COMMISSION REGULATION (EU) 2021/1891

of 26 October 2021

amending Annexes XIV and XV to Regulation (EU) No 142/2011 as regards imports into and transit through the Union of animal by-products and derived products

(Text with EEA relevance)

THE EUROPEAN COMMISSION,

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

Having regard to Regulation (EC) No 1069/2009 of the European Parliament and of the Council of 21 October 2009 laying down health rules as regards animal by-products and derived products not intended for human consumption and repealing Regulation (EC) No 1774/2002 (Animal by-products Regulation) (¹), and in particular Article 41(3), first and third subparagraphs, and Article 42(2), points (a), (b) and (d), thereof,

Whereas:

- (1) Commission Regulation (EU) No 142/2011 (²) lays down implementing measures for the public and animal health rules for animal by-products and derived products laid down in Regulation (EC) No 1069/2009, including models of health certificates and the list of third countries authorised for imports into and transit through the Union of consignments of animal by-products and derived products.
- (2) In particular, Chapter II of Annex XIV to Regulation (EU) No 142/2011 sets out the specific requirements for the importation into and the transit through the Union of consignments of animal by-products and derived products for uses outside the feed chain for farmed animals other than fur animals. Such consignments are required to comply with, inter alia, the rules set out in Table 2 of Section 1 of that Chapter.
- (3) More specifically, row 14 of Table 2 sets out, inter alia, the list of third countries authorised for imports into and transit though the Union of consignments of animal by-products and derived products for uses outside the feed chain, including consignments of fur for the manufacture of derived products, category 3 materials, referred to in Article 10, point (n), of Regulation (EC) No 1069/2009. Certain Member States have requested that row 14 of Table 2 be amended so as to include a list of third countries authorised for imports into the Union of fur for the manufacture of derived products. There is not a list of third countries authorised for imports into the Union of products of fur animals, but Commission Implementing Regulation (EU) 2021/404 (³) sets out a list of third countries, territories or zones thereof authorised for the entry into the Union of consignments of fresh meat of ungulates. Following an evaluation of the request by the Member States, it is appropriate to include a list of third countries from which fur for the manufacture of derived products may be imported into the Union in row 14 of Table 2. As the third countries from which the entry into the Union of fresh meat of ungulates is authorised for mythematic for the entry into the Union in row 14 of Table 2. As the third countries from which the entry into the Union of fresh meat of ungulates is authorised for mythematic for the entry into the Union of fresh meat of ungulates is authorised for the entry into the Union of fresh meat of third countries from which the further of products and protection of public and animal health, it is opportune to allow imports of fur for the manufacture of derived products from those third countries.

⁽¹⁾ OJ L 300, 14.11.2009, p. 1.

⁽²⁾ Commission Regulation (EU) No 142/2011 of 25 February 2011 implementing Regulation (EC) No 1069/2009 of the European Parliament and of the Council laying down health rules as regards animal by-products and derived products not intended for human consumption and implementing Council Directive 97/78/EC as regards certain samples and items exempt from veterinary checks at the border under that Directive (OJ L 54, 26.2.2011, p. 1).

^{(&}lt;sup>3</sup>) Commission Implementing Regulation (EU) 2021/404 of 24 March 2021 laying down the lists of third countries, territories or zones thereof from which the entry into the Union of animals, germinal products and products of animal origin is permitted in accordance with Regulation (EU) 2016/429 of the European Parliament and the Council (OJ L 114, 31.3.2021, p. 1).

- (4) Annex XIV to Regulation (EU) No 142/2011 should therefore be amended accordingly.
- In addition, Chapter 3(F) and Chapter 8 of Annex XV to Regulation (EU) No 142/2011 set out models of health (5) certificates for imports into, or transit through, the Union, of animal by-products for the manufacture of petfood and for those used for purposes outside the feed chain or for trade samples, respectively. Those model health certificates contain among others, the requirement that the animals from which animal by-products are derived must have stayed in a single holding for 40-days before slaughter. From an animal health point of view, such a 40-day pre-slaughter residency period ensures the safety of unprocessed animal by-products when they are imported into the Union. Freedom from foot-and-mouth-disease without practicing vaccination is the most favourable animal health status in accordance with standards of the World Organisation for Animal Health (OIE), and third countries with that animal health status are authorised for imports into the Union and transit through the Union of consignments of fresh meat without such a 40-day residency period, provided that they comply with all other animal health requirements. Certain third countries that are free of foot-and-mouth disease without practicing vaccination have asked the Commission to be authorised for imports into the Union of unprocessed animal by-products without the 40-day pre-slaughter residency period. Animal health conditions for imports of animal by-products should be aligned with animal health requirements for entry into the Union of fresh meat laid down in and Commission Implementing Regulation (EU) 2021/404.
- (6) The model health certificates for animal by-products for the manufacture of petfood, set out in Chapter 3(F) of Annex XV to Regulation (EU) No 142/2011, and for animal by-products to be used for purposes outside the feed chain or for trade samples, set out in Chapter 8 of Annex XV to that Regulation, should therefore be amended accordingly.
- (7) Furthermore, Chapter V of Annex VIII to Regulation (EU) No 142/2011 provides that derived products of Category 1 or Category 2 material should be permanently marked with a chemical marker to prevent their entry into the feed chain and to ensure official controls of feed. Marking with a chemical marker is not required for Category 3 rendered fats. It is therefore necessary to correct the erroneous wording in the model health certificate set out in Chapter 10(B) of Annex XV to that Regulation for imports of 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 the Union.
- (8) Annex XV to Regulation (EU) No 142/2011 should therefore be amended accordingly.
- (9) In order to avoid any disruption of trade, this Regulation should provide for a transitional period during which the commodities concerned by the amendments made to Regulation (EU) No 142/2011, by this Regulation, should continue to be accepted for importation into and transit through the Union, provided that those commodities comply with the requirements laid down in Regulation (EU) No 142/2011 before the amendments made by this Regulation.
- (10) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on Plants, Animals, Food and Feed,

HAS ADOPTED THIS REGULATION:

Article 1

Annexes XIV and XV to Regulation (EU) No 142/2011 are amended in accordance with the Annex to this Regulation.

Article 2

For a transitional period until 31 May 2022, consignments of animal by-products and of derived products accompanied by a health certificate duly completed and signed in accordance with the appropriate model health certificate set out in Chapter 3(F), Chapter 8 and Chapter 10(B) of Annex XV to Regulation (EU) No 142/2011, in the version applicable before the amendments provided for by Article 1 of this Regulation, shall continue to be accepted for importation into or transit through the Union, provided that such health certificates were duly completed and signed no later than 31 March 2022.

Article 3

This Regulation shall enter into force on the twentieth day following that of its publication in the Official Journal of the European Union.

This Regulation shall be binding in its entirety and directly applicable in all Member States.

Done at Brussels, 26 October 2021.

For the Commission The President Ursula VON DER LEYEN ANNEX

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

1. in Annex XIV, Chapter II, Section 1, Table 2, row 14, column 'third countries' list', the following point (d) is added:

'(d) In the case of fur for the manufacture of derived products:

Third countries listed in Part 1 of Annex XIII to Commission Implementing Regulation (EU) 2021/404 (*) from which the entry into the Union of fresh meat of ungulates is authorised.

- (*) Commission Implementing Regulation (EU) 2021/404 of 24 March 2021 laying down the lists of third countries, territories or zones thereof from which the entry into the Union of animals, germinal products and products of animal origin is permitted in accordance with Regulation (EU) 2016/429 of the European Parliament and the Council (OJ L 114, 31.3.2021, p. 1).'.
- 2. Annex XV is amended as follows:

(a) Chapter 3(F) is replaced by the following:

'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	NTRY:				Veterinary certificate to EU			
	I.1. Consignor		I.2. Certificate refe	erence No	I.2.a.			
	Name		I.3. Central competent authority					
	Address		I.4. Local compete					
				autionty				
	Tel.							
Jent	I.5. Consignee		I.6. Person respon	sible for the load	in EU			
ignn	Name		Name	Name				
ons	Address		Address					
ed c								
atch	Postcode Tel.		Postcode					
lispa	3 199900	LO Deview of Code	Tel.	100 and a	110 Deview of Code			
ofo	I.7. Country of ISO code origin	I.8. Region of Code origin	e I.9. Country of destination	ISO code	I.10. Region of Code destination			
tails	l	l	destination	I	destination			
Ď								
Part I: Details of dispatched consignment	I.11. Place of origin		I.12. Place of destin	nation				
•	Name Address	Approval number	Nama		om warehouse			
	Name	Approval number	Name Address	Appr	oval number			
	Address Name	Approval number	Postcode					
	Address							
	I.13. Place of loading		I.14. Date of depart	ure				
	I.15. Means of transport		I.16. Entry BIP in El	U				
		p □ Railway wagon □						
		ier 🗆	1.17.					
	Identification Documentation reference	•						
	Documentation reference	5						
	I.18. Description of commodity			I.19. Commo	odity code (HS code)			
					I.20. Quantity			
	I.21. Temperature of product				I.22. Number of packages			
	Ambient	Chilled	Frozen					
	I.23. Seal/Container No				I.24. Type of packaging			
	I.25. Commodities certified for:	:						
	Manufacture of petfood	J Fu	ther process \Box		Technical use 🗆			
	I.26. For transit through EU to	third country	I.27. For impor	t or admission int	o EU 🗆			
	-							
	Third country	ISO code						
	I.28. Identification of the comm	odities						
		Approva	I number of establishme	ents				
	Species (Scientific name)	Nature of commodity	lanufacturing plant N	Number of packag	es Net weight Batch number			

	COUNTRY	Animal by-products for the manufacture of petfood								
	н.	Health info	ormation	II.a. Certificate refere	nce No	II.b.				
		1069/2009 142/2011(1	I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council ^(1a) and Commission Regulation (EU) No 142/2011 ^(1b) , and in particular Chapter II of Annex XIV thereto, and certify that the animal by-products described above:							
		II.1.1. consist of animal by-products that satisfy the animal health requirements below;								
ication		II.1.2. (²)either								
rtifi		(²)or								
Part II: Certification		(²)or	that are anir			or Lagomorpha, aquatic animals or				
Ра	(²)either	[II.1.3.	eradicate any	epizootic disease, and whi		ch were not slaughtered or killed to				
			(a) come	rom holdings where						
			(i)	mplementing Regulation (of rinderpest, Newcastle d he period of the precedin during the period of the pre vicinity within a 10 km radi	(EU) 2018/18 isease or hig ig 30 days, n eceding 40 da us, during the	animals are listed in accordance with 82, there has been no case/outbreak hly pathogenic avian influenza during or of classical or African swine fever ys; nor in the holdings situated in their period of the preceding 30 days; and				
			(ii)	period of the preceding 60) days, nor in	of foot-and-mouth disease during the the holdings situated in their vicinity of the preceding 30 days; and				
		(²)either	[(b) have remained in their holdings of origin for a period of at least 40 days before the date of departure and which have been transported directly to the slaughterhouse without any contact with other animals which did not comply with the same health conditions;]]							
		(²) or	[(b) have remained on holdings under veterinary supervision in the third country or part of the territory of the third country–of origin from which imports of fresh meat of ungulates are authorised without any restrictions in accordance with Implementing Regulation (EU) 2021/404, and at the slaughterhouse							
			(i)		ghter and hav	spection during the period of 24 hours re shown no evidence of the diseases are susceptible; and				
			(ii)	with the relevant provision	s of Union leg	ne of slaughter or killing in accordance islation and have met requirements at napters II and III of Council Regulation				
	(²)or	[11.1.3.		tained or produced from use, and which	animals whi	ch were not killed to eradicate any				
			(a) have	een captured and killed in	the wild in ar	area:				
			(i)	following diseases for w Implementing Regulation rinderpest, Newcastle dise	hich the ani n (EU) 20 ease or highly days, nor of	s been no case/outbreak of any of the mals are listed in accordance with 18/1882: foot-and-mouth disease, pathogenic avian influenza during the classical or African swine fever during d				
			(ii)	of a country not authorised during the preceding 30 d days; and	for export to t ays or of pore	rom any country or part of the territory he European Union of poultry material cine material during the preceding 40				
			chillin		re and immed	riod of 12 hours following the killing for diately afterwards to a game-handling establishment;]				

COUNTRY			Animal by-products for the manufacture of petfood				
П.	Health inform	nation	II.a. Certificate reference No II.b.				
	II.1.4.	been no cas susceptible o the preparati only after th	btained in an establishment around whic e/outbreak of the diseases referred to in luring the period of the preceding 30 day on of raw material for exportation to the ne removal of all meat, and the tota at under the control of an official veterina	point II.1.3 for which the animals are s or, in the event of a case of disease European Union has been authorised al cleaning and disinfection of the			
	II.1.5.	comply with	btained and prepared without contact w the conditions required above, and it n with pathogenic agents;				
	II.1.6.	containers be	packed in new packaging preventing a paring the label indicating 'RAW MATERI DD' and the name and address of the nion;	AL ONLY FOR THE MANUFACTURE			
	II.1.7.	consist only	of the following animal by-products:				
	(²)either	of an	ases and parts of animals slaughtered or imals killed which were deemed fit for hu n legislation until irreversibly declared a ons;]	uman consumption in accordance with			
	(²)and/or	slaug cons of an	ases and the following parts originating ghtered in a slaughterhouse and were of umption following an ante-mortem inspe nimals from game killed for human con lation:	considered fit for slaughter for humar ction or bodies and the following parts			
		(i)	carcases or bodies and parts of animals consumption in accordance with Unio any signs of disease communicable to	on legislation, but which did not show			
		(ii) (iii)	heads of poultry; hides and skins, including trimmings including the phalanges and the carpu metatarsus bones;				
		(iv) (v)	pig bristles; feathers;]				
	(²)and/or	cons	al by-products arising from the produc umption, including degreased bone, g ge from milk processing;]				
	(²)and/or	are r prob	ucts of animal origin, or foodstuffs conta to longer intended for human consumpt lems of manufacturing or packaging defe iblic or animal health arise;]	ion for commercial reasons or due to			
	(²)and/or		tic animals, and parts of such animals, any signs of diseases communicable to				
	(²)and/or	-	al by-products from aquatic animals orig ufacturing products for human consumpl				
	(²)and/or	-	ollowing material originating from anim ase communicable through that material shells from shellfish with soft tissue or the following originating from terrestria – hatchery by-products, – eggs, – egg by-products, including egg sh	to humans or animals: flesh; Il animals:			
		(iii)	day-old chicks killed for commercial re	asons;]			
	(²)and/or	-	al by-products from aquatic or terrestr ogenic to humans or animals;]	ial invertebrates, other than specie			

COUNTRY		Animal by-products for the manufacture of petfood				
II. Health inform	mation	II.a. Certificate re	eference No	II.b.		
(²)and/or	- except Regula	ategory 1 material	as referred to in 7 009 and Category	Article 8, point (a)(iii), (iv) and (v), of 2 material as referred to in Article 9,		
(²)and/or	prohib	d by Council Dire in accordance	ective 96/22/EC(4a)	ed with certain substances which are), the import of the material being point (a)(ii) of Regulation (EC) No		
II.1.8.	European Uni	legislation in such ant of destination i	n a way that they	been preserved in accordance with will not spoil between dispatch and nion or during the transit through the		
II.1.9.	substances pro	bited by Directive 9	6/22/EC for the ma	hich have been treated with certain inufacture of petfood, the import being Regulation (EC) No 1069/2009:		
	Union by frozen b into sepa Europea of each p	been marked in the third country before entry into the territory of the European by a cross of liquefied charcoal or activated carbon on each outer side of each block, or, when the raw material is transported in pallets which are not divided parate consignments during transport to the petfood plant of destination in the ean Union or during the transit through the European Union, on each outer side n pallet, in a way that the marking covers at least 70 % of the diagonal length of zen block and be of at least 10 cm width;				
	third cou liquefied	y before entry into	the territory of the ting charcoal powd	raw material has been marked in the e European Union by spraying it with ler in such a way that the charcoal is		
	referred	above and other ne		w material which has been treated as terial, all the raw materials have been		
(²)(⁵)[II.2.	Specific requir	ents				
(²)(⁶)[II.2.1.	referred to in p	nt (II.1.2), where va	accination program	als that have been kept in the territory mes against foot-and-mouth disease in domestic bovine animals.]		
(²)(⁷)[II.2.2.	offal of domes than + 2 °C for	ruminants, which period of at least th	have maturated hree hours, or in the	mal by-products derived from trimmed at an ambient temperature of more case of masseter muscles of bovine period of at least 24 hours.]]		
(²)[II.3.	the animal by-	ne animal by-products for the manufacture of petfood contains or is obtained from animal- y products of ruminant origin and:				
(²)either	spongiform e	ephalopathy (BSE		ified as posing a negligible bovine dance with Commission Decision enous BSE case;]]		
(²)or	with Decision 2 by-products or the ban on the ruminants, as Health Code, h	7/453/EC in which erived products we beding of ruminant ined in the World been effectively e	there has been an re derived from an ts with meat-and-b Organisation for An nforced in that cou			
	(²)or [e derived from bov I slaughtered in a	ine, ovine or caprir	ovine, ovine or caprine animals.]]] ne animals born, continuously reared classified as posing a negligible BSE 53/EC.]]]		
		not contain: specified risk m	naterial as defined	in point 1 of Annex V to Regulation n Parliament and of the Council(9);		

COUNTRY

Animal by-products for the manufacture of petfood

II.	Health information	II.a. Certificate reference No II.b.
	(b	caprine animals, except from those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC, in which there has been no indigenous BSE case,
Note		
Part		
_	Box reference I.6: Person responsible	e for the consignment in the European Union: this box is to be filled in only if ; it may be filled in if the certificate is for a commodity to be imported into the
_		ion: this box is to be filled in only if it is a certificate for a transit commodity. d in free zones, free warehouses and custom warehouses.
-	5	per (railway wagons or container and lorries), flight number (aircraft) or name n the case of unloading and reloading in the European Union.
-	Box reference I.19: use the appropriat 23.01; 41.01.	te Harmonized System (HS) code: 05.04; 05.06; 05.07; 05.11.91 or 05.11.99;
—	Box reference I.23: for bulk containers	s, the container number and the seal number (if applicable) must be included.
-	Box reference I.25: technical use: an production or manufacturing of petfor	y use other than feeding of farmed animals, other than fur animals, and the od.
—	Box reference I.26 and I.27: fill in acc	ording to whether it is a transit or an import certificate.
—	Box reference I.28:	
		wing: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or Suidae, a, invertebrates other than Mollusca and Crustacea;
	 manufacturing plant: provide 	the veterinary control number of the approved establishment.
Part	П:	
(1a)	OJ L 300, 14.11.2009, p. 1.	
(^{1b})	OJ L 54, 26.2.2011, p. 1.	
(1c)	The name and ISO code number of t	ne exporting country as laid down in:
		sion Implementing Regulation (EU) 2021/404 (OJ L 114, 31.3.2021, p. 1);
		enting Regulation (EU) 2021/404, and
		n Regulation (EC) No 119/2009 (OJ L 39, 10.2.2009, p. 12).
	species concerned) must be included	
(^{1d})	Only for countries from which game authorised for importation into the Eu	e meat intended for human consumption of the same animal species is ropean Union.
(2)	Delete as appropriate.	
(3)	Excluding raw blood, raw milk, hides certificates in that Annex for the impo	and skins, hooves and horn, pig bristles and feathers (see relevant specific rt of these products).
(4)	OJ L 303, 18.11.2009, p. 1.	
(^{4a})	OJ L 125, 23.5.1996, p. 3.	
(5)	South American or South African con domestic ruminants for human cons masseter muscles of bovine animals	vided when the material of domestic ruminants originated in the territory of a untry or part thereof from where only maturated and deboned fresh meat of sumption is permitted for exportation to the European Union. The whole s, incised in accordance with Section IV, Chapter I, Part B.1, of Annex I to European Parliament and of the Council (OJ L 139, 30.4.2004, p. 206), are

Animal by-products for the manufacture of petfood

п.	Health information	II.a.	Certificate reference No	II.b.							
(6)											
(7)	Only for certain South American and South African countries.										
(8)	OJ L 172, 30.6.2007, p. 84.										
(9)	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 must accompany the consignment until it reaches the border inspection post of the European Union. 										
Offic	ial veterinarian/Official inspector										
	Name (in capital letters):			Qualification and title:							
	Date:			Signature:							
	Stamp:										
Ĺ											
;											

(b) Chapter 8 is replaced by the following:

'CHAPTER 8

Health certificate

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

COUNTRY:

Veterinary certificate to EU

	I.1. Consignor	I.2. Certificate reference No	l.2.a.				
	Name Address	I.3. Central competent authority					
	Tel.	I.4. Local competent authority					
Part I: Details of dispatched consignment	I.5. Consignee Name Address Postcode Tel.	I.6. Person responsible for the load in EU Name Address Postcode					
che	101.	Tel.					
of dispat	I.7. Country of ISO code I.8. Region of Code origin origin	I.9. Country of ISO destination code	I.10. Region of Code destination				
ails c	I.11. Place of origin	I.12. Place of destination					
Part I: Det	Name Approval number Address Name Approval number		ustom warehouse D oproval number				
	Address Name Approval number Address	Postcode					
	I.13. Place of loading	I.14. Date of departure					
	I.15. Means of transport	I.16. Entry BIP in EU					
	Aeroplane 🛛 Ship 🗆 Railway wagon 🗆						
	Road vehicle Other Identification	l.17.					
	Documentation references I.18. Description of commodity	1.19. Com	modity code (HS code)				
			I.20. Quantity				
	I.21. Temperature of product Ambient Chilled	Frozen 🛛	I.22. Number of packages				
	I.23. Seal/Container No	I.24. Type of packaging					
	I.25. Commodities certified for:						
	Technical use 🗆						
	I.26. For transit through EU to third country	I.27. For import or admission	into EU 🛛				
	Third country ISO code						
	I.28. Identification of the commodities						
		ber of establishments uring plant Number of packages	Net weight Batch number				

EN

	COUNTRY				al by-products to be used for purpos chain or for trade samples ⁽²⁾	es outside the			
	н.	Health info	ormatio	n	II.a. Certificate reference No	II.b.			
Part II: Certification		1069/2009	of the l o), and i above [are ti analy Regu CONS	European Parliament and n particular Chapter II of , rade samples which cons ses as referred to in the lation (EU) No 142/2011, SUMPTION'.]	are that I have read and understood R of the Council(^{1a}), and Commission R Annex XIV thereto, and certify that the a st of animal by-products intended for pa e definition of trade samples in point that bear the label 'TRADE SAMPLE N ements set out in point II.1.];	egulation (EU) No Inimal by-products articular studies or 39 of Annex I to			
o :≡	II.1.	The animal	by proc	ducts described above					
Part	II.1.1	The animal by products described above have been							
		(²)either	[(a)	obtained from materials imported from a third country, territory or p thereof: (3) authorised to export fresh meat to European Union;]					
		(²)and/or	[(b)	obtained in the	exporting third country, terr (³) from animals that	itory or part			
				fresh meat to the preceding three r	that third country, territory or part there European Union since birth or for a pe nonths before the date of slaughter; and wild in that third country, territory or par	riod of at least the l/or			
		(²)and/or	[(c)		rodents, lagomorphs, or aquatic anima				
	(²)[II.1.2.		lerived from eggs, milk, rodents, lagomo ebrates and unprocessed furs, have b						
		(²)either	[(a)	there has not b disease, Newcas period of the pre during the period	llowing diseases for which the animal een any case/outbreak of rinderpest de disease or highly pathogenic avian ir ceding 30 days, nor of classical or A of the preceding 40 days; nor in the h n a 10 km radius, during the period of	, swine vesicular Ifluenza during the frican swine fever oldings situated in			
				during the period their vicinity with days; and	not been any case/outbreak of foot-a of the preceding 60 days, nor in the h n a 25 km radius, during the period of	oldings situated in			
			(b)	which: (i) were not killed to [(²)either	eradicate any epizootic disease;				
				(ii) remained on thei the date of de slaughterhouse w the same health of	r holdings of origin for a period of at lea parture and which were transported ithout contact with other animals which o conditions;	d directly to the			
				or part of the terri meat of ungulates	n holdings under veterinary supervision ory of the third country-of origin from whi are authorised without any restrictions ementing Regulation (EU) 2021/404]	ch imports of fresh			
				(iii) at the slaughterho period of 24 hour	ouse, passed the ante-mortem health ins s before the time of slaughter and show	ed no evidence of			
				(iv) were handled in killing in accorda complied with re	rred to above for which the animals are he slaughterhouse before and at the ti nce with the relevant provisions of Uni equirements at least equivalent to th I of Council Regulation (EC) No 1099/2	me of slaughter or on legislation and ose laid down in			

COUNTRY

II.	Health information	on		II.a. Certificate reference No	II.b.
	(²)or	[(a)	following disease disease, rinderpe during the period fever during the p (ii) that is situated at another territory of	e wild in an area: km radius there has been no case/outbre s for which the animals are susceptible: st, Newcastle disease or highly pathogenic of the preceding 30 days nor of classical of eriod of the preceding 40 days; and a distance that exceeds 20 km from the bor of a third country or part thereof, which is n e exportation of such material to the Europ	foot-and-mouth avian influenza or African swine ders separating ot authorised at
		(b)		ransported within a period of 12 hours for on nmediately afterwards to a game established t:]]	
(²)[II.1.3.	been obta case/outbr of the prec of raw mate	ined in eak of d eding 3 erial for	erials other than materials an establishment around iseases referred to in point 0 days or, in the event of exportation to the Europea	derived from fish or invertebrates caught in I which, within a radius of 10 km, there II.1.2 for which the animals are susceptible a case/outbreak of one of those diseases, In Union was authorised only after the rem- establishment under the control of an offici	e has been no during a period the preparation oval of all meat,
II.1.4.	have been	obtaine	ed and prepared without c	ontact with other material which does not on handled so as to avoid contamination w	comply with the
II.1.5.	have been cleaned an in containe 'ANIMAL E OUTSIDE	d disinf ers seal 3Y-PRO THE FI	ected before use and, in th ed under the responsibilit DUCTS ONLY FOR THE	 prevents any leakage or in packaging vectors case of consignments shipped other than of the competent authority, bearing the MANUFACTURE OF DERIVED PRODUC e and address of the establishment of destance 	via parcel post, label indicating TS FOR USES
II.1.6.	European		following animal by produ	ata	
11.1.0.	(²)either	[-	of animals killed which w	nimals slaughtered or, in the case of game, rere deemed fit for human consumption in a reversibly declared as animal by-products	accordance with
	(²)and/or	[-	slaughtered in a slaugh consumption following a of animals from game legislation: (i) carcases or bodi human consumpl show any signs c (ii) heads of poultry; (iii) hides and skins,	wing parts originating either from anim renhouse and were considered fit for slaug in ante-mortem inspection or bodies and the killed for human consumption in accorda es and parts of animals which were rejec- tion in accordance with Union legislation, b f disease communicable to humans or anim including trimmings and splitting thereof, langes and the carpus and metacarpus bo s;	hter for human of following parts nce with Union sted as unfit for ut which did not mals; horns and feet,
	(²)and/or	[-	animal by-products from referred to in Article 1	n poultry and lagomorphs slaughtered c (3), point (d), of Regulation (EC) No 8 nd of the Council ^(2a) , which did not shor o humans or animals:1	53/2004 of the
	(²)and/or	[-	blood of animals which blood to humans or anir a slaughterhouse after	did not show any signs of disease commu nals, obtained from animals that have been having been considered fit for slaugh an ante-mortem inspection in accordar	n slaughtered in hter for human
	(²)and/or	[-	animal by-products aris	ing from the production of products inten degreased bone, greaves and centrifug sing:]	

II. Healt	th informati	on		II.a.	Certificate reference No	II.b.
	(²)and/or	[-	are no longer intended	for hur	odstuffs containing products of anin nan consumption for commercial re packaging defects or other defects	asons or due to
	(²)and/or	[-	risk to public or animal h petfood and feedingstuf products or derived p commercial reasons or	nealth a ffs of a roducts due to	rises;] nimal origin, or feedingstuffs contai s, which are no longer intended problems of manufacturing or packa	ning animal by- for feeding for
	(²)and/or	[-	blood, placenta, wool, fe	eathers	k to public or animal health arises;] , hair, horns, hoof cuts and raw milk y signs of any disease communical	0 0
	(²)and/or	[-	aquatic animals, and pa	arts of	such animals, except sea mammals	
	(²)and/or	[-	animal by-products from	n aquat	mmunicable to humans or animals; ic animals originating from establish	
	(²)and/or	[-	disease communicable (i) shells from shellf (ii) the following orig — hatchery by — eggs; — egg by-prod	originati througl ish with inating -products, in	ng from animals which did not sho n that material to humans or animals n soft tissue or flesh; from terrestrial animals: cts; ncluding egg shells;	
	(²)and/or	[-		n aqua	commercial reasons;] atic or terrestrial invertebrates, othe als:1	er than species
	(²)and/or	[-	animals and parts there except Category 1 mate	eof of t erial as 9/2009	ne zoological orders of Rodentia ar referred to in Article 8, point (a)(iii) and Category 2 material as referre	, (iv) and (v), of
	(²)and/or	[-	furs originating from dea	ad anir	nals that did not show clinical signs oduct to humans or animals;]	of any disease
ll.1.7.	Union legis	slation in	rozen at the plant of origi	in or h	ave been preserved in accordance spoil between the time of dispatch	
(²)(⁸)	territory or disease ar	part thei e regula	reof referred to in point II.1 rly carried out and officiall	.1, whe ly contr	orm animals that have been obtained are vaccination programmes against olled in domestic bovine animals.]] of animal by-products derived from o	foot-and-mouth
(²)[II.1.9.	the animal (²)either (²)or	[are deri [are deri			ovine, ovine or caprine animals.]] ne animals and does not contain an	d is not derived
		from: (²) eithei	born, continuously posing a negligit	reared	ne materials other than those derived and slaughtered in a country or reg vine spongiform encephalopathy sion Decision 2007/453/EC(⁹).]]	ion classified as
	(2))or	(ÈC) No 9 (b) mechanica or caprine continuous as posing 2007/453/ (c) animal by	99/200 ally sep e anim sly rea g a no 'EC, in 'EC, in	terial as defined in point 1 of Annex 1 of the European Parliament and of parated meat obtained from bones ials, except from those animals t red and slaughtered in a country or r egligible BSE risk in accordance which there has been no indigenous to redrived product obtained from in which have been killed, after stunning	the Council(10); of bovine, ovine hat were born, region classified with Decision s BSE case, bovine, ovine or
			of the cen instrumen injected ir born, con	itral ne t introc nto the tinuous as po	rvous tissue by means of an elonga duced into the cranial cavity, or by cranial cavity, except for those an sly reared and slaughtered in a co psing a negligible BSE risk in a	ted rod-shaped means of gas imals that were puntry or region

COUNTRY

II. Health inf	formation		II.a. Certificate reference No	II.b.
	(b) or (c) (b) or (c) (c) (c) (c) (c) (c) (c) (c) (c) (c)	tain milk or milk prod farmed animals, oth k or milk products o animals, other than e derived from ovine classical scr an awarene classical scr of an awarene classical scr of an awarene classical scr of an awarene classical scr of the confir o ovine and ca destroyed; the feeding greaves, as Organisation banned and least the pre ginate from holding spicion of TSE; ginate from holding spicion of TSE; ginate from holding spicion of TSE; ginate from holding spicion of TSE; ginate from holding spicion of the se of classical scrap either [all ovine ar destroyed o genotype, bi allele and ot or [all animals and destroy least two yes scrapie cas negative rei laboratory n Regulation (over the ag genotype: - animals and - animals	ctions apply to holdings of ovine or ca spicion of transmissible spongiform en- mation of classical scrapie; aprine animals affected with classical so to ovine and caprine animals of mea defined in the Terrestrial Animal Health n for Animal Health (OIE), of ruminal effectively enforced in the whole coun- ceeding seven years; gs where no official restrictions are s where no case of classical scrapie h e preceding seven years or, following the r slaughtered, except for breeding ran reeding ewes carrying at least one ARF her ovine animals carrying at least one in which classical scrapie was confirm ed, and the holding has been subjected ears since the date of confirmation of sults for the presence of TSE in ac nethods set out in Chapter C, point EC) No 999/2001, of all of the following e of 18 months, except ovine animal s which have been slaughtered for hu s which have died or been killed on the not killed in the framework of a co	re intended for feed cts: n kept continuously filled: tem is in place for prine animals in the cephalopathy (TSE) crapie are killed and t-and-bone meal on h Code of the World int origin has been try for a period of at imposed due to a as been diagnosed he confirmation of a as been diagnosed he confirmation of a twe been killed and ns of the ARR/ARR allele and no VRG e ARR allele;] ed have been killed of the last classica cluding testing with ccordance with the 3.2, of Annex X to g animals which are is of the ARR/ARR uman consumption: e holding but which
in only if it is certificate is for the establishm Box reference been issued b Box reference Box reference a transi wareho — product the com — Box reference (ship) is to be	a certificate for a cor or a commodity to be e 1.11: In the case of content only. e 1.11 and 1.12: Appro- by the competent auther e 1.12: Place of destin is for the manufacture it commodity. Product uses. is for trade samples of the trade samples of the trade samples of the trade samples of the trade s	nmodity to be transi imported into the EL onsignments for trac wal number: the reg nority. ation: this box is to b of derived products ts in transit may on or analyses: the plar re appropriate. mber (railway wagor of unloading and re	de samples or analyses: indicate the na istration number of the establishment pe filled in for the following products: for uses outside the feed chain: only if ly be stored in free zones, free waref at in the European Union indicated in as or container and lorries), flight numb loading in the European Union, the co	ay be filled in if the ame and address o or plant, which has f it is a certificate fo nouses and custom the authorisation o er (aircraft) or name

П.	Health information	II.a.	Certificate reference No	II.b.					
_	Box reference I.23: for bulk containers, the container nu Box reference I.25:	ımber a	and the seal number (if applicable) mus	t be included.					
	 technical use: any use other than feeding of farm manufacturing of petfood; for the purposes of the certificate, 'technical use' 			production or					
-	Box reference 1.26 and 1.27: except for trade samples, which are not sent in transit, fill in according to whether it is a transit or an import certificate.								
-	 Box reference I.28: products for the manufacture of derived products for uses outside the feed chain: Manufacturing plant: provide the veterinary control number of the approved establishment; products for the particular technological studies or analyses: the plant in the European Union indicated in the authorisation of the competent authority where appropriate; species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or Suidae, PESCA, Mollusca, Crustacea, invertebrates other than Mollusca and Crustacea. 								
Part									
(1a)	OJ L 300, 14.11.2009, p. 1.								
(1b)	OJ L 54, 26.2.2011, p. 1.								
(2)	Delete as appropriate.								
(2a) (3)	OJ L 139, 30.4.2004, p. 55. The name and ISO code number of the exporting cour	thu an I	aid down in:						
	 Part 1 of Annex XIII to Commission Implementing F Part 1 of Annex XIV to Implementing Regulation (E Annex I to Commission Regulation (EC) No 119/20 In addition the ISO code of territories and parts there (EU) 2021/404 and to Regulation (EC) No 119/2009 regulation 	Regulat U) 202 09 (OJ of refer eferred	ion (EU) 2021/404 (OJ L 114, 31.3.20 1/404, and L 39, 10.2.2009, p. 12). rred to in the Annexes to Implementir	g Regulation					
(4)	species concerned) must be included where applicable								
(4)	Only for countries from where the game meat intended authorised for importation into the European Union. OJ L 303, 18.11.2009, p. 1.	ed for h	numan consumption of the same anin	nal species is					
(6)	Supplementary guarantees to be provided where the r a South American or South African country or part ther domestic ruminants for human consumption is author masseter muscles of bovine animals, incised in accorr B.1, of Annex I to Regulation (EC) No 854/2004 of 30.4.2004, p. 206), are also permitted.	eof fror orised f dance v	n where only maturated and deboned or exportation to the European Union with the requirements of Section IV, C	fresh meat of n. The whole hapter I, Part					
(7)	Only for certain South American countries.								
(8)	Only for certain South American and South African cou	untries.							
(9)	OJ L 172, 30.6.2007, p. 84.								
(10)	OJ L 147, 31.5.2001, p. 1.								
_	The signature and the stamp must be in a different col Note for the person responsible for the consignment in purposes and must accompany the consignment until into the European Union.	n the E	uropean Union: this certificate is only						
Offici	al veterinarian/Official inspector								
	Name (in capital letters):		Qualification and title:						
	Date: Stamp:		Signature:						
<u>اــــــــــــــــــــــــــــــــــــ</u>									

(c) Chapter 10(B) is replaced by the following:

'CHAPTER 10(B)

Health certificate

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

COUNTRY:

Veterinary certificate to EU

	I.1. Consignor	I.2. Certificate reference No I.2.a.						
	Name	I.3. Central competent authority						
	Tel.	I.4. Local competent authority						
Part I: Details of dispatched consignment	I.5. Consignee Name Address Postal code Tel.	I.6. Person responsible for the load in EU Name Address Postal code Tel.						
f dispatche	I.7. Country of ISO code I.8. Region of Code origin origin	I.9. Country of ISO destination code	I.10. Region of Code destination					
ails o	I.11. Place of origin	I.12. Place of destination						
Part I: Deta	Name Approval number Address Name Approval number Address	Custom warehouse Name Approval number Address						
	Name Approval number Address	Postal code						
	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 Ambient D Chilled D	I.22. Number of packag						
	I.23. Seal/Container No		I.24. Type of packaging					
	I.25. Commodities certified for: Technical use □							
	I.26. For transit through EU to third country	I.27. For import or admission into EU						
	Third country ISO code							
	I.28. Identification of the commodities							
	Approval number of establishments Species Manufacturing plant (scientific name)	Number of packages Net we	ight Batch number					

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

	II.	Health infor	matior	1	II.a. Certificate reference No II.b.				
	-	I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/200 of the European Parliament and of the Council (^{1a}), and in particular Articles 8, 9 and 10 thereof, an Commission Regulation (EU) No 142/2011(^{1b}), and in particular Chapter II of Annex XIV thereto, and certif that the rendered fats described above:							
	II.1.	consist of rendered fats not intended for human consumption that satisfy the health requirements below;							
u	II.2.	have been prepared exclusively with the following animal by-products:							
Part II: Certification	(²)[II.2.1.	in the case of materials destined for the production of renewable fuels referred to in Chapter IV, Section 2, point L, of Annex IV to Regulation (EU) No 142/2011, biodiesel or oleochemical products, animal by-products referred to in Articles 8, 9 and 10 of Regulation (EC) No 1069/2009;]							
Part II: ((²)[II.2.2.	in the case of materials destined for the production of renewable fuels referred to in Chapter IV, Section 2, point J, of Annex IV to Regulation (EU) No 142/2011, the materials have been prepared exclusively from animal by-products referred to in Articles 9 and 10 of Regulation (EC) No 1069/2009;]							
	(²)[II.2.3.		e case of materials destined for purposes other than cosmetics, pharmaceuticals or medical devices, th erials have been prepared exclusively from:						
	_	(²)either	[-		esidues of authorised substances or contaminants referred to in Article 15(3) of Council Directive				
		(²)and/or	[-	products of animal origin which ha the presence of foreign bodies in t	ve been declared unfit for human consumption due to hose products;]				
		(²)and/or	[-	Regulation (EC) No 1069/2009,	ther than those referred to in Articles 8 and 10 of that died other than being slaughtered or killed for mals killed for disease control purposes;]				
		(²)and/or	[-	animals killed, and which are fit	aughtered or, in the case of game, bodies or parts of for human consumption in accordance with Union r human consumption for commercial reasons;]				
		(²)and/or	[-	slaughtered in a slaughterhouse consumption following an ante-me	ts originating either from animals that have been and were considered fit for slaughter for human ortem inspection or bodies and the following parts of an consumption in accordance with Union legislation:				
					arts of animals which are rejected as unfit for human e with Union legislation, but which did not show any able to humans or animals;				
				(ii) heads of poultry;					
					g trimmings and splitting thereof, horns and feet, nd the carpus and metacarpus bones, tarsus and				
				(iv) pig bristles;(v) feathers;]					
		(²)and/or	[-	to humans or animals obtained slaughterhouse after having been	w any signs of disease communicable through blood I from animals that have been slaughtered in a considered fit for slaughter for human consumption on in accordance with Union legislation;]				
		(²)and/or	[-		the production of products intended for human d bone, greaves and centrifuge or separator sludge				
		(²)and/or	[-	no longer intended for human con	stuffs containing products of animal origin, which are sumption for commercial reasons or due to problems fects or other defects from which no risk to public or				
		(²)and/or	[-	products or derived products, whi	nimal origin, or feeding stuffs containing animal by- ch are no longer intended for feeding for commercial nufacturing or packaging defects or other defects from ealth arises:]				

COUNTRY

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

II.	Health in	formation		II.a.	Certificate reference No	II.b.	
	(²)and/or	[-			horns, hoof cuts and raw milk origin y disease communicable through t	-	
	(²)and/or	[-	aquatic animals, and parts of s any signs of diseases commu		nimals, except sea mammals, which to humans or animals;]	n did not show	
	(²)and/or	[-	animal by-products from aqu manufacturing products for hu		imals originating from plants or e onsumption;]	stablishments	
	(²)and/or	[-	communicable through that m (i) shells from shellfish wit (ii) the following originating — hatchery by-produ — eggs, — egg by-products, i (iii) day-old chicks killed for	aterial h soft t g from t icts, ncludir r comm	issue or flesh; ærrestrial animals: ng egg shells, ærcial reasons;]		
	(2)and/or	[-	animals;]		s other than species pathogenic t		
	(²)and/or	[-	except Category 1 material a	as refei 09 and	pological orders of Rodentia and rred to in Article 8, point (a)(iii), (i Category 2 material as referred t	v) and (v), of	
	(²)and/or	[-	hides and skins, hooves, fea	thers,	wool, horns, hair and fur originati of disease communicable through t	0	
	(²)and/or	[-	through that material to h slaughterhouse and which we	umans ere con	id not show any signs of disease of or animals, which were slaug sidered fit for slaughter for human in accordance with Union legislatio	htered in a consumption	
(²)[II.2.4.			erials destined for purposes of	other th	nan the production of organic fert		
	improvers (²)either	s, cosmetic [-		fined ir	n Article 3(1), point (g), of Regula	tion (EC) No	
	(²)and/or	[-		anima	ls containing specified risk material		
(²)and/or [-		[-	Article 3(1), point (g), of Regulation (EC) No 999/2001 at the time of disposal;] animal by-products which have been derived from animals which have been submitted to illegal treatment as defined in Article 1(2), point (d), of Council Directive 96/22/EC ^(2c) or Article 2, point (b), of Directive 96/23/EC;]				
	(²)and/or	[-	animal by-products containin contaminants listed in Group	ng resi B(3) of iid dow	dues of other substances and e Annex I to Directive 96/23/EC, if n by Union legislation or, in the abs	such residues	
11.3.	the rende	red fats:	, ,				
	 have been subjected to processing in a method) as set out in Chapter III of An pathogenic agents, 						
	[(²)(b) of gl	Category yceroltrihe	1 and 2 materials have been		ed before dispatch to the Europea s minimum concentration of at least		
	(c) in		of rendered fats of ruminant or	igin, in	soluble impurities in excess of 0,1	5 % in weight	
	(d) ha (e) be	ave been tr	ansported under conditions wh on the packaging or cont		vent their contamination, and indicating "NOT FOR HUMAN	OR ANIMAL	

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

	nealthin	ormation		II.a. Certificate referen	ce No	II.b.
(²)[.4 .	in the case improvers (²)either (²)or	e of materials dest the rendered fats [are derived from [are derived from (²) either [b cc ne	described above other ruminants than b bovine, ovine or caprir ovine, ovine and caprir ontinuously reared and egligible bovine spong ommission Decision 20 a) specified risk mat No 999/2001;) mechanically sep caprine animals, or reared and slaug negligible BSE rist there has been no) animal by-product caprine animals w	ers, cosmetics, pharmaceu povine, ovine or caprine an ne animals and does not co ne materials other than those slaughtered in a country of jiform encephalopathy (B: 007/453/EC(³).] erial as defined in point 1 co parated meat obtained from except from those animals phered in a country or re- sk in accordance with Deco o indigenous BSE case, at or derived product obta- thich have been killed, after	ticals, medical de imals.] Intain and is not of se derived from a region classified SE) risk in acco of Annex V to Reg m bones of bovi that were born, egion classified cision 2007/453/f ained from bovir	evices or so derived from animals born d as posing ordance wit gulation (EC ne, ovine of continuous as posing EC, in whic ne, ovine of eration of th
			introduced into th cranial cavity, ex reared and slaug	ssue by means of an elon ne cranial cavity, or by me accept for those animals t phtered in a country or re k in accordance with Decis	eans of gas inject hat were born, egion classified	cted into th continuous as posing
Notes Part I:						
in	only if it is a o		mmodity to be transite	nt in the European Union: t d through the European U	•	
– Bo by	x reference I. ⁻ the competer	11: Approval numb nt authority.	•	pean Union. Iber of the establishment or	plant, which has	been issue
– Bo by – Bo - cor -	x reference I. the competer x reference I. approval nur mpetent author place of de Products in t	11: Approval numb nt authority. 12: mber: the registrati prity; estination: this bo transit may only be	er: the registration num on number of the estal ox is to be filled in e stored in free zones, f		as been issued e for a transit m warehouses.	by th
- Bo by - Bo - cor - Bo (sh the	x reference I. the competer x reference I. approval nur mpetent author place of de Products in t x reference I. nip) is to be pre border inspe	11: Approval numb nt authority. 12: mber: the registratio prity; estination: this bo transit may only be 15: Registration nu ovided. In the case ection post of the p	er: the registration num on number of the estal ox is to be filled in estored in free zones, f mber (railway wagons e of unloading and relo point of entry into the Eu	ber of the establishment or olishment or plant, which h only if it is a certificat free warehouses and custo or container and lorries), fli ading in the European Unio	as been issued e for a transit im warehouses. ght number (airci on, the consignor	by th commodit raft) or nam r must infor
- Bo by - Bo - - - Bo (sh the - Bo - Bo - Bo	x reference I. the competer approval nur mpetent author place of de Products in t x reference I. hip) is to be pr border inspe x I.19: use the .03; 15.04; 15 x reference I. x reference I.	11: Approval numb nt authority. 12: mber: the registratio prity; estination: this be transit may only be 15: Registration nu ovided. In the case ection post of the p e appropriate Harr 5.05; 15.06; 15.16 23: for bulk contain 25: technical use:	er: the registration num on number of the estal ox is to be filled in e stored in free zones, f mber (railway wagons e of unloading and relo point of entry into the Eu nonized System (HS) of or 15.18. ers, the container num any use other than fe	ber of the establishment or olishment or plant, which h only if it is a certificat iree warehouses and custo or container and lorries), fli ading in the European Unio uropean Union. code under the following h ber and the seal number (if reding of farmed animals,	as been issued e for a transit om warehouses. ght number (airci on, the consignor eadings: 04.05; 1 applicable) must	by th commodit raft) or nam r must infor 15.01, 15.0 be include
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Rendered fats not intended for human consumption for certain purposes outside the feed chain

П.	Health information	II.a.	Certificate reference No	II.b.					
-	 The signature and the stamp must be in a different colour to that of the printing. Note for the person responsible for the consignment in the European Union: this certificate is only for veterinary purposes and must accompany the consignment until it reaches the border inspection post of the point of entry into the European Union. 								
Offic	ial veterinarian/Official inspector Name (in capital letters): Date: Stamp:		Qualification and title: Signature:						
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