to Regulation (EU) No 605/20	10.				
( <sup>5</sup> )	OJ L 147, 31.5.2001, p. 1.						
( <sub>6</sub> )	OJ L 94, 1.4.2006, p. 28.						
— <sup>1</sup>	The signature and the seal must be in a different colour from that o	f the printing.					
	Note for the importer: this certificate is only for veterinary purposes a post of the European Union.	nd must accompany the consignment u	ntil it reaches the border inspection				
Offi	cial veterinarian/Official inspector						
	Name (in capital letters):	Qualification	and title:				
	Date:	Signature:					
	Stamp:						

# CHAPTER 2(B)

# Health certificate

For colostrum and colostrum products from bovine animals not intended for human consumption for dispatch to or transit through  $\binom{2}{2}$  the European Union

cou	COUNTRY							Veterinary certific	ate to EU
I.1. Consignor I.2.			1.2.	Certificat	e reference No	)	I.2.a.		
		Name			I.3. Central competent authority				
		Address		1.3.	Central	ompetent auth	ority		
		Tel.			Local co	mpetent author	ity		
¥	1.5.	Consignee	1.6.	Person r	esponsible for	the load	in EU		
me		Name		Name					
sign		Address			Address				
S		Destanda							
ped		Postcode Tel.			Postcode Tel.	,			
atcl									
disp	1.7.	Country of origin ISO code	I.8. Region of origin Code	1.9.	Country		code	I.10. Region of destination	Code
ğ		1			desinali			destination	
Part I: Details of dispatched consignment	111	Place of origin		112	Place of	destination			
ë.	''''	Tidoo or origin		1.12.	riace of	destination			
ar			Approval number		Name			Custom warehouse	
ď		Address Name	Approval number		Address			Approval number	
		Address	Approvai number		Postcode				
		Name Address	Approval number						
				I.14. Date of departure					
	interviews of reasoning								
	l.15.	Means of transport		I.16.	Entry BIF	in EU			
		Aeroplane Ship	Railway wagon						
		Road vehicle Other	]	I.17. Number(s) of CITES					
		Identification							
		Documentation references							
	I.18.	Description of commodity		I.19. Commodity code (HS code)					
							100 0	unatib.	
							1.20. Q	uanity	
	1.21.	Temperature of product					1.22. Nu	umber of packages	
		Ambient	Chilled	Froze	n 🗆				
	1.23.	Seal/Container No					I.24. Ty	pe of packaging	
	125	Commodities certified for:							
	1.23.								
		Animal feedingstuff	Further process	_				al use 🗆	
	1.26.	For transit through EU to third co	ountry	1.27.	For impo	rt or admission	into EU		
		Third country	SO code						
	1.28.	Identification of the commodities							
			number of establishments		Net weigh	nt		Batch number	
	(	Scientific name) Ma	anufacturing plant						

## COUNTRY

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

II	ı.	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 Parliamer and of the Council ( <sup>1a</sup> ), and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011 ( <sup>1b</sup> ), and in particular Annex X Chapter II, Section 4 and Annex XIV, Chapter I thereto, and certify that the colostrum ( <sup>2</sup> ) or the colostrum products ( <sup>2</sup> ) referred to in box I.2 comply with the following conditions:							
Part II: Certification	l.1.	they were produced and derived in							
Part II: C	1.2.	transmissible	they were produced from colostrum derived from animals which at the time of milking did not show clinical signs of any disease transmissible through colostrum to humans or animals, and which had been kept for at least 30 days prior to production on holdings that were not subject to official restrictions due to foot-and-mouth disease or rinderpest;						
ı	1.3.		they are colostrum or colostrum products of bovine animals that have been subject to High Temperature Short Time pasteurisation at 72 °C for at least 15 seconds, or an equivalent pasteurisation achieving a negative reaction to a phosphatase test in bovine milk, in combination with:						
		(2)(4) either		on that the colostrum or colostrum produces of FMD have been detected in the	lucts have been produced at least 21 cne exporting country;]	days before the shipping and in this			
		( <sup>2</sup> )( <sup>4</sup> ) or			produced on//, this date, in cognment is presented to a border inspe				
		and have been obtained from animals subject to regular veterinary inspections to ensure that they come from holdings on which all bovine herds are:							
			(2)(4) either	[recognised as officially tuberculosis	and brucellosis free (5),]				
			(²)(4) or	[not restricted under the national leg and brucellosis,]	islation of the third country of origin re	egarding eradication of tuberculosis			
		and	(2)(4) either	[recognised as official enzootic-bovin	e-leukosis free (5);]				
			(²)(4) or		e control of enzootic bovine leukosis a ag of this disease in the herd during th				
lı.	1.4.	every preca	ution was tak	en to avoid contamination of the colos	strum/colostrum product after processing	ng;			
lı.	1.5.	the colostru	m/colostrum p	product was packed:					
		(2) either	[in new cont	tainers,]					
		(2) or	[in vehicles	or bulk containers disinfected prior to	loading using a product approved by	the competent authority,]			
		and		ers are marked so as to indicate the na category 3 material and not intended for	uture of the colostrum/colostrum productor human consumption;	t and bear labels indicating that the			
l I	l.6.								
		(2) either  [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC No 999/2001 of the European Parliament and of the Council (6) or mechanically separated meat obtained from bones of bovine, ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunnin by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervou tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity.]							
		(²) or [the product does not contain and is not derived from bovine, ovine or 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 by a decision in accordance with Article 5(2) of Regulation (EC) No 999/2001;]							
II	1.7.	in addition a	as regards TS	EE:					
		(²) either							
			(i) it has be	een subject to regular official veterinar	y checks;				
L									

#### COUNTRY

(2) or

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

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

- (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or, following the confirmation of a classical scrapie case:
  - all animals in which classical scrapie was confirmed have been killed and destroyed, and
  - all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARR genotype and breeding ewes carrying at least one ARR allele and no VRQ allele;
- (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are introduced into the holding only if they come from a holding which complies with the requirements set out in points (i) and (ii).]
- [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin, and destined to a Member State listed in the Annex to Commission Regulation (EC) No 546/2006 (?), the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last seven years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfied the following requirements for the last seven years:
  - (i) it has been subject to regular official veterinary checks;
  - (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or, following the confirmation of a classical scrapie case:
    - all animals in which classical scrapie was confirmed have been killed and destroyed, and
    - all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARR genotype and breeding ewes carrying at least one ARR allele and no VRQ allele;
  - (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are introduced into the holding only if they come from a holding which complies with the requirements set out in points (i) and (ii).]

### Notes

### Part I:

- Box reference I.6: Person responsible for the load in the European Union: this box is to be filled in only if it is a certificate for transit commodity.
- 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
  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: 23.09.10, 23.09.90, 35.01, 35.02 or 35.04.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) must be included.
- Box reference I.25: technical use: any use other than for animal consumption.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: 'Manufacturing plant': provide the registration number of treatment or processing establishment.

### Part II:

- (1a) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.
- (9) For completion if the authorisation to import into the European Union is restricted to certain regions of the third country concerned.
- (4) this condition applies only to third countries listed in column 'A' of Annex I to Commission Regulation (EU) No 605/2010.
- (5) Officially tuberculosis and brucellosis free herd as laid down in Annex A to Council Directive 64/432/EEC; and officially enzootic-bovine-leukosis free herd as laid down in Chapter I of Annex D to Council Directive 64/432/EEC.
- (6) OJ L 147, 31.5.2001, p. 1.
- (7) OJ L 94, 1.4.2006, p. 28.

COUN	TRY Colostrum and	colostrum products from bovine animals not for human consumption							
II.	Health information	II.a. Certificate reference No	II.b.						
_ No	<ul> <li>The signature and the seal must be in a different colour from that of the printing.</li> <li>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.</li> </ul>								
Officia	al veterinarian/Official inspector								
	lame (in capital letters):	Qualification	and title:						
	Date:	Signature:							
s	stamp:								

# CHAPTER 3(A)

# **Health certificate**

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

COUNTRY Veterinary certificate					
	l.1.	Consignor	I.2. Certificate reference No I.2.a.		
		Name	LO Combrel commentant sufficients		
		Address	I.3. Central competent authority		
		Tel.	I.4. Local competent authority		
뉱	1.5.	Consignee	I.6. Person responsible for the load in EU		
l me		Name	Name		
sign		Address	Address		
2		Postcode	Postcode		
ped		Tel.	Tel.		
of dispatched consignment					
disp	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10. Region of Code destination		
6			destination destination		
Part I: Details	111	Place of origin	I.12. Place of destination		
e .		That of origin	1.12. Place of destination		
Έ		Name Approval number	Name Custom warehouse		
ية ا		Address Name Approval number	Address Approval number		
		Address	Postcode		
		Name Approval number Address	roscode		
	l.13.	Place of loading	I.14. Date of departure		
	l.15.	Means of transport	I.16. Entry BIP in EU		
		Aeroplane Ship Railway wagon			
		Road vehicle Other	1.17.		
		Identification			
		Documentation references			
	l.18.	Description of commodity	I.19. Commodity code (HS code) 23.09.10		
			I.20. Quantity		
	1.21.	Temperature of product	I.22. Number of packages		
		Ambient Chilled	Frozen		
	1.23.	Seal/Container No	I.24. Type of packaging		
	1.25.	Commodities certified for:	-		
		Animal feedingstuff ☐ Technical use ☐			
	1.26.	For transit through EU to third country	I.27. For import or admission into EU		
		Third country ISO code			
	1.28.	Identification of the commodities			
		Species Approval number of establishme (Scientific name) Manufacturing plant	ents Net weight Batch number		

COUNTRY Canned Petfood

_		···						
	II.	Health inf	ormation	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 Pa and of the Council ( <sup>1a</sup> ) and in particular Articles 8 and 10 thereof, and Commission Regulation (EU) No 142/2011 ( <sup>1b</sup> ), and in p Annex XIII, Chapter II and Annex XIV, Chapter II, thereof and certify that the petfood described above:						
Part II: Certification	II.1.	II.1. has been prepared and stored in an establishment or plant approved and supervised by the competent authority in accordance with 24 of Regulation (EC) No 1069/2009;						
: Certif	II.2.	II.2. has been prepared exclusively with the following animal by-products:						
Part II		(2) either	[- carcases and parts of animals slaughtered or, in human consumption in accordance with Union reasons;]					
		(²) and/or [- carcases and the following parts originating either from animals that have been slaughtered in a slaughterhouse and we considered fit for slaughter for human consumption following an ante-mortem inspection or bodies and the following parts animals from game killed for human consumption in accordance with Union legislation:						
			<ul><li>(i) carcases or bodies and parts of animals white legislation, but which did not show any signs</li></ul>					
		(ii) heads of poultry;						
	<ul><li>(iii) hides and skins, including trimmings and splitting thereof, horns and feet, including the phalanges and the carpus a metacarpus bones, tarsus and metatarsus bones, of animals, other than ruminants;</li></ul>							
			(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, obtain animals other than ruminants that have been slaughtered in a slaughterhouse after having been considered fit for sl 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 greaves and centrifuge or separator sludge from milk processing;]							
		<ul> <li>(²) and/or [- products of animal origin, or foodstuffs containing products of animal origin, which are no longer intended for consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defect which no risk to public or animal health arise;]</li> <li>(²) and/or [- petfood and feedingstuffs of animal origin, or feedingstuffs containing animal by-products or derived products, which longer intended for feeding for commercial reasons or due to problems of manufacturing or packaging defects or defects from which no risk to public or animal health arises;]</li> </ul>						
	(2) and/or [- blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk originating from live animals that did not any disease communicable through that product to humans or animals;]				animals that did not show signs of			
	(2) and/or [- aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of diseases commun to humans or animals;]  (2) and/or [- animal by-products from aquatic animals originating from plants or establishments manufacturing products for hononumption;]							
		(²) and/or	[- the following material originating from animals wh to humans or animals:	ich did not show any signs of disease	communicable through that material			
			(i) shells from shellfish with soft tissue or flesh;					
			(ii) the following originating from terrestrial anima	als:				
			<ul><li>hatchery by-products,</li></ul>					
			— eggs,					
			<ul> <li>egg by-products, including egg shells;</li> </ul>					
			(iii) day-old chicks killed for commercial reasons;	:]				

COUNTRY	Canned Petfood

I.a. Certificate reference No					***************************************		
(°) and/or [- material from animals which have been treated with certain substances which are prohibited pur 98/22/EC. the import of the material being permitted in accordance with Article 35(a)(ii) of No 1089/2009s.]  II.3. has been subjected to heat treatment to a minimum Fo value of 3 in hermetically sealed containers;  III.4. was analysed by a random sampling of at least five containers from each processed batch by laboratory diagnostic or adequate heat treatment of the whole consignment as foreseen under point III.3;  III.5. has undergone all precautions to avoid contamination with pathogenic agents after treatment.  III.6. (°) either III.6. (°) e	II.	Health informat	ion	II.a. Certificate reference No	II.b.		
96/22/EC, the import of the material being permitted in accordance with Article 35(a)(ii) of No 1098/20091.  II.3. has been subjected to heat treatment to a minimum Fo value of 3 in hermetically sealed containers;  III.4. was analyzed by a random sampling of at least five containers from each processed batch by laboratory diagnostic or adequate heat treatment of the whole consignment as foreseen under point II.3;  III.5. has undergone all precautions to avoid contamination with pathogenic agents after treatment.  III.6.  (**) either III.6 product does not contain and is not derived from specified risk material as defined in Annax V to 999/2001 of the European Parliament and of the Council (**) or mechanically separated meat obtain boving, ovine or caprire animals; and the animals from which this product is derived have not been stunning by means of gas injected into the cranial cavity or field by the same method or slaughtere central nervous issue by means of an elongated rod-shaped instrument introduced into the cranial (**) or III. The product does not contain and is not derived from bovine, ovine or caprine materials other than at animals born, continuously reared and salughtered in a country or region classified as posing a neglig decision in accordance with Article 5(2) of Regulation (EC) No 999/2001;]  III.7. in addition as regards TSE:  (**) either III nease of animal by-products intended for feeding ruminants and containing milk or milk products origin, the ovine and caprine animals from which these products are derived have been kept continuor for the last three years on a holding where no official movement restriction is imposed due to a susy which has satisfied the following requirements for the last three years:  (i) It has been subject to regular official veterinary checks;  (ii) no classical scrapic case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, ha or, following the confirmation of a classical scrapie was confirmed have been kelled and destroyed, and genotype and an anim		(²) and/or [	- animal by-products from aquatic or terrestrial	invertebrates other than species path	ogenic to humans or animals;]		
II.4. was analysed by a random sampling of at least five containers from each processed batch by laboratory diagnostic radequate heat treatment of the whole consignment as foreseen under point II.3;  II.5. has undergone all precautions to avoid contamination with pathogenic agents after treatment.  II.6.  (**) either		( <sup>2</sup> ) and/or [	96/22/EC, the import of the material being				
adequate heat freatment of the whole consignment as foreseen under point II.3;  II.5. has undergone all precautions to avoid contamination with pathogenic agents after treatment.  II.6.  (**) either**  (**) the product does not contain and is not derived from specified risk material as defined in Annex V to No 999/2001 of the European Parliament and of the Council (**) or mechanically separated meat obtabovine, ovine or caprine animals; and the animals from which this product is derived have not been sturning by means of gas injected into the cranial cavity or killed by the same method or slauphtere central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity or region classified as posing a neglig decision in accordance with Article 5(2) of Regulation (EC) No 999/2001;  II.7. in addition as regards TSE:  (**) either**  (**) either**  (**) case of animal by-products intended for feeding ruminants and containing milk or milk products or origin, the ovine and caprine animals from which these products are derived have been kept continuo for the last three years on a holding where no official movement restriction is imposed due to a susy which has statisted the following requirements for the last three years:  (i) it has been subject to regular official veterinary checks;  (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, ha or, following the confirmation of a classical scrapie case:  — all animals in which classical scrapie was confirmed have been killed and destroyed, and  — all goats and sheep on the holding have been killed and destroyed, except for breeding rams genotype and breeding ewes carrying at least one ARR allele and no VRO allele,  (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are in holding only if they come from a holding which complies with the requirements set out in points and caprine animals from which these products are derived have been kept continuously s	II.3.	has been su	bjected to heat treatment to a minimum Fc value	ue of 3 in hermetically sealed contain	ers;		
III.6.  (**) either Roduct does not contain and is not derived from specified risk material as defined in Annex V t No 999/2001 of the European Parliament and of the Council (**) or mechanically separated meat obtain bovine, ovine or caprine animals; and the animals from which this product is derived have not been stunning by means of gas injected into the crainal cavity or killed by the same method or slaughtere central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial of the product does not contain and is not derived from bovine, ovine or caprine materials other than the animals born, continuously reared and slaughtered in a country or region classified as posing a neglig decision in accordance with Article 5(2) of Regulation (EC) No 999/2001:]  III.7. In addition as regards TSE:  (**) either (In case of animal by-products intended for feeding ruminants and containing milk or milk products or origin, the ovine and caprine animals from which these products are derived have been kept continuo for the last three years on a holding where no official movement restriction is imposed due to a susy which has satisfied the following requirements for the last three years:  (i) it has been subject to regular official veterinary checks;  (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, ha or, following the confirmation of a classical scrapie case:  — all animals in which classical scrapie was confirmed have been killed and destroyed, and  — all goats and sheep on the holding have been killed and destroyed, except for breeding rams genotype and breeding ewes carrying at least one ARR allele and no VRQ allele,  (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are in holding only if they come from a holding which complies with the requirements set out in points and caprine animals from which these products are derived have been ket ontentions (EC) No 548/2 and caprine animals from which thes	II.4.						
(a) either Ithe product does not contain and is not derived from specified risk material as defined in Annax V No 999/2001 of the European Parliament and of the Council (b) or mechanically separated meat obtain bovine, owine or caprine animals; and the animals from which this product is derived have not been stuming by means of gas injected into the cranial cavity or killed by the same method or slaughters central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial of the cranial cavity or region classified as posing a neglig decision in accordance with Article 5(2) of Regulation (EC) No 999/2001;]  III.7. In addition as regards TSE:  (a) either In case of animal by-products intended for feeding ruminants and containing milk or milk products or origin, the ovine and caprine animals from which these products are derived have been kept continuo for the last three years on a holding where no official movement restriction is imposed due to a susy which has satisfied the following requirements for the last three years:  (i) It has been subject to regular official veterinary checks;  (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, ha or, following the confirmation of a classical scrapie case:  — all animals in which classical scrapie was confirmed have been killed and destroyed, and  — all goats and sheep on the holding have been killed and destroyed, except for breeding rams genotype and breeding ewes carrying at least one ARR allele and no VRQ allele,  (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are in holding only if they come from a holding which complies with the requirements set out in points origin, and destined to a Member State Isled in the Annax to Commission Regulation (EC) No 546/2 and caprine animals from which these products are derived have been kept continued or years on a holding where no official veterinary checks;  (i) It has been subject to regular official vete	II.5.	has undergone all precautions to avoid contamination with pathogenic agents after treatment.					
No 999/2001 of the European Parliament and of the Council (*) or mechanically separated meat obtain bovine, ovine or caprine animals; and the animals from which this product is derived have not been stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtere central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity or killed by the same method or slaughtere central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity or region classified as posing a neglig decision in accordance with Article 5(2) of Regulation (EC) No 999/2001;]  II.7. in addition as regards TSE:  (*) either  [in case of animal by-products intended for feeding ruminants and containing milk or milk products or origin, the ovine and caprine animals from which these products are derived have been kept continue for the last three years on a holding where no official movement restriction is imposed due to a susy which has satisfied the following requirements for the last three years:  (i) it has been subject to regular official veterinary checks;  (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, ha or, following the confirmation of a classical scrapie was confirmed have been killed and destroyed, and  — all goats and sheep on the holding have been killed and destroyed, except for breeding rams genotype and breeding ewes carrying at least one ARR allele and no VRQ allele,  (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are in holding only if they come from a holding which complies with the requirements set out in points origin, and destined to a Member State listed in the Annex to Commission Regulation (EC) No 546/2 and caprine animals from which these products are derived have been kept continuously since birth or years on a holding where no official movement restriction is imposed due to a suspicion of TSE and where followin	II.6.						
animals born, continuously reared and slaughtered in a country or region classified as posing a neglig decision in accordance with Article 5(2) of Regulation (EC) No 999/2001;]  II.7. In addition as regards TSE:  (**) either		( <sup>2</sup> ) either	No 999/2001 of the European Parliament and bovine, ovine or caprine animals; and the an stunning by means of gas injected into the co	of the Council (3) or mechanically sep imals from which this product is deriv ranial cavity or killed by the same me	arated meat obtained from bones of ed have not been slaughtered after thod or slaughtered by laceration of		
(c) either  [in case of animal by-products intended for feeding ruminants and containing milk or milk products or origin, the ovine and caprine animals from which these products are derived have been kept continuor for the last three years on a holding where no official movement restriction is imposed due to a susy which has satisfied the following requirements for the last three years:  (i) it has been subject to regular official veterinary checks;  (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, ha or, following the confirmation of a classical scrapie case:  — all animals in which classical scrapie was confirmed have been killed and destroyed, and  — all goats and sheep on the holding have been killed and destroyed, except for breeding rams genotype and breeding ewes carrying at least one ARR allele and no VRQ allele,  (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are in holding only if they come from a holding which complies with the requirements set out in points  (c) or  [in case of animal by-products intended for feeding ruminants and containing milk or milk products or origin, and destined to a Member State listed in the Annex to Commission Regulation (EC) No 548/2 and caprine animals from which these products are derived have been kept continuously since birth or years on a holding where no official movement restriction is imposed due to a suspicion of TSE and we the following requirements for the last seven years:  (i) it has been subject to regular official veterinary checks;  (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, ha or, following the confirmation of a classical scrapie case:  — all animals in which classical scrapie was confirmed have been killed and destroyed, and  — all goats and sheep on the holding have been killed and destroyed, except for breeding rams.		( <sup>2</sup> ) or	animals born, continuously reared and slaught	tered in a country or region classified			
origin, the ovine and caprine animals from which these products are derived have been kept continuo for the last three years on a holding where no official movement restriction is imposed due to a susy which has satisfied the following requirements for the last three years:  (i) it has been subject to regular official veterinary checks;  (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, ha or, following the confirmation of a classical scrapie case:  — all animals in which classical scrapie was confirmed have been killed and destroyed, and  — all goats and sheep on the holding have been killed and destroyed, except for breeding rams genotype and breeding ewes carrying at least one ARR allele and no VRQ allele,  (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are in holding only if they come from a holding which complies with the requirements set out in points  (2) or [in case of animal by-products intended for feeding ruminants and containing milk or milk products or origin, and destined to a Member State listed in the Annex to Commission Regulation (EC) No 546/2 and caprine animals from which these products are derived have been kept continuously since birth or years on a holding where no official movement restriction is imposed due to a suspicion of TSE and we the following requirements for the last seven years:  (i) it has been subject to regular official veterinary checks;  (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, ha or, following the confirmation of a classical scrapie case:  — all animals in which classical scrapie was confirmed have been killed and destroyed, and  — all goats and sheep on the holding have been killed and destroyed, except for breeding rams.	II.7.	in addition a	s regards TSE:				
<ul> <li>(ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, ha or, following the confirmation of a classical scrapie case:  — all animals in which classical scrapie was confirmed have been killed and destroyed, and  — all goats and sheep on the holding have been killed and destroyed, except for breeding rams genotype and breeding ewes carrying at least one ARR allele and no VRQ allele,</li> <li>(iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are in holding only if they come from a holding which complies with the requirements set out in points</li> <li>(²) or [in case of animal by-products intended for feeding ruminants and containing milk or milk products or origin, and destined to a Member State listed in the Annex to Commission Regulation (EC) No 546/2 and caprine animals from which these products are derived have been kept continuously since birth or years on a holding where no official movement restriction is imposed due to a suspicion of TSE and we the following requirements for the last seven years:  (i) it has been subject to regular official veterinary checks;  (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, ha or, following the confirmation of a classical scrapie case:  — all animals in which classical scrapie was confirmed have been killed and destroyed, and  — all goats and sheep on the holding have been killed and destroyed, except for breeding rams.</li> </ul>		(²) either	origin, the ovine and caprine animals from wh for the last three years on a holding where no	ich these products are derived have boofficial movement restriction is impo	een kept continuously since birth or		
or, following the confirmation of a classical scrapie case:  — all animals in which classical scrapie was confirmed have been killed and destroyed, and  — all goats and sheep on the holding have been killed and destroyed, except for breeding rams genotype and breeding ewes carrying at least one ARR allele and no VRQ allele,  (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are in holding only if they come from a holding which complies with the requirements set out in points  (²) or [in case of animal by-products intended for feeding ruminants and containing milk or milk products or derived have been kept continuously since birth or years on a holding where no official movement restriction is imposed due to a suspicion of TSE and we the following requirements for the last seven years:  (i) it has been subject to regular official veterinary checks;  (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, ha or, following the confirmation of a classical scrapie case:  — all animals in which classical scrapie was confirmed have been killed and destroyed, and  — all goats and sheep on the holding have been killed and destroyed, except for breeding rams.			(i) it has been subject to regular official vete	erinary checks;			
— all goats and sheep on the holding have been killed and destroyed, except for breeding rams genotype and breeding ewes carrying at least one ARR allele and no VRQ allele,  (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are in holding only if they come from a holding which complies with the requirements set out in points  (2) or  [in case of animal by-products intended for feeding ruminants and containing milk or milk products or origin, and destined to a Member State listed in the Annex to Commission Regulation (EC) No 546/2 and caprine animals from which these products are derived have been kept continuously since birth or years on a holding where no official movement restriction is imposed due to a suspicion of TSE and we the following requirements for the last seven years:  (i) it has been subject to regular official veterinary checks;  (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has or, following the confirmation of a classical scrapie case:  — all animals in which classical scrapie was confirmed have been killed and destroyed, and  — all goats and sheep on the holding have been killed and destroyed, except for breeding rams.					No 999/2001, has been diagnosed		
genotype and breeding ewes carrying at least one ARR allele and no VRQ allele,  (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are in holding only if they come from a holding which complies with the requirements set out in points  (2) or [in case of animal by-products intended for feeding ruminants and containing milk or milk products or origin, and destined to a Member State listed in the Annex to Commission Regulation (EC) No 546/2 and caprine animals from which these products are derived have been kept continuously since birth or years on a holding where no official movement restriction is imposed due to a suspicion of TSE and withe following requirements for the last seven years:  (i) it has been subject to regular official veterinary checks;  (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has or, following the confirmation of a classical scrapie case:  — all animals in which classical scrapie was confirmed have been killed and destroyed, and  — all goats and sheep on the holding have been killed and destroyed, except for breeding rams			— all animals in which classical scrapie	was confirmed have been killed and o	destroyed, and		
holding only if they come from a holding which complies with the requirements set out in points  (2) or  [in case of animal by-products intended for feeding ruminants and containing milk or milk products or origin, and destined to a Member State listed in the Annex to Commission Regulation (EC) No 546/2 and caprine animals from which these products are derived have been kept continuously since birth or years on a holding where no official movement restriction is imposed due to a suspicion of TSE and with the following requirements for the last seven years:  (i) it has been subject to regular official veterinary checks;  (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has or, following the confirmation of a classical scrapie case:  — all animals in which classical scrapie was confirmed have been killed and destroyed, and  — all goats and sheep on the holding have been killed and destroyed, except for breeding rams.							
origin, and destined to a Member State listed in the Annex to Commission Regulation (EC) No 548/2 and caprine animals from which these products are derived have been kept continuously since birth or years on a holding where no official movement restriction is imposed due to a suspicion of TSE and with the following requirements for the last seven years:  (i) it has been subject to regular official veterinary checks;  (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has or, following the confirmation of a classical scrapie case:  — all animals in which classical scrapie was confirmed have been killed and destroyed, and  — all goats and sheep on the holding have been killed and destroyed, except for breeding rams							
<ul> <li>(ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, ha or, following the confirmation of a classical scrapie case:</li> <li>— all animals in which classical scrapie was confirmed have been killed and destroyed, and</li> <li>— all goats and sheep on the holding have been killed and destroyed, except for breeding rams</li> </ul>		( <sup>2</sup> ) or	origin, and destined to a Member State listed and caprine animals from which these product years on a holding where no official movemen	in the Annex to Commission Regulat is are derived have been kept continuous at restriction is imposed due to a suspi	ion (EC) No 546/2006 (4), the ovine ously since birth or for the last seven		
or, following the confirmation of a classical scraple case:  — all animals in which classical scraple was confirmed have been killed and destroyed, and  — all goats and sheep on the holding have been killed and destroyed, except for breeding rams			(i) it has been subject to regular official vete	erinary checks;			
<ul> <li>— all goats and sheep on the holding have been killed and destroyed, except for breeding rams</li> </ul>					No 999/2001, has been diagnosed		
			- all animals in which classical scrapie	was confirmed have been killed and o	destroyed, and		
(iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are ir holding only if they come from a holding which complies with the requirements set out in points							

COUNTRY Canned Petfo				
II. Health information	II.a. Certificate reference No	II.b.		
Notes				
Part I:				
Box reference I.6: Person responsible for the consignment in the Eucommodity; it may be filled in if the certificate is for import commod		n only if it is a certificate for transit		
<ul> <li>Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses.</li> </ul>				
Box reference I.15: Registration number (railway wagons or containe provided in the event of unloading and reloading.	er and lorries), flight number (aircraft) o	or name (ship); information is to be		
Box reference I.23: for bulk containers, the container number and	e seal number (if applicable) should b	e given.		
Box reference I.25: technical use: any use other than for animal cor	sumption.			
Box reference I.26 and I.27: fill in according to whether it is a transi	t or an import certificate.			
Part II:				
( <sup>1a)</sup> OJ L 300, 14.11.2009, p. 1. ( <sup>1b)</sup> OJ L 54, 26.2.2011, p. 1.				
(2) Delete as appropriate.				
(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 Eurol accompany the consignment until it reaches the border inspection p		or veterinary purposes and has to		
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  $\binom{2}{2}$  the European Union

cou	DUNTRY Veterinary certificate to EU				
	l.1.	Consignor	I.2. Certificate reference No I.2.a.		
		Name	LO Control compatent authority		
		Address	I.3. Central competent authority		
		Tel.	I.4. Local competent authority		
뉱	1.5.	Consignee	I.6. Person responsible for the load in EU		
l e		Name	Name		
l iĝ		Address	Address		
Š					
b		Postcode Tel.	Postcode Tel.		
dispatched consignment		161.	161.		
ispa	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10. Region of Code		
of d			destination destination		
ils					
Part I: Details	l.11.	Place of origin	I.12. Place of destination		
art I		Name Approval number Address	Name Custom warehouse ☐ Address Approval number		
-		Name Approval number Address	Address Approval number		
		Name Approval number	Postcode		
		Address			
	I.13.	Place of loading	I.14. Date of departure		
	I.15.	Means of transport	I.16. Entry BIP in EU		
		Aeroplane Ship Railway wagon			
		Road vehicle Other I	1.17.		
		Documentation references			
	110	Description of commodity	Lite Comment's made (US and a)		
	1.10.	Description of commonly	I.19. Commodity code (HS code) 23.09.10		
			I.20. Quantity		
			,		
	1.21.	Temperature of product	I.22. Number of packages		
		Ambient Chilled Chilled	Frozen		
	1.23.	Seal/Container No	I.24. Type of packaging		
	1.25.	Commodities certified for:			
		Animal feedingstuff Technical use			
	1.26.	For transit through EU to third country	I.27. For import or admission into EU		
		Third country ISO code			
	1.28.	Identification of the commodities			
		Species Approval number of establishme (Scientific name) Manufacturing plant	nts Net weight Batch number		

## COUNTRY

# Processed petfood other than canned petfood

	II.	Health information	II.a. Certificate reference No	II.b.				
		I, the undersigned official veterinarian, declare that I have real and of the Council ( <sup>Ia</sup> ) and in particular Articles 8 and 10 th Annex XIII, Chapter II and Annex XIV, Chapter II thereof and	ereof, and Commission Regulation (EU)	No 142/2011 (1b), and in particular				
tion	II.1.	has been prepared and stored in a plant approved and supervised by the competent authority in accordance with Article 24 of Regulatio (EC) No 1069/2009;						
II: Certification	II.2.	has been prepared exclusively with the following animal by-products:						
Part II: Ce		(2) either [- carcases and parts of animals slaughtered or, in the case of game, bodies or parts of animals killed, and which are fit human consumption in accordance with Union legislation, but are not intended for human consumption for commercing 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:						
		<ul> <li>(i) carcases or bodies and parts of animals which are rejected as unfit for human consumption in accordance with Unic legislation, but which did not show any signs of disease communicable to humans or animals;</li> </ul>						
	(ii) heads of poultry;							
	(iii) hides and skins, including trimmings and splitting thereof, horns and feet, including the phalanges and the carpus a metacarpus bones, tarsus and metatarsus bones, of animals, other than ruminants;							
	(iv) pig bristles;							
		(v) feathers;]						
	(²) and/or [- blood of animals which did not show any signs of disease communicable through blood to humans or animals, obtained animals other than ruminants that have been slaughtered in a slaughterhouse after having been considered fit for slau 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 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 his consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects which no risk to public or animal health arise;]							
		(²) and/or [- petfood and feedingstuffs of animal origin, or fe longer intended for feeding for commercial re defects from which no risk to public or animal	asons or due to problems of manufactu					
		(²) and/or [- blood, placenta, wool, feathers, hair, horns, ho any disease communicable through that produ		animals that did not show signs of				
		(²) and/or [- aquatic animals, and parts of such animals, exc to humans or animals;]	ept sea mammals, which did not show a	ny signs of diseases communicable				
		(²) and/or [- animal by-products from aquatic animals original consumption;]	inating from plants or establishments i	manufacturing products for human				
		(²) and/or [- the following material originating from animals to humans or animals:	hich did not show any signs of disease	communicable through that material				
		(i) shells from shellfish with soft tissue or fles	h;					
		(ii) the following originating from terrestrial ani	mals:					
		<ul> <li>hatchery by-products,</li> </ul>						
		— eggs,						
		- egg by-products, including egg shells,						
		(iii) day-old chicks killed for commercial reason	s;]					

#### COUNTRY

#### Processed petfood other than canned petfood

COUNTRY				Processed petfood other than canned petfood		
II.	Health inf	ormati	ion II.a. Certificate refere	nce No	II.b.	
	(²) and/or	[- ani	imal by-products from aquatic or terrestrial invertebrates other than sp	pecies pathogen	nic to humans or animals;]	
	(²) and/or		aterial from animals which have been treated with certain substances we import of the material being permitted in accordance with Article 35			
II.3.						
	(2) either	[wa	as subjected to a heat treatment of at least 90 °C throughout its subs	stance;]		
	(²) or	[wa	as produced as regards ingredients of animal origin using exclusively	products which	had been:	
		(a)	) in the case of animal by-products or derived products from meat or least 90 °C throughout its substance;	meat products	subjected to a heat treatment of at	
		(b)	) in the case of milk and milk based products,			
			(i) if they are from third countries or parts of third countries listed (EU) No 605/2010 (³) submitted to a pasteurisation treatment s			
			<ul><li>(ii) with a pH reduced to less than 6 from third countries or parts Decision 2004/438/EC, first submitted to a pasteurisation treatrest;</li></ul>			
			(iii) if they are from third countries or parts of third countries lis No 605/2010, submitted to a sterilisation process or a double in to produce a negative phosphatase test on its own;			
			(iv) if they are from third countries or parts of third countries lis No 605/2010, where there has been an outbreak of foot-and vaccination against foot-and-mouth disease has been carried or	d-mouth disease	in the last 12 months or where	
			either			
			- a sterilisation process whereby an Fc value equal or greater	r than 3 is achie	eved	
			or			
			<ul> <li>an initial heat treatment with a heating effect at least equal least 72 °C for at least 15 seconds and sufficient to produce by</li> </ul>			
			either			
			<ul> <li>a second heat treatment with a heating effect at least equal which would be sufficient to produce a negative reaction to milk, or dried milk-based products by a drying process</li> </ul>			
			or			
			- an acidification process such that the pH has been maintain	ned at less than	6 for at least one hour;	
		(c)	) in the case of gelatine, produced using a process that ensures that treatment with acid or alkall, followed by one or more rinses with su necessary repeated, extraction by heat, followed by purification by	ubsequent adjus	tment of the pH and subsequent, if	
		(d)	) in the case of hydrolysed protein produced using a production procontamination of raw Category 3 material, and, in the case of hydroly hides and skins produced in a processing plant dedicated only to hydroly a molecular weight below 10000 Dalton and a process involving the liming and intensive washing followed by:	ysed protein enti drolysed protein	irely or partly derived from ruminant production, using only material with	

(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

#### COUNTRY

#### Processed petfood other than canned petfood

ii. Head in the made is the second of the se	II.	Health information	II.a. Certificate reference No	II.b.
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- (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 of the European Parliament and of the Council (4);
- (f) in the case of collagen submitted to a process ensuring that unprocessed Category 3 material is subjected to a treatment involving washing, pH adjustment using acid or alkali followed by one or more rinses, filtration and extrusion, the use of preservatives other than those permitted by 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;
- (k) in the case of fishmeal submitted to any of the processing methods or to a method and parameters which ensure that the
  products complies with the microbiological standards for derived products set out in Chapter I of Annex X to Regulation
  (EU) No 142/2011;
- (I) 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 remaining total insoluble impurities does not excess 0,15 % in weight;
- (m) 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 under (i), applies a treatment of the obtained phosphoric liquor with lime, resulting in a precipitate of dicalcium phosphate at pH 4 to 7; and
  - (iii) finally, air dries the precipitate of dicalcium phosphate with inlet temperature of 65 °C to 325 °C and end temperature between 30 °C and 65 °C;
- (n) in the case of tricalcium phosphate produced by a process that ensures
  - (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 °C;
- (o) 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 under 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, be 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 (5):

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

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

#### COUNTRY

#### Processed petfood other than canned petfood

II.	Health information	II.a. Certificate reference No	II.b.
II.5.	has undergone all precautions to avoid contamination with path	ogenic 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';

11.7.

(2) either

(2) or

(2) either

[the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001 of the European Parliament and of the Council (§) or mechanically separated meat obtained from bones of bovine, ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity:]

[the product does not contain and is not derived from bovine, ovine or 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 by a decision in accordance with Article 5(2) of Regulation (EC) No 999/2001.]

- II.8. in addition as regards TSE:
  - [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin, the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last three years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfied the following requirements for the last three years:
    - (i) it has been subject to regular official veterinary checks;
    - (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or, following the confirmation of a classical scrapie case:
      - all animals in which classical scrapie was confirmed have been killed and destroyed, and
      - all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARR genotype and breeding ewes carrying at least one ARR allele and no VRQ allele;
    - (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are introduced into the holding only if they come from a holding which complies with the requirements set out in points (i) and (ii).]
  - (²) or [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin, and destined to a Member State listed in the Annex to Commission Regulation (EC) No 546/2006 (²), the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last seven years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfied the following requirements for the last seven years:
    - (i) it has been subject to regular official veterinary checks;
    - (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or, following the confirmation of a classical scrapie case:
      - all animals in which classical scrapie was confirmed have been killed and destroyed, and
      - all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARR genotype and breeding ewes carrying at least one ARR allele and no VRQ allele;
    - (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are introduced into the holding only if they come from a holding which complies with the requirements set out in points (i) and (ii).]

### Notes

### Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only
  be stored in free zones, free warehouses and custom warehouses.

Processed petfood other than canned petfood

COUNTRY

Status: This is the original version (as it was originally adopted).

П.	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 event of unloading and reloading.						
— I	Box reference I.23: for bulk containers, the container number and the	e seal number (if applicable) should b	e given.				
— I	Box reference I.25: technical use: any use other than for animal consumption.						
— I	Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.						
Par	Part II:						
( <sup>1a</sup> )	OJ L 300, 14.11.2009, p. 1.						
(1b)	OJ L 54, 26.2.2011, p. 1.						
( <sup>2</sup> )	Delete as appropriate.						
(3)	OJ L 175, 10.7.2010, p. 1.						
(4)	OJ L 139, 30.4.2004, p. 55.						
( <sup>5</sup> )	Where:						
	n = number of samples to be tested;						
	$\boldsymbol{m} = \text{threshold}$ value for the number of bacteria; the result is consident;	ered satisfactory if the number of bacte	eria in all samples does not exceed				
	M = maximum value for the number of bacteria; the result is consider or more; and	lered unsatisfactory if the number of ba	cteria in one or more samples is M				
	c = number of samples the bacterial count of which may be betwee count of the other samples is m or less.	een m and M, the sample still being co	nsidered acceptable if the bacterial				
( <sup>6</sup> )	OJ L 147, 31.5.2001, p. 1.						
(7)	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 Europaccompany the consignment until it reaches the border inspection p		or veterinary purposes and has to				
Offic	Official veterinarian/Official inspector						
	Name (in capital letters):	Qualification	and title:				
	Date:	Signature:					
	Stamp:						

# CHAPTER 3(C)

# Health certificate

For dogchews intended for dispatch to or for transit through  $\binom{2}{2}$  the European Union

cou	NTRY	,			Veterinary certificate to EU	
	l.1.	Consignor		I.2. Certificate reference No	I.2.a.	
		Name				
		Address		I.3. Central competent authority		
		Tel.		I.4. Local competent authority		
뉱	1.5.	Consignee		I.6. Person responsible for the load i	n EU	
me.		Name		Name		
sign		Address		Address		
Ö		Destands				
pa		Postcode Tel.		Postcode Tel.		
atch				10		
of dispatched consignment	1.7.	Country of origin ISO code	I.8. Region of origin Code	I.9. Country of destination	I.10. Region of Code destination	
Part I: Details	1.11.	. Place of origin		I.12. Place of destination		
art I:		Name Address	Approval number		Custom warehouse	
-		Name Address	Approval number		Approval number	
	Name Approval number Address			Postcode		
	l.13.	Place of loading		I.14. Date of departure		
	l.15.	. Means of transport		I.16. Entry BIP in EU		
		Aeroplane Ship	Railway wagon			
		Road vehicle Other		1.17.		
		Identification				
		Documentation references				
	I.18.	. Description of commodity		I.19. Commodity code (		
				I.20. Qu	antity	
	1.21.	. Temperature of product		1.22. Nu	mber of packages	
		Ambient	Chilled	Frozen 🗆		
	1.23.	. Seal/Container No		I.24. Ty	pe of packaging	
	1.25.	. Commodities certified for:				
		Animal feedingstuff	Technical use			
	1.26.	. For transit through EU to third of	country	I.27. For import or admission into EU		
		Third country	ISO code			
	1.28.	. Identification of the commodities	•			
	(		number of establishments anufacturing plant	Net weight	Batch number	

COUNTRY

Health information II.a. Certificate reference No II.b. I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council (<sup>1a</sup>), and in particular Article 10, and Commission Regulation (EU) No 142/2011 (<sup>1b</sup>), and in particular Annex XIII, Chapter II and Annex XIV, Chapter II thereof, and certify that the dogchews described above: have been prepared exclusively with the following animal by-products: Part II: Certification (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, of animals other than ruminants; (iv) pig bristles; (v) feathers;] (2) and/or [- blood of animals which did not show any signs of disease communicable through blood to humans or animals, obtained from animals other than ruminants that have been slaughtered in a slaughterhouse after having been considered fit for slaughter for human consumption following an 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 [- animal by-products from aquatic animals originating from plants or establishments manufacturing products for human consumption:1 (2) and/or [- material from animals which have been treated with certain substances which are prohibited pursuant to Directive 96/22/EC, the import of the material being permitted in accordance with Article 35(a)(ii) of Regulation (EC) No 1069/2009;] have been subjected [in the case of dogchews made from hides and skins of ungulates or from fish, to a treatment sufficient to destroy pathogenic (2) either 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;] 11.3. 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): absence in 25 g: n = 5, c = 0, m = 0, M = 0, Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram; have undergone all precautions to avoid contamination with pathogenic agents after treatment: 11.4. 11.5. were packed in new packaging;

COUNTRY

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

II.6.

(2) either [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001 of the European Parliament and of the Council (4) or mechanically separated meat obtained from bones of bovine, ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity:]

(2) or [the product does not contain and is not derived from bovine, ovine or 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 by a decision in accordance with Article 5(2) of Regulation (EC) No 999/2001;]

#### II.7. in addition as regards TSE:

(e) either [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin, the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last three years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfied the following requirements for the last three years:

- (i) it has been subject to regular official veterinary checks;
- (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or, following the confirmation of a classical scrapie case:
  - all animals in which classical scrapie was confirmed have been killed and destroyed, and
  - all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARR genotype and breeding ewes carrying at least one ARR allele and no VRQ allele;
- (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are introduced into the holding only if they come from a holding which complies with the requirements set out in points (i) and (ii).]
- (2) or [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin, and destined to a Member State listed in the Annex to Commission Regulation (EC) No 546/2006 (5), the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last seven years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfied the following requirements for the last seven years:
  - (i) it has been subject to regular official veterinary checks;
  - (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or, following the confirmation of a classical scrapie case:
    - all animals in which classical scrapie was confirmed have been killed and destroyed, and
    - all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARR genotype and breeding ewes carrying at least one ARR allele and no VRQ allele;
  - (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are introduced into the holding only if they come from a holding which complies with the requirements set out in points (i) and (ii).]

### Notes

### Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit
  commodity; it may be filled in if the certificate is for import commodity.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only
  be stored in free zones, free warehouses and custom warehouses.

-	MINI		Dogenew				
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 event of unloading and reloading.						
_	Box reference I.19: Alternatively, commodity codes 2309 and 4101 n	nay be chosen.					
_	Box reference I.23: for bulk containers, the container number and the	e seal number (if applicable) should b	e given.				
-	Box reference I.25: technical use: any use other than for animal con-	sumption.					
-	Box reference I.26 and I.27: fill in according to whether it is a transit	or an import certificate.					
Par	t II:						
(1a)	OJ L 300, 14.11.2009, p. 1.						
(1b)	OJ L 54, 26.2.2011, p. 1.						
( <sup>2</sup> )	Delete as appropriate.						
(3)	Where:						
	n = number of samples to be tested;						
	m = threshold value for the number of bacteria; the result is conside m;	ered satisfactory if the number of bact	eria in all samples does not exceed				
	M = maximum value for the number of bacteria; the result is consider or more; and	ered unsatisfactory if the number of ba	octeria in one or more samples is M				
	c = number of samples the bacterial count of which may be betwee count of the other samples is m or less.	en m and M, the sample still being co	nsidered acceptable if the bacterial				
(4)	OJ L 147, 31.5.2001, p. 1.						
( <sup>5</sup> )	OJ L 94, 1.4.2006, p. 28.						
-	The signature and the stamp must be in a different colour to that of	the printing.					
	<ul> <li>Note for the person responsible for the consignment in the European Union: This certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.</li> </ul>						
Offi	icial 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

cou	NTR	1	Veterinary certificate to EU				
	l.1.	Consignor	I.2. Certificate reference No I.2.a.				
		Name	I.3. Central competent authority				
		Address					
		Tel.	I.4. Local competent authority				
뉱	1.5.	Consignee	I.6. Person responsible for the load in EU				
ine i		Name	Name				
sigr		Address	Address				
l o		Postcode	Postcode Tel.				
dispatched consignment		Tel.					
patc	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO I.10. Region of Code				
dis		Todamary or original locations and an original country or original	destination code destination				
s of							
l: Details	l.11.	Place of origin	I.12. Place of destination				
:D		Name Approval number	Name Custom warehouse □				
Part		Address	Address Approval number				
-		Name Approval number Address					
		Name Approval number	Postcode				
		Address					
	I.13.	Place of loading	I.14. Date of departure				
	145	Manua of transport	I.16. Entry BIP in EU				
	1.15.	Means of transport	1.10. Entry bir in Eo				
		Aeroplane Ship Railway wagon Road vehicle Other Ship					
		Identification	1.17.				
		Documentation references					
	I.18.	Description of commodity	I.19. Commodity code (HS code)				
			I.20. Quantity				
	L21.	Temperature of product	I.22. Number of packages				
		Ambient ☐ Chilled ☐	Frozen				
	1.23.	Seal/Container No	I.24. Type of packaging				
	1.25.	Commodities certified for:					
		Animal feedingstuff Technical use					
	1.26.	For transit through EU to third country	I.27. For import or admission into EU				
		Third country ISO code					
	1.28.	Identification of the commodities					
			umber of establishments Net weight Batch number nufacturing plant				

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

#### COUNTRY

Part II:

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

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

- II.1. consist of animal by-products that satisfy the health requirements below;
- II.2. consist of animal by-products:

  (a) derived from meat which s

   Commission Regulation
  - (a) derived from meat which satisfies the relevant animal and public health requirements laid down in:

    - and/or Commission Regulation (EC) No 798/2008 (4), and provided the animals from which the meat is derived come from a territory
      or part of a territory ........................ (ISO code) 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 the animals from which the meat is derived come from a territory or part of a territory .................. (ISO code) 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 last 12 months and where no vaccination has taken place during that time (only as relevant for the susceptible species);
  - (b) derived from animals that, at the slaughterhouse, have passed the ante-mortem health inspection during the 24 hours before the slaughter and have shown no evidence of the diseases referred in the Regulations above for which the animals are susceptible; and
  - (c) derived from animals that have been treated in the slaughterhouse before and at the time of slaughter or killing in accordance with the relevant provisions of Council Directive 93/119/EC (6) on the protection of animals at the time of slaughter or killing;
  - II.3. consist only of the following animal by-products:
    - (a) parts of slaughtered animals, which were fit for human consumption in accordance with Union legislation, but are not intended for human consumption for commercial reasons, and
    - (b) parts of slaughtered animals, which are rejected as unfit for human consumption but are not affected by any signs of diseases communicable to humans or animals and derive from carcases that are fit for human consumption in accordance with Union legislation;
  - II.4. have been obtained and prepared without contact with other material not complying with the conditions required in the Regulations above, and it has been handled so as to avoid contamination with pathogenic agents;
  - II.5. have been packed in final packaging which bear labels indicating 'RAW PET FOOD NOT FOR HUMAN CONSUMPTION' or 'ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS NOT FOR HUMAN CONSUMPTION' and then in leak-proof and officially sealed boxes/containers or in new packaging preventing any leakage and officially sealed boxes/containers which bear labels indicating 'RAW PET FOOD NOT FOR HUMAN CONSUMPTION' or 'ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS NOT FOR HUMAN CONSUMPTION', the name and the address of the establishment of destination;
  - II.6. in the case of raw petfood:
    - (a) have been prepared and stored in a plant approved and supervised by the competent authority in accordance with Article 24 of Regulation (EC) No 1069/2009; and
    - (b) were examined by random sampling of at least five samples from each batch taken during storage (before dispatch) and complies with the following standards (7):

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

Enterobacteriaceae: n = 5, c = 2, m = 10, M = 5000 in 1 gram;

II.7.

(2) either
[the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001 of the European Parliament and of the Council (8) or mechanically separated meat obtained from bones of bovine, ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity.]

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

### COUNTRY

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

(²) or [the product does not contain and is not derived from bovine, ovine or 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 by a decision in accordance with Article 5(2) of Regulation (EC) No 999/2001;]

#### II.8. in addition as regards TSE:

- (2) either [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin, the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last three years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfied the following requirements for the last three years:
  - (i) it has been subject to regular official veterinary checks;
  - (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or, following the confirmation of a classical scrapie case:
    - all animals in which classical scrapie was confirmed have been killed and destroyed, and
    - all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARR genotype and breeding ewes carrying at least one ARR allele and no VRQ allele;
  - (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are introduced into the holding only if they come from a holding which complies with the requirements set out in points (i) and (ii).]
- (2) or [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin, and destined to a Member State listed in the Annex to Commission Regulation (EC) No 546/2006 (9), the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last seven years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfied the following requirements for the last seven years:
  - (i) it has been subject to regular official veterinary checks;
  - (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or, following the confirmation of a classical scrapie case:
    - all animals in which classical scrapie was confirmed have been killed and destroyed, and
    - all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARR genotype and breeding ewes carrying at least one ARR allele and no VRQ allele;
  - (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are introduced into the holding only if they come from a holding which complies with the requirements set out in points (i) and (ii).]

## Notes

### Part I

- Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit
  commodity; it may be filled in if the certificate is for import commodity.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only
  be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in the event of unloading and reloading.
- Box reference I.19: use the appropriate HS code: 05.11.91; 05.11.99 or 23.09.90.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be given.

ou	INTRY	Raw petfood for direct sale or animal by- products to be fed to fur animals			
II.	Health information	II.a. Certificate reference No	II.b.		
_	Box reference I.25: technical use: any use other than for animal co	nsumption.			
_	Box reference I.26 and I.27: fill in according to whether it is a trans	it or an import certificate.			
_	Box reference I.28: Nature of commodity: select raw petfood or ani	mal by-product.			
Par	t II:				
( <sup>1a</sup> )	OJ L 300, 14.11.2009, p. 1.				
( <sup>1b</sup> )	OJ L 54, 26.2.2011, p. 1.				
( <sup>2</sup> )	Delete as appropriate.				
( <sup>3</sup> )	OJ L 73, 20.3.2010, p. 1.				
( <sup>4</sup> )	OJ L 226, 23.8.2008, p. 1.				
( <sup>5</sup> )	OJ L 39, 10.2.2009, p. 12.				
( <sup>6</sup> )	OJ L 340, 31.12.1993, p. 21.				
( <sup>7</sup> )	Where:				
	n = number of samples to be tested;				
	m = threshold value for the number of bacteria; the result is consider m;n	ered satisfactory if the number of bacte	eria in all samples does not exceed		
	$\mbox{\bf M} = \mbox{\bf maximum value}$ for the number of bacteria; the result is consider or more; and	ered unsatisfactory if the number of ba	cteria in one or more samples is M		
	c = number of samples the bacterial count of which may be betwee count of the other samples is m or less.n	en m and M, the sample still being co	nsidered acceptable if the bacterial		
( <sup>8</sup> )	OJ L 147, 31.5.2001, p. 1.				
( <sup>9</sup> )	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 Europaccompany the consignment until it reaches the border inspection po		or veterinary purposes and has to		
Offic	cial veterinarian/Official inspector				
	Name (in capital letters):	Qualification and	d 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  $\binom{2}{2}$  the European Union

COUNTRY Veterinary certificate to							ficate to EU				
	l.1.	Consignor				I.2. Certificate reference No I.2.a.					
		Name				I.3. Central competent authority					
		Address						ompetent a	utilionty		
						1.4.	Local cor	mpetent aut	hority		
		Tel.				_					
ent	1.5.	Consignee		1.6.	Person re	esponsible 1	or the load	in EU			
Ē		Name					Name				
ısig		Address					Address				
Ö	Postcode						Postcode				
ped		Tel.					Tel.	,			
dispatched consignment		T									
isp	1.7.	Country of origin	ISO code	I.8. Region of origin	Code	1.9.	Country of destination		ISO code	I.10. Region of destination	Code
o d			l		I		destinatio	"' I		destination	1
ls o											
l: Details	1.11.	Place of origin				l.12.	Place of	destination			
<u>:</u>		Name		Approval number			Name			Custom warehouse [	¬
Part		Address					Address			Approval number	_
۵.		Name		Approval number							
		Address					Postcode	,			
		Name Address		Approval number							
	L13	Place of loading				1.14	Date of d	leparture			
		riaco or localing					Date 0. 0	.opartaro			
	1.15.	Means of transport				116	Entry BIP	in FII			
	·					1.10.	Linky Dir	III E0			
		Aeroplane  Road vehicle	Ship  Other	, ,	ш	L17.					
		Identification	Other	_							
		Documentation refe	erences								
	1.18.	Description of com				I.19. Commodity code (HS code)					
			,			1.15. Commonly code (115 code)					
						I.20. Quantity					
	1.01	Temperature of pro	duat								
	1.21.			Ob. 111 - 11 - 17		I.22. Number of packages Frozen □					
		Ambient		Chilled							
	1.23.	Seal/Container No				I.24. Type of packaging					
	1.25.	Commodities certifi	ed for:								
		Animal feedingstuff		Technical use	. 🗆						
	• –					1.07	For impo	t or admiss	ion into Ell		
	I.26. For transit through EU to third country				1.27.	For impor	t or admiss	ion into EU			
	Third country ISO code										
	1.28.	Identification of the	commoditie	s							
		Species	Net	ure of commodity	Annroyal re-	mhor	of actablic	hmonto	Not .	weight Batch	n number
	Species Nature of commodity Approval nu (Scientific name) Man					ring plant	milents	IARL /	weigin battr	i iluliibei	

c	OUN	TRY		Flavouring innards for use in the manufacture of petfo							
ſ	II.		Health infe	orn	nation	II.a. Certificate reference No	II.b.				
		I, the undersigned official veterinarian, declare that I ha and of the Council ( <sup>1a</sup> ) and in particular Article 8 and Annex XIII, Chapter III and Annex XIV, Chapter II the			ouncil (1a) and in particular Article 8 and 10 there	of, and Commission Regulation (EU)	No 142/2011 (1b), and in particular				
	II.1. consist of animal by-products that satisfy the animal he			ani	mal by-products that satisfy the animal health requ	uirements below;					
	ertifica	2.	have been	en prepared including the following animal by-products which are exclusively:							
	Part II: Certification		(²) either	[-		ered or, in the case of game, bodies or parts of animals killed, and which are fit fo th Union legislation, but are not intended for human consumption for commercia					
			(2) and/or	[-	carcases and the following parts originating either considered fit for slaughter for human consumption animals from game killed for human consumption	n following an ante-mortem inspection					
	$\frac{1}{2}$				(i) carcases or bodies and parts of animals whic legislation, but which did not show any signs						
					(ii) heads of poultry;						
				(iii) hides and skins, including trimmings and split metacarpus bones, tarsus and metatarsus bo		the phalanges and the carpus and					
	(iv) pig bristles;				(iv) pig bristles;						
		(v) feathers;]		(v) feathers;]							
	animals other than ruminants that have be			[-	animals other than ruminants that have been slaug	gns of disease communicable through blood to humans or animals, obtained from an slaughtered in a slaughterhouse after having been considered fit for slaughter mortern inspection in accordance with Union legislation;]					
			(2) and/or	[-	animal by-products arising from the production of greaves and centrifuge or separator sludge from		mption, including degreased bone,				
			( <sup>2</sup> ) and/or	[-	products of animal origin, or foodstuffs containing consumption for commercial reasons or due to which no risk to public or animal health arise;]						
			( <sup>2</sup> ) and/or	[-	petfood and feedingstuffs of animal origin, or feed longer intended for feeding for commercial reason defects from which no risk to public or animal here.	ons or due to problems of manufactur					
			(2) and/or	[-	blood, placenta, wool, feathers, hair, horns, hoof cany disease communicable through that product t		animals that did not show signs of				
		(²) and/or [- aquatic animals, and parts of such animals, to humans or animals;]				t sea mammals, which did not show a	ny signs of diseases communicable				
			(²) and/or	[-	animal by-products from aquatic animals original consumption;]	ating from plants or establishments r	manufacturing products for human				
(²) and/or [- the following material originating from an to humans or animals:			[-	the following material originating from animals whice to humans or animals:	ch did not show any signs of disease of	communicable through that material					

(i) shells from shellfish with soft tissue or flesh;

## COUNTRY

# Flavouring innards for use in the manufacture of petfood

COUN			Flavouring innards for use in the manufacture of petroo			
II.	Health inf	formation	II.a. Certificate reference No	II.b.		
		(ii) the following originating from terrestrial animals	:			
		<ul> <li>hatchery by-products,</li> </ul>				
		— eggs,				
		<ul> <li>egg by-products, including egg shells;</li> </ul>				
		(iii) day-old chicks killed for commercial reasons;]				
	(2) and/or	[- animal by-products from aquatic or terrestrial invert	rebrates other than species pathoger	nic to humans or animals;]		
	(2) and/or	[- material from animals which have been treated with the import of the material being permitted in accord				
II.3.	have beer agents;	n subjected to processing in accordance with Annex XIII				
II.4.		n examined by the competent authority taking a random standards $(^3)$ :	n sample immediately prior to dispat	tch and found it to comply with the		
	Salmonella	a: absence in 25g: n = 5, c = 0, m = 0, M	1 = 0,			
	Enterobac	teriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gra	m;			
II.5.	the end p	roduct was:				
	(2) either	[packed in new or sterilised bags,]				
	(²) or	[transported in bulk in containers or other means of transported by the competent authority before use,]	ansport that were thoroughly cleaned	d and disinfected with a disinfectant		
	and which	bear labels indicating 'NOT FOR HUMAN CONSUMP'	TION';			
II.6.	the end p	roduct was stored in enclosed storage;				
II.7.	the produc	ct has undergone all precautions to avoid contamination	with pathogenic agents after treatment	nent;		
II.8.						
	(²) either	[the product does not contain and is not derived from 999/2001 of the European Parliament and of the Coulovine or caprine animals; and the animals from whice means of gas injected into the cranial cavity or killed by means of an elongated rod-shaped instrument into	ncil (4) or mechanically separated mathematics and like this product is derived have not lightly the same method or slaughtered by	eat obtained from bones of bovine, been slaughtered after stunning by		
	(²) or	[the product does not contain and is not derived from I born, continuously reared and slaughtered in a count accordance with Article 5(2) of Regulation (EC) No 9:	ry or region classified as posing a n			
II.9.	in addition	as regards TSE:				
	(2) either	[in case of animal by-products intended for feeding ru the ovine and caprine animals from which these prod three years on a holding where no official movement r the following requirements for the last three years:	ucts are derived have been kept con	ntinuously since birth or for the last		
		(i) it has been subject to regular official veterinary cl	hecks;			
		(ii) no classical scrapie case, as defined in point 2(g following the confirmation of a classical scrapie cases		999/2001, has been diagnosed or,		
		all animals in which classical scrapie was con	firmed have been killed and destroy	ed, and		
		— all goats and sheep on the holding have be				
		genotype and breeding ewes carrying at least	one ARR allele and no VRQ allele;			

## COUNTRY

#### Flavouring innards for use in the manufacture of petfood

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

(iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are introduced into the holding only if they come from a holding which complies with the requirements set out in points (i) and (ii).]

(2) or [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin, and destined to a Member State listed in the Annex to Commission Regulation (EC) No 546/2006 (5), the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last seven years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfied the following requirements for the last seven years:

- (i) it has been subject to regular official veterinary checks;
- (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or, following the confirmation of a classical scrapie case:
  - all animals in which classical scrapie was confirmed have been killed and destroyed, and
  - all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARR genotype and breeding ewes carrying at least one ARR allele and no VRQ allele;
- (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are introduced into the holding only if they come from a holding which complies with the requirements set out in points (i) and (ii).]

#### Notes

#### Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only
  be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be
  provided in the event of unloading and reloading.
- Box reference I.19: use the appropriate HS code: 05.04 or 05.11.91.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be given.
- Box reference I.25: technical use: any use other than for animal consumption.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: 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.
- (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.

COUNTRY	Flavouring innards for u	se in the manufacture of petfood		
II. Health information	II.a. Certificate reference No	II.b.		
(4) OJ L 147, 31.5.2001, p. 1.				
( <sup>5</sup> ) OJ L 94, 11.4.2006, p. 28.				
- The signature and the stamp must be in a different colour to that of	the printing.			
<ul> <li>Note for the person responsible for the consignment in the European Union: This certificate is only for veterinary purposes and ha accompany the consignment until it reaches the border inspection post.</li> </ul>				
Official veterinarian/Official inspector				
Name (in capital letters):	Qualification ar	nd title:		
Date:	Signature:			
Stamp:				

# CHAPTER 3(F)

# **Health certificate**

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

cou	NTR	(	Veterinary certificate to EU	
	l.1.	Consignor	I.2. Certificate reference No I.2.a.	
		Name	I.3. Central competent authority	
		Address	1.3. Central competent additionty	
		Tel.	I.4. Local competent authority	
<u>۔</u>	1.5.	Consignee	I.6. Person responsible for the load in EU	
l er		Name	Name	
l ig		Address	Address	
l Suc				
5		Postcode	Postcode	
) ě		Tel.	Tel.	
dispatched consignment	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10. Region of Code	
dis			destination destination	
6				
Part I: Details of	l.11.	Place of origin	I.12. Place of destination	
Ü		Name Approval number	Name Custom warehouse	
<u>F</u>		Address	Address Approval number	
ية		Name Approval number	Bostonia	
		Address	Postcode	
		Name Approval number		
		Address		
	I.13.	Place of loading	I.14. Date of departure	
	I.15.	Means of transport	I.16. Entry BIP in EU	
	Aeroplane ☐ Ship ☐ Railway wagon ☐			
		Road vehicle Other O		
		Identification	1.17.	
		Documentation references		
	I.18.	Description of commodity	I.19. Commodity code (HS code)	
			42.06	
			I.20. Quantity	
	1.01	Temperature of product	I.22. Number of packages	
	1.21.	Ambient Chilled	Frozen   Frozen	
	1.23.	Seal/Container No	I.24. Type of packaging	
	1.25.	Commodities certified for:	<u>'</u>	
		Animal feedingstuff ☐ Further p	rocess Technical use	
	1.26.	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 Nature of commodity Approval number of (Scientific name) Manufacturing		

ç	ou	NTRY			Animal by-products	for the manufacture of petfood
		II.	Health inf	ormation	II.a. Certificate reference No	II.b.
			Parliament	ersigned official veterinarian, declare that I have t and of the Council ( <sup>1a</sup> ) and Commission Regulati that the animal by-products described above:		
	ے	II.1.1.	consist of	animal by-products that satisfy the animal health	requirements below;	
	Part II: Certification	II.1.2.	have been	n obtained in the territory of:	(1	c) from animals:
	: Cert		(2) either	[(a) that have remained in this territory since bit	rth or for at least the last three months	s before slaughter;]
	Part		(2) or	[(b) killed in the wild in this territory $(^{1d})$ ;]		
		II.1.3.	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 been neither case/outbreak rinderpest, swine vesicular disease, Newcastle disease or highly pathogenic avian influenza during the prior 3 days, nor of classical or African swine fever during the prior 40 days; nor in the holdings situated in their vicinii within 10 km, during the prior 30 days; and				
				(ii) where there has been neither case/ou holdings situated in their vicinity within		ing the prior 60 days, nor in the
	(b) which:					
	<ul> <li>(i) were not killed to eradicate any epizootic disease;</li> <li>(ii) have remained in their holdings of origin for at least 40 days before departure and which have been transpedirectly to the slaughterhouse without contact with other animals which did not comply with the same honditions;</li> <li>(iii) at the slaughterhouse, have passed the ante-mortem health inspection during the 24 hours before the slaughterhouse shown no evidence of the diseases referred to above for which the animals are susceptible;</li> </ul>					
	(iv) have been treated in the slaughterhouse before and at the time of slaughter or killing in accordance with relevant provisions of Council Directive 93/119/EC (4) on the protection of animals at the time of slaughte killing]					
			(2) or	[(a) captured and killed in the wild in an area:		
<ul> <li>(i) in which within 25 km there has been no case/outbreak of any of the following diseases for which the an susceptible: foot-and-mouth disease, rinderpest, Newcastle disease or highly pathogenic avian influenz the prior 30 days, nor of classical or African swine fever during the prior 40 days; and</li> </ul>					pathogenic avian influenza during	
				(ii) that is situated at a distance that exceed thereof, which is not authorised at these		
				(b) which after killing were transported within 12 to a game establishment, or directly to a game		centre and immediately afterwards
		II.1.4.	referred to preparation	n obtained in an establishment around which, wo in point II.1.3 for which the animals are suscept of raw material for exportation to the European and disinfection of the establishment under the co	otible during the prior 30 days or, in the Union has been authorised only after	e event of a case of disease, the

COUNTRY		Animal by-products for the manufacture of petfoc				
II.	Health inf	rmation II.a. Certificate reference No II.b.				
II.1.5.	have been obtained and prepared without contact with other material not complying with the conditions required above, and it has been handled so as to avoid contamination with pathogenic agents;					
II.1.6.		have been packed in new packaging preventing any leakage and in officially sealed containers bearing the label indicating 'RAW MATERIAL ONLY FOR THE MANUFACTURE OF PET FOOD' and the name and address of the EU establishment of destination;				
II.1.7.	consist only of the following animal by-products:					
	(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;]				
	(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:					
		<ul> <li>(i) carcases or bodies and parts of animals which are rejected as unfit for human consumption in accordance wit Union legislation, but which did not show any signs of disease communicable to humans or animals;</li> </ul>				
		(ii) heads of poultry;				
		(iii) hides and skins, including trimmings and splitting thereof, horns and feet, including the phalanges and the carpu and metacarpus bones, tarsus and metatarsus bones, of animals other than ruminants;				
		(iv) pig bristles;				
	(v) feathers;]					
	(2) and/or [- animal by-products arising from the production of products intended for human consumption, including degrease 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 hum consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects frowhich no risk to public or animal health arise;]					
	(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 huma consumption;]				
	(2) and/or	[- the following material originating from animals which did not show any signs of disease communicable through the 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,				
	<ul><li>— egg by-products, including egg shells;</li></ul>					
		(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	[- material from animals which have been treated with certain substances which are prohibited pursuant to Directiv 96/22/EC, the import of the material being permitted in accordance with Article 35(a)(ii) of Regulation (EC No 1069/2009;]				

COUNTRY			Animal by-produc	ts for the manufacture of petfood	
II.	Health info	rmation	II.a. Certificate reference No	II.b.	
II.1.8.		deep-frozen at the plant of origin or have been p en dispatch and delivery to the plant of destina		ation in such a way that they will not	
II.1.9.		of raw material derived from animals which h. //22/EC for the manufacture of petfood, the impr 09:			
	(a) it has been marked in the third country before entry into the territory of the Union by a cross of liquefied charcoal or ac carbon on each outer side of each frozen block, or, when the raw material is transported in pallets which are not divid separate consignments during transport to the petfood plant of destination, on each outer side of each pallet, in a way to marking covers at least 70 % of the diagonal length of the frozen block and be of at least 10 cm width;				
		of material which is not frozen, the raw material y spraying it with liquefied charcoal or by apply ; and			
		ase the animal by-products are made up of ra raw material, all the raw materials have been n			
( <sup>2</sup> ) ( <sup>5</sup> ) [II.2.	Specific rec	uirements			
(²) ( <sup>6</sup> ) II.2.1.		ducts in this consignment come from animals programmes against foot-and-mouth disease a			
(²) ( <sup>7</sup> ) II.2.2.	maturated a	ducts in this consignment consist only of animal int an ambient temperature of more than $+ 2  ^{\circ}$ C dideboned meat of domestic animals, for at lea	for at least three hours, or in the c		
II.3.					
	(²) either	[the product does not contain and is not derived 999/2001 of the European Parliament and of bovine, ovine or caprine animals; and the animals stunning by means of gas injected into the cracentral nervous tissue by means of an elongate	the Council (8) or mechanically sepa- mals from which this product is deriv- anial cavity or killed by the same met	rated meat obtained from bones of ed have not been slaughtered after thod or slaughtered by laceration of	
	( <sup>2</sup> ) or	[the product does not contain and is not derive animals born, continuously reared and slaughte decision in accordance with Article 5(2) of Reg	ered in a country or region classified a		
II.4.	in addition	as regards TSE:			
	(²) either	[in case of animal by-products intended for fee origin, the ovine and caprine animals from whic for the last three years on a holding where no which has satisfied the following requirements	ch these products are derived have be official movement restriction is impo	een kept continuously since birth or	
		(i) it has been subject to regular official veter	inary checks;		
		(ii) no classical scrapie case, as defined in po or, following the confirmation of a classical		No 999/2001, has been diagnosed	
		- all animals in which classical scrapie w	as confirmed have been killed and d	lestroyed, and	
		<ul> <li>all goats and sheep on the holding have genotype and breeding ewes carrying a</li> </ul>			
		(iii) ovine and caprine animals, with the exception holding only if they come from a holding w			

#### COUNTRY

#### Animal by-products for the manufacture of petfood

II.a. Certificate reference No II.b.
feeding ruminants and containing milk or milk products of ovine or caprine d in the Annex to Commission Regulation (EC) No 546/2006 (9), the ovine cts are derived have been kept continuously since birth or for the last seven int restriction is imposed due to a suspicion of TSE and which has satisfied years:
erinary checks;
point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed cal scrapie case:
was confirmed have been killed and destroyed, and
ave been killed and destroyed, except for breeding rams of the ARR/ARR g at least one ARR allele and no VRQ allele;
eption of sheep of the ARR/ARR prion genotype, are introduced into the which complies with the requirements set out in points (i) and (ii).]
Euroepan Union: this box is to be filled in only if it is a certificate for transit dity.
nly if it is a certificate for transit commodity. The products in transit can only i.
ner and lorries), flight number (aircraft) or name (ship); information is to be
11.99.
the seal number (if applicable) should be included.
onsumption.
sit or an import certificate.
ntrol number of the approved establishment.
down in:

In addition the ISO code of regionalisation in this Annex (where applicable for the susceptible species concerned) should be included.

the Annex to Regulation (EC) No 798/2008, andthe Annex to Regulation (EC) No 119/2009.

cou	INTRY	Animal by-produc	ts for the manufacture of petfood	
II.	Health information	II.a. Certificate reference No	II.b.	
( <sup>1d</sup> )	(1d) Only for countries from where game meat intended for human consumption of the same animal species is authorised for importation European Union.			
(²)	Delete as appropriate.			
(3)	Excluding raw blood, raw milk, hides and skins, hooves and horn, pithese products).	g bristles and feathers (see relevant	specific certificates for the import of	
(4)	OJ L 340, 31.12.1993, p. 21.			
(5)	Supplementary guarantees to be provided when the material of don African country or part thereof from where only maturated and debon for exportation to the European Union. The whole masseter muscl Chapter I, Part B(1) of Regulation (EC) No 854/2004 of the European permitted.	ed fresh meat of domestic ruminants les of bovine animals, incised in acc	for human consumption is permitted cordance with Annex I, Section IV,	
(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.			
( <sup>9</sup> )	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 Europaccompany the consignment until it reaches the border inspection po		or veterinary purposes and has to	
Offi	cial veterinarian/Official inspector			
	Name (in capital letters):	Qualification and	title:	
	Date:	Signature:		
	Stamp:			

# CHAPTER 4(A)

# **Health certificate**

For the import of blood and blood products from equidae to be used outside the feed chain, for dispatch to or for transit through  $(^2)$  the European Union

cou	NTR	1	Veterinary certificate to EU
	l.1.	Consignor	I.2. Certificate reference No I.2.a.
		Name	I.3. Central competent authority
		Address	no. Soma compount additionly
		Tel	I.4. Local competent authority
		Tel.	<del> </del>
ent	1.5.	Consignee	I.6. Person responsible for the load in EU
트		Name	Name
Isig		Address	Address
8		Postcode	Postcode
)ed		Tel.	Tel.
dispatched consignment		Our to divide 100 and 100 Burlow divide 0 and	10 0 140 Police ( 0.4
lisb	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO I.10. Region of Code destination code
of			
ails	1.11.	Place of origin	I.12. Place of destination
Det			_
Part I: Details		Name Approval number Address	Name Custom warehouse ☐ Address Approval number
_		Name Approval number Address	Postcode
		Name Approval number	1 0310000
		Address	
	I.13.	Place of loading	I.14. Date of departure
	l.15.	Means of transport	I.16. Entry BIP in EU
		Aeroplane Ship Railway wagon	
		Road vehicle Other I	l.17.
		Documentation references	
	1.18.	Description of commodity	I.19. Commodity code (HS code) 30.02
			I.20. Quantity
			1.20. Quantity
	I.21.	Temperature of product	I.22. Number of packages
		Ambient Chilled C	Frozen
	1.23.	Seal/Container No	I.24. Type of packaging
	1.25.	Commodities certified for:	
		Technical use □	
	1.26.	For transit through EU to third country	I.27. For import or admission into EU
		Third country ISO code	
	1.28.	Identification of the commodities	
		Species	Approval number of establishments
		(Scientific name)	Manufacturing plant

# Blood and blood products from equidae for purposes outside the

COUNTRY feed chain Health information II.a. Certificate reference No II.b. I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council (<sup>1a</sup>) and in particular Article 8(c) and Article 8(d) and Article 10 thereof, and Commission Regulation (EU) No 142/2011 (<sup>1b</sup>), and in particular Annex XIII, Chapter IV thereof, and certify that the blood or blood products of equidae described above: II.1. consist of blood or blood products from equidae that satisfy the health requirements below: Certification 11.2. consist exclusively of blood or blood products of equidae not intended for human nor animal consumption; have been obtained from animals that originate from a third country, territory or part thereof listed in the column 'third countries' lists' of row No 3 of Table 2 in Section 1 of Chapter II of Annex XIV to Regulation (EU) No 142/2011 where the following diseases are compulsorily II.3. notifiable: African horse sickness, dourine, glanders (Burkholderia mallei), equine encephalomyelitis (all types including Venezuelan equine encephalomyelitis), equine infectious anaemia, vesicular stomatitis, rabies, anthrax; ≝ Part have been derived from blood which was collected under the supervision of a veterinarian, from equidae, which on inspection at the time of collection were free from clinical signs of infectious disease: (2) either [in slaughterhouses approved in accordance with Regulation (EC) No 853/2004 of the European Parliament and of the Council (3);] (2) or [in slaughterhouses approved and supervised by the competent authority of the country of export;] (2) or lin facilities approved and supervised by the competent authority of the country of export for the purpose of collecting blood from equidae for the production of blood products for purposes other than feeding for farmed animals;] II.5. have been derived from blood which was collected from equidae. II.5.1. which on inspection on the date of blood collection did not show clinical signs of any of the compulsorily notifiable diseases listed in Annex A to Council Directive 2009/156/EC (4), and of equine influenza, equine piroplasmosis, equine rhinopneumonitis and equine viral arteritis listed in point 4 of Article 1.2.3 of the Terrestrial Animal Health Code of the World Organisation for Animal Health (OIE), 2010 edition; which have been kept for at least 30 days prior to the date of and during blood collection on holdings under veterinary supervision which were not subject to a prohibition order pursuant to Article 4(5) or restrictions for African horse sickness in accordance with Article 5 of Directive 2009/156/EC; which had no contact with equidae from a holding which was subject to a prohibition order for animal health reasons pursuant to Article 4(5) of Directive 2009/156/EC; II.5.4. for which the period for the prohibition order referred to in points II.5.2. and II.5.3 has been determined as followed: (2) either [where not all the animals of species susceptible to the disease located on the holding have been slaughtered and the premises disinfected the period of prohibition has been: - six months in the case of glanders (Burkholderia mallei), beginning on the date on which the equidae infected with the disease are slaughtered, - six months in the case of equine encephalomyelitis of any type, including Venezuelan equine encephalomyelitis, beginning on the date on which the equidae infected with the disease are slaughtered, in the case of equine infectious anaemia, until the date on which, the infected animals having been slaughtered, and the
remaining animals have shown a negative reaction to two Coggins tests carried out three months apart, - during six months from the date of the last recorded case of vesicular stomatitis, - during one month from the date of the last recorded case of rabies.

- during 15 days from the date of the last recorded case of anthrax:

COUNT	RY			Blood and blood products from e feed chain	quidae for purposes outside the
II.	Health inf	ormation		II.a. Certificate reference No	II.b.
	(²) or	disinfected, the period of prohibition	n shall be 30 day	sease located on the holding have b s, beginning on the date on which the , where the period of prohibition shall	animals were slaughtered and the
II.6.		ucts must come from a establishment nditions set out in Articles 23 or 24		ed or registered by the competent auth C) No 1069/2009;	ority of the third country meeting the
II.7.	blood prod	lucts have been produced from blood	d which fulfils the	e conditions referred in II.4 and II.5 ar	nd
	(2) either	birth if less than three months old,	prior to the date	dae which have been kept for a perio e of collection on holdings under vete of blood collection has been free of:	
		(a) African horse sickness for two	years;		
		(b) Venezuelan equine encephalor	myelitis for a peri	iod of at least two years;	
		(c) glanders;			
		(2) either [for a period of three	e years;]		
		slaughterhouse refer	rred to in II.4, in s and sinuses a	he animals have passed the post-moncluding a careful examination of much their ramifications, after splitting to	cous membranes from the trachea,
		(d) in the case of blood products	other than serum	n, vesicular stomatitis for six months;]	
	(²) or	possible causative pathogens for A	frican horse sick	ng treatments, followed by an effective kness, equine encephalomyelitis of all ular stomatitis and glanders ( <i>Burkhold</i>	types including Venezuelan equine
		(2) either [heat treatment at a tem	perature of 65°C	for at least three hours;]]	
		(2) or [irradiation at 25 kGy by	gamma rays;]]		
		(2) or [change in pH to pH 5 f	or two hours;]]		
		(2) or [heat treatment of at least	st 80°C througho	out their substance;]]	
II.8.	all precauti and packa		nination of the blo	ood and blood products with pathogenic	c agents during production, handling
II.9.		blood products were packed in PTION' and bearing the approval nur		neable containers clearly labelled 'I blishment of collection;	NOT FOR HUMAN OR ANIMAL
II.10.	the produc	ts were stored in enclosed storage.			
Notes					
Part I:					
		6: Person responsible for the consignay be filled in if the certificate is for		ropean Union: this box is to be filled ty.	n only if it is a certificate for transit

Box reference I.11 and I.12: Approval number: the registration number of the establishment or plant, which has been issued by the competent

Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses.

COUNT	RY	Blood and blood products from e feed chain	quidae for purposes outside the	
II.	Health information	II.a. Certificate reference No	II.b.	
	reference I.15: Registration number (railway wagons or containe vided in case of unloading and reloading.	r and lorries), flight number (aircraft) o	or name (ship); information is to be	
— Вох	reference I.23: for bulk containers, the container number and the	e seal number (if applicable) must be	included.	
— Вох	— Box reference I.25: technical use: any use other than for animal consumption.			
— Вох	reference I.26 and I.27: fill in according to whether it is a transit	t or an import certificate.		
— Вох	reference I.28: Manufacturing plant: provide the veterinary control	ol number of the registered establishm	ent of collection.	
Part II:				
( <sup>1a</sup> ) OJ	J L 300, 14.11.2009, p. 1.			
(1b) OJ	L 54, 26.2.2011, p. 1.			
(²) De	elete as appropriate.			
(3) OJ	L 139, 30.4.2004, p. 55.			
(4) OJ	L 192, 23.7.2010, p. 1.			
— The	signature and the stamp must be in a different colour to that of	the printing.		
	e for the person responsible for the consignment in the European consignment until it reaches the border inspection post.	Union: this certificate is only for veterin	ary purposes and must accompany	
Official	veterinarian/Official inspector			
Na	ame (in capital letters):	Qualification and	d title:	
Da	ate:	Signature:		
St	amp:			

# 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  $\binom{2}{1}$  the European Union

cou	NTRY	,	Veterinary certificate to El
	I.1.	Consignor	I.2. Certificate reference No I.2.a.
		Name	I.3. Central competent authority
	Address		no. Solidar compotent additionly
		Tel.	I.4. Local competent authority
			LO. Borrow was the feet to FII
e l	I.5.	Consignee	I.6. Person responsible for the load in EU
툂		Name	Name Address
nsi		Address	Address
8		Postcode	Postcode
) š		Tel.	Tel.
of dispatched consignment	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO I.10. Region of Code destination code destination
<b>b</b>			
etails	l.11.	Place of origin	I.12. Place of destination
Part I: Details		Name Approval number Address	Name Custom warehouse ☐ Address Approval number
		Name Approval number Address	Postcode
		Name Approval number Address	
	I.13.	Place of loading	I.14. Date of departure
	l.15.	Means of transport	I.16. Entry BIP in EU
		Aeroplane Ship Railway wagon	
		Road vehicle Other	1.17.
		Identification	
		Documentation references	
	I.18.	Description of commodity	I.19. Commodity code (HS code)
			I.20. Quantity
	I.21.	Temperature of product	I.22. Number of packages
		Ambient Chilled Chilled	Frozen
	1.23.	Seal/Container No	I.24. Type of packaging
	1.25.	Commodities certified for:	
		Animal feedingstuff Technical use	
	1.26.	For transit through EU to third country	I.27. For import or admission into EU
		Third country ISO code	
	1.28.	Identification of the commodities	
		Species Nature of commodity (Scientific name)	Approval number of establishments Batch number  Manufacturing plant

# Blood products not intended for human consumption that could be

#### COUNTRY used as feed material Health information II.a. Certificate reference No. II.b. I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council (<sup>1a</sup>) and Commission Regulation (EU) No 142/2011 (<sup>1b</sup>) and certify that the blood products described above: II.1. consist of blood products that satisfy the health requirements below; II.2. consist exclusively of blood products not intended for human consumption; Certification II.3. have been prepared and stored in a plant, approved, validated and supervised by the competent authority in accordance with Article 24 of Regulation (EC) No 1069/2009; have been prepared exclusively with the following animal by-products: Part (2) either [blood of slaughtered animals, which is fit for human consumption in accordance with Union legislation, but is not intended for human consumption for commercial reasons;] (2) and/or [blood of slaughtered animals, which is rejected as unfit for human consumption in accordance with Union legislation, but which did not show any signs of diseases communicable to humans or animals, derived from carcases that have been slaughtered in a slaughterhouse and were considered fit for human consumption following an ante-mortem inspection in accordance with Union legislation;] II.5. have been submitted (2) either No 142/2011] [to a method and parameters which ensure that the product complies with the microbiological standards set in Chapter I of Annex X to Regulation (EU) No 142/2011,] (2) or in order to kill pathogenic agents; 11.6. have been examined under the responsibility of the competent authority taking a random sample immediately prior to dispatch and found it 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.7. the end product was: (2) either [packed in new or sterilised bags:] (2) or Itransported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectant approved by the competent authority before use, ] and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'; II.8. the end product was stored in enclosed storage: II.9. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment; II.10. (2) either (the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/ [the product does not contain and is not derived from specified risk material as defined in Annex V to Hegulation (EC) No 999/ 2001 of the European Parliament and of the Council (5) or mechanically separated meat obtained from bones of bovine, ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity.]

[the product does not contain and is not derived from bovine, ovine or 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 by a decision in accordance with Article 5(2) of Regulation (EC) No 999/2001.]

(2) or

Stamp:

Status: This is the original version (as it was originally adopted).

# Blood products not intended for human consumption that could be COUNTRY used as feed material Health information II.a. Certificate reference No II.b. Notes Part I: - Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity - Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses. - Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading. Box reference I.19: use the appropriate HS code: 05.11.91 or 05.11.99. Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included. Box reference I.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. 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 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 = maximum value for the number of bacteria; the result is considered unsatisfactory if the number of bacteria in one or more samples is M or more; and c = number of samples the bacterial count of which may be between m and M, the sample still being considered acceptable if the bacterial count of the other samples is m or less. (5) OJ L 147, 31.5.2001, p. 1. 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 has to accompany the consignment until it reaches the border inspection post. Official veterinarian/Official inspector Name (in capital letters): Qualification and title: Signature: Date:

# CHAPTER 4(C)

# Health certificate

For untreated blood products, excluding 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

cou	NTRY	,	Veterinary certificate to El
	l.1.	Consignor	I.2. Certificate reference No I.2.a.
		Name	I.3. Central competent authority
	Address		
		Tel.	I.4. Local competent authority
¥	1.5.	Consignee	I.6. Person responsible for the load in EU
me		Name	Name
sign		Address	Address
Ö			
ped.		Postcode Tel.	Postcode Tel.
of dispatched consignment			
disp	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10. Region of Code destination
ō			
Part I: Details	l.11.	Place of origin	I.12. Place of destination
:		Name Approval number	Name Custom warehouse □
Part		Address	Address Approval number
		Name Approval number Address	Postcode
		Name Approval number Address	. 000000
	I.13.	Place of loading	I.14. Date of departure
	l.15.	Means of transport	I.16. Entry BIP in EU
	Aeroplane ☐ Ship ☐ Railway wagon ☐		
		Road vehicle Other	I.17.
		Identification	
		Documentation references	
	I.18.	Description of commodity	I.19. Commodity code (HS code) 30.02
			I.20. Quantity
	I.21.	Temperature of product	I.22. Number of packages
		Ambient Chilled C	Frozen
	1.23.	Seal/Container No	I.24. Type of packaging
	1.25.	Commodities certified for:	
		Technical use	
	1.26.	For transit through EU to third country	I.27. For import or admission into EU
		Third country ISO code	
	1.28.	Identification of the commodities	
		Species Nature of commodity (Scientific name)	Approval number of establishments Batch number Manufacturing plant

Untreated blood products, excluding of equidae, for the manufacture of derived products for purposes outside the feed chain for furmed entirely

... (3) where no case of foot-and-mouth disease has been

recorded for 12 months and in which vaccination programmes against foot-and-mouth disease are being officially carried out and controlled in domestic ruminant animals for at least 12 months (4);]]

# COUNTRY chain for farmed animals 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 Annex XIV, Chapter II thereof, and certify that: II.1. the blood products described above consist of blood products that satisfy the requirements below; Certification 11.2. they consist exclusively of blood products not intended for human or animal consumption; II.3. they have been prepared and stored in a plant supervised by the competent authority or in the establishment of collection (2), Part II: exclusively with the following animal by-products: (2) either [- blood of slaughtered animals, which is fit for human consumption in accordance with Union legislation, but is not intended for human consumption for commercial reasons;] (2) and/or [- blood of slaughtered animals, which is rejected as unfit for human consumption in accordance with Union legislation, but which did not show any signs of diseases communicable to humans or animals, derived from carcases that have been slaughtered in a slaughterhouse and were considered fit for human consumption following an ante-mortem inspection in accordance with Union legislation;] (2) and/or [- blood of slaughtered animals, which did not show any signs of diseases communicable to humans or animals, obtained from animals other than ruminants that have been slaughtered in a slaughterhouse after having been considered fit for human consumption following an ante-mortem inspection in accordance with Union legislation: (2) and/or [- blood and blood products derived from the production of products intended for human consumption;] (2) and/or [- blood and blood products originating from live animals that did not show signs of any disease communicable through that product to humans or animals:1 (²) and/or [- material from animals which have been treated with certain substances which are prohibited pursuant to Directive 96/22/EC, the import of the material being permitted in accordance with Article 35(a)(ii) of Regulation (EC) No 1069/2009;] the blood from which such products are manufactured has been collected: (2) either [in slaughterhouses approved in accordance with Union legislation;] (2) or [in slaughterhouses approved and supervised by the competent authority of the third country;] (2) or [from live animals in facilities approved and supervised by the competent authority of the third country.] (2) [II.5. in the case of blood products derived from animals belonging to the taxa Artiodactyla, Perissodactyla and Proboscidea, including their crossbreds, the products come: from a country where no case of rinderpest, peste des petits ruminants and Rift Valley fever has been recorded for 12 months and in which vaccination has not been carried out against those diseases for at least 12 months; (2) II.5.2. either [from the territory of a country or region with code ..... .... (3) where no case of foot-and-mouth disease has been recorded for 12 months and in which vaccination has not been carried out against this disease for at least 12 months;]

[from the territory of a country or region with code ....

or

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

# COUNTRY chain for farmed animals Health information II.a. Certificate reference No II.b. (2) [II.5.3. In addition, in case of animals other than Suidae and Tayassuidae: (2) either [in the country or region of origin no case of vesicular stomatitis and bluetongue (2) (including the presence of seropositive animals) has been recorded for 12 months and in which vaccination has not been carried out against those diseases for at least 12 months;] (in the country or region of origin vesicular stomatitis and bluetongue (2) seropositive animals are present (4);]] (2) [II.5.4. In addition, in case of animals other than Suidae and Tayassuidae: [in the country or region of origin no case of swine vesicular disease, classical swine fever and African swine fever has been recorded for at least 12 months and vaccination has not been carried out against those diseases for at least 12 months in the susceptible either [in the country or region of origin no case of vesicular stomatitis and bluetongue (²) (including the presence of seropositive animals) has been recorded for 12 months and in which vaccination has not been carried out against those diseases for at (2) [II.5.4.2. least 12 months:1 (2) [II.5.4.2. [in the country or region of origin vesicular stomatitis seropositive animals are present (4);]]] (2) [II.6. in the case of blood products derived from poultry or other avian species the animals and the products come from the territory of a the OIE. which for at least 12 months has not carried out vaccination against avian influenza, where the animals from which the products derive have not been vaccinated against Newcastle disease with vaccines prepared from a Newcastle disease master strain showing a higher pathogenicity than lentogenic virus strains;] II.7. the products were: (2) either [packed in new or sterilised bags or bottles.] (2) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectant approved by the competent authority before use,] the outer packaging or containers bear labels indicating 'NOT FOR HUMAN OR ANIMAL CONSUMPTION'; II.8. the products were stored in enclosed storage: the products have undergone all precautions to avoid contamination with pathogenic agents during transport; 11.10 (2) either [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001 of the European Parliament and of the Council (<sup>6</sup>) or mechanically separated meat obtained from bones of bovine, ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity.] [the product does not contain and is not derived from bovine, ovine or 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 by a decision in accordance with Article 5(2) of Regulation (EC) No 999/2001.] (2) or Notes Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity.
- 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. The products in transit can only be stored in free zones, free warehouses and custom warehouses.

Untreated blood products, excluding of equidae, for the

Status: This is the original version (as it was originally adopted).

COUNTRY	1	chain for farmed animals				
II.	Health information	II.a. Certificate reference No	II.b.			
	eference I.15: Registration number (railway wagons or contained led in case of unloading and reloading.	r and lorries), flight number (aircraft) or	name (ship); information is to be			
— Box re	eference I.23: for bulk containers, the container number and the	e seal number (if applicable) should be	e included.			
— Box re	eference I.25: technical use: any use other than for animal con	sumption.				
— Box re	eference I.26 and I.27: fill in according to whether it is a transit	t or an import certificate.				
Part II:						
( <sup>1a</sup> ) OJ I	L 300, 14.11.2009, p. 1.					
( <sup>1b</sup> ) OJ I	L 54, 26.2.2011, p. 1.					
(²) Dele	ete as appropriate.					
(3) Cod	e of the territory as it appears in Part 1 of Annex II to Regulati	ion (EU) No 206/2010.				
	is case following the border check provided for in Directive 97/ Directive, the products must be transported directly to the plan		ditions laid down in Article 8(4) of			
( <sup>5</sup> ) Cod	e of the territory as it appears in Part 1 of Annex II to Decision	n 2006/696/EC.				
(6) OJ I	L 147, 31.5.2001, p. 1.					
— The s	ignature and the stamp must be in a different colour to that of	the printing.				
— Note	for the person responsible for the consignment in the Europ npany the consignment until it reaches the border inspection po	rean Union: this certificate is only for ost.	veterinary purposes and has to			
Official ve	eterinarian/Official inspector					
Name	(in capital letters):	Qualification an	d title:			
Date:		Signature:				
Stamp	:					

# CHAPTER 4(D)

# Health certificate

For treated blood products, excluding 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  $\binom{2}{1}$  the European Union

cou	NTRY	•	Veterinary certificate to EU					
	I.1.	Consignor	I.2. Certificate reference No I.2.a.					
		Name	I.3. Central competent authority					
		Address	I.d. Local competent outhority					
			I.4. Local competent authority					
	1.6	Tel.	LC. Deven responsible for the lead in EU					
ieu	1.5.	Consignee	I.6. Person responsible for the load in EU					
igi		Name	Name					
ous		Address	Address					
ğ		Postcode	Postcode					
dispatched consignment		Tel.	Tel.					
isp	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10. Region of Code					
7			destination destination					
ais			140 51 - 4 1 5 5					
Det	1.11.	Place of origin	I.12. Place of destination					
Part I: Details		Name Approval number Address	Name Custom warehouse ☐ Address Approval number					
		Name Approval number Address	Postcode					
		Name Approval number Address						
	I.13.	Place of loading	I.14. Date of departure					
			140 5 : 000 : 50					
	1.15.	Means of transport	I.16. Entry BIP in EU					
		Aeroplane  Ship  Railway wagon  Railway						
		Road vehicle Other Identification	1.17.					
		Documentation references						
	I.18.	Description of commodity	I.19. Commodity code (HS code)					
			30.02					
			I.20. Quantity					
	I.21.	Temperature of product	I.22. Number of packages					
		Ambient Chilled Chilled	Frozen					
	1.23.	Seal/Container No	I.24. Type of packaging					
	1.25.	Commodities certified for:						
		Technical use						
	1.26.	For transit through EU to third country	I.27. For import or admission into EU					
		Third country ISO code						
	1.28.	Identification of the commodities						
		Species Nature of commodity (Scientific name)	Approval number of establishments Batch number Manufacturing plant					

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

### COUNTRY

	II.	Health information		II.a. Certificate reference No II.b.						
		and of th		d and understood Regulation (EC) No 1069/2009 of the European Parliament d Article 8(d) and Article 10 thereof, and Commission Regulation (EU) ereof, and certify that:						
	II.1.	the blood	products described above consist of blood products	that satisfy the requirements below;						
ation	II.2.	they consi	st exclusively of blood products not intended for hun	nan or animal consumption;						
ertific	II.3.	they have	been prepared and stored in a plant supervised by t	he competent authority exclusively with	the following animal by-products:					
Part II: Certification		(²) either	[- blood of slaughtered animals, which is fit for hum for human consumption for commercial reasons;]		nion legislation, but is not intended					
		( <sup>2</sup> ) and/or	[- blood of slaughtered animals, which is rejected a which did not show any signs of diseases comm slaughtered in a slaughterhouse and were considenced accordance with Union legislation;]	nunicable to humans or animals, deriv	ed from carcases that have been					
		(²) and/or [- blood of slaughtered animals, which did not show any signs of diseases communicable to humans or animals, obtained fr animals other than ruminants that have been slaughtered in a slaughterhouse after having been considered fit for hum consumption following an ante-mortem inspection in accordance with Union legislation;]								
		(2) and/or	[- blood and blood products originating from live anir these products to humans or animals;]	mals that did not show clinical signs of a	any disease communicable through					
		(2) and/or	[- material from animals which have been treated with the import of the material being permitted in account.]							
	II.4.	the blood	from which such products are manufactured has bee	en collected:						
		(2) either	[in slaughterhouses approved in accordance with U	Inion legislation,]						
		( <sup>2</sup> ) or	[in slaughterhouses approved and supervised by the	e competent authority of the third cour	ntry,]					
		( <sup>2</sup> ) or	[from live animals in facilities approved and supervi	ised by the competent authority of the	third country.]					
	(²) [II.5.	Tayassuid		g treatments, guaranteeing the absence	Proboscidea including their crossbreeds, other than Suidae and ents, guaranteeing the absence of pathogens of foot-and-mouth t Valley fever and bluetongue:					
		(2) either	[heat treatment at a temperature of 65 °C for at lea	st three hours, followed by an effectiveness check;]						
		(2) or	[irradiation at 25 kGy by gamma rays, followed by	an effectiveness check;]						
		( <sup>2</sup> ) or	[change in pH to pH 5 for two hours, followed by a	an effectiveness check;]						
		( <sup>2</sup> ) or	[heat treatment of at least 80 °C throughout their su	ubstance, followed by an effectiveness	check.]]					
	(²) [II.6.	following to	e of blood products derived from Suidae, Tayassuidae reatments guaranteeing the absence of pathogens of t disease, classical swine fever, African swine fever, No ss:	the following diseases: foot-and-mouth of	disease, vesicular stomatitis, swine					
		(2) either	[heat treatment at a temperature of 65 °C for at lea	ast three hours, followed by an effective	eness check;]					
		(²) or	[irradiation at 25 kGy by gamma rays, followed by	an effectiveness check;]						
		(²) or	[heat treatment of at least 80 $^{\circ}\text{C}$ for Suidae/Taya throughout their substance, followed by an effective		oultry and other avian species (2)					

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

#### COUNTRY

(2) [II.7. In the case of blood products derived from species other than listed under II.5 or II.6 the products have undergone of the following

- II.8. The products were:

Health information

- (2) either [packed in new or sterilised bags or bottles,]
- (2) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectant approved by the competent authority before use;] and

II.a. Certificate reference No

the outer packaging or containers bear labels indicating 'NOT FOR HUMAN OR ANIMAL CONSUMPTION';

- II.9. the products were stored in enclosed storage;
- II.10. the products have undergone all precautions to avoid contamination with pathogenic agents after treatment;

II.11.

- (e) either [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001 of the European Parliament and of the Council (e) or mechanically separated meat obtained from bones of bovine, ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity.]
- (²) or [the product does not contain and is not derived from bovine, ovine or 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 by a decision in accordance with Article 5(2) of Regulation (EC) No 999/2001.]

#### Notes

#### Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit
  commodity; it may be filled in if the certificate is for import commodity.
- 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. The products in transit can only be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.
- Box reference I.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.

#### Part II

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

co	JNTRY	Treated blood products, excluding of equidae, for the manufactur of derived products for purposes outside the feed chain for farme animals					
П.	Health information	II.a. Certificate reference No	II.b.				
(2)	Delete as appropriate.						
(3)	OJ L 147, 31.5.2001, p. 1.						
-	The signature and the stamp must be in a different colour to that of the printing.						
-	<ul> <li>Note for the person responsible for the consignment in the European Union: this certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.</li> </ul>						
Off	Official veterinarian/Official inspector						
	Name (in capital letters):	Qualification an	nd title:				
	Date: Signature:						
	Stamp:						

# CHAPTER 5(A)

# Health certificate

For fresh or chilled hides and skins of ungulates, intended for dispatch to or for transit through  $\binom{2}{2}$  the European Union

cou	NTRY	1	Veterinary certificate to EU			
	l.1.	Consignor	I.2. Certificate reference No I.2.a.			
		Name	I.3. Central competent authority			
		Address	,			
		Tel.	I.4. Local competent authority			
	1.5.	Consignee	I.6. Person responsible for the load in EU			
nent	1.0.	Name	Name			
ignr		Address	Address			
ons		Post of	Postcode			
ed c		Postcode Tel.	Tel.			
dispatched consignment	1.7	Country of origin 100 and 10 Barrian of origin Code	LO Country of ICO and I LO Borion of Code			
dsip	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10. Region of Code destination			
οţο						
etails	l.11.	Place of origin	I.12. Place of destination			
Part I: Details of		Name Approval number Address	Name Custom warehouse ☐ Address Approval number			
ď		Name Approval number Address	Postcode			
		Name Approval number Address				
	I.13.	Place of loading	I.14. Date of departure			
	l.15.	Means of transport	I.16. Entry BIP in EU			
		Aeroplane ☐ Ship ☐ Railway wagon ☐				
		Road vehicle Other	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	I.22. Number of packages			
		Ambient Chilled Chilled	Frozen			
	1.23.	Seal/Container No	I.24. Type of packaging			
	1.25.	Commodities certified for:	·			
		Animal feedingstuff ☐ Technical use ☐				
	1.26.	For transit through EU to third country	I.27. For import or admission into EU			
		Third country ISO code	_			
	1.28.	Identification of the commodities				
		Species Approval number (Scientific name) Manufactu				

COUNTRY Fresh or chilled hides and skins of ungulates II.a. Certificate reference No II. Health information I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council (<sup>1a</sup>) and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011 (<sup>1b</sup>), and in particular Annex XIV, Chapter II thereof, and certify that the hides and skins described above: have been obtained from animals that: Certification (2) either [- were slaughtered and their carcases are fit for human consumption in accordance with Union legislation;] [- were slaughtered in a slaughterhouse, after undergoing ante-mortem inspection, and were considered fit, as a result of such inspection, for slaughter for human consumption in accordance with Union legislation;] (2) or Part II: originate from a country or, in the case of regionalisation in accordance with Union legislation, from a part of a country from which imports 11.2. of all categories of fresh meat of the corresponding species are authorised and which: for at least 12 months before dispatch, has been free from the following diseases (3): (a) [- classical swine fever, and African swine fever;] [- rinderpest;] and (b) has been free for at least 12 months before dispatch from foot-and-mouth disease and where, for 12 months before dispatch, no vaccination has been carried out against foot-and-mouth disease (3); II.3. have been obtained from: [animals that have remained in the territory of the country of origin for at least three months before being slaughtered or since birth in the case of animals less that three months old:1 [in the case of hides and skins from bi-ungulates, animals that come from holdings in which there has been no outbreak of foot-and-mouth disease in the previous 30 days, and around which within a radius of 10 km there has been no case of foot-and-mouth disease for 30 [in the case of hides and skins from swine, animals that come from holdings in which there has been no outbreak of swine vesicular disease in the previous 30 days, or of classical or African swine fever in the previous 40 days, and around which within a radius of 10 km there has been no case of these diseases for 30 days;] [animals that have shown no evidence of [foot-and-mouth disease], [rinderpest], [classical swine fever], [African swine fever] or [swine vesicular disease] (3) during ante-mortem health inspection at the slaughterhouse during the 24 hours before slaughter;] II 4 have undergone all precautions to avoid contamination with pathogenic agents. Notes Part I: Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity. 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. The products in transit can only be stored in free zones, free warehouses and custom warehouses Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in the event of unloading and reloading.

Box reference I.19: use the appropriate HS code: 41.01; 41.02 or 41.03.

COUNTRY	Fresh or	chilled hides and skins of ungulates				
II. Health information	II.a. Certificate reference No	II.b.				
Box reference I.23: for bulk containers, the container number and	the seal number (if applicable) show	uld 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 tra	Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.					
Part II:						
( <sup>1a)</sup> OJ L 300, 14.11.2009, p. 1.						
( <sup>1b</sup> ) OJ L 54, 26.2.2011, p. 1.						
(²) Delete as appropriate.						
(3) Delete diseases not applicable to the species concerned.						
— The signature and the stamp must be in a different colour to that	t of the printing.					
Note for the person responsible for the consignment in the Eu accompany the consignment until it reaches the border inspection.		ly for veterinary purposes and has to				
Official veterinarian/Official inspector						
Name (in capital letters):	Qualification and	d title:				
Date: Signature:						
Stamp:						

# CHAPTER 5(B)

# **Health certificate**

For treated hides and skins of ungulates, intended for dispatch to or for transit through  $\binom{2}{2}$  the European Union

cou	UNTRY Veterinary certificate to EU								
	I.1.	Consignor	1.2.	Certificate	e reference N	lo	1.:	2.a.	
		Name	I.3. Central competent authority						
		Address		Ochtrar o	ompotont dat	inonty			
			1.4.	Local cor	mpetent author	ority			
		Tel.	_						
鸉	I.5.	Consignee	1.6.	Person r	esponsible fo	r the loa	d in E	U	
Ĕ		Name		Name					
igi		Address		Address					
[ 등		Destroyle		Deeterde					
8		Postcode Tel.		Postcode Tel.	,				
of dispatched consignment			_						
sps	1.7.	Country of origin ISO code I.8. Region of origin Code	1.9.	Country destination		ISO code	I.10.	Region of destination	Code
Ē				destinatio	on I	code		destination	
<u>s</u>			-						
Part I: Details	l.11.	Place of origin	1.12.	Place of	destination				
≘		Name Approval number		Name			Cu	stom warehouse	ı
ᆲ		Address		Address			Ap	proval number	.
		Name Approval number							
		Address		Postcode	•				
		Name Approval number Address							
	112	Place of loading	114	Date of o	donarturo				
	1.10.	Place of loading	1.14.	Date of t	Jopanuio				
Н	L 15	Means of transport	I.16. Entry BIP in EU						
		•		Linay Dii	20				
		Aeroplane Ship Railway wagon Railway wagon							
		Road vehicle Other I	I.17. Number(s) of CITES						
		Documentation references							
	1.40				140.0		1- 016	S 4-3	
	1.18.	Description of commodity	I.19. Commodity code (HS code)						
						1.20.	Quan	tity	
	101	Tamasandana of acadast	I.22. Number of packages						
	1.21.	Temperature of product	F			1.22.	MUMB	e or packages	
			Frozer						
	1.23.	Seal/Container No				1.24.	Туре	of packaging	
	1.25.	Commodities certified for:							
		Animal feedingstuff ☐ Technical use ☐							
		_	_						
	1.26.	For transit through EU to third country	1.27.	For impo	rt or admission	on into E	U		,
		Third country ISO code							
	1.65	International Control of the account of the control							
	1.28.	Identification of the commodities							
		Species Approval number			ts			Net weig	ght
		(Scientific name) Manufactu	ring pl	ant					

COUNTRY Treated hides and skins of ungulates Health information 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 10 thereof, and Commission Regulation (EU) No 142/2011 (1b), and in particular Annex XIV, Chapter II thereof, and certify that the hides and skins described above: II.1. have been obtained from animals that: Part II: Certification (2) either [- were slaughtered and their carcases are fit for human consumption in accordance with Union legislation;] (2) or [- were slaughtered in a slaughterhouse, after undergoing ante-mortem inspection, and were considered fit, as a result of such inspection, for slaughter for human consumption in accordance with Union legislation;] (2) or [- did not show any clinical signs of any disease communicable to humans or animals through the hide or skin, and were not killed to eradicate any epizootic disease;] [II.2. come from animals originate from a third country or, in the case of regionalisation in accordance with Union legislation, from a (2) either part of a third country listed in Part 1 of Annex II to Commission Regulation (EU) No 206/2010 (3) from which imports of fresh meat of the corresponding species are authorised and have been: (2) either [dried:1 (2) or [dry-salted or wet-salted for at least 14 days prior to dispatch;] (2) or [dry-salted or wet-salted on the following date ... .. and according to the declaration of the transporter, the hides and skins will be transported by ship and the duration of transport will be such that they will have undergone a minimum of 14 days of salting before they reach the EU border inspection post;] (2) or [salted for seven days in sea salt with the addition of 2 % of sodium carbonate:] (2) or of transport will be such that they will have undergone a minimum of seven days of salting before they reach the EU border inspection post.]] (2) or [II.2. come from animals originate from a third country or, in the case of regionalisation in accordance with Union legislation, from a part of a third country listed in Part 1 of Annex II to Regulation (EU) No 206/2010 from which imports of fresh meat of the corresponding species are NOT authorised and have been: (2) either [salted for seven days in sea salt with the addition of 2 % of sodium carbonate;] (2) or [salted in sea salt with the addition of 2 % of sodium carbonate on the following date and according to the declaration of the transporter, the hides and skins will be transported by ship and the duration of transport will be such that they will have undergone a minimum of seven days of salting before they reach the EU border inspection post;] (2) or [dried for 42 days at a temperature of at least 20 °C;]] II.3. the consignment has not been in contact with other animal products or with live animals presenting a risk of spreading a serious Notes Part I: Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity.

Treated hides and skins of ungulates

COUNTRY

Status: This is the original version (as it was originally adopted).

П.	Health information	II.a. Certificate reference No II.b.					
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.							
	<ul> <li>Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses.</li> </ul>						
	<ul> <li>Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) and information is to be provided in the event of unloading and reloading.</li> </ul>						
— Во	— Box reference I.19: use the appropriate HS code: 41.01; 41.02 or 41.03.						
— Во	x reference I.23: for bulk containers, the container number and the	e seal number (if applicable) should be	e given.				
— Во	x reference I.25: technical use: any use other than for animal con-	sumption.					
— Во	x reference I.26 and I.27: fill in according to whether it is a transit	or an import certificate.					
Part I	l:						
( <sup>1a</sup> ) O	J L 300, 14.11.2009, p. 1.						
( <sup>1b</sup> ) O	J L 54, 26.2.2011, p. 1.						
(²) D	elete as appropriate.						
( <sup>3</sup> ) O	J L 73, 20.3.2010, p. 1.						
( <sup>4</sup> ) O	J L 147, 31.5.2001, p. 1.						
— Th	e signature and the stamp must be in a different colour to that of	the printing.					
	ote for the person responsible for the consignment in the Europ company the consignment until it reaches the border inspection po		r veterinary purposes and has to				
Officia	al veterinarian/Official inspector						
\ \	lame (in capital letters):	Qualification and	i title:				
0	Date:	Signature:					
s	Stamp:						

# CHAPTER 5(C)

# Official declaration

For treated hides and skins of ruminants and of equidae that are intended for dispatch to or for transit through (1) the European Union and have been kept separate for 21 days or will undergo transport for 21 uninterrupted days before importation

cou	OUNTRY Veterinary certificate to EU							
	l.1.	Consignor	I.2. Certificate reference No I.2.a.					
		Name	I.3. Central competent authority					
		Address	- Company					
		Tel.	I.4. Local competent authority					
_	15	Consignee	I.6. Person responsible for the load in EU					
le l	1.0.	Name	Name					
l g		Address	Address					
Suc								
Ö		Postcode	Postcode					
dispatched consignment		Tel.	Tel.					
spa	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO I.10. Region of Code					
of di			destination code destination					
lls o		Place of orbits	Liab Share of the first first					
Part I: Details	1.11.	Place of origin	I.12. Place of destination					
=		Name Approval number	Name Custom warehouse					
Part		Address	Address Approval number					
_		Name Approval number Address						
		Name Approval number	Postcode					
		Address						
	I.13.	Place of loading	I.14. Date of departure					
	l.15.	Means of transport	I.16. Entry BIP in EU					
		Aeroplane ☐ Ship ☐ Railway wagon ☐						
		Road vehicle Other	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	I.22. Number of packages					
		Ambient Chilled C	Frozen					
	1.23.	Seal/Container No	I.24. Type of packaging					
	1.25.	Commodities certified for:	·					
		Animal feedingstuff Technical use						
	1.26.	For transit through EU to third country	I.27. For import or admission into EU					
		Third country ISO code						
	1.28.	Identification of the commodities						
			ber of establishments Net weight acturing plant					

# Treated hides and skins of ruminants and of equidae that have been kept separate for 21 days or will undergo transport for 21 uninterrupted days before importation

cou	INTRY				kept separate for 21 days or uninterrupted days before importar	will undergo transport for 21
	II.	Healt	h informatio	on	II.a. Certificate reference No	II.b.
			I, the under	rsigned declare that the hides and skins	described above:	
		II.1.	have been	obtained from animals that:		
			(1) either	[- were slaughtered and their carcase	s are fit for human consumption in a	accordance with Union legislation;]
cation			( <sup>1</sup> ) or	[- were slaughtered in a slaughterhous result of such inspection, for slaughternoons and statement of the st	e, after undergoing ante-mortem inspec er for human consumption in accorda	
Part II: Certification			( <sup>1</sup> ) or	[- did not show any clinical signs of ar and were not killed to eradicate any		r animals through the hide or skin,
art		II.2.	have been:			
٦			(1) either	[- dried;]		
			( <sup>1</sup> ) or	[- dry-salted or wet-salted for at least	14 days prior to dispatch;]	
			( <sup>1</sup> ) or	[- salted for seven days in sea salt wit	th the addition of 2 % of sodium carbo	onate;]
		II.3.	have not by transmissible	peen in contact with other animal pro- le disease;	ducts or with live animals presenting	ng a risk or spreading a serious
	(2) either	[II.4.	have been under point	kept separate immediately before disparate II.2.]	atch for 21 days under official supervi	ision after the treatment described
	(²) or	[II.4.	following th	e declaration of the transporter, the dur	ation of the transport period is foresee	en to be at least 21 days.]
	Notes					
	Part I:					
				sponsible for the consignment in the Eu n if the certificate is for import commod		n only if it is a certificate for transit
	<ul> <li>Box refer authority.</li> </ul>		1 and I.12: /	Approval number: the registration number	er of the establishment or plant, which	has been issued by the competent
				destination: this box is to be filled in only warehouses and custom warehouses.	if it is a certificate for transit commod	ity. The products in transit can only
				ion number (railway wagons or containe ading and reloading.	r and lorries), flight number (aircraft) o	or name (ship); information is to be
	— Box refer	ence I.	9: use the a	appropriate HS code: 41.01; 41.02 or 4	1.03.	
	— Box refer	ence I.2	3: for bulk o	containers, the container number and th	e seal number (if applicable) should b	e given.
	— Box refer	ence I.2	25: technical	use: any use other than for animal cor	sumption.	
	— Box refer	ence I.2	26 and I.27:	fill in according to whether it is a transi	t or an import certificate.	
	Part II:					
	(1) Delete as	appro	oriate.			
	— The signs	ature ar	d the stamp	must be in a different colour to that of	the printing.	
				sible for the consignment in the Europe until it reaches the border inspection p		or veterinary purposes and has to

COUNTRY	Treated hides and skins of ruminants and of equidae that have been kept separate for 21 days or will undergo transport for 21 uninterrupted days before importation
II. Health information	II.a. Certificate reference No II.b.
Official veterinarian/Official inspector	
Name (in capital letters):	Qualification and title:
Date:	Signature:
Stamp:	

# CHAPTER 6(A)

# Health certificate

For treated game trophies and other preparations of birds and ungulates, being solely bones, horns, hooves, claws, antlers, teeth, hides or skins, for dispatch to or for transit through  $\binom{2}{}$  the European Union

COUNTRY Veterinary certificate to						ate to EU		
	I.1.	Consignor	1.2.	Certificate	reference N	o	I.2.a.	
		Name	1.3.	I.3. Central competent authority				
		Address	nor contract during					
		Tel.	I.4. Local competent authority					
	1.5		10	D			4 in FII	
len	1.5.	Consignee	1.6.		esponsible fo	r the loa	d in EU	
l m		Name Address		Name Address				
nsi		Addiess		Address				
8		Postcode		Postcode	•			
che		Tel.		Tel.				
dispatched consignment	1.7.	Country of origin ISO code I.8. Region of origin Code	1.9.	Country destination		ISO code	I.10. Region of destination	Code
o d				dostinatio	"'	code	destination	
l: Details	l.11.	Place of origin	I.12.	Place of	destination			
Part I: D		Name Approval number Address		Name Address			Custom warehouse  Approval number	
۵		Name Approval number Address		Postcode				
		Name Approval number Address						
	I.13.	Place of loading	l.14.	Date of o	departure			
	l.15.	Means of transport	I.16. Entry BIP in EU					
		Aeroplane						
		Road vehicle Other	I.17. Number(s) of CITES					
		Identification  Documentation references						
							1- (10 1-)	
	1.18.	Description of commodity			1.19. Commo	odity cod	de (HS code)	
						1.20.	Quantity	
	I.21.		I.22. Number of packages					
	1.23.	Seal/Container No	I.24. Type of packaging					
	1.25.	Commodities certified for:						
		Technical use						
	1.26.	For transit through EU to third country	I.27. For import or admission into EU			ı		
		Third country ISO code						
	1.28.	Identification of the commodities						
		Species Nature of (Scientific name)	comm	odity			Number of packa	ges

Treated game trophies and other preparations of birds and ungulates, being solely bones, horns, hooves, claws, antlers, COUNTRY teeth, hides or skins Health information 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 (<sup>1a</sup>) and Commission Regulation (EU) No 142/2011 (<sup>1b</sup>), and in particular Annex XIV, Chapter II thereof, and certify that the game trophies described above: II.1. have been packaged, immediately after treatment, without being in contact with other products of animal origin likely to contaminate them, in individual, transparent and closed packages so as to avoid any subsequent contamination; Certification (2) either [II.2. in the case of game trophies or other preparations consisting solely of hides or skin: (2) either [have been dried:] Part II: (2) or [have been dry-salted or wet-salted for a minimum of 14 days before dispatch;] (2) or (date) and, according to the declaration of the trans-[were dry-salted or wet-salted on .. porter, will be transported by ship and the duration of the transport will be such that they will have undergone a minimum of 14 days salting before they reach the EU border inspection post;]] (2) or [II.2. in the case of game trophies or other preparations consisting solely of bone, horns, hooves, claws, antlers or teeth: (a) have been immersed in boiling water for an appropriate time so as to ensure that any matter other than bone, horns, hooves, (b) have been disinfected with a product authorised by the competent authority, in particular with hydrogen peroxide where parts consisting of bone are concerned.] II.3. (2) either If the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001 of the European Parliament and of the Council (3) or mechanically separated meat obtained from bones of bovine, ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial (2) or [the product does not contain and is not derived from bovine, ovine or 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 by a decision in accordance with Article 5(2) of Regulation (EC) No 999/2001.] Notes Part I: Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity. Box reference I.11 and I.12: Approval number: the registration number of the establishment or plant, which has been issued by the competent

Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only

Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be

Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.

authority.

be stored in free zones, free warehouses and custom warehouses.

provided in case of unloading and reloading

Treated game trophies and other preparations of birds and

Status: This is the original version (as it was originally adopted).

COUNTRY	teeth, hides or skins	norns, nooves, claws, antiers,				
II. Health information	II.a. Certificate reference No	II.b.				
— Box reference I.19: use the appropriate HS code: 05.05; 05.06; 05.07 or 97.05.						
— 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.						
<ul> <li>Box reference I.28: for nature of commodity, specify choosing one or more possibilities among the following: [bones], [horns], [hooves], [claws [antiers], [teeth], [hides] or [skins].</li> </ul>						
Part II:						
( <sup>1a</sup> ) OJ L 300, 14.11.2009, p. 1.						
( <sup>1b</sup> ) OJ L 54, 26.2.2011, p. 1.						
(²) Delete as appropriate.						
(³) OJ L 147, 31.5.2001, p. 1.						
— The signature and the stamp must be in a different colour to that of the printing.						
<ul> <li>Note for the person responsible for the consignment in the European Union: this certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.</li> </ul>						
Official veterinarian/Official inspector						
Name (in capital letters):	Qualification and	d title:				
Date:	Signature:					
Stamp:						

# CHAPTER 6(B)

# Health certificate

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

cou	COUNTRY Veterinary certificate to EU										
	l.1.	1. Consignor				I.2. Certificate reference No I.2.a.					
		Name			I.3. Central competent authority						
		Address			1.3. Central competent authority						
						I.4. Local competent authority					
		Tel.									
ent	1.5.	i. Consignee			I.6. Person responsible for the load in EU						
Е	Name			Name							
Isig		Address				Address					
8		Postcode Tel.			Postcode Tel.						
Je d											
dispatched consignment		0	Busham of salata							0	
lsb	1.7.	Country of origin ISO code I.8	. Region of origin	Code	1.9.	Country destination	of on	ISO code	1.10.	Region of destination	Code
of							···				I
l: Details	1.11.	Place of origin			1.12	Place of	destination				I
Deta		r lass or origin				1 1000 01	doomaton				
<b>=</b>			proval number						Custom warehouse  Approval number		
Part		Address				Addioss			API	orovar mamber	
	Name Approval number Address Name Approval number					Postcode					
						rosicode	,				
		Address									
	I.13.	Place of loading			I.14.	Date of	departure				
	1.15.	I.15. Means of transport				I.16. Entry BIP in EU					
	Aeroplane Ship Railway wagon										
	Road vehicle Other Identification Documentation references  I.18. Description of commodity				I.17. Number(s) of CITES						
					I.19. Commodity code (HS code)						
								1.20.	Quant	ity	
	I.21.							1.22.	Numb	er of packages	
	I.23. Seal/Container No I.25. Commodities certified for:					ı			I.24. Type of packaging		
		Technical use									
	Teominal ass										
	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					Number of packages					
		(Scientific name)									

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

	II.	н	ealth	nformati	on	II.a. Certificate reference No	II.b.						
			Parlia	ament an	gned official veterinarian, declare that I have d of the Council ( <sup>1a</sup> ) and Commission Regu ertify that the game trophies described above:	ılation (EU) No 142/2011 ( <sup>1b</sup> ), and in	c) No 1069/2009 of the European particular Annex XIV, Chapter II						
_	( <sup>2</sup> )	either	[II.1.	with res	pect to game trophies or other preparations o	f cloven-hoofed animals, excluding swi	ine:						
Part II: Certification		(a) (region) has been free from foot-and-mouth disease and rinderpest for the previous 12 months, and due the same period, no vaccination against any of those diseases has taken place; and											
t ⊪.				(b) the	ne trophies or other preparations described above:								
Par				ť	vere obtained from animals which were killed in the corresponding susceptible domestic specie estrictions because of outbreaks of diseases	es and where, during the last 60 days,	there have been no animal health						
					originated from animals that were killed at a dis a third country not authorised to export untreate								
	( <sup>2</sup> )	or	[II.1.	with res	pect to game trophies or other preparations o	f wild swine:							
				dise	ase, foot-and-mouth disease and porcine enter add out against any of those diseases during t	roviral encephalmiyelitis (Teschen disea	African swine fever, swine vesicular ise) and no vaccinations have been						
				(b) the	game trophies or other preparations described	d above:							
				" (	vere obtained from animals which were killed corresponding susceptible domestic species a estrictions because of outbreaks of diseases	and where, during the last 60 days, t	there have been no animal health						
					originated from animals that were killed at a dis a third country not authorised to export untreal								
(2) or [II.1. with respect to game trophies or other preparations of solipeds, the game trophies or other preparations described about obtained from wild solipeds that were killed in the territory of the exporting country mentioned above;]													
	(2) or [II.1. with respect to game trophies or other preparations of game birds:												
	(a) (region) is free from highly pathogenic avian influenza and Newcastle disease; and												
	<ul> <li>(b) the game trophies or other preparations described above were obtained from wild game birds that were killed in that read where during the last 30 days there have been no animal health restrictions because of outbreaks of disease to the wild birds are susceptible;]</li> <li>II.2. The game trophies or other preparations described above have been packaged without being in contact with other production animal origin likely to contaminate them, in individual, transparent and closed packages so as to avoid any subsequent contamination.</li> </ul>												
			II.3.										
				( <sup>2</sup> ) either	[the product does not contain and is not der No 999/2001 of the European Parliament at of bovine, ovine or caprine animals; and the after stunning by means of gas injected laceration of central nervous tissue by me cavity.]	nd of the Council (3) or mechanically se ne animals from which this product is a into the cranial cavity or killed by the	eparated meat obtained from bones derived have not been slaughtered a same method or slaughtered by						
(²) or [the product does not contain and is not derived from bovine, ovine or caprine materials other than those derived animals born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE a decision in accordance with Article 5(2) of Regulation (EC) No 999/2001.]													

cour	ITRY	Game trophies or o consisting of entire pa			and ungulates
II.	Health information	II.a. Certificate referenc	e No	II.b.	
Note	s				
Part	l:				
	ox reference I.6: Person responsible for the consignment in the Euronmodity; it may be filled in if the certificate is for import commodity		s to be filled in	only if it is a certi	ficate for transit
	iox reference I.11 and I.12: Approval number: the registration number uthority.	er of the establishment or	r plant, which I	has been issued by	the competent
	ox reference I.12: Place of destination: this box is to be filled in only e stored in free zones, free warehouses and custom warehouses.	if it is a certificate for tra	ansit commodi	ty. The products in	transit can only
	ox reference I.15: Registration number (railway wagons or container rovided in case of unloading and reloading.	r and lorries), flight numl	ber (aircraft) o	r name (ship); infor	rmation is to be
— в	ox reference I.19: use the appropriate HS code: 05.05; 05.06 or 05	5.07.			
— в	ox reference I.23: for bulk containers, the container number and the	e seal number (if applica	able) should be	e included.	
— в	ox reference I.25; technical use; any use other than for animal con-	sumption.			
— в	tox reference I.26 and I.27: fill in according to whether it is a transit	or an import certificate.			
Part	II:				
( <sup>1a</sup> )	OJ L 300, 14.11.2009, p. 1.				
( <sup>1b</sup> )	OJ L 54, 26.2.2011, p. 1.				
(²)	Delete as appropriate.				
(3)	OJ L 147, 31.5.2001, p. 1.				
— т	he signature and the stamp must be in a different colour to that of	the printing.			
	lote for the person responsible for the consignment in the European Une consignment until it reaches the border inspection post.	Jnion: this certificate is or	nly for veterina	ry purposes and ha	s to accompany
Offic	ial veterinarian/Official inspector				
	Name (in capital letters):	Qu	ualification and	title:	
	Date:	Sig	gnature:		
	Stamp:				

# CHAPTER 7(A)

# Health certificate

For pig bristles from third countries or regions thereof that are free from African swine fever, intended for dispatch to or for transit through  $\binom{2}{1}$  the European Union

cou	OUNTRY Veterinary certificate to EU				
	I.1.	Consignor	I.2. Certificate reference No I.2.a.		
		Name	I.3. Central competent authority		
		Address			
		Tel.	I.4. Local competent authority		
ᇦ	1.5.	Consignee	I.6. Person responsible for the load in EU		
mer	1.5.	Name	Name		
igu		Address	Address		
l oi					
be		Postcode	Postcode		
tch		Tel.	Tel.		
ispa	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO I.10. Region of Code		
of d			destination code destination		
ils					
Deta	1.11.	Place of origin	I.12. Place of destination		
Part I: Details of dispatched consignment		Name Approval number Address	Name Custom warehouse ☐ Address Approval number		
Par		Name Approval number	The state of the s		
		Address	Postcode		
		Name Approval number Address	Tostcode		
	l.13.	Place of loading	I.14. Date of departure		
	l.15.	Means of transport	I.16. Entry BIP in EU		
		Aeroplane Ship Railway wagon			
		Road vehicle Other	1.17.		
		Identification			
		Documentation references			
	I.18.	Description of commodity	I.19. Commodity code (HS code) 05.02		
			I.20. Quantity		
	I.21.	Temperature of product	I.22. Number of packages		
		Ambient Chilled Chilled	Frozen		
	I.23.	Seal/Container No	I.24. Type of packaging		
	1.25.	Commodities certified for:			
		Animal feedingstuff ☐ Technical use ☐			
	1.26.	For transit through EU to third country	I.27. For import or admission into EU		
		Third country ISO code			
	1.28.	Identification of the commodities			
		Approval number of establishments Nun Manufacturing plant	nber of packages Net weight		

# Pig bristles from third countries or regions thereof that are free from COUNTRY African swine fever Health information II.a. Certificate reference No II.b. I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council (<sup>1a</sup>) and in particular Article 10(b)(iv) thereof, and Commission Regulation (EU) No 142/2011 (<sup>1b</sup>), and in particular Annex XIV, Chapter II thereof, and certify that: II.1. the pig bristles described above have been obtained from pigs originating, and slaughtered in a slaughterhouse, in the country of origin; the pigs, from which the pig bristles have been obtained, did not show during inspection, carried out at the time of slaughtering, signs of diseases communicable to humans or animals and were not killed to eradicate any epizootic disease; 11.2. Certification II.3. the country of origin or, in case of regionalisation according to Union legislation, the region of origin, has been free from African swine fever for at least 12 months; ≝ II.4. the pig bristles are dry and securely enclosed in packaging. Part Notes Part I: Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity. 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. The products in transit can only be stored in free zones, free warehouses and custom warehouses. Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading. - Box reference I.23; for bulk containers, the container number and the seal number (if applicable) should be included. Box reference I.25: technical use: any use other than for animal consumption. Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate. - Box reference I.28: Manufacturing plant: provide the veterinary control number of the registered 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. - 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 has to 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 7(B)

# **Health certificate**

For pig bristles from third countries or regions thereof that are not free from African swine fever, intended for dispatch to or for transit through  $\binom{2}{1}$  the European Union

COU	DUNTRY Veterinary certificate to EU					
	l.1.	Consignor	I.2. Certificate reference No I.2.a.			
		Name				
		Address	I.3. Central competent authority			
		71001000	I.d. Local comments to the other			
		Tel.	I.4. Local competent authority			
_	1.5.	Consignee	I.6. Person responsible for the load in EU			
Jen	1.5.	•				
Ē		Name	Name			
ısić		Address	Address			
8		Postcode	Postcode			
eq		Tel.	Tel.			
달		10	10.1			
of dispatched consignment	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO I.10. Region of Code			
ä			destination code destination			
5						
Part I: Details	I.11.	Place of origin	I.12. Place of destination			
Det			_			
Ξ		Name Approval number	Name Custom warehouse ☐ Address Approval number			
Par		Address	Approval number			
		Name Approval number Address				
			Postcode			
		Name Approval number Address				
			Late Date of James 1			
	1.13.	Place of loading	I.14. Date of departure			
	l.15.	Means of transport	I.16. Entry BIP in EU			
		Aeroplane ☐ Ship ☐ Railway wagon ☐				
		Road vehicle Other	1.5			
		Identification	1.17.			
		Documentation references				
	I.18.	Description of commodity	I.19. Commodity code (HS code) 05.02			
			I.20. Quantity			
	1.01	Temperature of product	I.22. Number of packages			
	1.21.					
		Ambient Chilled C	Frozen			
	I.23.	Seal/Container No	I.24. Type of packaging			
	1.25.	Commodities certified for:				
		Animal feedingstuff ☐ Technical use ☐				
		Animal reedingston				
	1.26.	For transit through EU to third country	I.27. For import or admission into EU			
		Third country ISO code				
	1.28.	Identification of the commodities	1			
		Approval number of establishments Num	ber of packages Net weight			
		Manufacturing plant	ivor or packages			

# Pig bristles from third countries or regions thereof that are not free from African swine fever

II.b.

### COUNTRY

I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council (<sup>1a</sup>) and in particular Article 10(b)(iv) thereof, and Commission Regulation (EU) No 142/2011 (<sup>1b</sup>), and in particular Annex XIV, Chapter II thereof, and certify that:

II.a. Certificate reference No

II.1. the pig bristles described above have been obtained from pigs originating, and slaughtered in a slaughterhouse, in the country of origin;

II: Certification

Part

II.2.

the pigs from which the pig bristles have been obtained did not show during inspection, carried out at the time of slaughtering, signs of diseases communicable to humans or animals and were not killed to eradicate any epizootic disease;

II.3. the pig bristles mentioned above have been:

(2) either [boiled;]

Health information

(2) or [dyed;]

(2) or [bleached;]

II.4. the pig bristles are dry and securely enclosed in packaging.

Notes

### Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit
  commodity; it may be filled in if the certificate is for import commodity.
- 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. The products in transit can only be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.
- Box reference I.25: technical use: any use other than for animal consumption.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: Manufacturing plant: provide the veterinary control number of the registered 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.
- 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 has to accompany
  the consignment until it reaches the border inspection post.

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

# CHAPTER 8

# Health certificate

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

cou	OUNTRY Veterinary certificate to EU				
	I.1.	Consignor	I.2. Certificate reference No I.2.a.		
		Name	I.3. Central competent authority		
		Address	no. Soma composite authority		
		Tel.	I.4. Local competent authority		
_	1.5.	Consignee	I.6. Person responsible for the load in EU		
neu	1.5.	Name	Name		
dispatched consignment		Address	Address		
			,		
		Postcode	Postcode		
tche		Tel.	Tel.		
spa	I.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO I.10. Region of Code		
of di			destination code destination		
ls o					
Part I: Details	l.11.	Place of origin	I.12. Place of destination		
<u></u>		Name Approval number	Name Custom warehouse ☐		
art		Address	Address Approval number		
-		Name Approval number Address			
		Name Approval number	Postcode		
		Address			
	I.13.	Place of loading	I.14. Date of departure		
	I.15.	Means of transport	I.16. Entry BIP in EU		
		Aeroplane ☐ Ship ☐ Railway wagon ☐			
		Road vehicle Other	1.17.		
		Identification			
		Documentation references			
	I.18.	Description of commodity	I.19. Commodity code (HS code)		
			I.20. Quantity		
	I.21.	Temperature of product	I.22. Number of packages		
		Ambient ☐ Chilled ☐	Frozen 🗆		
	1.23.	Seal/Container No	I.24. Type of packaging		
	1.25.	Commodities certified for:	'		
		Technical use □			
			_		
	1.26.	For transit through EU to third country	I.27. For import or admission into EU		
		Third country ISO code			
	1.28.	Identification of the commodities			
		Species Nature of commodity Approval number of e Manufacturing			

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

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.	
		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council ( <sup>1a</sup> ) and Commission Regulation (EU) No 142/2011 ( <sup>1b</sup> ), and in particular Annex XIV, Chapter II thereof, and certify that the animal by-products described above:	
_	II.1.	are trade samples which consist of animal by-products intended for particular studies or analyses as referred to in definition No 39 of Annex I to Commission Regulation (EU) No 142/2011, that are bearing the label 'TRADE SAMPLE NOT FOR HUMAN CONSUMPTION'; or	
Certification	II.2.	satisfy the animal health requirements below;	
Certif	II.2.1.	have been obtained in the territory of:(3) from animals:	
Part II:		(2) either [(a) that have remained in this territory since birth or for at least the last three months before slaughter;]	
"		(2) or [(b) killed in the wild in this territory (4);]	
	II.2.2.	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 been neither case/outbrer rinderpest, swine vesicular disease, Newcastle disease or highly pathogenic avian influenza during the pric		
		<ul><li>(ii) where there has been neither case/outbreak of foot-and-mouth disease during the prior 60 days, nor in the holdings situated in their vicinity within 25 km, during the prior 30 days; and</li></ul>	
		(b) which:	
		(i) were not killed to eradicate any epizootic disease;	
		<ul><li>(ii) have remained in their holdings of origin for at least 40 days before departure and which have been transported directly to the slaughterhouse without contact with other animals which did not comply with the same health conditions;</li></ul>	
		(iii) at the slaughterhouse, have passed the ante-mortem health inspection during the 24 hours before the slaughter and have shown no evidence of the diseases referred to above for which the animals are susceptible; and	
		<ul><li>(iv) have been treated in the slaughterhouse before and at the time of slaughter or killing in accordance with the relevant provisions of Council Directive 93/119/EC (<sup>5</sup>) on the protection of animals at the time of slaughter or killing]</li></ul>	
		(a) or [(a) captured and killed in the wild in an area:	
		(i) in which within 25 km there has been no case/outbreak of any of the following diseases for which the animals are susceptible: foot-and-mouth disease, rinderpest, Newcastle disease or highly pathogenic avian influenza during the prior 30 days nor of classical or African swine fever during the prior 40 days; and	
		(ii) that is situated at a distance that exceeds 20 km from the borders separating another territory of a country or part thereof, which is not authorised at these dates for exporting this material to the European Union; and	
		(b) which after killing were transported within 12 hours for chilling either to a collection centre and immediately afterwards to a game establishment, or directly to a game establishment;]	
	II.2.3.	have been obtained in an establishment around which, within a radius of 10 km, there has been no case/outbreak of diseases referred to in point II.2.2 for which the animals are susceptible during the prior 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 removal of all meat, and the total cleaning and disinfection of the establishment under the control of an official veterinarian;	
	II.2.4. have been obtained and prepared without contact with other material not complying with the conditions required above, and it has been handled so as to avoid contamination with pathogenic agents;		

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

COUNTRY	j	or for trade samples (2)	
II.	Health information	II.a. Certificate reference No	II.b.
II.2.5.	have been packed in new packaging preventing any leak and in containers sealed under the responsibility of the co ONLY FOR THE MANUFACTURE OF DERIVED PRODUC of the EU establishment of destination;	empetent authority, bearing the label i	ndicating 'ANIMAL BY-PRODUCTS
II.2.6.	consist only of the following animal by-products:		
	<ul> <li>(²) either [- carcases and parts of animals slaughtered or for human consumption in accordance with commercial reasons;]</li> </ul>		
	(2) and/or [- carcases and the following parts originating were considered fit for slaughter for human following parts of animals from game killed f	n consumption following an ante-mo	rtem inspection or bodies and the
	(i) carcases or bodies and parts of animals Union legislation, but which did not show		
	(ii) heads of poultry;		
	<ul><li>(iii) hides and skins, including trimmings and and metacarpus bones, tarsus and meta</li></ul>		
	(iv) pig bristles;		
	(v) feathers;]		
	(2) and/or [- animal by-products arising from the product bone, greaves and centrifuge or separator s		consumption, including degreased
	(²) and/or [- products of animal origin, or foodstuffs contaconsumption for commercial reasons or due which no risk to public or animal health arise	to problems of manufacturing or pack	
	(²) and/or [- aquatic animals, and parts of such animals, enicable to humans or animals;]	except sea mammals, which did not s	how any signs of diseases commu-
	(²) and/or [- animal by-products from aquatic animals original consumption;]	ginating from establishments or plants	manufacturing products for human
	(²) and/or [- the following material originating from anima material to humans or animals:	Is which did not show any signs of o	disease communicable through that
	(i) shells from shellfish with soft tissue or flo	esh;	
	(ii) the following originating from terrestrial a	nimals:	
	<ul><li>hatchery by-products;</li></ul>		
	— eggs;		
	<ul> <li>egg by-products, including egg shells</li> </ul>	;	
	(iii) day-old chicks killed for commercial reas	sons;]	
	(²) and/or [- fur originating from dead animals that did not humans or animals;]	show clinical signs of any disease co	mmunicable through that product to
II.2.7.	have been deep-frozen at the plant of origin or have been not spoil between dispatch and delivery to the plant of de		gislation in such a way that they will

# Animal by-products to be used for purposes outside the feed chain

### COUNTRY or for trade samples (2) Health information II.a. Certificate reference No. II.b. (2) (6) [II.2.8. Specific requirements

- (²) (ĭ) [II.2.8.1. The by-products in this consignment come from animals that have been obtained in the territory mentioned under (II.2.1), where vaccination programmes against foot-and-mouth disease are being regularly carried out and officially controlled in domestic bovine
- (2) (8) [II.2.8.2. The by-products in this consignment consist of animal by-products derived from offal or deboned meat.]

### II.2.9.

- (2) either [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001 of the European Parliament and of the Council (9) or mechanically separated meat obtained from bones of bovine, ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity;]
- [the product does not contain and is not derived from bovine, ovine or 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 by a decision in accordance with Article 5(2) of Regulation (EC) No 999/2001;]

### II.2.10. in addition as regards TSE:

- (2) either [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin, the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last three years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfied the following requirements for the last three years:
  - (i) it has been subject to regular official veterinary checks;
  - (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or, following the confirmation of a classical scrapie case:
    - all animals in which classical scrapie was confirmed have been killed and destroyed, and
    - all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARR genotype and breeding ewes carrying at least one ARR allele and no VRQ allele;
  - (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are introduced into the holding only if they come from a holding which complies with the requirements set out in points (i) and (ii).]
- (2) or fin case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin, and destined to a Member State listed in the Annex to Commission Regulation (EC) No 546/2006 (<sup>10</sup>), the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last seven years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfied the following requirements for the last seven years:
  - (i) it has been subject to regular official veterinary checks;
  - (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or, following the confirmation of a classical scrapie case:
    - all animals in which classical scrapie was confirmed have been killed and destroyed, and
    - all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARR genotype and breeding ewes carrying at least one ARR allele and no VRQ allele;
  - (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are introduced into the holding only if they come from a holding which complies with the requirements set out in points (i) and (ii).]

### Notes

## Part I:

Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity

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

### COUNTRY

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

- Box reference I.11: In case of consignments for the particular technological studies or analyses: indicate name and address of establishment
- Box reference I.11 and I.12: Approval number: the registration number of the establishment or plant, which has been issued by the competent
- 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 transit commodity. The products
    in transit can only be stored in free zones, free warehouses and custom warehouses.
  - products for the particular technological studies or analyses: the EU plant indicated in authorisation of competent authority.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading.
- Box reference I.19: use the appropriate HS code: 05.11.91; 05.11.99 or 30.01.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.
- Box reference I.25: technical use: any use other than for animal consumption.
- Box reference I.25: for the purposes of the certificate, 'technical use' includes use as a trade sample.
- Box reference I.26 and I.27: except for trade samples, which are not sent in transit, fill in according to whether it is a transit or an import certificate.
- Box reference I.28:
  - products for the manufacture of derived products for uses outside the feed chain: Manufacturing plant: provide the veterinary control number
    of the approved establishment.
  - products for the particular technological studies or analyses: the EU plant indicated in authorisation of competent authority.

### 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) 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.
  - the Annex to Regulation (EC) No 798/2008, and
  - the Annex to Regulation (EC) No 119/2009.
  - In addition the ISO code of regionalisation in this Annex (where applicable for the susceptible species concerned) should be included.
- (4) Only for countries from where game meat intended for human consumption of the same animal species is authorised for importation into the European Union.
- (5) OJ L 340, 31.12.1993, p. 21.

COUN		Animal by-products to be used for or for trade samples (2)	purposes outside the feed chain
II.	Health information	II.a. Certificate reference No	II.b.
f	Supplementary guarantees to be provided when the material of don African country or part thereof from where only maturated and debon or exportation to the European Union. The whole masseter muscles on Part B(1) of Regulation (EC) No 854/2004 of the European Parliar	ed fresh meat of domestic ruminants for of bovine animals, incised in accordance	or human consumption is permitted e with Annex I, Section IV, Chapter
(7) (	Only for certain South American countries.		
(8) (	Only for certain South American and South African countries.		
(9) (	OJ L 147, 31.5.2001, p. 1.		
(10)	OJ L 94, 1.4.2006, p. 28.		
_ т	he signature and the stamp must be in a different colour to that of	the printing.	
	ote for the person responsible for the consignment in the European Unie consignment until it reaches the border inspection post.	Inion: this certificate is only for veterinal	ry purposes and has to accompany
Offici	ial veterinarian/Official inspector		
	Name (in capital letters):	Qualification and	I title:
	Date:	Signature:	
	Stamp:		

# CHAPTER 9

# Health certificate

For fish oil 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  $\binom{2}{2}$  the European Union

cou	NTR	•	Veterinary certificate	to EU	
	l.1.	Consignor	I.2. Certificate reference No I.2.a.		
		Name	I.3. Central competent authority		
		Address	I.d. Local compostant sutherity		
		Tel.	I.4. Local competent authority		
턽	1.5.	Consignee	I.6. Person responsible for the load in EU		
dispatched consignment		Name	Name		
nsig		Address	Address		
8		Postcode	Postcode		
tche		Tel.	Tel.		
ispa	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO I.10. Region of Code destination	ode	
₽			destination code destination		
l: Details	l.11.	Place of origin	I.12. Place of destination		
Part I: D		Name Approval number Address	Name Custom warehouse Address Approval number		
_		Name Approval number Address	Postcode		
		Name Approval number Address			
	I.13.	Place of loading	I.14. Date of departure		
	l.15.	Means of transport	I.16. Entry BIP in EU		
		Aeroplane Ship Railway wagon			
		Road vehicle Other I	1.17.		
		Documentation references			
	I.18.	Description of commodity	I.19. Commodity code (HS code)		
			I.20. Quantity		
	I.21.	Temperature of product	I.22. Number of packages		
		Ambient Chilled Chilled	Frozen		
	1.23.	Seal/Container No	I.24. Type of packaging		
	1.25.	Commodities certified for:			
		Animal feedingstuff Technical use			
	1.26.	For transit through EU to third country	I.27. For import or admission into EU		
		Third country ISO code			
	1.28.	Identification of the commodities			
		Nature of commodity Approval number of establishments Manufacturing plant	Number of packages Net weight Batch num	nber	

# Fish oil not intended for human consumption to be used as feed

COU	INTRY
	Ш

### material or for purposes outside the feed chain 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 Annex XIV, Chapter II thereof, and certify that the fish oil described above:

- II.1. consists of fish oil that satisfies the health requirements below;
- II.2. contains exclusively fish oil not intended for human consumption;
- II: Certification has been prepared and stored in a dedicated fish plant approved, validated and supervised by the competent authority in accordance with Article 24 of Regulation (EC) No 1069/2009; II.3.
- 11.4. has been prepared exclusively with the following animal by-products: Part
  - (2) either [- animal by-products arising from the production of products intended for human consumption;]
  - 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;]
  - aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of diseases commu-(2) and/or [nicable to humans or animals:1
  - (2) and/or [animal by-products from aquatic animals originating from plants or establishments manufacturing products for human
  - II.5. the fish oil:
    - (a) has been subjected to processing in accordance with Annex X, Chapter II, Section 3 of Regulation (EU) No 142/2011, in order to kill pathogenic agents;
    - (b) has not been in contact with other types of oils including rendered fats from any species of terrestrial animals, and
    - (2) either [(c) is packaged in new containers or in containers that have been cleaned and disinfected if necessary for the prevention of contamination and all precautions taken to prevent their contamination,]
    - [(c) where bulk transport is intended, the pipe, pumps and bulk tanks and any other bulk container or bulk road tanker used in (2) or the transportation of the product from the manufacturing plant either directly on to the ship or into shore tanks or directly to plants have been inspected and found to be clean before use,]
    - (d) which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'. and

### Notes

### Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit commodity: it may be filled in if the certificate is for import commodity.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading
- Box reference I.19: use the appropriate HS code: 15.04 or 15.18.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.
- Box reference I.25: technical use: any use other than for animal consumption.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28; Manufacturing plant; provide the registration number of the treatment/processing establishment.

COUNTRY	Fish oil not intended for human c material or for purposes outside the	
II. Health information	II.a. Certificate reference No	II.b.
Part II:		
( <sup>1a</sup> ) OJ L 300, 14.11.2009, p. 1.		
( <sup>1b</sup> ) OJ L 54, 26.2.2011, p. 1.		
(²) Delete as appropriate.		
- The signature and the stamp must be in a different colour to that of	the printing.	
<ul> <li>Note for the person responsible for the consignment in the European Union: this certificate is only for veterinary purposes and ha accompany the consignment until it reaches the border inspection post.</li> </ul>		
Official veterinarian/Official inspector		
Name (in capital letters):	Qualification and	d title:
Date:	Signature:	
Stamp:		

# 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

cou	COUNTRY Veterinary certificate to EU					
	l.1.	Consignor	2. Certificate reference No	) I.2.a.		
		Name	I.3. Central competent authority			
		Address	Solida compotent durining			
		Tel.	<ol> <li>Local competent author</li> </ol>	ity		
dispatched consignment	I.5.	Consignee	6. Person responsible for	the load in EU		
ᄩ		Name	Name			
nsić		Address	Address			
8		Postcode	Postcode			
hed		Tel.	Tel.			
patc	17	Country of origin ISO code I.8. Region of origin Code	9. Country of I	SO I.10. Region of Code		
dis	1.7.	Country of origin 150 code 1.8. Region of origin Code		ode destination		
5						
Part I: Details of	l.11.	Place of origin	12. Place of destination			
<u>:</u>		Name Approval number	Name	Custom warehouse		
art		Address	Address	Approval number		
-		Name Approval number Address	Postcode			
		Name Approval number Address				
	I.13.	Place of loading	14. Date of departure			
	I.15.	Means of transport	16. Entry BIP in EU			
		Aeroplane ☐ Ship ☐ Railway wagon ☐				
		Road vehicle Other	17.			
		Identification				
		Documentation references				
	I.18.	Description of commodity	I.19. Commod	dity code (HS code)		
				I.20. Quantity		
	I.21.	Temperature of product		I.22. Number of packages		
		Ambient Chilled C	ozen 🗌			
	1.23.	Seal/Container No		I.24. Type of packaging		
	1.25.	Commodities certified for:				
		Animal feedingstuff ☐ Technical use ☐				
	126	For transit through EU to third country	27. For import or admission	n into EU		
			27. Tor import or durinosion			
		Third country ISO code				
	1.28.	Identification of the commodities				
		Species Nature of commodity Approval number of (Scientific name) Manufacturing		packages Net weight Batch number		

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

### COUNTRY

≝ Part

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

I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council (<sup>1a</sup>) and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011 (<sup>1b</sup>), and in particular Annex XIV, Chapter II thereof, and certify that the rendered fats described above:

- II.1. consist of rendered fats that satisfy the health requirements below:
- II.2. consist of rendered fats not intended for human consumption;
- Certification have been prepared and stored in a plant approved, validated 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 II.3. Council (3), in order to kill pathogenic agents;
  - II.4. 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
    - (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;
      - (v) feathers;]
    - (2) and/or [- blood of animals which did not show any signs of disease communicable through blood to humans or animals, obtained from animals other than ruminants that have been slaughtered in a slaughterhouse after having been considered fit for slaughter for human consumption following an ante-mortem inspection in accordance with Union legislation:]
    - (2) and/or [- animal by-products arising from the production of products intended for human consumption, including degreased bone, greaves and centrifuge or separator sludge from milk processing;]
    - (2) and/or [- products of animal origin, or foodstuffs containing products of animal origin, which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects from which no risk to public or animal health arise:1
    - (2) and/or [- petfood and feedingstuffs of animal origin, or feedingstuffs containing animal by-products or derived products, which are no longer intended for feeding for commercial reasons or due to problems of manufacturing or packaging defects or other defects from which no risk to public or animal health 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:1
    - (2) and/or [- the following material originating from animals which did not show any signs of disease communicable through that material
      - (i) shells from shellfish with soft tissue or flesh;

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

COUN	ΓRY	Rendered fats not intended for human consumption to be used a feed material
II.	Health info	
		(ii) the following originating from terrestrial animals:
		— hatchery by-products,
		— eggs,
		egg by-products, including egg shells;
		(iii) day-old chicks killed for commercial reasons;]
II.5.	(2) either	[- in the case of material of porcine origin, come from a country or part of a territory free from foot-and-mouth disease for the
		previous 24 months and free from classical swine fever and African swine fever for the previous 12 months;]
	(²) and/or	[- in the case of material of poultry origin, come from a country or part of a territory free from Newcastle disease and aviar influenza for the previous 6 months;]
	(²) and/or	[- in the case of material of ruminant origin, come from a country or part of a territory free from foot-and-mouth disease for the previous 24 months and free from rinderpest for the previous 12 months;]
	(²) and/or	[- where there has been an outbreak of one of the abovementioned diseases during the relevant period mentioned above, and where the rendered fats are derived from a susceptible species, have been subjected to a heat treatment for at least 70 °C for 30 minutes or at least 90 °C for at least 15 minutes, and
		details of the critical control points are recorded and maintained so that the owner, operator or their representative and, as necessary, the competent authority can monitor the operation of the plant; the information must include the particle size, critical temperature and, as appropriate, the absolute time, pressure profile, raw material feed rate and fat recycling rate.
II.6.	if derived f 0,15 % in	rom ruminant animals were purified in such way that the maximum levels of remaining total insoluble impurities does not exceed weight;
II.7.	the render	ed fats:
		(a) have been subjected to processing in accordance with Annex X, Chapter II, Section 3 of Regulation (EU) No 142/2011, or treatment in accordance with Section XII of Annex III to Regulation (EC) No 853/2004, in order to kill pathogenic agents; and
	(²) either	[(b) are packaged in new containers or in containers that have been cleaned and disinfected if necessary for the prevention of contamination, and all precautions taken to prevent their contamination;]
	(²) or	[(b) where bulk transport is intended, the pipe, pumps and bulk tanks and any other bulk container or bulk road tanker used in the transportation of the product from the manufacturing plant either directly on to the ship or into shore tanks or directly to plants have been checked under the responsibility of the competent authority and found to be clean before use;]
	and which	bear labels indicating 'NOT FOR HUMAN CONSUMPTION';
II.8.		
	(²) either	[the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) No. 999/2001 of the European Parliament and of the Council (4) or mechanically separated meat obtained from bones of bovine, ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity:]
	(²) or	[the product does not contain and is not derived from bovine, ovine or 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 by a decision in accordance with Article 5(2) of Regulation (EC) No 999/2001;]
II.9.	in addition	as regards TSE:
	( <sup>2</sup> ) either	[in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin, the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last three years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfied the following requirements for the last three years:
		(i) it has been subject to regular official veterinary checks;

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

II.b.

### COUNTRY

## Health information II.a. Certificate reference No

- (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or, following the confirmation of a classical scrapie case:
  - all animals in which classical scrapie was confirmed have been killed and destroyed, and
  - all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARR genotype and breeding ewes carrying at least one ARR allele and no VRQ allele;
- (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are introduced into the holding only if they come from a holding which complies with the requirements set out in points (i) and (ii),]
- (2) or [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin, and destined to a Member State listed in the Annex to Commission Regulation (EC) No 546/2006 (5), the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last seven years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfied the following requirements for the last seven years:
  - (i) it has been subject to regular official veterinary checks;
  - (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or, following the confirmation of a classical scrapie case:
    - all animals in which classical scrapie was confirmed have been killed and destroyed, and
    - all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARR genotype and breeding ewes carrying at least one ARR allele and no VRQ allele;
  - (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are introduced into the holding only if they come from a holding which complies with the requirements set out in points (i) and (ii).]

### Notes

### Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit
  commodity; it may be filled in if the certificate is for import commodity.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only
  be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading.
- Box reference I.19: use the appropriate HS code: 15.02; 15.03; 15.04; 15.05; 15.06; 15.16.10; 15.17 or 15.18.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.
- Box reference I.25: technical use: any use other than for animal consumption.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: Manufacturing plant: provide the registration number of the treatment/processing establishment.

### Part II:

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

COUNTRY		feed material		
II. Health information		II.a. Certificate reference No	II.b.	
( <sup>4</sup> ) OJ L 147, 31.5.2001, p. 1.				
( <sup>5</sup> ) 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 t accompany the consignment until it real			or veterinary purposes and has to	
Official veterinarian/Official inspector				
Name (in capital letters):		Qualification a	nd title:	
Date:		Signature:		
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  $\binom{2}{1}$  the European Union

cou	NTRY	1	Veterinary certificate to EU		
	l.1.	Consignor	I.2. Certificate reference No I.2.a.		
		Name Address	I.3. Central competent authority		
			I.4. Local competent authority		
	_	Tel.			
ment	1.5.	Consignee Name	I.6. Person responsible for the load in EU		
sign		Address	Name Address		
8		P. 1. 1.			
ched		Postcode Tel.	Postcode Tel.		
of dispatched consignment	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of destination ISO code destination Code		
l: Details	l.11.	Place of origin	I.12. Place of destination		
Part I: D		Name Approval number Address	Name Custom warehouse ☐ Address Approval number		
•		Name Approval number Address	Postoodo		
		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 Ship			
		Identification	1.17.		
		Documentation references			
	I.18.	Description of commodity	I.19. Commodity code (HS code)		
			I.20. Quantity		
	1.21.	Temperature of product	I.22. Number of packages		
		Ambient Chilled Chilled	Frozen 🗆		
	1.23.	Seal/Container No	I.24. Type of packaging		
	1.25.	Commodities certified for:			
		Technical use			
	1.26.	For transit through EU to third country	I.27. For import or admission into EU		
		Third country ISO code			
	1.28.	Identification of the commodities			
		Species Nature of commodity Approval number of (Scientific name) Manufacturi			

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

COUNTRY				Rendered fats not intended for human consumption to be used for certain purposes outside the feed chain			
	II.	Health in	formation	II.a. Certificate reference No	II.b.		
		Parliamen	lersigned official veterinarian, declare that I have t and of the Council ( <sup>1a</sup> ) and in particular <i>i</i> D11 ( <sup>1b</sup> ), and in particular Annex XIV, Chapter II the	Articles 8, 9 and 10 thereof, and	Commission Regulation (EU)		
	II.1.	consist of	rendered fats not intended for human consumption	that satisfy the health requirements b	elow;		
ution	II.2.	have been	n prepared exclusively with the following animal by-	products:			
Part II: Certification	II.2.1.		e of materials destined for the production of biodie 069/2009;	sel, animal by-products referred to in A	Articles 8, 9 and 10 of Regulation		
ar ::	II.2.2.	in the cas	e of materials destined for other purposes:				
<u> </u>		(²) either	[- animal by-products containing residues of author to in Article 15(3) of Directive 96/23/EC;]	rised substances or contaminants exceed	eding the permitted levels referred		
		(2) and/or	[- products of animal origin which have been declar those products;]	red unfit for human consumption due to	the presence of foreign bodies in		
		(²) and/or	[- animals and parts of animals, other than those re other than being slaughtered or killed for humans.]				
		(²) and/or	[- carcases and parts of animals slaughtered or, in human consumption in accordance with Union reasons;]				
		(²) and/or	[- carcases and the following parts originating eithe considered fit for slaughter for human consumpt of animals from game killed for human consumptions.]	ion following an ante-mortem inspectior	or bodies and the following parts		
			(i) carcases or bodies and parts of animals whi legislation, but which did not show any sign				
			(ii) heads of poultry;				
			(iii) hides and skins, including trimmings and spl metacarpus bones, tarsus and metatarsus b				
			(iv) pig bristles;				
			(v) feathers;]				
		(²) and/or	[- blood of animals which did not show any signs from animals other than ruminants that have be slaughter for human consumption following an a	en slaughtered in a slaughterhouse af	ter having been considered fit for		
		(2) and/or	[- animal by-products arising from the production of greaves and centrifuge or separator sludge from		nption, including degreased bone,		
		(²) and/or	[- products of animal origin, or foodstuffs contain consumption for commercial reasons or due to which no risk to public or animal health arise;]				
		(²) and/or	[- petfood and feedingstuffs of animal origin, or fe no longer intended for feeding for commercial re defects from which no risk to public or animal it	asons or due to problems of manufactu			
		(2) and/or	[- blood, placenta, wool, feathers, hair, horns, hoof any disease communicable through that produc		animals that did not show signs of		
		(2) and/or	[- aquatic animals, and parts of such animals, ex- nicable to humans or animals;]	cept sea mammals, which did not sho	w any signs of diseases commu-		
		(2) and/or	[- animal by-products from aquatic animals origin consumption;]	nating from plants or establishments m	nanufacturing products for human		

Rendered fats not intended for human consumption to be used for COUNTRY certain purposes outside the feed chain Health information II.a. Certificate reference No II.b. II. (e) 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 [- taquatic 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) and Category 2 material as referred to in Article 9(a) to (g) of Regulation (EC) No 1069/2009:1 (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;] in the case of materials destined for purposes other than the production of organic fertilisers or soil improvers: (2) either [- specified risk material as defined in Article 3(1)(g) of Regulation (EC) No 999/2001 of the European Parliament and of the Council (3):] (2) and/or [- entire bodies or parts of dead animals containing specified risk material as defined in Article 3(1)(g) of Regulation (EC) No 999/2001 at the time of disposal;] (²) and/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 Directive 96/22/EC or Article 2(b) of Directive 96/23/EC;] (²) and/or [- animal by-products containing residues of other substances and environmental contaminants listed in Group B(3) of Annex I to Directive 96/23/EC, if such residues exceed the permitted levels laid down by Union legislation or, in the absence thereof, by legislation of the Member State of importation;] II.3. the rendered fats: (a) have been subjected to processing in accordance with method ....... as laid down in Chapter III of Annex IV to Regulation (EU) No 142/2011, in order to kill pathogenic agents (b) have been marked before shipment to the European Union with glyceroltriheptanoate (GTH), so that a homogenous minimum concentration of at least 250 mg GTH per kilogram fat is achieved; (c) in the case of rendered fats of ruminant origin, insoluble impurities in excess of 0.15 % in weight have been removed: (d) have been transported under conditions which prevent their contamination; and

(e) bear labels on the packaging or container indicating 'NOT FOR HUMAN OR ANIMAL CONSUMPTION';

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

# COUNTRY certain purposes outside the feed chain П. Health information II.a. Certificate reference No II.b. 11.4 in the case of materials destined for organic fertilisers or soil improvers: [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001 or mechanically separated meat obtained from bones of bovine, ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity.] [the product does not contain and is not derived from bovine, ovine or 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 by a decision in accordance with Article 5(2) of Regulation (EC) No 999/2001.] (2) or Notes Part I: Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity. Box reference I.11 and I.12: Approval number: the registration number of the establishment or plant, which has been issued by the competent Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading. Box reference I.19: use the appropriate HS code: 15.02; 15.03; 15.04; 15.05; 15.06; 15.16.10; 15.17 or 15.18. Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included. Box reference I.25: technical use: any use other than for animal consumption. - Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate. - Box reference I.28: Manufacturing plant: provide the registration number of the treatment/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, - 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 has to accompany the consignment until it reaches the border inspection post. Official veterinarian/Official inspector Qualification and title: Name (in capital letters): Date: Signature: Stamp:

## CHAPTER 11

## Health certificate

For gelatine and collagen not intended for human consumption to be used as feed material or for purposes outside the feed chain, intended for dispatch to or for transit through  $\binom{2}{1}$  the European Union

cou	DUNTRY Veterinary certificate to EU						
	l.1.	Consignor	I.2. Certificate reference No I.2.a.				
		Name Address	I.3. Central competent authority				
		Tel.	I.4. Local competent authority				
뒽	1.5.	Consignee	I.6. Person responsible for the load in EU				
me		Name	Name				
ığ		Address	Address				
Si o							
ğ		Postcode	Postcode				
ţ		Tel.	Tel.				
Part I: Details of dispatched consignment	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of destination ISO code l.10. Region of destination				
ls o							
Detai	l.11.	Place of origin	I.12. Place of destination				
art I:		Name Approval number Address	Name Custom warehouse ☐ Address Approval number				
•		Name Approval number Address	5				
		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 _	L.17.				
		Road vehicle Other O					
		Identification  Documentation references					
	I.18.	Description of commodity	I.19. Commodity code (HS code)				
			I.20. Quantity				
			1.20. Quantity				
	I.21.	Temperature of product	I.22. Number of packages				
		Ambient Chilled Chilled	Frozen				
	1.23.	Seal/Container No	I.24. Type of packaging				
	1.25.	Commodities certified for:					
		Animal feedingstuff Technical use					
	1.26.	For transit through EU to third country	I.27. For import or admission into EU				
		Third country ISO code					
	1.28.	Identification of the commodities					
		Species Nature of commodity Approval number of (Scientific name) Manufacturin					

OUN	ITRY			Gelatine and collagen not intender used as feed material or for purpo				
	II.	Health info	ormation	II.a. Certificate reference No	II.b.			
		Parliament	ersigned official veterinarian, declare that I have and of the Council ( <sup>1a</sup> ) and in particular Article Annex XIV, Chapter I thereof, and certify that the g	10 thereof, and Commission Regulatio				
II.1. consists of gelatine/collagen (2) that satisfy the health requirements below;								
<u>ا</u>	II.2.	consist exc	clusively of gelatine/collagen (2) not intended for hu	man consumption;				
Part II: Certification	II.3.	has been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance with Article 24 of Regulation (EC) No 1069/2009, in order to kill pathogenic agents;						
ਜ਼ੂ ਜ਼ੂ	II.4.	has been p	prepared exclusively with the following animal by-p	roducts:				
(2) either [- carcases and parts of animals slaughtered or, in the case of game, bodies or parts of animals killed, human consumption in accordance with Union legislation, but are not intended for human consumptions;]								
		(2) and/or	[- carcases and the following parts originating eith considered fit for slaughter for human consump of animals from game killed for human consumption of animals from game killed for human consumptions.]	tion following an ante-mortem inspection	or bodies and the following parts			
			(i) carcases or bodies and parts of animals w legislation, but which did not show any sig					
			(ii) heads of poultry;					
			(iii) hides and skins, including trimmings and s metacarpus bones, tarsus and metatarsus					
			(iv) pig bristles;					
			(v) feathers;]					
		(2) and/or	[- animal by-products arising from the production greaves and centrifuge or separator sludge from		nption, including degreased bone,			
		(²) and/or	[- products of animal origin, or foodstuffs contail consumption for commercial reasons or due to which no risk to public or animal health arise;]	o problems of manufacturing or packag				
		(²) and/or	[- petfood and feedingstuffs of animal origin, or fellonger intended for feeding for commercial readefects from which no risk to public or animal	asons or due to problems of manufactur				
		(²) and/or	[- aquatic animals, and parts of such animals, e nicable to humans or animals;]	xcept sea mammals, which did not sho	w any signs of diseases commu-			
		(2) and/or	[- animal by-products from aquatic animals orig consumption;]	inating from plants or establishments n	nanufacturing products for human			
	II.5.	the gelatine	e/collagen (²)					
			(a) was wrapped, packaged, stored and transpo packaging took place in a dedicated room, a					

Wrappings and packages containing gelatine/collagen  $(^2)$  carry the words 'GELATINE/COLLAGEN  $(^2)$  SUITABLE FOR ANIMAL CONSUMPTION'; and

# Gelatine and collagen not intended for human consumption to be

COUNTRY used as feed material or for purposes outside the feed chain II. Health information II.a. Certificate reference No II.b [(b) in the case of gelatine, has been produced by a process that is ensuring that unprocessed Category 3 material is subjected to a treatment with acid or alkali, followed by one or more rinses, involving pH adjustment, extraction by (2) either 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, has been produced by a process that 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, in order to kill pathogenic agents;] II.6. in the case of gelatine from materials other than hides and skins: [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001 of the European Parliament and of the Council (3) or mechanically separated meat obtained from bones of bovine, ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning by (2) either means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity;] (2) or Ithe product does not contain and is not derived from bovine, ovine or 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 by a decision in accordance with Article 5(2) of Regulation (EC) No 999/2001;] 11.7. in the case of gelatine from materials other than hides and skins: in addition as regards TSE: [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin, the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last (2) either three years on a holding where no official movement restriction is imposed due to a suspicion of TŚE and which has satisfied the following requirements for the last three years: (i) it has been subject to regular official veterinary checks; (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or, following the confirmation of a classical scrapie case: - all animals in which classical scrapie was confirmed have been killed and destroyed, and - all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARR genotype and breeding ewes carrying at least one ARR allele and no VRQ allele, (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are introduced into the holding only if they come from a holding which complies with the requirements set out in points (i) and (ii).] (2) or fin case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin, and destined to a Member State listed in the Annex to Commission Regulation (EC) No 546/2006 (4), the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last seven years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfied the following requirements for the last seven years: (i) it has been subject to regular official veterinary checks; (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or, following the confirmation of a classical scrapie case: - all animals in which classical scrapie was confirmed have been killed and destroyed, and - all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARR

genotype and breeding ewes carrying at least one ARR allele and no VRQ allele;

(iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are introduced into the holding only if they come from a holding which complies with the requirements set out in points (i) and (ii).]

Stamp:

co	UNTRY	Gelatine and collagen not intende used as feed material or for purp	
II.	Health information	II.a. Certificate reference No	II.b.
No	tes		
Pa	rt I:		
_	Box reference I.6: Person responsible for the consignment in the Eucommodity; it may be filled in if the certificate is for import commod		in only if it is a certificate for trans
-	Box reference I.12: Place of destination: this box is to be filled in only be stored in free zones, free warehouses and custom warehouses.	r if it is a certificate for transit commod	dity. The products in transit can only
_	Box reference I.15: Registration number (railway wagons or containe provided in case of unloading and reloading.	r and lorries), flight number (aircraft)	or name (ship); information is to be
-	Box reference I.19: use the appropriate HS code: 35.03 or 35.04.		
_	Box reference I.23: for bulk containers, the container number and the	e seal number (if applicable) should	be included.
_	Box reference I.25: technical use: any use other than for animal cor	nsumption.	
_	Box reference I.26 and I.27: fill in according to whether it is a transi	t or an import certificate.	
-	Box reference I.28: Nature of commodity: select gelatine or collager Manufacturing plant: provide the registration number of treatment/pro		
Pa	rt II:		
( <sup>1a</sup>	OJ L 300, 14.11.2009, p. 1.		
( <sup>1b</sup>	OJ L 54, 26.2.2011, p. 1.		
( <sup>2</sup> )	Delete as appropriate.		
( <sup>3</sup> )	OJ L 147, 31.5.2001, p. 1.		
( <sup>4</sup> )	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 Europaccompany the consignment until it reaches the border inspection p		or veterinary purposes and has to
Of	icial veterinarian/Official inspector		
	Name (in capital letters):	Qualification and	d title:

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

cou	UNTRY Veterinary certificate to EU					
	l.1.	Consignor	I.2. Certificate reference No I.2.a.			
		Name	I.3. Central competent authority			
		Address	no. Solida compotent dationly			
		Tel.	I.4. Local competent authority			
_	1.5.	Consignee	I.6. Person responsible for the load in EU			
neu	1.5.	Name	Name			
gu		Address	Address			
onsi			7.11.11.000			
Ö		Postcode	Postcode			
tche		Tel.	Tel.			
of dispatched consignment	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO I.10. Region of Code			
ğ			destination code destination			
ls o						
l: Details	1.11.	Place of origin	I.12. Place of destination			
<u>:</u>		Name Approval number	Name Custom warehouse □			
Part		Address	Address Approval number			
-		Name Approval number Address				
		Name Approval number	Postcode			
		Address				
	I.13.	Place of loading	I.14. Date of departure			
	I.15.	Means of transport	I.16. Entry BIP in EU			
		Aeroplane ☐ Ship ☐ Railway wagon ☐				
		Road vehicle Other	1.17.			
		Identification				
		Documentation references				
	I.18.	Description of commodity	I.19. Commodity code (HS code)			
			I.20. Quantity			
	I.21.	Temperature of product	I.22. Number of packages			
		Ambient ☐ Chilled ☐	Frozen			
	1.23.	Seal/Container No	I.24. Type of packaging			
	1.25.	Commodities certified for:	'			
		Animal feedingstuff ☐ Technical use ☐				
	1.26.	For transit through EU to third country	I.27. For import or admission into EU			
		Third country ISO code	_			
		Third country ISO code				
	1.28.	Identification of the commodities				
		Species Nature of commodity Approval number of e (Scientific name) Manufacturing				

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

### COUNTRY

COU		JNTRY		1		for uses outside the feed chain		
		II.	Health info	orn	nation	II.a. Certificate reference No	II.b.	
			and of the	Co	ned official veterinarian, declare that I have read a uncil ( <sup>1a</sup> ) and in particular Article 10 thereof, and Co reof, and certify that the hydrolysed protein/dicalciu	ommission Regulation (EU) No 142/20	11 (1b), and in particular Annex XIV,	
Dart II. Cortification	ے	II.1.	consists of	hy	drolysed protein/dicalcium phosphate/tricalcium ph	nosphate (2) that satisfy the health requ	irements below;	
	li.2. consists exclusively of hydrolysed protein/dicalcium phosphate/trical						ricalcium phosphate (2) not intended fo	or human consumption;
	t II: Cert	II.3.	has been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance with Article Regulation (EC) No 1069/2009, in order to kill pathogenic agents;					
	Par	II.4.	has been p	ore	pared exclusively with the following animal by-proc	ducts:		
		II.4.1.	in the case	e of	dicalcium phosphate derived from defatted bones	<b>3:</b>		
		carcases and parts of animals slaughtered or, in the case of game, bodies or parts of animals killed, and which are fit consumption in accordance with Union legislation, but are not intended for human consumption for commercial reasons;						
L	$\dashv$	II.4.2.	in case of	oth	er materials:			
(²) either [- carcases and parts of animals slaughtered or, in the case of game, bodies or parts of human consumption in accordance with Union legislation, but are not intended for reasons:]								
			(2) and/or	[-	carcases and the following parts originating either considered fit for slaughter for human consumption animals from game killed for human consumption	n following an ante-mortem inspection		
					(i) carcases or bodies and parts of animals whic legislation, but which did not show any signs			
					(ii) heads of poultry;			
					(iii) hides and skins, including trimmings and split metacarpus bones, tarsus and metatarsus bo			
					(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 of animals other than ruminants 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  [- animal by-products arising from the production of products intended for human consumption, including degree greaves and centrifuge or separator sludge from milk processing;]					ng been considered fit for slaughter			
					mption, including degreased bone,			
(2) and/or [- products of animal origin, or foodstuffs containing products of animal origin, or consumption for commercial reasons or due to problems of manufacturing or which no risk to public or animal health arise;]  (2) and/or [- petfood and feedingstuffs of animal origin, or feedingstuffs containing animal byplonger intended for feeding for commercial reasons or due to problems of manufacturing or 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 any disease communicable through that product to humans or animals;]								
						ons or due to problems of manufactur		
							animals that did not show signs of	

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

COUNTRY for uses outside the feed chain Health information II.a. Certificate reference No (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:1 (2) and/or [- animal by-products from aquatic animals originating from plants or establishments manufacturing products for human consumption:1 (2) and/or [- the following material originating from animals which did not show any signs of disease communicable through that material (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;] 11.5. the hydrolysed protein/dicalcium phosphate/tricalcium phosphate (2): (a) was wrapped and packaged in packaging which bear labels indicating 'NOT FOR HUMAN CONSUMPTION' and stored and transported under satisfactory hygiene conditions, and in particular wrapping and packaging took place in a dedicated room, and only preservatives permitted under Union legislation were used; and (2) either [(b) in the case of hydrolysed protein, has been produced by a process involving appropriate measures to minimise contamination of raw Category 3 material. In the case of hydrolysed proteins entirely or partly derived from ruminants hides and skins, has been produced in a processing plant dedicated only to hydrolysed proteins production, using a process involving the preparation of the 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 3 hours at 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.] (2) or [(b) in the case of dicalcium phosphate, has been 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) followed by treatment of the obtained phosphoric liquor with lime, resulting in a precipitate of dicalcium phosphate at pH 4 to 7, and (iii) finally air-dries this precipitate, with inlet temperature of 65 °C to 325 °C and end temperature between 30 °C and (2) or [(b) in the case of tricalcium phosphate, has been produced by a process ensuring: (i) that all Category 3 bone-material is finely crushed and degreased in counter-flow with hot water (bone chips less than (ii) continuous cooking with steam at 145 °C during 30 minutes at 4 bars, (iii) separation of the protein broth from the hydroxyapatite (tricalcium phosphate) by centrifugation, and (iv) granulation of the tricalcium phosphate after drying in a fluidised bed with air at 200 °C.]

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

### COUNTRY

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

II.6.

(2) either

[the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001 of the European Parliament and of the Council (3) or mechanically separated meat obtained from bones of bovine, ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity;]

(2) or [the product does not contain and is not derived from bovine, ovine or 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 by a decision in accordance with Article 5(2) of Regulation (EC) No 999/2001;]

- II.7. in addition as regards TSE:
  - either [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin, the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last three years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfied the following requirements for the last three years:
    - (i) it has been subject to regular official veterinary checks;
    - (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or, following the confirmation of a classical scrapie case:
      - all animals in which classical scrapie was confirmed have been killed and destroyed, and
      - all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARR genotype and breeding ewes carrying at least one ARR allele and no VRQ allele;
    - (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are introduced into the holding only if they come from a holding which complies with the requirements set out in points (i) and (ii).]
  - (²) or [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin, and destined to a Member State listed in the Annex to Commission Regulation (EC) No 546/2006 (\*), the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last seven years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfied the following requirements for the last seven years:
    - (i) it has been subject to regular official veterinary checks;
    - (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or, following the confirmation of a classical scrapie case:
      - all animals in which classical scrapie was confirmed have been killed and destroyed, and
      - all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARR genotype and breeding ewes carrying at least one ARR allele and no VRQ allele;
    - (iii) ovine and caprine animals, with the exception of sheep of the ARR/ARR prion genotype, are introduced into the holding only if they come from a holding which complies with the requirements set out in points (i) and (ii).]

### Notes

### Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only
  be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading.
- Box reference I.19: use the appropriate HS code: 28.35 or 35.04.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.

COUNTRY	Hydrolysed protein, dicalcium pho not intended for human consumpti for uses outside the feed chain				
II. Health information	II.a. Certificate reference No	II.b.			
— Box reference I.25: technical use: any use other than for animal con	sumption.				
Box reference I.26 and I.27: fill in according to whether it is a transit	t or an import certificate.				
Box reference I.28: Nature of commodity: specify if hydrolysed prote	in, dicalcium phosphate or tricalcium p	phosphate.			
Manufacturing plant: provide the registration number of treatment/pro	cessing establishment.				
Part II:					
( <sup>1a)</sup> OJ L 300, 14.11.2009, p. 1.					
( <sup>1b</sup> ) OJ L 54, 26.2.2011, p. 1.					
(²) Delete as appropriate.					
(³) OJ L 147, 31.5.2001, p. 1.					
( <sup>4</sup> ) 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 Europaccompany the consignment until it reaches the border inspection personal control of the consignment until it reaches the border inspection personal control of the consignment of the consignment in the Europaccompany the consignment of the consignment in the Europaccompany the Europacc		r veterinary purposes and has to			
Official veterinarian/Official inspector					
Name (in capital letters):	Qualification and	i title:			
Date: Signature:					
Stamp:					

# **CHAPTER 13**

# Health certificate

For apiculture by-products intended exclusively for use in apiculture, intended for dispatch to or for transit through  $\binom{2}{1}$  the European Union

cou	NTRY	Veterinary certificate to EU			
	I.1.	Consignor	I.2. Certificate reference No I.2.a.		
		Name	I.3. Central competent authority		
		Address	1.5. Certifal competent authority		
		Tel.	I.4. Local competent authority		
aut	1.5.	Consignee	I.6. Person responsible for the load in EU		
Ě		Name	Name		
nsig		Address	Address		
8		Postcode	Postcode		
hed		Tel.	Tel.		
dispatched consignment	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10. Region of Code		
ğ			destination destination		
ls o					
Detai	I.11.	Place of origin	I.12. Place of destination		
Part I: Details of		Name Approval number Address	Name Custom warehouse ☐ Address Approval number		
۵ ا		Name Approval number			
		Address Name Approval number	Postcode		
		Address			
	I.13.	Place of loading	I.14. Date of departure		
	l.15.	Means of transport	I.16. Entry BIP in EU		
		Aeroplane Ship Railway wagon			
		Road vehicle Other	1.17.		
		Identification			
	110	Documentation references	L10 Commodity and (US ands)		
	1.18.	Description of commodity	I.19. Commodity code (HS code)		
			I.20. Quantity		
	I.21.	Temperature of product	I.22. Number of packages		
			Frozen 🗆		
	1.23.	Seal/Container No	I.24. Type of packaging		
	1.25.	Commodities certified for:			
		Technical use □			
	1.26.	For transit through EU to third country	I.27. For import or admission into EU		
		Third country ISO code			
	1.28.	Identification of the commodities			
		Species Nature of commodity (Scientific name)	Approval number of establishments Net weight Manufacturing plant		

COL	JNTRY			Apiculture by-products intended	exclusively for use in apiculture	
	II.	Health info	ormation	II.a. Certificate reference No	II.b.	
		and of the	rsigned official veterinarian, declare that I have read at Council ( <sup>1a</sup> ) and in particular Article 10 thereof, and Cothereof, and certify that the apiculture by-products de	ommission Regulation (EU) No 142/201		
۰	II.1.	come from with:	come from an area where the diseases mentioned below are officially notifiable and which is not subject to any restrictions associated with:			
		(a) Americ	an foulbrood (Paenibacillus larvae larvae);			
ation		(b) Acarios	sis (Acarapis woodi (Rennie));			
ertific		(c) Small h	nive beetle (Aethina tumida); and			
Part II: Certification		(d) Tropilaelaps mites ( <i>Tropilaelaps</i> spp.);				
Part	II.2.	have been				
		(2) either	[subjected to a temperature of - 12 $^{\circ}\text{C}$ or lower for	at least 24 hours.]		
		(²) or	[in the case of wax refined or processed in accordance IV to Regulation (EU) No 142/2011]	ance with processing method 1-2-3-4-	.5-7 (2) as set out in Chapter III of	
	Notes					
	Part I:					
			6: Person responsible for the consignment in the Eurapy be filled in if the certificate is for import commoditions.		n only if it is a certificate for transit	
		reference I. ority.	eference I.11 and I.12: Approval number: the registration number of the establishment or plant, which has been issued by the competent rity.			
		reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only tored in free zones, free warehouses and custom warehouses.				
			eference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be ed in the event of unloading and reloading.			
	— Вох	reference I.19: use the appropriate HS code: 05.11.99 and specify the commodity as listed under note Box reference I.28.				
	— Вох	reference I.23: for bulk containers, the container number and the seal number (if applicable) should be given.				
	— Вох	reference I.25: technical use: any use other than for animal consumption.				
	— Вох	reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.				
	— Вох	r reference I.28: Nature of commodity: means honey, beeswax, royal jelly, propolis or pollen used in bee-keeping;				
	Part II:					
	( <sup>1a</sup> ) O	J L 300, 14.	11.2009, p. 1.			
	(1b) O.	J L 54, 26.2	.2011, p. 1.			
	(²) De	elete as appropriate.				
	— The	signature a	signature and the stamp must be in a different colour to that of the printing.			
			erson responsible for the consignment in the Europ consignment until it reaches the border inspection po		r veterinary purposes and has to	
Official veterinarian/Official inspector						
	Na	me (in capita	al letters):	Qualification and	title:	
	Dat	te:		Signature:		
	Sta	mp:				

# CHAPTER 14(A)

# **Health certificate**

For fat derivatives not intended for human consumption to be used outside the feed chain, intended for dispatch to or for transit through  $\binom{2}{1}$  the European Union

COUNTRY Veterinary certificate to EV			
	l.1.	Consignor	I.2. Certificate reference No I.2.a.
		Name Address	I.3. Central competent authority
		Tel.	I.4. Local competent authority
ent	1.5.	Consignee	I.6. Person responsible for the load in EU
ᇤ		Name	Name
onsi		Address	Address
of dispatched consignment		Postcode Tel.	Postcode Tel.
of disp	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of destination ISO code destination Code
etails	l.11.	Place of origin	I.12. Place of destination
Part I: Details		Name Approval number Address	Name Custom warehouse ☐ Address Approval number
اتا		Name Approval number Address	
		Name Approval number Address	Postcode
	I.13.	Place of loading	I.14. Date of departure
	l.15.	Means of transport	I.16. Entry BIP in EU
		Aeroplane Ship Railway wagon	
		Road vehicle Other I	1.17.
		Documentation references	
	I.18.	Description of commodity	I.19. Commodity code (HS code)
			15.16.10
			I.20. Quantity
	I.21.	Temperature of product	I.22. Number of packages
		Ambient Chilled Chilled	Frozen
	1.23.	Seal/Container No	I.24. Type of packaging
	1.25.	Commodities 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	
	1.28.	Identification of the commodities	
		Species Nature of commodity Approval number of (Scientific name) Manufacture	and the same

# Fat derivatives not intended for human consumption to be used

#### COUNTRY outside the feed chain Health information II.a. Certificate reference No. II.b. I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council (<sup>1a</sup>) and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011 (<sup>1b</sup>), and in particular Annex XIV, Chapter II thereof, and certify that the fat derivatives described above: II.1. consist of fat derivatives that satisfy the health requirements below; II.2. consist of fat derivatives intended for purposes outside the feed chain, other than in cosmetics, pharmaceuticals and medical devices; Certification have been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance with Article 24 of Regulation (EC) No 1069/2009, in order to kill pathogenic agents; II.3. 11.4. have been prepared from rendered fats exclusively produced from the following materials: art II.4.1. in case the fat derivatives are intended for uses outside the feed chain, other than in organic fertilisers, soil improvers, cosmetics, pharmaceuticals and medical devices, the following Category 1 materials: (2) either [- animal by-products which have been derived from animals which have been submitted to illegal treatment as defined in Article 1(2)(d) of Directive 96/22/EC or Article 2(b) of Directive 96/23/EC;] (2) and/or [- animal by-products containing residues of other substances and environmental contaminants listed in Group B(3) of Annex I to Directive 96/23/EC, if such residues exceed the permitted levels laid down by Union legislation or, in the absence thereof, by legislation of the Member State of importation;] II.4.2. in case the fat derivatives are intended for use in organic fertilisers or soil improvers or other uses outside the feed chain, other than in cosmetics, pharmaceuticals and medical devices, the following Category 2 materials: (²) either [- animal by-products containing residues of authorised substances or contaminants exceeding the permitted levels referred to in Article 15(3) of Directive 96/23/EC;] (2) and/or [- products of animal origin which have been declared unfit for human consumption due to the presence of foreign bodies in those products: (2) and/or [- animals and parts of animals, other than those referred to in Articles 8 and 10 of Regulation (EC) No 1069/2009, that died other than being slaughtered or killed for human consumption, including animals killed for disease control purposes;] II.4.3. Category 3 materials: (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, of animals, other than ruminants; (iv) pig bristles; (v) feathers;]

(2) and/or [- blood of animals which did not show any signs of disease communicable through blood to humans or animals, obtained from animals other than ruminants, that have been slaughtered in a slaughterhouse after having been considered fit for slaughter for human consumption following an 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,

(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;]

greaves and centrifuge or separator sludge from milk processing;]

# Fat derivatives not intended for human consumption to be used outside the feed chain

#### COUNTRY

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

- (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;]
- II.5. in case of fat derivatives produced from animal by-products referred to in point II.4.1 and point II.4.2:
  - (a) have been produced using the following methods:
    - (2) either [transesterification or hydrolysis at least 200 °C, under corresponding appropriate pressure, for 20 minutes (glycerol, fatty acids and esters)]
    - (2) or [saponification with NaOH 12M (glycerol and soap):
      - (2) either [in a batch process at 95 °C for three hours;]
      - (2) or [in a continuous process at 140 °C, 2 bars (2000 hPa) for eight minutes;]]
    - $(^2)$  or [hydrogenation at 160 °C at 12 bars (12000 hPa) pressure for 20 minutes;]
  - (b) are packaged in new containers or in containers that have been cleaned, and all precautions are taken to prevent its contamination which bear labels indicating 'NOT FOR HUMAN OR ANIMAL CONSUMPTION';
- II.6. in case of fat derivatives produced from animal by-products referred to in point II.4.3, the fat derivatives have been produced in accordance with one of the processing methods 1-2-3-4-5-6-7 (²) referred to in Chapter III of Annex IV to Regulation (EU) No 142/2011.

#### Notes

#### Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity.
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.
- Box reference I.25: technical use: any use other than for animal consumption.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: Manufacturing plant: provide the registration number of treatment/processing establishment.

COUNTRY		Fat derivatives not intended for human consumption to be used outside the feed chain			
II.	Health information	II.a. Certificate reference No	II.b.		
Part	II:				
( <sup>1a</sup> ) (	( <sup>1a</sup> ) OJ L 300, 14.11.2009, p. 1.				
(1b) (	OJ L 54, 26.2.2011, p. 1.				
(2) [	Delete as appropriate.				
— т	The signature and the stamp must be in a different colour to that of the printing.				
	<ul> <li>Note for the person responsible for the consignment in the European Union: this certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.</li> </ul>				
Offic	ial veterinarian/Official inspector				
1	Name (in capital letters):	Qualification and	title:		
1	Date: Signature:				
8	Stamp:				

# CHAPTER 14(B)

#### Health certificate

For fat derivatives not intended for human consumption to be used as feed or outside the feed chain, intended for dispatch to or for transit through  $\binom{2}{1}$  the European Union

cou	COUNTRY Veterinary certificate to EU				
	I.1.	Consignor	I.2. Certificate reference No I.2.a.		
		Name	I.3. Central competent authority		
		Address			
		Tel.	I.4. Local competent authority		
펕	1.5.	Consignee	I.6. Person responsible for the load in EU		
E	1.5.	Name	Name		
sigi		Address	Address		
8			,		
pe		Postcode	Postcode		
of dispatched consignment		Tel.	Tel.		
disb	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10. Region of Code		
5			destination destination		
Part I : Details		Discover of solution	I 40 Discount destination		
Det	1.11.	Place of origin	I.12. Place of destination		
ا ٿِ ا		Name Approval number Address	Name Custom warehouse ☐ Address Approval number		
Par		Name Approval number	Addiess Approval Humber		
	Address Name Approval number		Postcode		
		Address	144.5.		
	1.13.	Place of loading	I.14. Date of departure		
	l.15.	Means of transport	I.16. Entry BIP in EU		
		Aeroplane ☐ Ship ☐ Railway wagon ☐			
		Road vehicle Other	1.17.		
		Identification			
	140	Documentation references			
	1.18.	Description of commodity	I.19. Commodity code (HS code) 15.16.10		
			I.20. Quantity		
	I.21.	Temperature of product	I.22. Number of packages		
		Ambient Chilled	Frozen 🗆		
	1.23.	Seal/Container No	I.24. Type of packaging		
	1.25.	Commodities certified for:			
		Animal feedingstuff ☐ Technical use ☐			
	126	For transit through EU to third country	I.27. For import or admission into EU		
	1.20.	. –	1.27. For import or admission into Eo		
		Third country ISO code			
	1.28.	Identification of the commodities			
		Species Nature of commodity Approval number of			
		(Scientific name) Manufacturin	ng plant packages		

# Fat derivatives not intended for human consumption to be used as

cou	COUNTRY		feed or outside the feed chain		
	II.	Health infe	ormation	II.a. Certificate reference No	II.b.
		Parliament	ersigned official veterinarian, declare that I have and of the Council ( <sup>1a</sup> ) and in particular Article Annex XIV, Chapter II thereof, and certify that the f	10 thereof, and Commission Regulation	
	II.1.	consist of	fat derivatives that satisfy the health requirements	below;	
cation	II.2. consist of fat derivatives not intended for human consumption;				
II.2. consist of fat derivatives not intended for human consumption;   II.3. have been prepared and stored in a plant approved, validated and supervised by the competent author of Regulation (EC) No 1069/2009, in order to kill pathogenic agents;   II.4. have been prepared from rendered fats exclusively produced from the following Category 3 materials				ority in accordance with Article 24	
Part	II.4. have been prepared from rendered fats exclusively produced from the following Category 3 materials:				
		(²) either	[- carcases and parts of animals slaughtered or, in human consumption in accordance with Union reasons;]		
		(²) and/or	[- carcases and the following parts originating either considered fit for slaughter for human consumpti of animals from game killed for human consumptions.]	ion following an ante-mortem inspection	or bodies and the following parts
			(i) carcases or bodies and parts of animals whi legislation, but which did not show any sign		
			(ii) heads of poultry;		
		(iii) hides and skins, including trimmings and splitting thereof, horns and feet, including the phalanges and the carpus metacarpus bones, tarsus and metatarsus bones, of animals, other than ruminants;			
		(iv) pig bristles;			
			(v) feathers;]		
		(²) and/or	[- blood of animals which did not show any signs from animals other than ruminants that have be slaughter for human consumption following an a	en slaughtered in a slaughterhouse at	ter having been considered fit for
		(2) and/or	[- animal by-products arising from the production of greaves and centrifuge or separator sludge from		nption, including degreased bone,
		(²) and/or	<ul> <li>[- products of animal origin, or foodstuffs contain consumption for commercial reasons or due to which no risk to public or animal health arise;]</li> </ul>		
		(²) and/or	[- petfood and feedingstuffs of animal origin, or fer no longer intended for feeding for commercial re- defects from which no risk to public or animal h	asons or due to problems of manufactu	
		(2) and/or	[- blood, placenta, wool, feathers, hair, horns, hoof any disease communicable through that product		animals that did not show signs of
		(²) and/or	[- aquatic animals, and parts of such animals, exc nicable to humans or animals;]	cept sea mammals, which did not sho	w any signs of diseases commu-
		(²) and/or	[- animal by-products from aquatic animals origin consumption;]	nating from plants or establishments n	nanufacturing products for human
		(²) and/or	[- the following material originating from animals material to humans or animals:	which did not show any signs of dis	ease communicable through that
			(i) shells from shellfish with soft tissue or flesh;		

	Fat derivatives not intended for human consumption to be used as feed or outside the feed chain		
II. Health information	II.a. Certificate reference No	II.b.	
(ii) the following originating from terrestrial animals:			

- hatchery by-products,
- eggs.
- egg by-products, including egg shells;
- (iii) day-old chicks killed for commercial reasons;]
- II.5. are packaged in new containers or in containers which bear labels indicating 'NOT FOR HUMAN CONSUMPTION', that have been cleaned, and all precautions are taken to prevent its contamination.

Notes

#### Part I:

- Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity.
- Box reference I.11 and I.12: Approval number: the registration number of the establishment or plant, which has been issued by the competent
- Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses.
- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.
- Box reference I.25: technical use: any use other than for animal consumption.
- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
- Box reference I.28: Manufacturing plant: provide the registration number of treatment/processing establishment.

- (1a) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.
- The signature and the stamp must be in a different colour to that of the printing.
- Note for the person responsible for the consignment in the European Union: this certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.

Official veterinarian/Official inspector

Qualification and title: Name (in capital letters):

Date: Signature:

Stamp:

### CHAPTER 15

# **Health certificate**

For egg products not intended for human consumption that could be used as feed material, intended for dispatch to or for transit through  $\binom{2}{1}$  the European Union

cou	DUNTRY Veterinary certificate to E			
	I.1.	Consignor	I.2. Certificate reference No I.2.a.	
		Name	I.3. Central competent authority	
		Address	I de la collection de l	
		Tel.	I.4. Local competent authority	
ŧ	I.5.	Consignee	I.6. Person responsible for the load in EU	
me		Name	Name	
nsig		Address	Address	
8		Postcode	Postcode	
chec		Tel.	Tel.	
spat	I.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO I.10. Region of Code	
đ Ģ			destination code destination	
ils o		Place of a late	LAC Element testing	
Part I: Details of dispatched consignment	1.11.	Place of origin	I.12. Place of destination	
Ξ		Name Approval number	Name Custom warehouse ☐ Address Approval number	
Par		Address Name Approval number	Approval Humber	
		Address	Postcode	
		Name Approval number Address	. 30.033	
	I.13.	Place of loading	I.14. Date of departure	
	l.15.	Means of transport	I.16. Entry BIP in EU	
		Aeroplane Ship Railway wagon		
		Road vehicle Other O	1.17.	
		Identification  Documentation references		
	I.18.	Description of commodity	I.19. Commodity code (HS code)	
			35.02	
			I.20. Quantity	
	I.21.	Temperature of product	I.22. Number of packages	
		Ambient Chilled C	Frozen	
	1.23.	Seal/Container No	I.24. Type of packaging	
	1.25.	Commodities certified for:		
	Animal feedingstuff Technical use			
	1.26.	For transit through EU to third country	I.27. For import or admission into EU	
		Third country ISO code		
	1.28.	Identification of the commodities		
		Species Nature of commodity Approval number of ex (Scientific name) Manufacturing		

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

#### COUNTRY

Certification

Part II:

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

I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council (<sup>1a</sup>) and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011 (<sup>1b</sup>), and in particular Annex XIV, Chapter I thereof, and certify that the egg products described above:

- II.1. consist of egg products that satisfy the health requirements below:
- II.2. consist exclusively of egg products not intended for human consumption;
- II.3. have been prepared and stored in a plant, approved, validated and supervised by the competent authority in accordance with Article 24 of Regulation (EC) No 1069/2009 or Article 4(2) of Regulation (EC) No 853/2004 of the European Parliament and of the Council (3), in order to kill pathogenic agents;
- II.4. have been prepared (derived) exclusively with the following animal by-products:
  - (2) either [- animal by-products arising from the production of products intended for human consumption;]
  - (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 [- the following material originating from terrestrial animals which did not show any signs of disease communicable through that material to humans or animals:
    - hatchery by-products,
    - eggs,
    - egg by-products, including egg shells;]
- II.5. have been subjected to processing:

  - (3) or [in accordance to a method and parameters which ensure that the products complies with the microbiological standards set in Chapter I of Annex X, to Regulation (EU) No 142/2011;]
  - (3) or [in accordance with Section X, Chapters I to III of Annex III to Regulation (EC) No 853/2004;]
- II.6. have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply with the following standards (5):

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

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

- II.7. meet Union standards on residues of substances that are harmful or might alter the organoleptic characteristics of the product or make its use as feed dangerous or harmful to animal health;
- II.8. the end product was:
  - (3) either [packed in new or sterilised bags,]
  - (3) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectant approved by the competent authority before use,]

and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION';

- II.9. the end product was stored in enclosed storage;
- II.10. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment.

Notes

#### Part I:

Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit
commodity; it may be filled in if the certificate is for import commodity.

		Egg products not intended for hur used as feed	man consumption that could be		
II.	Health information	II.a. Certificate reference No	II.b.		
_	Box reference I.12: Place of destination: this box is to be filled in only be stored in free zones, free warehouses and custom warehouses.	y if it is a certificate for transit commodi	ty. The products in transit can only		
_	Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in case of unloading and reloading.				
_	Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.				
_	Box reference I.25: technical use: any use other than for animal cor	nsumption.			
_	Box reference I.26 and I.27: fill in according to whether it is a trans	it or an import certificate.			
Par	t II:				
( <sup>1a</sup> )	OJ L 300, 14.11.2009, p. 1.				
( <sup>1b</sup> )	OJ L 54, 26.2.2011, p. 1.				
( <sup>2</sup> )	Delete as appropriate.				
(3)	OJ L 139, 30.4.2004, p. 55.				
(4)	Insert method 1 to 5 or 7 as applicable.				
( <sup>5</sup> )	Where:				
	n = number of samples to be tested;				
	$\mbox{\it m} = \mbox{\it threshold}$ value for the number of bacteria; the result is conexceed $\mbox{\it m};$	nsidered satisfactory if the number of	bacteria in all samples does not		
	$M=\mbox{maximum}$ value for the number of bacteria; the result is consider or more; and	ered unsatisfactory if the number of bac	cteria in one or more samples is M		
	c = number of samples the bacterial count of which may be betwe count of the other samples is m or less.	en m and M, the sample still being cor	nsidered acceptable if the bacterial		
- 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 U the consignment until it reaches the border inspection post.	Inion: this certificate is only for veterinal	ry purposes and has to accompany		
Offic	cial veterinarian/Official inspector				
	Name (in capital letters):	Qualification and	title:		
	Date:	Signature:			
	Stamp:				

#### **CHAPTER 16**

#### **Model declaration**

Declaration by the importer of bones and bone products (excluding bone meal), horns and horn products (excluding horn meal) and hooves and hoof products (excluding hoof meal) intended for use other than as feed material, organic fertilisers or soil improvers for dispatch to the European Union

Note for the importer: This declaration is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post.

I, the undersigned, declare that the following products (1):
(a) bones and bone products (excluding bone meal);
(b) horns and horn products (excluding horn meal);
(c) hooves and hoof products (excluding hoof meal);
are intended to be imported by me into the Union, and I declare that these products will not be diverted at any stage for any use in food, feed material, organic fertilisers or soil improvers and will be conveyed directly for the purpose of further processing or treatment to:
Name:
Furthermore, I declare that the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001 or mechanically separated meat obtained from bones of bovine, ovine or caprine animals.
The importer:
Name:
Done at
Signature
Reference number as indicated on the Common Veterinary Entry Document (CVED) provided for in Annex III to Commission Regulation (EC) No 136/2004:
Official stamp of the border inspection post of entry into the EU (2)
Signature: (Signature of the official veterinarian of the border inspection post) (2)
Name: (Name in capital letters)

<sup>(1)</sup> Delete as appropriate.

 $<sup>(^2)</sup>$  The signature and the stamp must be in a different colour to that of the printing.

### CHAPTER 17

# **Health certificate**

For processed manure, derived products from processed manure and guano from bats intended for dispatch to or for transit through  $(^2)$  the European Union

cou	OUNTRY Veterinary certificate to				
	l.1.	Consignor	I.2. Certificate reference No I.2.a.		
		Name	I.3. Central competent authority		
		Address	n.o. Central competent authority		
			I.4. Local competent authority		
		Tel.			
ĭ	1.5.	Consignee	I.6. Person responsible for the load in EU		
Ē		Name	Name		
sign		Address	Address		
lo co		P			
eq		Postcode Tel.	Postcode Tel.		
tch		10.1	10.1		
of dispatched consignment	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO I.10. Region of Code		
f di			destination code destination		
s o					
Part I: Details	l.11.	Place of origin	I.12. Place of destination		
Ξ		Name Approval number	Name Custom warehouse Address Approval number		
Par		Address	Address Approval number		
		Name Approval number Address			
		Name Approval number	Postcode		
		Address			
	I.13.	Place of loading	I.14. Date of departure		
		· ·	, ,		
	I.15.	Means of transport	I.16. Entry BIP in EU		
		Aeroplane ☐ Ship ☐ Railway wagon ☐			
		Road vehicle Other			
		Identification	1.17.		
		Documentation references			
	110	Description of commodity	I.19. Commodity code (HS code)		
	1.10.	Description of commodity	1.19. Commodity code (115 code)		
			I.20. Quantity		
	1 21	Temperature of product	I.22. Number of packages		
	1.21.				
		Ambient Chilled C	Frozen		
	1.23.	Seal/Container No	I.24. Type of packaging		
	1.25.	Commodities certified for:			
		Technical use			
			LOT For house the advisor late FU		
	1.26.	For transit through EU to third country	I.27. For import or admission into EU		
		Third country ISO code			
	1.28.	Identification of the commodities			
		Species Nature of commodity	Approval number of establishments		
		(Scientific name)	Manufacturing plant		

(1b) OJ L 54, 26.2.2011, p. 1.

Status: This is the original version (as it was originally adopted).

# Processed manure, derived products from processed manure and COUNTRY guano from bats Health information II.a. Certificate reference No II.b. I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council (<sup>1a</sup>) and in particular Article 9 thereof, and Commission Regulation (EU) No 142/2011 (<sup>1b</sup>), and in particular Annex XIV, Chapter II thereof, and certify that the processed manure, the derived products from processed manure and the guano from bats described come from a plant for the manufacture of products for purposes other than feeding to farmed animals, a biogas plant or a composting plant approved by the competent authority of the third country meeting the special conditions laid down in Regulation (EC) No 1069/2009 and in Regulation (EU) No 142/2011; II.1. Certification II.2.(2) have been subjected to: [a heat treatment process of at least 70 °C for at least 60 minutes;] or ≝ Part [an equivalent treatment validated and authorised by the importing Member State in accordance with the specific conditions laid down in Regulation (EC) No 1069/2009 and in Regulation (EU) No 142/2011 as follows: II.3. (a) free from Salmonella (no salmonella in 25 g treated product); (b) free from Escherichia coli or from Enterobacteriaceae (based on the aerobic count: less than 1 000 cfu per gram of treated product); have been subjected to reduction in spore-forming bacteria and toxin formation; II.4. are securely enclosed in: (a) well-sealed and insulated containers; or (b) properly sealed packs (plastic bags or 'big bags'). Notes Part I: Box reference I.6: Person responsible for the consignment in the European Union: this box is to be filled in only if it is a certificate for transit commodity; it may be filled in if the certificate is for import commodity. Box reference I.11 and I.12: Approval number: the registration number of the establishment or plant, which has been issued by the competent Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit can only be stored in free zones, free warehouses and custom warehouses. Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in the event of unloading and reloading. - Box reference I.23; for bulk containers, the container number and the seal number (if applicable) should be given. Box reference I.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.31: Nature of commodity: enter if processed manure, derived products from processed manure or guano from bats. (1a) OJ L 300, 14.11.2009, p. 1.

COUNTRY	guano from bats				
II. Health information	II.a. Certificate reference No	II.b.			
(²) Delete as appropriate.					
- The signature and the stamp must be in a different colour to that of	— The signature and the stamp must be in a different colour to that of the printing.				
	<ul> <li>Note for the person responsible for the consignment in the European Union: This certificate is only for veterinary purposes and has accompany the consignment until it reaches the border inspection post.</li> </ul>				
Official veterinarian/Official inspector					
Name (in capital letters):	Qualification and	I title:			
Date:	Signature:				
Stamp:					

### CHAPTER 18

# **Health certificate**

For horns and horn products, excluding horn meal, and hooves and hoof products, excluding hoof meal, intended for the production of organic fertilisers or soil improvers intended for dispatch to or for transit through  $\binom{2}{}$  the European Union

cou	COUNTRY Veterinary certificate to EU					
	l.1.	Consignor	I.2. Certificate reference No I.2.a.			
		Name Address	I.3. Central competent authority			
		Tel.	I.4. Local competent authority			
Ħ	I.5.	Consignee	I.6. Person responsible for the load in EU			
me		Name	Name			
sigr		Address	Address			
o						
eq		Postcode Tel.	Postcode Tel.			
atch			101.			
ispa	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10. Region of Code destination			
s of dispatched consignment			desination			
Detail	l.11.	Place of origin	I.12. Place of destination			
Part I: Details		Name Approval number Address	Name Custom warehouse ☐ Address Approval number			
_		Name Approval number Address	Postcode			
		Name Approval number Address				
	I.13.	Place of loading	I.14. Date of departure			
	l.15.	Means of transport	I.16. Entry BIP in EU			
		Aeroplane  Ship  Railway wagon  Railway				
		Road vehicle Other O	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	I.22. Number of packages			
			Frozen _			
		Seal/Container No	I.24. Type of packaging			
	1.25.	Commodities certified for:				
		Further process				
	1.26.	For transit through EU to third country	I.27. For import or admission into EU			
		Third country ISO code				
	1.28.	Identification of the commodities				
		Species Approval number of establishments (Scientific name) Manufacturing plant	Net weight Batch number			

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

#### COLINTRY

- Box reference I.28: Nature of commodity.

COUNTRY		JNTRY			organic fertilisers or soil improvers	
		II.	Health info	ormation	II.a. Certificate reference No	II.b.
uoi			and of the	rsigned official veterinarian, declare that I have read a Council ( <sup>1a</sup> ), and Commission Regulation (EU) No 142 and horn products, excluding horn meal, and hooves	2/2011 (1b), and in particular Annex XIV	, Chapter II thereof, and certify that
	ou	II.1.	(²) either	[originate from animals that were slaughtered in a slavesult of such inspection, for slaughter for human or		ortem inspection, and were fit, as a
	II: Certification		(²) or	[originate from animals that did not show clinical sanimals;]	signs of any disease communicable t	hrough that product to humans or
	Part II: C	II.2.	horns, horn 80 °C;	products, hooves and hoof products must have und	dergone a heat treatment for one hou	r at a core temperature of at least
	٦	II.3.	horns must have been removed without opening the cranial cavity;			
		II.4.	at any stag	e of processing, storage or transport every precaution	on shall be taken to avoid cross-contain	mination.
		II.5.	the horns a	and horn products, excluding horn meal, and hooves	and hoof products, excluding hoof me	eal, were packed:
	$\Box$		(2) either	[in new packaging or containers;]		
			(²) or	[in vehicles or bulk containers disinfected prior to lo	ading using a product approved by th	e competent authority;]
			and	[the packaging or containers are marked so as to ind FOR HUMAN AND ANIMAL CONSUMPTION' and to		
		II.6.				
999/2001 of the European Parliament and of the ovine or caprine animals; and the animals from means of gas injected into the cranial cavity or kil			(2) either	[the product does not contain and is not derived for 999/2001 of the European Parliament and of the Co ovine or caprine animals; and the animals from wh means of gas injected into the cranial cavity or killed by means of an elongated rod-shaped instrument in	uncil (4) or mechanically separated me ich this product is derived have not b by the same method or slaughtered by	eat obtained from bones of bovine, been slaughtered after stunning by
			(²) or	[the product does not contain and is not derived from born, continuously reared and slaughtered in a cour accordance with Article 5(2) of Regulation (EC) No	ntry or region classified as posing a n	
		Notes				
		Part I:				
				6: Person responsible for the consignment in the Euray be filled in if the certificate is for import commodit		n only if it is a certificate for transit
			reference I. ority.	11 and I.12: Approval number: the registration numbe	er of the establishment or plant, which	has been issued by the competent
	<ul> <li>Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. The products in transit be stored in free zones, free warehouses and custom warehouses.</li> </ul>			y. The products in transit must only		
	<ul> <li>Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information provided in the event of unloading and reloading.</li> </ul>			or name (ship); information is to be		
		— Вох	reference I.	23: for bulk containers, the container number and the	e seal number (if applicable) must be	given.
		— Вох	reference I.	25: technical use: any use other than for animal con-	sumption.	
		— Вох	reference I.	26 and I.27: fill in according to whether it is a transit	or an import certificate.	

COUNTRY	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.					
Part II:							
( <sup>1a</sup> ) OJ L 300, 14.11.2009, p. 1.	( <sup>1a</sup> ) OJ L 300, 14.11.2009, p. 1.						
( <sup>1b</sup> ) OJ L 54, 26.2.2011, p. 1.							
(²) Delete as appropriate.							
(3) Type of product: horns, horn products, hooves, hoof products.							
( <sup>4</sup> ) OJ L 147, 31.5.2001, p. 1.							
- The signature and the stamp must be in a different colour to that	t of the printing.						
Note for the person responsible for the consignment in the Europe the consignment until it reaches the border inspection post.	an Union: this certificate is only for veteri	nary purposes and must accompany					
Official veterinarian/Official inspector							
Name (in capital letters):	Qualification and	d title:					
Date:	Signature:						
Stamp:							

### CHAPTER 19

# **Health certificate**

For gelatine not intended for human consumption to be used by the photographic industry, intended for dispatch to the European Union

cou	COUNTRY Veterinary certificate to EU						
	l.1.	Consignor	I.2. Certificate reference No I.2.a.				
		Name					
		Address	I.3. Central competent authority				
	Tel.		I.4. Local competent authority				
<b>#</b>	1.5.	Consignee	I.6. Person responsible for the load in EU				
la e		Name	Name				
ign		Address	Address				
ő							
ğ		Postcode	Postcode Tel.				
tch		Tel.					
dispatched consignment	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10. Region of Code destination				
s of							
Part I: Details of	l.11.	Place of origin	I.12. Place of destination				
벁		Name Approval number	Name Custom warehouse ☐				
Pa		Address	Address Approval number				
		Name Approval number Address					
		Name Approval number Address	Postcode				
	I.13.	Place of loading	I.14. Date of departure				
	l.15.	Means of transport	I.16. Entry BIP in EU				
		Aeroplane ☐ Ship ☐ Railway wagon ☐					
		Road vehicle Other	I.17. Number(s) of CITES				
		Identification					
		Documentation references					
	1.18.	Description of commodity	I.19. Commodity code (HS code)				
		,	35.03				
			I.20. Quantity				
	1.21.	Temperature of product Ambient ☐ Chilled ☐	I.22. Number of packages				
	1.00	Seal/container No	I.24. Type of packaging				
	1.23.	Seal/Corrainer No	1.24. Type or packaging				
	I.25. Commodities certified for:						
	Technical use						
	1.26.		1.27. For import or admission into EU				
	I.28. Identification of the commodities						
		Oncoins Annual number of antablish	ments Net weight Batch number				
		Species Approval number of establish (Scientific name) Manufacturing plant	ments Net weight Batch number				

#### Gelatine not intended for human consumption to be used by the photographic industry

#### COUNTRY

Part 11.4.

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

I, the undersigned official, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council (<sup>1a</sup>) and in particular Articles 8 and 10 thereof, and Commission Regulation (EU) No 142/2011 (<sup>1b</sup>), and in particular Annex XIV, Chapter II thereof, and certify that the photographic gelatine described above:

- II.1. consists exclusively of photographic gelatine for photographic uses and is not intended for any other purpose;
- has been prepared and stored in a plant registered and supervised by the competent authority in accordance with Article 23 of Regulation (EC) No 1069/2009, which does not produce gelatine for food, feed or other uses intended for dispatch to the European II.2. Certification
- II.3. has been prepared with Category 3 animal by-products and/or bovine vertebral column classified as Category 1 material;
  - has been wrapped, packaged in new containers, stored and transported in sealed, leak-proof labelled containers in a vehicle under satisfactory hygiene conditions;
- II.5. has been produced by a process ensuring that the raw material is:
  - (3) either treated by pressure sterilisation as referred to in definition No 19 of Article 3 of Regulation (EC) No 1069/2009 (2);
  - (3) or subjected to:
    - (i) treatment with acid for at least two days, washing with water and treatment with an alkaline solution for at least 20 days; the pH must be adjusted and the material purified by means of filtration and sterilised at 138-140 °C for 4 seconds; or
    - (ii) treatment with alkali for at least two days, washing with water and treatment with an acid solution for 10-12 hours; the pH must be adjusted and the material purified by means of filtration and sterilised at 138-140 °C for 4 seconds.
- has been wrapped and packaged in wrappings and packages carrying the words 'PHOTOGRAPHIC GELATINE FOR THE PHOTOGRAPHIC INDUSTRY ONLY'. II.6.

#### Notes

#### Part I:

- Box reference I.5: The intended destination of the photographic gelatine can only be the Czech Republic, the Netherlands or the United
- Box reference I.9: Country of destination: only applicable for the Czech Republic, the Netherlands or the United Kingdom.
- 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.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in the event of unloading and reloading.
- Box reference I.23: Identification of container/seal number: only where applicable.
- Box reference I.25; technical use; any use other than for animal consumption.

#### Part II:

- (1a) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26,2,2011, p. 1.
- (2) Pressure sterilisation (method 1) is also referred to in Chapter III of Annex IV to Regulation (EU) No 142/2011 as follows:

#### 'Reduction

1. If the particle size of the animal by-products to be processed is more than 50 millimetres, the animal by-products must be reduced in size using appropriate equipment, set so that the particle size after reduction is no greater than 50 millimetres. The effectiveness of the equipment must be checked daily and its condition recorded. If checks disclose the existence of particles larger than 50 millimetres, the process must be stopped and repairs made before the process is resumed.

C	DUNTRY	Gelatine not intended for human consumption to be used by the photographic industry				
	II. Health information	II.a. Certificate reference No	II.b.			
Time, temperature and pressure						
	2. The animal by-products with the particle size of no greater than 50 millimetres must be heated to a core temperature of more than 133 °C for at least 20 minutes without interruption at a pressure (absolute) of at least 3 bars. The pressure must be produced by the evacuation of all air in the sterilisation chamber and the replacement of the air by steam ("saturated steam"); the heat treatment may be applied as the sole process or as a pre- or post-process sterilisation phase.					
	3. The processing may be carried out in batch or continuous systematical systems.	ems.'				
(	3) Delete as appropriate.					
-	- The signature and the stamp must be in a different colour to that of	the printing.				
-	<ul> <li>Note for the person responsible for the load in the European Union: consignment until it reaches the factory of destination from the border</li> </ul>		rposes and has to accompany the			
0	Official veterinarian/Official inspector					
	Name (in capital letters):	Qualification an	d title:			
	Date: Signature:					
	Stamp:					

# **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, in vitro diagnostics and laboratory reagents

COUNTRY Veterinary certificate to EU											
	l.1.	Consignor			1.2.	Certificat	e reference	No	1.2	2.a.	
		Name			1.3.	Central o	competent a	uthority			
		Address					ompotont a	attriority			
		Tel.		I.4. Local competent authority							
eut	1.5.	. Consignee			I.6. Person responsible for the load in EU						
ᄩ	Name			Name							
nsić	Address			Address							
8		Postcode				Postcode	е				
þe		Tel.				Tel.					
dispatched consignment	17	Country of origin ISO code	LO Basian of avisin	Code	1.9.	Carratar		ISO	140	Degien of	Code
disb	1.7.	Country of origin 180 code	I.8. Region of origin	Code	1.9.	Country	on	code	1.10.	Region of destination	Code
5											
l: Details	l.11.	Place of origin			I.12.	Place of	destination				
ä		Name	Approval number			Name			Cus	stom warehouse	1
Part		Address	Approvar namber			Address				proval number	
٩.		Name	Approval number								
	Address  Name Approval number Address					Postcode	е				
	I.13.	Place of loading			l.14.	Date of	departure				
	I.15.	5. Means of transport			I.16. Entry BIP in EU						
		Aeroplane ☐ Ship ☐ Railway wagon ☐									-
		Road vehicle Other			1.17.						
		Identification									
		Documentation references									
	I.18.	Description of commodity					I.19. Comr	nodity co	de (HS	S code)	
								1.20.	Quant	tity	
	I.21.	Temperature of product			I.22. Number of packages						
	Ambient ☐ Chilled ☐				Frozer						
	1.23.	I.23. Seal/Container No			I.24. Type of packaging						
	I.25. Commodities certified for:										
	Technical use □										
	_										
	1.26.	.26. For transit through EU to third country			I.27. For import or admission into EU			J			
	Third country ISO code										
	I.28. Identification of the commodities										
	Species Approval number of establish				ments			Net weig	ht	Batch	number
	(Scientific name) Manufacturing plant										
	1										

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

#### COUNTRY

Part

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 the Union and satisfy the definition provided for in point 35 of Annex I of 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,]
- (2) and/or [- active implantable medical devices,]
- (2) and/or [- in vitro diagnostic medical devices,]
- (2) and/or [- laboratory reagents;]
- (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 handling or transformation such as mixing, coating, assembling, packaging or labelling to make it suitable for placing on the market or putting into service as medicinal products, veterinary medicinal products, active implantable medical device, medical devices or in vitro diagnostic medical device in accordance with the Union legislation (1b) applicable to those products or as laboratory reagents;
- (3) it has been derived from the following material which may have originated from animals which have been submitted to illegal treatment as defined in Article 1(2)(d) of Council Directive 96/22/EC or Article 2(b) of Council Directive 96/23/EC (2):
- (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, of animals other than ruminants;
  - (iv) pig bristles;
  - (v) feathers;]
- (2) and/or [- blood of animals which did not show any signs of disease communicable through blood to humans or animals obtained from animals other than ruminants that have been slaughtered in a slaughterhouse after having been considered fit for slaughter for human consumption following an 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 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;]

II.a. Certificate reference No

Status: This is the original version (as it was originally adopted).

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

#### COUNTRY

Г.				
1 (2)	) and/or [- aquatic animals, and parts of such animals	event cas mammale	which did not show any signs of a	diseases communicable to humans

- (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.

Health information

- 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 die 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/ACTIVE IMPLANTABLE MEDICAL DEVICES/IN VITRO DIAGNOSTIC MEDICAL DEVICES/LABORATORY REAGENTS ONLY' and it is not intended to be diverted at any stage within the Union for any other use;
- (5) the consignment will be transported directly to the place of destination as indicated under point I.12 of this declaration, that is:
  - an establishment or plant for the production of medicinal products, veterinary medicinal products, medical devices, in vitro diagnostics
    or laboratory reagents, 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 shall only be dispatched to an establishment or plant referred to in the preceding subpoint of (5).

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

COUNTRY		diagnostics and laboratory reagents				
II.	Health information	II.a. Certificate reference No	II.b.			
Not	es					
-	Box reference I.25: technical use: any use other than for animal con	sumption.				
( <sup>1a</sup> )	( <sup>1a</sup> ) OJ L 54, 26.2.2011, p. 1.					
(1b)	(1b) Directive 2001/82/EC of the European Parliament and of the Council of 6 November 2001 on the Community code relating to veterinar medicinal products (OJ L 311, 28.11.2001), 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 199 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), as appropriate.					
(2)	Delete as appropriate.					
The	importer					
	Name (in capital letters):	Address:				
	Date:	Signature:				