Commission Regulation (EU) 2019/319 of 6 February 2019 amending Annex IX to Regulation (EC) No 999/2001 of the European Parliament and of the Council and Annex XV to Commission Regulation (EU) No 142/2011 as regards health certification at import into the Union concerning transmissible spongiform encephalopathies (Text with EEA relevance)

# COMMISSION REGULATION (EU) 2019/319

of 6 February 2019

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

(Text with EEA relevance)

### THE EUROPEAN COMMISSION,

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

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

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

# Whereas:

- (1) Regulation (EC) No 999/2001 lays down rules for the prevention, control and eradication of transmissible spongiform encephalopathies ('TSEs') in bovine, ovine and caprine animals. It applies to the production and placing on the market of live animals and products of animal origin and in certain specific cases to exports thereof. That Regulation also provides a legal basis for the classification, as laid down in Commission Decision 2007/453/EC<sup>(3)</sup>, of Member States and third countries or regions thereof according to their disease status for bovine spongiform encephalopathy (BSE) into those with a negligible BSE risk, a controlled BSE risk and an undetermined BSE risk.
- (2) Annex IX to Regulation (EC) No 999/2001 set outs the requirements for the importation into the Union of live animals, embryos, ova and products of animal origin. More particularly, Chapter B of that Annex sets out the requirements for imports of bovine animals, which takes into account the BSE status of the third countries or regions. In addition, Chapter D of that Annex lays down requirements for the provision of

- an attestation concerning the TSE related risk in the health certificate required for the importation into the Union of certain animal by-products and derived products, including, inter alia, processed animal protein.
- (3) Chapter B of Annex IX to Regulation (EC) No 999/2001, as amended by Commission Regulation (EU) 2016/1396<sup>(4)</sup>, requires that live bovine animals imported into the Union must not have been exposed to BSE cases or their cohort. Taking into account the fact that the main transmission route of BSE is through feed contaminated with the BSE prion, that requirement should be amended to provide that live bovine animals imported into the Union may not be BSE cases or their cohort. Chapter B of Annex IX to Regulation (EC) No 999/2001 should therefore be amended accordingly.
- (4) Regulation (EC) No 1069/2009 lays down public health and animal health rules for animal by-products and derived products in order to prevent and minimise risks to public and animal health arising from those products. Commission Regulation (EU) No 142/2011<sup>(5)</sup> lays down implementing measures for the public and animal health rules for animal by-products and derived products laid down in Regulation (EC) No 1069/2009, including certain requirements for the importation of animal by-products and derived products from third countries.
- (5) Annex I to Regulation (EU) No 142/2011 lists certain definitions to be used for the purposes of that Regulation. Article 31 of Regulation (EU) No 142/2011 provides that consignments of animal by-products and derived products for importation into or transit through the Union are to be accompanied by health certificates and declarations, in accordance with the models set out in Annex XV thereto.
- Organisation for Animal Health ('OIE Code')<sup>(6)</sup> recommends that meat-and-bone meal and greaves derived from ruminants, as defined in the OIE Code, and commodities containing such products, which originate from countries or zones with a negligible BSE risk status in which there has been an indigenous BSE case, may enter international trade only if the products were derived from cattle born after the date of the effective implementation, in the country, of the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants, as defined in the OIE Code. Point 2 of that Article recommends that meat-and-bone meal and greaves derived from ruminants, as defined in the OIE Code, and commodities containing such products, should not enter international trade if they originate from countries or zones with a controlled or undetermined BSE risk status.
- (7) The OIE Code defines meat-and-bone meal as the solid protein products obtained when animal tissues are rendered, including any intermediate protein product other than peptides of a molecular weight less than 10 000 daltons and amino-acids. Thus, meat-and-bone meal as defined in the OIE Code covers both the definition of meat-and-bone meal set out in point 27 of Annex I to Regulation (EU) No 142/2011 and the definition of processed animal protein set out in point 5 of that Annex.
- (8) In accordance with Article 41(2)(c) of Regulation (EC) No 1069/2009, imports into the Union of meat-and-bone meal, as defined in Union legislation, may only take place if implementing rules have been adopted setting out the conditions for such importation.

Since no such implementing rules have been adopted, the importation into the Union of meat-and-bone meal, derived from Category 1 or Category 2 material, is currently not allowed. However, imports into the Union of processed animal protein, as defined in Union legislation, may take place, subject to compliance with the TSE related import conditions laid down in Section B of Chapter D of Annex IX to Regulation (EC) No 999/2001, as well as with the conditions on the import of processed animal protein laid down in Regulation (EU) No 142/2011.

- (9) In order to align the TSE conditions for imports into the Union, laid down in Regulation (EC) No 999/2001, with the recommendations included in the BSE Chapter of the OIE Code, it is appropriate to amend Section B of Chapter D of Annex IX to Regulation (EC) No 999/2001 so that the requirement laid down in that Section take account of the recommendations of Article 11.4.13 of the OIE Code. However, since the use of processed animal protein derived from ruminants in the manufacturing of petfood is authorised in the Union, in order not to apply a discriminatory treatment towards imports compared to European Union production, the recommendations of Article 11.4.13 of the OIE Code should not be followed for the importation of petfood containing processed animal protein derived from ruminants, provided that such petfood is processed and labelled in accordance with Union legislation.
- (10) Section B of Chapter D of Annex IX to Regulation (EC) No 999/2001 should therefore be amended accordingly.
- (11)Products of animal origin may be required to be declared animal by-products by Union law, or by the decision of the responsible operator. When an operator decides that products of animal origin are to be declared as animal by-products, that decision is irreversible. Such animal by-products are excluded from use for human consumption. Certain animal by-products have the same Combined Nomenclature (CN) customs codes as animal products intended for human consumption which are laid down in Annex I to Council Regulation (EEC) No 2658/87<sup>(7)</sup>. For the classification in the CN customs codes the customs authorities in Member States need to be able to clearly differentiate between products which are fit for human consumption and those which are unfit for human consumption. In order to avoid any confusion for the purpose of that classification, the health guarantees referred to in the import certificates of unprocessed animal by-products should clarify that, although the animal by-products originate from animal products which were fit for human consumption at a former stage, they are now classified and treated as animal by-products which are permanently excluded from the food chain. The model health certificates set out in Chapters 3(D), 3(F) and 8 of Annex XV to Regulation (EU) No 142/2011 should therefore be amended accordingly.
- In addition, the TSE attestation in the model certificates for imports of and transit through the Union of certain animal by-products set out in Chapters 1, 1a, 2(A), 2(B), 3(A), 3(B), 3(C), 3(D), 3(E), 3(F), 4(B), 4(C), 4(D), 6(B), 8, 10(A), 10(B), 11, 12 and 18 of Annex XV to Regulation (EU) No 142/2011 should be amended to take account of the requirements of Chapter D of Annex IX to Regulation (EC) No 999/2001, as amended by Commission Regulation (EU) No 630/2013<sup>(8)</sup>, by Regulation (EU) 2016/1396 and by this Regulation.

- (13) The import conditions for processed animal protein referred to in the model health certificate set out in Chapter 1 of Annex XV to Regulation (EU) No 142/2011 require the absence of blood from ruminants in processed animal proteins imported from third countries. However, the new TSE attestation set out in point II.7. of that model health certificate, as amended by this Regulation, provides for adequate guarantees to mitigate the TSE risk in such products. Therefore, the wording 'other than ruminants' should be deleted in all the model health certificates set out in Annex XV to Regulation (EU) No 142/2011 that are to be amended by this Regulation.
- (14) Chapters 1, 1a, 2(A), 2(B), 3(A), 3(B), 3(C), 3(D), 3(E), 3(F), 4(B), 4(C), 4(D), 6(B), 8, 10(A), 10(B), 11, 12 and 18 of Annex XV to Regulation (EU) No 142/2011 should therefore be amended accordingly.
- (15) In addition, consignments of intermediate products intended for the manufacturing of cosmetic and pharmaceutical products are required to be accompanied by a declaration completed in accordance with the model set out in Chapter 20 of Annex XV to Regulation (EU) No 142/2011 when presented at a border inspection post ('BIP') for the purpose of veterinary checks. Intermediate products may consist of or may contain animal by-products. The existing model declaration indicates only a limited number of appropriate HS codes which are to be used by the operator to notify the product to the customs authorities in the Member States. It is not possible to set out an exhaustive list of HS codes in advance in the model declaration which would cover all combinations of animal by-products in the intermediate products. Therefore, it is appropriate to replace the existing HS codes in order that the person responsible for the consignment may declare intermediate products to the BIP by an appropriate HS code in accordance with Commission Decision 2007/275/EC<sup>(9)</sup>. Chapter 20 of Annex XV to Regulation (EU) No 142/2011 should be amended accordingly.
- (16) In order to avoid any disruption of trade, this Regulation should provide for a transitional period during which time the commodities concerned by the amendments made to Regulation (EU) No 142/2011 should continue to be accepted for importation into and transit through the Union, provided that those commodities comply with the requirements laid down in Regulation (EU) No 142/2011 before they were amended by this Regulation.
- (17) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on Plants, Animals, Food and Feed,

# HAS ADOPTED THIS REGULATION:

# Article 1

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

# Article 2

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

#### Article 3

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

#### Article 4

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

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

Done at Brussels, 6 February 2019.

For the Commission

The President

Jean-Claude JUNCKER

#### ANNEX I

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

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

### SECTION B

### Health certificate requirements

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

posing a negligible BSE risk in accordance with Decision 2007/453/EC;

or

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

or

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

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

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

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

or

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

#### ANNEX II

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

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

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

Documentation references

Document Generated: 2023-08-31

COL	INTRY	<b>′</b> :								Vete	rinary certific	cate to E	ΞU
	l.1.	Consignor					1.2.	Certificate refere	nce No	1.2	2.a.		
		Name Address					1.3.	Central compete	nt authority				
		Tel.					1.4.	Local competent	authority				
	1.5.	Consignee					1.6.	Person responsil	ble for the loa	ad in El	J		
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Part I : Details of dispatched consignment	l.11.	Place of origi	'n				I.12.	Place of destinat	ion				
Par		Name	Α	pprova	l number				Custo	om war	ehouse		
		Address						Name	Appro	oval nu	mber		
		Name	А	oprova	l number			Address					
		Address											
		Name	Α	pprova	l number			Postcode					
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	I.13.	Place of load	ing				1.14.	Date of departure	e				
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		Aeroplane			Railway wa	gon 🗖							
		Road vehicle		П			1.17.						

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

# COUNTRY

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

	II.	Health infor	mation		II.a.	Certificate reference No	II.b.
		the Europea	n Parliamen	it and of the	Counci	re that I have read and understood Rei I (¹a) and in particular Article 10 thereo ion 1 of Chapter II of Annex X, and Chap	f, and Commission Regulation
ıtion	II.1.			orotein or pro sumption that:	duct o	described above contains exclusively	processed animal protein not
Part II: Certification						establishment or plant approved and of Regulation (EC) No 1069/2009, and	supervised by the competent
art II:		(b) has be	een prepare	d exclusively	vith the	e following animal by-products:	
ď		(²) eitf	ner [-	animals kill	ed, an	s of animals slaughtered or, in the cas d which are fit for human consumption not intended for human consumption for	on in accordance with Union
		(²) and	d/or [-	slaughtered consumption	in a n follov	e following parts originating either for slaughterhouse and were considered wing an ante-mortem inspection or bod a killed for human consumption in accord	fit for slaughter for human lies and the following parts of
				consu	mption	bodies and parts of animals which ar in accordance with Union legislation, ase communicable to humans or animal	but which did not show any
				(ii) heads	of pou	ultry;	
					nalang	rins, including trimmings and splitting the es and the carpus and metacarpus b	
				(iv) pig br	stles;		
				(v) feather	rs;]		
		(²) and	d/or [-	to humans slaughterho	or ar use af	which did not show any signs of disease nimals, obtained from animals that l ter having been considered fit for slau nortem inspection in accordance with Ur	nave been slaughtered in a ghter for human consumption
		(²) and	d/or [-		n, inclu	ts arising from the production of puding degreased bone, greaves and cong;]	
		(²) and	d/or [-	longer inten	ded fo	origin, or foodstuffs containing products r human consumption for commercial r packaging defects or other defects fro ;]	easons or due to problems of
		(²) and	d/or [-		did n	ool, feathers, hair, horns, hoof cuts and ot show signs of any disease commun s;]	
		(²) and	d/or [-			and parts of such animals, except sea roses communicable to humans or animals	
		(²) and	d/or [-			s from aquatic animals originating fr ducts for human consumption;]	om establishments or plants

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

# COUNTRY

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

						containing such protein
II.	Health information	on		II.a.	Certificate reference No	II.b.
	(²) and/or	[-		-	erial originating from animals which did ough that material to humans or animals	, ,
			(i) shells	from s	shellfish with soft tissue or flesh;	
			(ii) the fo	lowing	originating from terrestrial animals:	
			- 1	atcher	ry by-products,	
			_ •	eggs,		
			_ •	gg by-	-products, including egg shells;	
			(iii) day-o	d chicl	ks killed for commercial reasons;]	
	(²) and/or	[-	aquatic and and other th		trial invertebrates other than species pa ects;]	thogenic to humans or animals
	(²) and/or	[-	Category 1	materi	thereof of the zoological orders of Rod al as referred to in Article 8(a)(iii), (iv) a ticle 9(a) to (g) of Regulation (EC) No 10	nd (v) and Category 2 material
	and					
	(c) has been s	ubjecte	d to the follow	ing pr	ocessing standard:	
	(²) either	at a	pressure (al	solute	erature of more than 133°C for at least e) of at least 3 bars produced by satura of more than 50 millimetres;]	
	(²) or			(indica	nmalian protein other than fishmeal, the ate the processing method) as set out 2/2011;]	
	(²) or	(ind			the processing method 1-2-3-4-5-6-7 g method) as set out in Chapter III of	
	(²) or	(ind No	icate the pro	cessing ere in	olood, the processing method 1-2-3-4-5-7 g method) as set out in Chapter III of case of method 7 a heat treatment of a e;]	Annex IV to Regulation (EU)
II.2.	the competent au following standard		examined a r	andom	sample immediately prior to dispatch a	and found it to comply with the
	Salmonella:		Abser	ce in 2	25 g: n = 5, c = 0, m = 0, M = 0	
	Enterobacteriacea	ie:	n = 5,	c = 2,	m = 10, M = 300 in 1g;	
II.3.	the product has u	ndergo	ne all precaut	ons to	avoid recontamination with pathogenic	agents after treatment;
II.4.	the end product:					
	(²) either [was	oacked	in new or ste	ilised l	bags,]	

(i)

# COUNTRY

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

II.	Health informa			II - O - III - I - I - I - I - I - I - I							
	nealth illionna	ition		II.a. Certificate reference No	II.b.						
	. ,	s transported in infected before u		containers or other means of transpo	ort that were thoroughly cleaned and						
	which bear labe	els indicating 'NO	OT FOR	HUMAN CONSUMPTION';							
II.5.	the end produc	t was stored in er	nclosed	storage;							
(²) [II.6.	the processed ruminant origin		or pro	duct described above contains or is	derived from animal-by products of						
	(²) either		e with	country or region, which is classified Decision 2007/453/EC, and in which							
	(²) or	with Decisi by-product ban on th ruminants,	ion 200 t or der he feed , as defi	country or region classified as posing 7/453/EC in which there has been an i ved product were derived from animal ing of ruminants with meat-and-bonned in the OIE Terrestrial Animal Healthegion, and]	ndigenous BSE case, and the animal ls born after the date from which the le meal and greaves derived from						
	(²) either [is derived from other ruminants than bovine, ovine or caprine animals.]										
	(²) or	[is derived	from bo	vine, ovine or caprine animals and doe	es not contain and is not derived from:						
		(²) either	continu	[bovine, ovine and caprine materials other than those derived from animals born continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]							
		(²) or	[ (a)	specified risk material as defined in p No 999/2001 of the European Parliame							
			(b)	mechanically separated meat obtain caprine animals, except from those a reared and slaughtered in a countrinegligible BSE risk in accorda 2007/453/EC (5), in which there has be	animals that were born, continuously ry or region classified as posing a ance with Commission Decision						
			(c)		ed, after stunning, by laceration of the an elongated rod-shaped instrument by means of gas injected into the sthat were born, continuously reared classified as posing a negligible BSE						
II.7.	the processed a	animal protein or	r produc	t described above:							
		es not contain m ned animals, othe		ilk products of ovine or caprine anima fur animals.]	I origin or is not intended for feed for						
	• •	(²) or [contains milk or milk products of ovine or caprine animal origin and is intended for feed for farmed animals, other than fur animals, and the milk or milk products:									
	(a)	are derived fro	om ovir	a and consine enimals which have be	en kept continuously since birth in a						

classical scrapie is compulsorily notifiable;

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

#### COUNTRY

Health information

II.

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

										-				
	(ii)	an awar	eness	, surve	illance	and mo	onitorin	ng sys	tem is	in plac	ce for o	classic	al scrapi	e;
	(iii)	official re									anima	ls in t	he case	of a
	(iv)	ovine an	d cap	rine an	imals a	ffected	with c	lassic	al scra	pie ar	e killed	d and o	destroye	d;
	(v)	the feed defined Health ( whole co	in the OIE),	Terres of run	strial A ninant	nimal F origin f	lealth nas be	Code en ba	of the anned	World and e	d Orga effectiv	anisatio	on for A	nimal
(b)	originate fro	m holding	s wh	ere no	official	restricti	ons ar	e imp	osed d	lue to	a susp	icion o	of TSE;	
(c)	originate fro at least the													
	(²) either	[all oving slaughte carrying at least o	red, at lea	except ast one	for bre	eding	rams (	of the	ARR/	/ARR	genoty	pe, bi	reeding	ewes
	( <sup>2</sup> ) or	[all anim and the of confir including laborator No 999// except of	holdir matic testi ry me 2001,	ng has on of t ng with thods s of all	been s he last negati set out of the	ubjecte classi ve resu in poin followir	ed for a ical so ilts for it 3.2 o	a perion crapie the proof Cha mals v	od of a case resenc apter C which a	t least to int e of T	t two y ensifie SE in a nnex X	rears sed TSI accord to Re	since the E monite lance wite gulation	date oring, th the (EC)
		— anim	nals w	hich h	ave bee	n slauç	ghtered	d for h	numan	consu	mption	n; and		
		— anim		hich haw							ut whi	ch wer	re not kil	led in
II.8. the processed a ruminant origin a											nimal-l	by pro	ducts of	non-
(²) either [not	intended for t	he produ	ction	of feed	for farr	ned ani	mals,	other	than fu	ır anim	nals.]			
Con resu	nded for the signor has ur lts of the ana ulation (EC) N	ndertaken ilyses car	to er	nsure thout in a	nat the	Border	Inspe	ection	Post o	of entry	y will b	e prov	ided wit	the

II.a. Certificate reference No

### Notes

# Part I:

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

**Health information** 

Document Generated: 2023-08-31

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

# COUNTRY

II.

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

_	Boxı	reference I.19: use the appropriate HS code: 05.05; 05.06; 05.07; 05.11; 23.01 or 23.09.								
_		reference I.25: technical use: any use other than feeding of farmed animals, other than fur animals, and the uction or manufacturing of pet food.								
_	Box	reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.								
_	Suida	reference I.28: Species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or ae, Pesca, Mollusca, Crustacea, invertebrates other than Mollusca and Crustacea. In the case of farmed fish, specify cientific name of the fish.								
Part	II:									
( <sup>1a</sup> )	OJ L	300, 14.11.2009, p. 1.								
(1b)	OJ L	54, 26.2.2011, p. 1.								
(²)	Delet	te as appropriate.								
(3)	Whe	re:								
	n = number of samples to be tested;									
	m =	threshold value for the number of bacteria; the result is considered satisfactory if the number of bacteria in all samples does not exceed m;								
	M =	maximum value for the number of bacteria; the result is considered unsatisfactory if the number of bacteria in one or more samples is M or more; and								
	c =	number of samples the bacterial count of which may be between m and M, the sample still being considered acceptable if the bacterial count of the other samples is m or less.								
(4)	OJ L	147, 31.5.2001, p. 1.								
( <sup>5</sup> )	OJ L	172, 30.6.2007, p. 84.								
( <sup>6</sup> )	desc than (EC) resul	ribed in this health certificate is intended to be used for the production of feed for non-ruminant farmed animals, other fur animals, the consignment must be analysed, in accordance with the methods set out in Annex VI to Regulation No 152/2009, in order to verify the absence of unauthorised constituents of animal origin. The information on the tof such analysis must be attached to this health certificate when presenting the consignment at an EU border								
( <sup>7</sup> )	OJ L	54, 26.2.2009, p. 1.								
_	The	signature and the stamp must be in a different colour to that of the printing.								
_										
Offic	ial ve	terinarian/Official inspector								
	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.  OJ L 147, 31.5.2001, p. 1.  OJ L 172, 30.6.2007, p. 84.  The Person responsible for the load referred to in Box I.6 must ensure that, if the processed animal protein or product described in this health certificate is intended to be used for the production of feed for non-ruminant farmed animals, other than fur animals, the consignment must be analysed, in accordance with the methods set out in Annex VI to Regulation (EC) No 152/2009, in order to verify the absence of unauthorised constituents of animal origin. The information on the result of such analysis must be attached to this health certificate when presenting the consignment at an EU border inspection post.  OJ L 54, 26.2.2009, p. 1.  The signature and the stamp must be in a different colour to that of the printing.  Note for the person responsible for the consignment in the European Union: This certificate is only for veterinary purposes and must accompany the consignment until it reaches the border inspection post.  Qualification and title:									
	Date	: Signature:								
	Stam	pp:								

Certificate reference No

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

# COUNTRY:

# Veterinary certificate to EU

	l.1.	Consignor					1.2.	Certificate refere	ence No	I.2.a.			
		Name					1.3.	Central compete	ent authority				
		Address					1.4.	Local competent	t authority				
		Tel.											
	1.5.	Consignee					1.6.	Person responsi	ble for the load	in EU			
ent		Name						Name					
gnm		Address						Address					
onsi								Butterd					
þ		Postcode						Postcode					
che		Tel.						Tel.					
spat	1.7.	Country of origin	ISO code	1.8.	Region of origin	Code	1.9.	Country of destination	ISO code	I.10. Region of destination	Code		
ğ		or origin	1	I	origin	I		destillation		destillation			
ails	111	Place of or	igin				112	Place of destina	tion				
Det		r lace of of	igii i				112.						
Part I : Details of dispatched consignment		Name Approval number						Custor	n warehouse				
ď		Address						Name	Approv	/al number			
		Name						Address					
		Address											
		Name		Appro	val number			Postcode					
		Address											
	I.13.	Place of loa	ading				1.14.	Date of departur	re e				
	l.15.	Means of tr	ransport				1.16.	Entry BIP in EU					
								,					
		Aeroplane	☐ Ship		Railway wa	agon 🗆							
		Road vehic	cle 🔲 Othe	r 🗆			1.17.						
		Identification	on										
		Documentation references											
	I.18.	Description of commodity							I.19. Commo	odity code (HS code)			
		,											
										I.20. Quantity			
	I.21.	Temperatu	re of product							I.22. Number of pa	ckages		
		Ambient 🗆	l		Chilled	]		Frozen 🛭	]				
	1.23.	Seal/Conta	oient ☐ Chilled ☐ I/Container No						I.24. Type of packaging				

Commission Regula	ation (EU) 2019/31	9 of 6 February 2	2019 amending	Annex IX to I	Regulation (EC)
ANNEYI					

17

1.25.	Commodities certifie	ed for:				
	Animal feedingstuff		Technical use		Manufacture of petfo	ood 🗖
I.26.	For transit through B	EU to third country		I.27. For import o	r admission into EU	
	Third country	ISO code	е			
1.28.	Identification of the	commodities	Approval number	of establishments		
Sp	ecies (Scientific name)	Nature of commod	dity Manufactu	ıring plant	Net weight	Batch number

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

# COUNTRY

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

		and products other than pe	etfood 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 R the European Parliament and of the Council ( $^{1a}$ ) and in particular Article 10 there (EU) No 142/2011 ( $^{1b}$ ), and in particular Section 1 of Chapter II of Annex X, and Ch certify that:	of, and Commission Regulation
ation	II.1.	the processed animal protein derived from farmed insects or product describ processed animal protein not intended for human consumption that:	ed above contains exclusively
Part II: Certification		<ul> <li>has been prepared and stored in an establishment or plant approved an authority in accordance with Article 24 of Regulation (EC) No 1069/2009, and</li> </ul>	
Part II:		(b) has been prepared exclusively from farmed insects of the following species:	
		(²) either [- Black Soldier Fly (Hermetia illucens);]	
		(²) and/or [- Common Housefly (Musca domestica);]	
		(²) and/or [- Yellow Mealworm (Tenebrio molitor);]	
		(²) and/or [- Lesser Mealworm (Alphitobius diaperinus);]	
		(²) and/or [- House cricket (Acheta domesticus);]	
		(²) and/or [- Banded cricket (Gryllodes sigillatus);]	
		(²) and/or [- Field Cricket (Gryllus assimilis).]	
		and	
		(c) has been processed by method [1]-[2]-[3]-[4]-[5]-[7] ( <sup>2</sup> ) as set out in Chapt (EU) No 142/2011;	er III of Annex IV to Regulation
		and	
		(d) the substrate for the feeding of farmed insects may only contain product following products of animal origin of Category 3 material:	cts of non-animal origin or the
		— fishmeal;	
		<ul> <li>blood products from non-ruminants;</li> </ul>	
		<ul> <li>di and tricalcium phosphate of animal origin;</li> </ul>	
		<ul> <li>hydrolysed proteins from non-ruminants;</li> </ul>	
		<ul> <li>hydrolysed proteins from hides and skins of ruminants;</li> </ul>	
		<ul> <li>gelatine and collagen from non-ruminants;</li> </ul>	
		eggs and egg products;	
		<ul> <li>milk, milk based-products, milk-derived products, and colostrum;</li> </ul>	
		— honey;	
		<ul><li>rendered fats;</li></ul>	

# COUNTRY

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

												ontaining such	
II.	Health in	formation	1		П	.a. Ce	ertificate	e refere	nce No			II.b.	
	and												
	ma	aterials of		in than								in contact with an did not contain n	
II.2.		etent auth standards		ned a ra	andom s	ample	immedia	ately pr	ior to d	ispatch a	ind fou	and it to comply v	with the
	Salmonel	la:		Absenc	ce in 25	g: n = {	5, c = 0,	, m = 0,	M = 0				
	Enteroba	cteriaceae	:	n = 5, c	5, c = 2, m = 10, M = 300 in 1g;								
II.3.	the produ	ict has und	lergone all p	orecautio	ons to av	void rec	ontami	nation v	vith path	nogenic a	agents	after treatment;	
II.4.	the end p	roduct:											
	(²) either	[was pa	cked in new	or steril	ilised ba	gs,]							
	(²) or		ansported in ted before u		n contair	ners or	other r	means o	of trans	port that	were	thoroughly clean	ed and
	which bear labels indicating 'NOT FOR HUMAN CONSUMPTION/ PROCESSED INSECT PROTEIN – SHALL NOT BE USED IN FEED FOR FARMED ANIMALS EXCEPT AQUACULTURE AND FUR ANIMALS';												
II.5.	the end p	roduct was	s stored in e	enclosed	d storage	<b>;</b> ;							
(²) [II.6.		essed anir origin and:		or prod	duct des	scribed	above	contai	ns or i	s derived	d from	animal-by prod	ucts of
	(2)	either		e with [								a negligible BSE een no indigenou	
	(2)	or	with Decis by-product ban on the ruminants,	sion 2007 t or deri he feed , as defir	from a country or region classified as posing a negligible BSE risk in accordar on 2007/453/EC in which there has been an indigenous BSE case, and the animor derived product were derived from animals born after the date from which e feeding of ruminants with meat-and-bone meal and greaves derived from the one of the o						animal nich the d from		
	(2)	either	[is derived	from oth	her rumi	inants t	han bov	vine, ov	ine or c	aprine an	nimals.	11	
	(2)	or	[is derived	from bo	ovine, ov	ine or o	caprine	animals	s and do	oes not c	ontain	and is not derive	d from:
	(²) either [bovine, ovine and caprine materials other than those derived from animals be continuously reared and slaughtered in a country or region classified as positine negligible BSE risk in accordance with Decision 2007/453/EC.]]												
			(²) or	[(a)								ex V to Regulation Council (4);	on (EC)
				(b)	caprine reared negligib	anima and sole B	ils, exc laughte SE ris	ept fron red in sk in	n those a cour accor	animals ntry or r dance	that egion with	es of bovine, o were born, contin classified as po Commission D nous BSE case,	nuously

II.a. Certificate reference No

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

# COUNTRY

**Health information** 

II.

II.8.

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

•••	110414111111	••••••••••••••••••••••••••••••••••••••				This. Continuate reference rise			
				ca ce in cr ar	aprine entral troduc anial nd sla	al by-product or derived product obtained from bovine, ovine or ne animals which have been killed, after stunning, by laceration of the al nervous tissue by means of an elongated rod-shaped instrument luced into the cranial cavity, or by means of gas injected into the al cavity, except for those animals that were born, continuously reared shaughtered in a country or region classified as posing a negligible BSE in accordance with Decision 2007/453/EC.]]]			
II.7.	the processed animal protein or product described above:								
	(²) either		does not contain milk or milk products of ovine or caprine animal origin or is not intended for feed for armed animals, other than fur animals.]						
	(²) or					of ovine or caprine animal origin and is intended for feed for farmed and the milk or milk products:			
						caprine animals which have been kept continuously since birth in a conditions are fulfilled:			
			(i)	classical sci	rapie i	e is compulsorily notifiable;			
			(ii)	an awarene	SS, SL	surveillance and monitoring system is in place for classical scrapie;			
			(iii)			ons apply to holdings of ovine or caprine animals in the case of a E or the confirmation of classical scrapie;			
			(iv)	ovine and c	aprine	ne animals affected with classical scrapie are killed and destroyed;			
			(v)	defined in the	he Te :), of	ovine and caprine animals of meat-and-bone meal or greaves, as Terrestrial Animal Health Code of the World Organisation for Animal of ruminant origin has been banned and effectively enforced in the or a period of at least the preceding seven years;			
		(b)	originate fro	m holdings v	vhere	re no official restrictions are imposed due to a suspicion of TSE;			
						re no case of classical scrapie has been diagnosed during a period of a years or, following the confirmation of a case of classical scrapie:			
			(²) either	slaughtered	, exc east	caprine animals on the holding have been killed and destroyed or xcept for breeding rams of the ARR/ARR genotype, breeding ewes st one ARR allele and no VRQ allele and other ovine animals carrying R allele;]			
			( <sup>2</sup> ) or	and the hole of confirma including tes laboratory in No 999/200	ding hation of sting was the s	which classical scrapie was confirmed have been killed and destroyed, a has been subjected for a period of at least two years since the date of the last classical scrapie case to intensified TSE monitoring, g with negative results for the presence of TSE in accordance with the nods set out in point 3.2 of Chapter C of Annex X to Regulation (EC) of all of the following animals which are over the age of 18 months, nimals of the ARR/ARR genotype:			
				— animals	whic	nich have been slaughtered for human consumption; and			
						nich have died or been killed on the holding but which were not killed in ork of a disease eradication campaign.]]			

[the processed animal protein or product described above contains or is derived from animal-by products of non-

ruminant origin and is, according to the statement of the Consignor referred to in Box I.1,

Health information

Document Generated: 2023-08-31

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

#### COUNTRY

II.

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

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

Certificate reference No

#### Notes

#### Part I:

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

#### Part II:

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

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

# COUNTRY

п

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

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

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

Ambient  $\square$ 

I.23. Seal/Container No

Document Generated: 2023-08-31

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

col	JNTRY	<b>'</b> :						Veterinary certificat	e to EU	
	l.1.	Consignor			1.2.	Certificate referer	ice No	I.2.a.		
		Name			1.3.	I.3. Central competent authority				
		Address			I.4. Local competent authority					
						•				
		Tel.								
	1.5.	Consignee		1.6.	Person responsib	le for the load	d in EU			
ent		Name				Name				
gnm		Address				Address				
onsi										
ğ		Postcode				Postcode				
tche		Tel.				Tel.				
ispa	1.7.	Country ISO code of origin	I.8. Region of origin	Code	1.9.	Country of destination	ISO code	I.10. Region of destination	Code	
οę			Ī							
Part I : Details of dispatched consignment	I.11. Place of origin				1.12.	Place of destination	on			
: De										
art		Name A	pproval number					Custom warehouse		
ď		Address				Name		Approval number		
		Name A	pproval number			Address				
		Address								
		Name A	pproval 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 wag	gon 🗖						
		Road vehicle Other			l.17.	Number(s) of CIT	ES			
		Identification								
		Documentation references								
	I.18.	Description of commodity					I.19. Comm	odity code (HS code)		
								I.20. Quantity		
	1.21.	Temperature of product						I.22. Number of packages		

Frozen

I.24. Type of packaging

Chilled

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

# COUNTRY

# Milk, milk-based products and milk-derived products not

				, ,	for human consumption	,					
	II.	Health inforr	nation	II.a. Certificate reference No	II.b.						
		the Europear (EU) No 142/ certify that the	he undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2008 European Parliament and of the Council (1a), and in particular Article 10 thereof, and Commission Regular U) No 142/2011 (1b), and in particular Section 4 of Chapter II of Annex X, and Chapter I of Annex XIV thereto, trify that the milk (2), the milk-based products (2) and milk-derived products (2) referred to in box 1.28 comply be following conditions:								
on	II.1.			red in(inse							
Part II: Certification		listed in Part mouth diseas	I of Annex II to ( se (FMD) and rir	(insert name of region) (3), which is Commission Regulation (EU) No 605/2010 (4), and which has been free from foot-and-inderpest for a period of 12 months immediately prior to export and has not practised st during that period;							
Part II	II.2.	any disease	were produced from raw milk derived from animals which at the time of milking did not show clinical disease transmissible through milk to humans or animals, and which had been kept for a period of days prior to production on holdings that were not subject to official restrictions due to foot-and-mouth of erpest;								
	II.3.	they are milk	or milk products	that:							
		(²) either	[have undergo	ne one of the treatments or combinations thereof de	escribed in point II.4;]						
		(²) or		ey to be fed to animals of species susceptible to ected from milk subjected to one of the treatments d							
			(²) either	[the whey was collected at least 16 hours after clot	ting and has a pH below 6;]						
			(²) (⁵) or [the whey has been produced at least 21 days before the shipping period no cases of FMD have been detected in the exporting country;]								
			(²) ( <sup>5</sup> ) or	[the whey has been produced on//, this day voyage duration, being at least 21 days before the border inspection post of the European Union;]]							
	II.4.	they have be	en subject to one	e of the following treatments:							
		(²) either		ture short time pasteurisation at 72°C for at leachieving a negative reaction to a phosphatase to							
			(²) either	[a subsequent second high temperature short time 15 seconds or an equivalent pasteurisation which to a phosphatase test in bovine milk;]							
			(²) or	[a subsequent drying process that in the case combined with additional heating to 72°C or higher							
			(²) or	[a subsequent process by which the pH is reduced level below 6;]	l and kept for at least one hour at a						
			(²) ( <sup>5</sup> ) or	[the condition that the milk/milk product has been the date of shipping and during that period no cast the exporting country;]							
			(²) ( <sup>5</sup> ) or	[the milk/milk product has been produced on/ consideration of the foreseen voyage duration, bein that the consignment is presented to a border Union;]	ng at least 21 days prior to the date						
			(²) or	[sterilisation at a level of at least F <sub>0</sub> 3;]]							

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

# COUNTRY

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

								for human consumption
II.	Health info	rmation	II	.a.	. (	Certificate reference No		II.b.
	(²) or	[ultra high tem	perature tre	eatr	tme	ent at 132°C for at least one sec	ond i	n combination with:
		(²) either				drying process that in the c additional heating to 72°C or hig		of milk intended for feeding is
		(²) or	[a subsequentle su			process by which the pH is redu	ced a	and kept for at least one hour at a
		(²) ( <sup>5</sup> ) or		fsh	hip	ping and during that period no c		produced at least 21 days prior to s of FMD has been detected in the
		(²) ( <sup>5</sup> ) or	considerat	ion	n o	f the foreseen voyage duration,	being	(insert the date), this date, in g at least 21 days prior to the date nspection post of the European
II.5.	every preciprocessing;		n to avoid	COI	onta	amination of the milk/milk-base	ed p	roduct/milk-derived product after
II.6.	the milk/mill	k-based product/m	milk-derived	pro	rod	uct was packed:		
	(²) either	[in new contain	[in new containers;]					
	(²) or	[in vehicles of competent aut		ntai	ine	ers disinfected prior to loading	g us	ing a product approved by the
	and		bear labels					k/milk-based product/milk-derived 3 material and not intended for
II.7.	the milk, mi	lk-based products	and milk-d	eriv	ive	d products described above:		
	(²) either	[does not cont farmed animal					nal o	rigin or is not intended for feed for
	(²) or					s of ovine or caprine animal orig s, and the milk or milk products:	gin ar	nd is intended for feed for farmed
		(a)				n ovine and caprine animals wh y where the following conditions		nave been kept continuously since fulfilled:
			(i)		cla	assical scrapie is compulsorily n	otifia	able;
			(ii)			n awareness, surveillance and assical scrapie;	d mo	onitoring system is in place for
			(iii)			ficial restrictions apply to holdinase of a suspicion of TSE or the		of ovine or caprine animals in the irmation of classical scrapie;
			(iv)			vine and caprine animals affecte estroyed;	ed wi	th classical scrapie are killed and
			(v)		gr O ba	reaves, as defined in the Terres rganisation for Animal Health	strial (OIE	imals of meat-and-bone meal or Animal Health Code of the World E), of ruminant origin has been e whole country for a period of at
		(b)	originate for	rom	m h	noldings where no official restric	ctions	s are imposed due to a suspicion

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

#### COUNTRY

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

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

#### Notes

#### Part I:

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

#### Part II:

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

ANNEXI

Document Generated: 2023-08-31

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

# COUNTRY

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

II.	Health information	II.a.	Certificate reference No		II.b.	
(²)	Delete as appropriate.					
(3)	For completion if the authorisation to import into or transit through the European Union is restricted to certain regions of the third country concerned.					
(4)	OJ L 175, 10.7.2010, p. 1.					
( <sup>5</sup> )	this condition applies only to third countries I	isted i	n column 'A' of Annex I to R	egulation (l	EU) No 605/2010.	
( <sup>6</sup> )	OJ L 147, 31.5.2001, p. 1.					
_	The signature and the stamp must be in a di	fferent	colour to that of the printing	g.		
_	Note for the person responsible for the cons and must accompany the consignment until				te is only for veterinary purposes	
Offic	ial veterinarian/Official inspector					
	Name (in capital letters):			Qualificatio	n and title:	
	Date:			Signature:		
	Stamp:					

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

I.21. Temperature of product

Chilled

Ambient  $\square$ 

I.23. Seal/Container No

I.20. Quantity

Frozen

I.22. Number of packages

I.24. Type of packaging

Document Generated: 2023-08-31

CO	UNTRY	Y:								Vete	rinary certifica	te to EU
	l.1.	Consignor					1.2.	Certificate refere	ence No	1.2	2.a.	
		Name					1.3.	I.3. Central competent authority				
		Address					1.4.	Local competent	authority			
		Tol										
	1.5.	Tel. Consignee					1.6.	Person responsi	blo for the lea	ad in El		
	1.5.	Name					1.6.	Name	ble for the loa	au III E	U	
nen	Name						Address					
ign	Address					Address						
suos		Postcode						Postcode				
eq		Tel.										
tch	ļ. <u> </u>							Tel.				
ispa	1.7.	Country of origin	ISO code	1.8.	Region of origin	Code	1.9.	Country of destination	ISO code	I.10.	Region of destination	Code
s of c					·							
tails	1.11.	I.11. Place of origin				1.12.	Place of destination	tion				
Part I : Details of dispatched consignment		· ·										
Ę		Name		Appro	val number					Cust	tom warehouse	
ď		Address						Name		App	roval number	
		Name		Appro	val number			Address				
		Address										
		Name		Appro	val number			Postcode				
		Address										
	1.13.	Place of lo	ading				1.14.	Date of departur	е			
	145	Maana of t					1.16	Entry BIP in EU				
	I.15. Means of transport  Aeroplane □ Ship □ Railway wagon □						1.16.	CHILLY BIP III EU				
						agon $\square$						
		Road vehic			. idiirray we	-9-11	1.17	Number(s) of Cl	TES			
	Identification						,		5			
			ation referen	ces								
	1,18		n of commod						I.19. Comr	nodity (	code (HS code)	

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

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

#### COUNTRY

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

	II.	Health informati	on	II.a. Certificate reference No	II.b.					
		the European Pa (EU) No 142/201	arliament and of the 1 (1b), and in particu	n, declare that I have read and understood Council (¹a), and in particular Article 10 the lar Section 4 of Chapter II of Annex X and Cl lostrum products (²) referred to in box I.28 co	reof, and Commission Regulation napter I of Annex XIV thereto, and					
	II.1. they were produced and derived in									
Part II: Certification		(insert name of region) (3), which is listed in Annex I to Commission Regulation (EU) No 605/2010 (4), and which has been free from foot-and-mouth disease (FMD) and rinderpest for a period of 12 months immediately prior to export and has not practised vaccination against rinderpest during that period;								
Part II: Ce	II.2. they were produced from colostrum derived from animals which at the time of milking did not show clinical signs of any disease transmissible through colostrum to humans or animals, and which had been kept for a period of at least 30 days prior to the date of production on holdings that were not subject to official restrictions due to foot-and-mout disease or rinderpest;									
	II.3.	pasteurisation at	ney are colostrum or colostrum products of bovine animals that have been subject to high temperature short time asteurisation at 72°C for at least 15 seconds, or an equivalent pasteurisation achieving a negative reaction to a nosphatase test in bovine colostrum, in combination with:							
		(²) ( <sup>5</sup> ) either	(2) (5) either [the condition that the colostrum or colostrum products have been produced during a period at least 21 days before the date of shipping and during this period no cases of FMD have been detected in the exporting country,]							
		(²) ( <sup>5</sup> ) or	[the condition that the colostrum or colostrum products have been produced on// (inser the date), this date, in consideration of the foreseen voyage duration, being at least 21 days before the consignment is presented to a border inspection post of the European Union,]							
		and	have been obtained from animals subject to regular veterinary inspections to ensure that they come from holdings on which all bovine herds are:							
			(²) ( <sup>5</sup> ) either	[recognised as officially tuberculosis and bru	cellosis free (6),]					
			(²) ( <sup>5</sup> ) or	[not restricted under the national legislation of eradication of tuberculosis and brucellosis,]	of the third country of origin for the					
		and	(²) ( <sup>5</sup> ) either	[recognised as official enzootic-bovine-leuko	sis-free (6),]					
			(²) ( <sup>5</sup> ) or	[included in an official system for the control there has been no evidence as result of clir disease in the herd during the period of the $\rho$	nical and laboratory testing of this					
	II.4.	every precaution	has been taken to a	void contamination of the colostrum/colostrur	m product after processing;					
	II.5.	the colostrum or	colostrum product w	as packed:						
		(²) either	[in new containers	.]						
		(²) or	[in vehicles or bu	ilk containers disinfected prior to loading uty,]	sing a product approved by the					
		and the containers are marked so as to indicate the nature of the colostrum/colostrum product and bear labels indicating that the product is Category 3 material and not intended for human consumption;								
	II.6.	the colostrum or	colostrum product do	pes not contain milk or milk products of ovine	or caprine animal origin.					
	Notes	lotes								
	Part I:									

#### Part I:

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

ANNEX I
Document Generated: 2023-08-31

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

# COUNTRY

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

II.	Health Information	II.a.	Certificate reference No		II.D.				
_	Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity.								
_	Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to be provided. In the case of unloading and reloading in the European Union, the consignor must inform the border inspection post of the European Union.								
-	Box reference I.19: use the appropriate Harmonised System (HS) code of the World Customs Organisation: 04.04.90; 23.09.10, 23.09.90, 35.01, 35.02 or 35.04.								
_	Box reference I.23: for bulk containers, the container number and the seal number (if applicable) must be included.								
_	Box reference I.25: technical use: any use other than feeding of farmed animals, other than fur animals, and the production or manufacturing of pet food								
_	Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.								
_	Box reference I.28: 'Manufacturing plant': provide the registration number of the treatment or processing establishment.								
Par	t II:								
( <sup>1a</sup> )	OJ L 300, 14.11.2009, p. 1.								
(1b)	OJ L 54, 26.2.2011, p. 1.								
(2)	Delete as appropriate.								
(3)	For completion if the authorisation for introduction into the European Union is restricted to certain regions of the third country concerned.								
(4)	OJ L 175, 10.7.2010, p. 1.								
(5)	This condition applies only to third countries authorised in column 'A' of Annex I to Commission Regulation (EU) No 605/2010 (OJ L 175, 10.7.2010, p. 1).								
(6)	Officially tuberculosis-free and brucellosis-free herd as laid down in Annex A to Council Directive 64/432/EEC (OJ 121, 29.7.1964, p. 1977/64) and officially enzootic-bovine-leukosis-free herd as laid down in Chapter I of Annex D to that Directive.								
_	The signature and the seal must be in a different colour from that of the printing.								
_	Note for the importer: this certificate is only for veterinary purposes and must accompany the consignment until it reaches the border inspection post of the European Union.								
Offic	cial veterinarian/Official inspector								
	Name (in capital letters):			Qualification	on and title:				
	Date:			Signature:					
	Stamp:								

CHAPTERealth certificateFor canned petfood intended for dispatch to or for transit 3(A) through (2) the European Union

I.21. Temperature of product

Chilled

Ambient  $\square$ 

I.23. Seal/Container No

23.09 I.20. Quantity

Frozen 🗖

I.22. Number of packages

I.24. Type of packaging

Document Generated: 2023-08-31

COL	JNTRY	Y:								Vete	erinary certifica	te to EU
	I.1. Consignor				1.2.	Certificate refere	ence No	1.2	2.a.			
		Name					1.3.	Central compete	nt authority			
		Address					1.4.	Local competent	authority			
		Tel.										
ment	1.5.	Consignee					1.6.	Person responsi	hle for the lo	ad in Fl	11	
	1.5.	Name				1.0.	Person responsible for the load in EU     Name					
		Address				Address						
sign		Address						Address				
cons		Postcode						Postcode				
atched	Tel.				Tel.							
	1.7.	Country	ISO code	1.8.	Region of	Code	1.9.	Country of	ISO	I 10	Region of	Code
disp	""	of origin	100 0000	1.0.	origin	Out	1.0.	destination	code	1.10.	destination	Code
Part I : Details of dispatched consignment												
	1.11.	Place of or	rigin				I.12.	Place of destina	tion	•		
ă												
art		Name		Appro	val number					Cust	tom warehouse	
-		Address						Name		App	roval number	
		Name		Appro	val number			Address				
		Address										
		Name Approval number					Postcode					
		Address										
	I.13.	Place of lo	ading				1.14.	Date of departur	е			
	115	Means of t	ransport				116	Entry BIP in EU				
	1.10.	mound of t	ranoport				1.10.	Linay bir iii Lo				
		Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐										
						1.17.						
		Identification	on									
		Documenta	ation reference	ces								
	I.18.	Description	n of commodi	ity					I.19. Comr	nodity o	code (HS code)	

.25. Con	Commodities certified for:								
Pett	Petfood 🗆		Technical use □						
.26. For	For transit through EU to third country	, 🗆	I.27. For import or admission into EU						
Thir	Third country ISO co	de							
.28. Ider	Identification of the commodities								
	Approval number of establishments								
(Sc	Species Manuf (Scientific name)	facturing plant	Net weight	Batch number					
	Species Manut								

	COUNT	RY				Canned Petfood				
	п.	Health infor	mati	on	II.a. Certificate reference No	II.b.				
		the Europea Regulation (	an Pa EU) I	arliament and of the	an, declare that I have read and understood Regine Council (1a), and in particular Articles 8 and and in particular Chapter II of Annex XIII and Chapter II of An	10 thereof, and Commission				
tion	II.1.	has been prepared and stored in an establishment or plant approved and supervised by the competent auth accordance with Article 24 of Regulation (EC) No 1069/2009;								
rtifica	II.2.	has been pro	has been prepared exclusively with the following animal by-products:							
Part II: Certification		(²) either	[-	killed, and which	rts of animals slaughtered or, in the case of gam are fit for human consumption in accordance with an consumption for commercial reasons;]					
		(²) and/or	and/or   [- carcases and the following parts originating either from animals that have been slaughter slaughterhouse and were considered fit for slaughter for human consumption following mortem inspection or bodies and the following parts of animals from game killed for consumption in accordance with Union legislation:							
					carcases or bodies and parts of animals which are consumption in accordance with Union legislation cigns of disease communicable to humans or animates.	, but which did not show any				
				(ii) h	neads of poultry;					
				i	nides and skins, including trimmings and splitti ncluding the phalanges and the carpus and me netatarsus bones;					
				(iv) p	oig bristles;					
				(v) f	eathers;]					
		(²) and/or	[-	Article 1(3)(d) of	n the farm as referred to in lean Parliament and of the e to humans or animals]					
		(²) and/or	[-	<ul> <li>blood of animals which did not show any signs of disease communicable throu humans or animals, obtained from animals that have been slaughtered in a slaughter having been considered fit for slaughter for human consumption following an inspection in accordance with Union legislation;]</li> </ul>						
		(²) and/or	[-	animal by-products arising from the production of products intended for human including degreased bone, greaves and centrifuge or separator sludge from milk pro						
		(²) and/or	[-	<ul> <li>products of animal origin, or foodstuffs containing products of animal origin, which a intended for human consumption for commercial reasons or due to problems of man packaging defects or other defects from which no risk to public or animal health arise</li> </ul>						
		(²) and/or	[-	[- petfood and feedingstuffs of animal origin, or feedingstuffs containing animal by-pi derived products, which are no longer intended for feeding for commercial reasons problems of manufacturing or packaging defects or other defects from which no risk to animal health arise;]						
		(²) and/or	[-	<ul> <li>blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk originating from liver that did not show signs of any disease communicable through that product to hanimals;</li> </ul>						
		(²) and/or	[-	<ul> <li>aquatic animals, and parts of such animals, except sea mammals, which did not show a of diseases communicable to humans or animals;]</li> </ul>						
		(²) and/or	[-	<ul> <li>animal by-products from aquatic animals originating from plants or establishments manufactur products for human consumption;</li> </ul>						

instrument introduced into the cranial cavity, or by means of gas injected into the cranial cavity, except for those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]]

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

COUNTRY Canned Petfood Health information II.a. Certificate reference No (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: 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; 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;] animals and parts thereof of the zoological orders of Rodentia and Lagomorpha, except (2) and/or Category 1 material as referred to in Article 8(a)(iii), (iv) and (v) of Regulation (EC) No 1069/2009 and Category 2 material as referred to in Article 9(a) to (g) of that Regulation;] material from animals which have been treated with certain substances which are prohibited by Council Directive 96/22/EC (2b), the import of the material being permitted in accordance with (2) and/or Article 35(a)(ii) of Regulation (EC) No 1069/2009;] 11.3 has been subjected to heat treatment to a minimum Fc value of 3 in hermetically sealed containers; 11.4. was analysed by a random sampling of at least five samples from each processed batch by laboratory diagnostic method to ensure adequate 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. (2) [II.6. the petfood described above (2) either [is derived from other ruminants than bovine, ovine or caprine animals.] (2) or [is derived from bovine, ovine or caprine animals and does not contain and is not derived from: (2) either Ibovine, ovine and caprine materials other than those derived from animals born. continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]] specified risk material as defined in point 1 of Annex V to Regulation (EC) (2) or [(a) No 999/2001 of the European Parliament and of the Council (3); (b) mechanically separated meat obtained from bones of bovine, ovine or caprine animals, except from those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Commission Decision 2007/453/EC (4), in which there has been no indigenous BSE case, animal by-product or derived product obtained from bovine, ovine or caprine animals which have been killed, after stunning, by laceration of the central nervous tissue by means of an elongated rod-shaped **Health information** 

II.b.

**Canned Petfood** 

Document Generated: 2023-08-31

COUNTRY

II.

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

Note	es
Part	l:
_	Box reference I.6: Person responsible for the consignment in the European Union: this box is required to be filled in only if it is a certificate for a commodity to be transited through the European Union; it may be filled in if the certificate is for a commodity to be imported into the European Union.
_	Box reference I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity. Products in transit may only be stored in free zones, free warehouses and custom warehouses.
_	Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship); information is to be provided in the event of unloading and reloading in the European Union.
_	Box reference I.23: for bulk containers, the container number and the seal number (if applicable) must be given.
_	Box reference I.25: technical use: any use other than feeding of farmed animals, other than fur animals, and the production or manufacturing of pet food
_	Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.
_	Box reference I.28: Species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or Suidae, Pesca, Mollusca, Crustacea, invertebrates other than Mollusca and Crustacea.
Part	II:
( <sup>1a</sup> )	OJ L 300, 14.11.2009, p. 1.
(1b)	OJ L 54, 26.2.2011, p. 1.
(2)	Delete as appropriate.
( <sup>2a</sup> )	OJ L 139, 30.4.2004, p. 55.
(2b)	OJ L 125, 23.5.1996, p. 3.
(3)	OJ L 147, 31.5.2001, p. 1.
(4)	OJ L 172, 30.6.2007, p. 84.
_	The signature and the stamp must be in a different colour to that of the printing.
_	Note for the person responsible for the consignment in the European Union: This certificate is only for veterinary purposes and must accompany the consignment until it reaches the border inspection post.
Offic	ial veterinarian/Official inspector
JK	Name (in capital letters):  Qualification and title:
	Date: Signature:
	Stamp:

II.a. Certificate reference No

(CHAPTER 3(B)

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

col	JNTRY	<b>/</b> :								Veterinary certifica	te to EU
	l.1.	Consignor					1.2.	Certificate refere	nce No	I.2.a.	
		Name					1.3.	Central compete	nt authority		
		Address					1.4.	Local competent	authority		
								·	,		
		Tel.									
	1.5.	Consignee					1.6.	Person responsil	ole for the loa	d in EU	
ent		Name						Name			
gnm		Address						Address			
onsi											
ğ		Postcode						Postcode			
tche		Tel.						Tel.			
ispa	1.7.	Country of origin	ISO code	1.8.	Region of origin	Code	1.9.	Country of destination	ISO code	I.10. Region of destination	Code
of d		<b>-</b>									
Part I : Details of dispatched consignment	1.11.	Place of or	igin				1.12.	Place of destinat	ion		
: De			-								
art		Name		Appro	val number					Custom warehouse	
ď		Address						Name		Approval number	
		Name		Appro	val number			Address			
		Address									
		Name		Appro	val number			Postcode			
		Address									
	I.13.	Place of loa	ading				I.14.	Date of departure	Э		
	I.15.	Means of tr	ransport				I.16.	Entry BIP in EU			
			_	_		_					
		Aeroplane			Railway wa	agon 🎞					
		Road vehic		er 📙			I.17.				
		Identification									
	140		ation reference						140 Comm	andity and (HC anda)	
	1.10.	Description	of commodi	ity					1.19. Comii	nodity code (HS code)	
								L		I.20. Quantity	
	1.21.	Temperatu	re of product	t						I.22. Number of pa	ckages

Frozen 🗖

I.24. Type of packaging

Chilled

Ambient  $\square$ 

I.23. Seal/Container No

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

1.25.	Commodities certified for:								
	Petfood		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  Approval number of establishments								
	Species (Scientific name)	Manufacturing plant	Net weight	Batch number					

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

	COUNTRY				Processed petfoo	d other than canned petfood				
	II.	Health info	ormati	on	II.a. Certificate reference No	II.b.				
		the Europe Regulation and certify	ean Pa (EU) that th	arliament and of th No 142/2011 ( <sup>1b</sup> ), a e petfood describe		10 thereof, and Commission napter II of Annex XIV thereto,				
ation	II.1.			ed and stored in a ulation (EC) No 106	plant approved and supervised by the competer 9/2009;	nt authority in accordance with				
ertific	II.2.	has been p	repare	ed exclusively with t	the following animal by-products:					
Part II: Certification		(²) either	[-	killed, and which	rts of animals slaughtered or, in the case of gar are fit for human consumption in accordance wit an consumption for commercial reasons;]					
		(²) and/or	[-	slaughterhouse a mortem inspection	following parts originating either from animals the and were considered fit for slaughter for human on on or bodies and the following parts of animals accordance with Union legislation:	consumption following an ante-				
				consump	or bodies and parts of animals which are tion in accordance with Union legislation, but wh communicable to humans or animals;					
				(ii) heads of	poultry;					
					skins, including trimmings and splitting thereof, horns and feet, including the and the carpus and metacarpus bones, tarsus and metatarsus bones;					
				(iv) pig bristle	es;					
				(v) feathers;]						
		(²) and/or	[-	Article 1(3)(d) of	cts from poultry and lagomorphs slaughtered of Regulation (EC) No 853/2004 of the Euro h did not show any signs of disease communicab	pean Parliament and of the				
		(²) and/or	[-	humans or anima having been cor	which did not show any signs of disease co als, obtained from animals that have been slaugh nsidered fit for slaughter for human consumpt ordance with Union legislation;]	tered in a slaughterhouse after				
		(²) and/or	[-		cts arising from the production of products intelested bone, greaves and centrifuge or separator slu					
		(²) and/or	[-	intended for huma	al origin, or foodstuffs containing products of animal consumption for commercial reasons or due to sor other defects from which no risk to public or a	problems of manufacturing or				
		(²) and/or	[-	derived products	dingstuffs of animal origin, or feedingstuffs con , which are no longer intended for feeding for ufacturing or packaging defects or other defects e;]	commercial reasons or due to				
		(²) and/or	[-		wool, feathers, hair, horns, hoof cuts and raw milk originating from live animals w signs of any disease communicable through that product to humans or					
		(²) and/or	[-		and parts of such animals, except sea mammals nunicable to humans or animals;]	, which did not show any signs				

COUNT						•	od other than canned petfood
II.	Health info	rmati	on		II.a.	Certificate reference No	II.b.
	(²) and/or	[-		by-products cts for human		aquatic animals originating from plants or umption;]	establishments manufacturing
	(²) and/or	[-				originating from animals which did not nat material to humans or animals:	show any signs of disease
			(i)	shells from	shellfi	ish with soft tissue or flesh;	
			(ii)	the following	g origi	inating from terrestrial animals:	
				— hatch	ery by	y-products,	
				— eggs,			
				— egg b	-proc	ducts, including egg shells,	
			(iii)	day-old chic	ks kil	lled for commercial reasons;]	
	(²) and/or	[-		by-products s or animals;		aquatic or terrestrial invertebrates other	er than species pathogenic to
	(²) and/or	[-	Catego	ory 1 material	as re	eof of the zoological orders of Roder eferred to in Article 8(a)(iii), (iv) and (v) of as referred to in Article 9(a) to (g) of that F	Regulation (EC) No 1069/2009
	(²) and/or	[-	Counc	il Directive 9	6/22/E	nich have been treated with certain subst EC ( <sup>2b</sup> ), the import of the material being tion (EC) No 1069/2009;]	
II.3.							
	(²) either	[wa	s subjec	ted to a heat	treatr	ment of at least 90 °C throughout its subst	tance;]
	(²) or	[wa	s produc	ed as regard	s ingr	redients of animal origin using exclusively	products which had been:
		(a)				products or derived products from meat o t 90 °C throughout its substance;	r meat products subjected to a
		(b)	in the	case of milk a	nd m	ilk based products,	
			(i)	Commission	n Reg	hird countries or parts of third countries li gulation (EU) No 605/2010 (³) submitted uce a negative phosphatase test;	
			(ii)	column C o	f Ann	ed to less than 6 from third countries or p lex I to Regulation (EU) No 605/2010, firs ent to produce a negative phosphatase tes	st submitted to a pasteurisation
			(iii)	Regulation	(EU)	hird countries or parts of third countries li No 605/2010, submitted to a sterilisati each treatment was sufficient to produce	ion process or a double heat
			(iv)	Regulation disease in	(EU) the	hird countries or parts of third countries li No 605/2010, where there has been a preceding 12 months or where vaccin n carried out in the preceding12 months, s	n outbreak of foot-and-mouth nation against foot-and-mouth
				either			
				— a steri	lisatio	on process whereby an Fc value equal or	greater than 3 is achieved
				or			
				paster	ırisati	eat treatment with a heating effect at lea ion process of at least 72 °C for at leas negative reaction to a phosphatase test, fo	t 15 seconds and sufficient to

COUNTRY

### Processed petfood other than canned petfood

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

#### either

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

or

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

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

II.	Health infe	ormatio	on		II.a.	Certifi	cate refer	ence No			II.b
		(1)	in the	case of dical	ium p	hospha	te produc	ed by a p	process th	at	
			(i)		with	dilute hy	/drochlori	c acid (a	t a minimu		and degreased with hot wa centration of 4 % and a pH
			(ii)								nt of the obtained phospho hate at pH 4 to 7; and
			(iii)	finally, air of 325 °C and							inlet temperature of 65 °C
		(m)	in the	case of trical	ium p	hospha	te produc	ed by a	process th	ıat ensı	ıres
			(i)	that all Cat hot water (b					crushed	and de	greased in counter-flow w
			(ii)	continuous	cooki	ng with	steam at	145 °C d	uring 30 m	ninutes	at 4 bar;
			(iii)	separation centrifugation			ein broth	from th	e hydrox	yapatite	e (tricalcium phosphate)
			(iv)	granulation	of the	tricalcii	um phosp	hate afte	r drying in	n a fluid	bed with air at 200 °C ;
		(n)		ensure that							ent method and paramete cal standards referred to
	(²) or			ct to a treat authority;]	nent	such a	s drying	or ferme	entation, v	which I	has been authorised by t
	(²) or	anin	nals, ha	s been subje	ct to	a treatm	nent whicl	n has be	en authori	ised by	es pathogenic to humans the competent authority a d animal health;]
II.4.				om sampling plant and co						cessed	I batch taken during or af
	Salmonella	a:		absence in	25g: ı	n = 5, c	= 0, m = 0	O, M = O,			
	Enterobact	teriacea	ae:	n = 5, c = 2	m =	10, M =	300 in 1	gramme;			
II.5.	has underg	gone all	l precau	tions to avoid	cont	aminatio	on with pa	thogenic	agents af	fter trea	atment;
II.6.		that the	e conte								ackages on which it is clea icating "NOT FOR HUM/
(²) [II.7.	the petfood	d descri	ibed abo	ove							
	(²) either	[is d	erived fi	om other rur	ninan	ts than b	oovine, ov	vine or ca	prine anin	nals.]	
	(²) or	[is d	erived fi	rom bovine, o	vine	or caprir	ne animal	s and do	es not cor	ntain ar	nd is not derived from:
				The sine of			rina mat	oriala at	ner than	thoso	derived from onimals bo
		(²) e	ither		y rea	ared and	d slaught	ered in	a country	or re	derived from animals bo gion classified as posing C.]]

COUNTRY

Processed petfood other than canned petfood

COUN	IKI		Processed petrood other than canned petrood						
II.	Health information		II.a.	Certificate reference No		II.b.			
	(1	anim slaug acco	als, hter dan	cally separated meat obtained from those animals that red in a country or region classifice with Commission Decision 200 nous BSE case,	were bed as p	oorn, continuously reared and oosing a negligible BSE risk in			
	(1	anim nervo the o those or re	als in the state of the state o	y-product or derived product ob which have been killed, after s tissue by means of an elongated al cavity, or by means of gas inje imals that were born, continuousl classified as posing a negligible k/EC.]]]	tunning rod-sha ected int y reared	, by laceration of the central ped instrument introduced into to the cranial cavity, except for d and slaughtered in a country			
Notes									

#### Part I:

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

### Part II:

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

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

CO	UNTR	Υ	Processed petfood other than canned petfood								
II.		Health information	II.a.	Certificate reference No		II.b.					
(4)	Whe	re:									
	n =	number of samples to be tested;									
	m =	m = threshold value for the number of bacteria; the result is considered satisfactory if the number of bacteria in all samples does not exceed m;									
	M =	maximum value for the number of bac more samples is M or more; and	teria; th	ne result is considered uns	satisfactory if th	ne number of bacteria in one or					
	c =	number of samples the bacterial cou acceptable if the bacterial count of the			m and M, the	sample still being considered					
(5)	OJ L	147, 31.5.2001, p. 1.									
( <sup>6</sup> )	OJ L	172, 30.6.2007, p. 84.									
_	The	signature and the stamp must be in a di	fferent	colour to that of the printing	ng.						
_		for the person responsible for the cons must accompany the consignment until									
Offi	cial ve	terinarian/Official inspector									
	Nam	e (in capital letters):			Qualification a	and title:					
	Date: Signature:										
	Stan	np:									

CHAPTERealth certificateFor dogchews intended for dispatch to or for transit through 3(C) (2) the European Union

### COUNTRY:

000	MIKI	•					veterinary certificat	e to Eu
	l.1.	Consignor	1.	2.	Certificate referen	nce No	I.2.a.	
		Name	1.	.3.	Central competer	nt authority		
		Address	1.	.4.	Local competent	authority		
		Tel.						
	1.5.	Consignee	1.	.6.	Person responsib	ole for the loa	d in EU	
ent		Name			Name			
gnm		Address			Address			
onsi								
Ö		Postcode			Postcode			
tche		Tel.			Tel.			
ispa	1.7.	Country ISO code I.8. Region of Cod of origin	le I.	.9.	Country of destination	ISO code	I.10. Region of destination	Code
ofd								
ails	l.11.	Place of origin	1.	.12.	Place of destinati	ion		
. De								
Part I : Details of dispatched consignment		Name Approval number					Custom warehouse	
Δ.		Address			Name		Approval number	
		Name Approval number			Address			
		Address						
		Name Approval number			Postcode			
		Address						
	I.13.	Place of loading	1.	.14.	Date of departure	•		
	I.15.	Means of transport	1.	.16.	Entry BIP in EU			
		Aeroplane  Ship  Railway wagon	<b>-</b>					
		Road vehicle  Other	1.	.17.				
		Identification						
		Documentation references						
	I.18.	Description of commodity				I.19. Comm	odity code (HS code)	
							I.20. Quantity	
	I.21.	Temperature of product					I.22. Number of pac	kages
		Ambient Chilled Chilled			Frozen			
	1.23.	Seal/Container No					I.24. Type of package	ging

Commission Regulatio	n (EU)	2019/319 of	6 February	2019	amending	Annex IX to	Regulation (	EC)
4NNFYII								

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ANNEX II

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Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

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

	COUNT	RY							Dogchews
	II.	Health info	rmati	on	II	a. Certificate reference N	No	II.b.	
		the Europe Regulation	an Pa (EU)	rliament a No 142/20	and of the Co	leclare that I have read ar uncil ( <sup>1a</sup> ), and in particula in particular Chapter II of a above:	r Article 10 of t	hat Regulation,	and Commission
Ę	II.1.	have been	prepai	ed exclus	sively with the	following animal by-produc	cts:		
Part II: Certification		(²) either	[-	killed, a	nd which are	f animals slaughtered or, fit for human consumption onsumption for commercial	in accordance		
Part II: (		(²) and/or	[-	slaughte mortem	erhouse and vinspection o	owing parts originating eith were considered fit for slau bodies and the following dance with Union legislatio	ighter for huma g parts of anim	n consumption t	following an ante-
				.,	consumption	bodies and parts of an in accordance with Union nunicable to humans or an	legislation, but		
				(ii)	heads of poul	try;			
						ns, including trimmings ar d the carpus and metacarp			
				(iv)	pig bristles;				
				(v)	feathers;]				
		(²) and/or	[-	humans having	or animals, o been conside	ich did not show any sig btained from animals that red fit for slaughter for nce with Union legislation;]	have been slau human consum	ghtered in a sla	ughterhouse after
		(²) and/or	[-			rising from the production one, greaves and centrifu			
		(²) and/or	[-			parts of such animals, exp able to humans or animals		als, which did no	ot show any signs
		(²) and/or	[-		oy-products fr s for human c	om aquatic animals origina onsumption;]	ting from plants	or establishme	nts manufacturing
		(²) and/or	[-	Council	Directive 96/	which have been treated 22/EC ( <sup>2a</sup> ), the import of tulation (EC) No 1069/2009	the material bei		
	II.2.	have been	subjec	ted					
		(²) either				nade from hides and skins nisms (including salmonell			eatment sufficient
		(²) and/or				made from animal by-prod nt of at least 90°C through			s of ungulates or
	II.3.					f at least five samples fro olies with the following star		ssed batch take	n during or after
		Salmonella	:		absence in	25g: n = 5, c = 0, m = 0, M	= 0,		
		Enterobacte	eriace	ae:	n = 5, c = 2	m = 10, M = 300 in 1 gran	nme;		

**Dogchews** 

COUNTRY

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

II.	Health info	rmation		II.a. (	Certificate reference No	II.b.		
II.4.	have underg	gone all preca	autions to av	oid conta	amination with pathogenic agents after	treatment;		
II.5.	were packed	were packed in new packaging;						
(²) [II.6.	the dogchev	ws described	above					
	(²) either	[is derived f	from other ru	ıminants	than bovine, ovine or caprine animals	.]]		
	(²) or	[is derived f	from bovine,	ovine or	caprine animals and does not contain	and is not derived from:		
		(²) either	continuou	ovine and caprine materials other than those derived from animals born, usly reared and slaughtered in a country or region classified as posing at BSE risk in accordance with Decision 2007/453/EC.]]				
		(²) or		specified risk material as defined in point 1 of Annex V to Regulation (No 999/2001 of the European Parliament and of the Council $(^4)$ ;				
			an sla ac	imals, ex lughtered cordance	lly separated meat obtained from bookcept from those animals that were do not not not not not not not not not no	born, continuously reared and posing a negligible BSE risk in		
			an ne the the or	imals wherevous tise cranial ose animal	product or derived product obtained nich have been killed, after stunnin sue by means of an elongated rod-sh cavity, or by means of gas injected in als that were born, continuously reard lassified as posing a negligible BSE r (C.]]]	g, by laceration of the central laped instrument introduced into into the cranial cavity, except for ed and slaughtered in a country		

### Notes

### Part I:

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

### Part II:

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

ANNEX II
Document Generated: 2023-08-31

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

COL	JNTRY			Dogchews			
II.	Health information	II.a. Certifica	te reference No	II.b.			
(2)	Delete as appropriate.						
( <sup>2a</sup> )	OJ L 125, 23.5.1996, p. 3.						
(3)	Where:						
_	n = number of samples to be tested;						
_	m = threshold value for the number of b samples does not exceed m;	acteria; the res	ult is considered satisfactory	if the number of bacteria in all			
-	M = maximum value for the number of bac more samples is M or more; and	teria; the result	is considered unsatisfactory if	the number of bacteria in one or			
_	c = number of samples the bacterial count of which may be between m and M, the sample still being considered acceptable if the bacterial count of the other samples is m or less.						
(4)	OJ L 147, 31.5.2001, p. 1.						
(5)	OJ L 172, 30.6.2007, p. 84.						
_	The signature and the stamp must be in a c	ifferent colour to	that of the printing.				
_	Note for the person responsible for the conand must accompany the consignment until						
Offic	cial veterinarian/Official inspector						
	Name (in capital letters):		Qualification	n and title:			
	Date:		Signature:				
	Stamp:						

CHAPTERealth certificateFor raw petfood for direct sale or animal by-products to 3(D) be fed to fur animals, intended for dispatch to or for transit through (2) the European Union

Ambient  $\square$ 

I.23. Seal/Container No

Document Generated: 2023-08-31

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

col	JNTRY	<b>/</b> :								Veterinary certifica	ite to EU	
	l.1.	Consignor					1.2.	Certificate refere	nce No	I.2.a.		
		Name					I.3. Central competent authority					
		Address					1.4.	Local competent	authority			
		Tel.										
	1.5.	Consignee					1.6.	Person responsib	ole for the loa	ad in EU		
ent		Name						Name				
gnm		Address					Address					
onsi												
o pe		Postcode						Postcode				
tch		Tel.						Tel.				
lispa	1.7.	Country of origin	ISO code	1.8.	Region of origin	Code	1.9.	Country of destination	ISO code	I.10. Region of destination	Code	
ofc												
Part I : Details of dispatched consignment	1.11.	Place of or	igin				1.12.	Place of destinat	ion			
e .												
artl	Name Approval number									Custom warehouse		
<u> </u>		Address						Name		Approval number		
		Name		Appro	val number			Address				
		Address										
		Name		Appro	val number		Postcode					
		Address										
	I.13.	Place of loa	ading				1.14.	Date of departure	e			
	I.15.	Means of to	ransport				I.16.	Entry BIP in EU				
		Aeroplane	☐ Ship		Railway wa	agon 🗖						
		Road vehicle  Other  Identification  Documentation references						1.17.				
	I.18.	Description	of commodi	ty					I.19. Comm	nodity code (HS code)		
										1		
										I.20. Quantity		
	1.21.	Temperatu	re of product							I.22. Number of pa	ckages	

Chilled

Frozen

I.24. Type of packaging

1.25.	Commodities certified for:								
	Petfood			Tech	nical use 🗆				
1.26.	For transit through I	EU to third country		I.27. For import of	or admission into EU				
	Third country	ISO code							
1.28.	Identification of the		val number	of establishments					
(8	Species Scientific name)	Nature of commodity	Manufactu	uring plant	Net weight	Batch number			

Health information

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

### COUNTRY

II.

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

II.b.

II.a. Certificate reference No

I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of

		the European Parliament and of the Council ( <sup>1a</sup> ) and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011 ( <sup>1b</sup> ), and in particular Chapter II of Annex XIII and Chapter II of Annex XIV thereto, and certify that the raw petfood or animal by-products described above:
_	II.1.	consist of animal by-products that satisfy the health requirements below;
icatio	II.2.	consist of animal by-products:
Certif		(a) derived from meat which satisfies the relevant animal and public health requirements laid down in:
Part II: Certification		<ul> <li>Commission Regulation (EU) No 206/2010 (<sup>3</sup>) and provided that the animals from which the meat is derived come from the third countries, territories or parts thereof</li></ul>
		<ul> <li>and/or Commission Regulation (EC) No 798/2008 (4), and provided that the animals from which the meat is derived come from the third countries, territories or parts thereof</li></ul>
		— and/or Commission Regulation (EC) No 119/2009 (5), and provided that the animals from which the meat is derived come from the third countries, territories or parts thereof
		(b) derived from animals that, at the slaughterhouse, have passed the ante-mortem health inspection during the period of 24 hours before the time of slaughter and have shown no evidence of the diseases referred in the Regulations referred to in point (a) for which the animals are susceptible; and
		(c) derived from animals that have been handled in the slaughterhouse before and at the time of slaughter or killing in accordance with the relevant provisions of Union legislation and have met requirements at least equivalent to those laid down in Chapters II and III of Council Regulation (EC) No 1099/2009 ( <sup>6</sup> ); or
		(d) in the case of feed for fur animals, are derived from aquatic animals which satisfy the relevant animal and public health requirements laid down in Commission Decision 2006/766/EC (7), and come from countries or territories thereof (ISO code of the country) as listed in Annex II to that Decision;
	II.3.1.	consist only of the following animal by-products:
		<ul> <li>(a) carcases and parts of animals slaughtered or, in the case of game, bodies or parts of animals killed which were deemed fit for human consumption in accordance with Union legislation until irreversibly declared as animal by-products for commercial reasons;</li> </ul>
		(b) parts of slaughtered animals, which are rejected as unfit for human consumption but are not affected by any signs of diseases communicable to humans or animals and derived from carcases that are fit for human consumption in accordance with Union legislation;
	II.3.2.	in the case of feed for fur animals in addition to II.3.1. consist also of the following animal by-products:
		(²) either [- animal by-products from poultry and lagomorphs slaughtered on the farm as referred to in Article 1(3)(d) of Regulation (EC) No 853/2004 of the European Parliament and of the Council (²a), which did not show any signs of disease communicable to humans or animals;]
		(²) and/or [- blood of animals which did not show any signs of disease communicable through blood to humans or animals, obtained from animals that have been slaughtered in a slaughterhouse after having been considered fit for slaughter for human consumption following an ante-mortem inspection in accordance with Union legislation;]
		(²) and/or [- animal by-products arising from the production of products intended for human consumption, including degreased bone, greaves and centrifuge or separator sludge from milk processing;]

Certificate reference No

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

### COUNTRY

**Health information** 

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

	(²) and/or	[-	products of animal origin, or foodstuffs containing products of animal origin, which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects from which no risk to public or animal health arises;]
	(²) and/or	[-	petfood and feedingstuffs of animal origin, or feedingstuffs containing animal by-products or derived products, which are no longer intended for feeding for commercial reasons or due to problems of manufacturing or packaging defects or other defects from which no risk to public or animal health arise;]
	(²) and/or	[-	blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk originating from live animals that did not show signs of any disease communicable through that product to humans or animals;]
	(²) and/or	[-	aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of diseases communicable to humans or animals;]
	(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:
			<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 invertebrates other than species pathogenic to humans or animals;]
	(²) and/or	[-	animals and parts thereof of the zoological orders of Rodentia and Lagomorpha, except Category 1 material as referred to in Article 8(a)(iii), (iv) and (v) of Regulation (EC) No 1069/2009 and Category 2 material as referred to in Article 9(a) to (g) of that Regulation;]
II.4.			ned and prepared without contact with other material which does not comply with the conditions laid ulation (EC) No 1069/2009, and it has been handled so as to avoid contamination with pathogenic
II.5.	CONSUMP CONSUMP preventing	TION TION any l	ted in final packaging which bear labels indicating 'RAW PET FOOD — NOT FOR HUMAN or 'ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS — NOT FOR HUMAN and then placed in leak-proof and officially sealed boxes/containers or in new packaging eakage and officially sealed boxes/containers which bear labels indicating 'RAW PET FOOD —

- II.6. in the case of raw petfood:
  - has been prepared and stored in a plant approved and supervised by the competent authority in accordance with Article 24 of Regulation (EC) No 1069/2009 and

NOT FOR HUMAN CONSUMPTION OF 'ANIMAL BY-PRODUCTS FOR FEED FOR FUR ANIMALS — NOT FOR

HUMAN CONSUMPTION', and the name and the address of the establishment of destination;

was examined by random sampling of at least five samples from each batch taken during storage (before dispatch) and complies with the following standards  $(^8)$ : (b)

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

#### COUNTRY

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

II.	Health informati	ion		II.a. Certificate reference No II.b.				
	Salmonella:	ab	sence in 25 g	: n=5, c=0, m=0, M=0				
	Enterobacteriace	ae: n=	5, c=2, m=10	M=5000 in 1 gram;				
(²) [II.7.	[the petfood or a products of rumin			e fed to fur animals described above contains or is derived from animal-by				
	(²) either			untry or region, which is classified as posing a negligible BSE risk in sion 2007/453/EC, and in which there has been no indigenous BSE case,				
	(²) or	(²) or [originates from a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC in which there has been an indigenous BSE case, and the animal by product or derived product were derived from animals born after the date from which the ban of the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants, a defined in the OIE Terrestrial Animal Health Code, has been effectively enforced in that country or region, and]]						
	(²) either	[is derived fr	is derived from other ruminants than bovine, ovine or caprine animals.]]					
	(²) or	[is derived fr	om bovine, ov	ovine or caprine animals and does not contain and is not derived from:				
		(²) either	continuous	, ovine and caprine materials other than those derived from animals born, ously reared and slaughtered in a country or region classified as posing a ble BSE risk in accordance with Decision 2007/453/EC.]]				
		(²) or		ecified risk material as defined in point 1 of A 999/2001 of the European Parliament and o				
			cap and BS	chanically separated meat obtained from orine animals, except from animals that were d slaughtered in a country or region classing E risk in accordance with Commission De- ich there has been no indigenous BSE case	e born, continuously reared fied as posing a negligible cision 2007/453/EC (10), in			
			cap the ins into cor	mal by-product or derived product obtain orine animals which have been killed, after central nervous tissue by means of trument introduced into the cranial cavity, or the cranial cavity, except for those ntinuously reared and slaughtered in a cou- sing a negligible BSE risk in accordance with	stunning, by laceration of an elongated rod-shaped r by means of gas injected animals that were born, ntry or region classified as			

### Notes

### Part I:

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

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

### COUNTRY

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

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

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

CHAPTERealth certificateFor flavouring innards for use in the manufacture of 3(E) petfood, intended for dispatch to or for transit through (2) the European Union

COL	JNTRY	<b>/</b> :			Veterinary certificate to EU		
	l.1.	Consignor	1.2.	Certificate reference No	I.2.a.		
		Name	I.3. Central competent authority				
		Address	1.4.	Local competent authority			
		Tel.					
	1.5.	Consignee	1.6.	Person responsible for the	load in EU		
ent		Name		Name			
gnn		Address		Address			
onsi							
o pa		Postcode		Postcode			
atch		Tel.	10	Tel.	140 Desire of Orde		
disp	1.7.	Country ISO code I.8. Region of Code of origin origin	1.9.	Country of ISO destination code	I.10. Region of Code destination		
o of c							
Part I : Details of dispatched consignment	l.11.	Place of origin	I.12.	Place of destination			
. D							
artı		Name Approval number			Custom warehouse		
ъ.		Address		Name Approval number			
		Name Approval number		Address			
		Address					
		Name Approval number	Postcode				
		Address					
	I.13.	Place of loading	l.14.	Date of departure			
	I.15.	Means of transport	I.16.	Entry BIP in EU			
		Aeroplane					
		Road vehicle  Other	l.17.				
		Identification					
		Documentation references					
	I.18.	Description of commodity		I.19. Co	mmodity 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 certific	Commodities certified for:							
	Petfood			Technical use □					
1.26.	For transit through I	EU to third country		I.27. For import or admission in	to EU				
	Third country	ISO code							
1.28.	Identification of the	commodities							
		Appro	oval number	of establishments					
(8	Species Scientific name)	Nature of commodity	Manufactu	uring plant Net weight	Batch number				

### COUNTRY

### Flavouring innards for use in the manufacture

						of petfood			
	II.	Health info	ormati	on	II.a. Certificate reference No	II.b.			
		the Europe Regulation	ean Pa (EU) l	arliament and of the No 142/2011 (1b), and	declare that I have read and understood Reg Council (1a), and in particular Article 8 and I in particular Chapter III of Annex XIII and Cl roducts described above:	10 thereof, and Commission			
	II.1.	consist of a	nimal	by-products that satisf	fy the animal health requirements below;				
ation	II.2.	have been	prepar	ed and include the fol	lowing animal by-products which are exclusive	ely:			
Part II: Certification		(²) either	[-	killed, and which are	of animals slaughtered or, in the case of gar e fit for human consumption in accordance wit consumption for commercial reasons;]				
Parl		(²) and/or	[-	slaughterhouse and mortem inspection	llowing parts originating either from animals that have been slaughtered in a were considered fit for slaughter for human consumption following an anteor bodies and the following parts of animals from game killed for human ordance with Union legislation:				
				consumption	r bodies and parts of animals which are n in accordance with Union legislation, but wh nunicable to humans or animals;				
				(ii) heads of pou	ultry;				
					kins, including trimmings and splitting thereof nd the carpus and metacarpus bones, tarsus a				
				(iv) pig bristles;					
				(v) feathers;]					
	(²) and/or [- blood of animals w humans or animals, having been consid				hich did not show any signs of disease co obtained from animals that have been slaugh dered fit for slaughter for human consumpt ance with Union legislation;]	tered in a slaughterhouse after			
		(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	[-	intended for human	origin, or foodstuffs containing products of anin consumption for commercial reasons or due to r other defects from which no risk to public or a	problems of manufacturing or			
		(²) and/or	[-	derived products, w	gstuffs of animal origin, or feedingstuffs con hich are no longer intended for feeding for cturing or packaging defects or other defects	commercial reasons or due to			
		(²) and/or	[-		ol, feathers, hair, horns, hoof cuts and raw mi signs of any disease communicable throug				
		(²) and/or	[-		d parts of such animals, except sea mammals, which did not show any signs licable to humans or animals;]				
		(²) and/or	[-	animal by-products f products for human	from aquatic animals originating from plants or consumption;]	establishments manufacturing			
		(²) and/or	[-		rial originating from animals which did not show any signs of disease ugh that material to humans or animals:				
				(i) shells from s	shellfish with soft tissue or flesh;				

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

#### COUNTRY

### Flavouring innards for use in the manufacture

COUNTR	,						FI	avouring innard	ls for use in the manufa of po	acture etfood
II.	Health infor	rmatio	n		1	II.a. Certifica	ate reference	No	II.b.	
			(ii)	the follow	wing or	iginating from	n terrestrial an	imals:		
				– h	natcher	y by-products	5,			
				– е	eggs,	ys,				
				- е	egg by- <sub>l</sub>	by-products, including egg shells;				
			(iii)	day-old o	chicks I	ks killed for commercial reasons;]				
	(²) and/or	[-		by-produ s or anima		m aquatic o	r terrestrial in	vertebrates other	er than species pathoge	enic to
	(²) and/or	[-	Catego	ry 1 mate	erial as	referred to in	Article 8(a)(iii		itia and Lagomorpha, Regulation (EC) No 1069 Regulation;]	
	(²) and/or	[-	Counci	I Directive	e 96/22	2/EC ( <sup>2a</sup> ), the		e material being	ances which are prohibi permitted in accordance	
II.3.	have been subjected to processing in order to kill pathogenic agents;			in acc	ordance with	Chapter III of	Annex XIII to Re	egulation (EU) No 142/20	)11, in	
II.4.	was analysed by a random sampling storage at the processing plant and co					of at least five samples from each processed batch taken during or after applies with the following standards (3):				r after
	Salmonella: absence i			ce in 25	5g: n = 5, c =	0, m = 0, M =	0,			
	Enterobacteriaceae: n = 5, c =				c = 2, n	n = 10, M = 3	00 in 1 gramn	ne;		
II.5.	the end product was:									
	(²) either	either [packed in new or sterilis			rilised b	pags,]				
	(²) or							f transport that ent authority befo	were thoroughly cleane re use,]	d and
	and which be	ear lab	oels indi	cating 'No	OT FO	OT FOR HUMAN CONSUMPTION';				
II.6.	the end prod	duct wa	as store	d in enclo	osed sto	ed storage;				
II.7.	the product I	has ur	ndergon	e all preca	autions	to avoid con	tamination wit	h pathogenic ag	ents after treatment;	
(²) [II.8.	the flavourin	g inna	ırds prod	ducts des	cribed	ribed above				
	(²) either	[is de	erived fr	om other	rumina	ınts than bovi	ine, ovine or c	aprine animals.]		
	(2) or	[is de	erived fr	om bovin	ie, ovine	e or caprine a	animals and de	oes not contain a	and is not derived from:	
		(²) ei	ither	continuo	ously re	eared and sl	laughtered in		derived from animals region classified as pos EC.]]	
		(²) O	r					d in point 1 of iament and of the	Annex V to Regulation e Council (4);	ı (EC)
				s s	animals slaughte accorda	, except fror ered in a cou	m those anim untry or regior mmission Dec	nals that were b n classified as p	es of bovine, ovine or coorn, continuously reare osing a negligible BSE CC (5), in which there has	d and risk in
				a n tl tl	animals nervous the crar those a or regio	which have stissue by me nial cavity, or nimals that w	e been killed, eans of an eld by means of vere born, cor	after stunning, ongated rod-sha gas injected int ntinuously reared	rom bovine, ovine or come by laceration of the comped instrument introduced to the cranial cavity, except and slaughtered in a cook in accordance with Design	central ed into ept for country

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

### COUNTRY

### Flavouring innards for use in the manufacture

CO	OUNTRY Flavouring innards for use in the manufacture of petfood					
II.	Health information	II.a. Certificate reference No	II.b.			
Not	es					
Parl	El:					
_	Box reference I.6: Person responsible for the it is a certificate for a commodity to be transcommodity to be imported into the European	sited through the European Union; it may be				
_	Box reference I.12: Place of destination: this transit may only be stored in free zones, free		or transit commodity. Products i			
_	Box reference I.15: Registration number (rail information is to be provided in the event of u					
_	Box reference I.19: use the appropriate HS c	ode: 05.04; 05.06, 05.11 or 23.09 .				
_	Box reference I.23: for bulk containers, the co	ontainer number and the seal number (if app	licable) should be given.			
_	Box reference I.25: technical use: any use production or manufacturing of pet food.	e other than feeding of farmed animals, of	other than fur animals, and th			
_	Box reference I.26 and I.27: fill in according t	o whether it is a transit or an import certificat	te.			
_	Box reference I.28:					
	<ul> <li>species: select from the following: Ave Mollusca, Crustacea, Invertebrates oth</li> </ul>	es, Ruminantia, Suidae, Mammalia other tha er than Mollusca and crustacea	an Ruminantia or Suidae, Pesca			
	<ul> <li>define the innard product.</li> </ul>					
Part	t II:					
<sup>(1a</sup> )	OJ L 300, 14.11.2009, p. 1.					
( <sup>1b</sup> )	OJ L 54, 26.2.2011, p. 1.					
(²)	Delete as appropriate.					
( <sup>2a</sup> )	OJ L 125, 23.5.1996, p. 3.					
( <sup>3</sup> )	Where:					
	n = number of samples to be tested;					
	m = threshold value for the number of bacteria; the result is considered satisfactory if the number of bacteria in all samples does not exceed m;					
	M = maximum value for the number of bac or more samples is M or more; and	teria; the result is considered unsatisfactory	if the number of bacteria in on			
	c = number of samples the bacterial cou acceptable if the bacterial count of the	nt of which may be between m and M, thother samples is m or less.	ne sample still being considere			
( <sup>4</sup> )	OJ L 147, 31.5.2001, p. 1.					
<sup>(5</sup> )	OJ L 172, 30.6.2007, p. 84.					
_	The signature and the stamp must be in a dif	ferent colour to that of the printing.				
_	Note for the person responsible for the consignand must accompany the consignment until i		ate is only for veterinary purpose			
Offic	cial veterinarian/Official inspector					
	Name (in capital letters):	Qualificatio	n and title:			
	Date:	Signature:				
	Stamp:					

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

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

### COUNTRY:

### Veterinary certificate to EU

	l.1.	Consignor	1.2.	Certificate reference No	I.2.a.
		Name	1.3.	Central competent authority	
		Address	1.4.	Local competent authority	
		Tel.			
	1.5.	Consignee	1.6.	Person responsible for the lo	ad in EU
ent		Name		Name	
guu		Address		Address	
onsi					
ğ		Postcode		Postcode	
che		Tel.		Tel.	
Part I: Details of dispatched consignment	1.7.	Country ISO code I.8. Region of Code of origin	1.9.	Country of ISO code	I.10. Region of Code destination
s of					
etails	l.11.	Place of origin	I.12.	Place of destination	
ă					
art		Name Approval number			Custom warehouse
₾.		Address		Name	Approval number
		Name Approval number		Address	
		Address			
		Name Approval number		Postcode	
		Address			
	I.13.	Place of loading	1.14.	Date of departure	
_	115	Means of transport	116	Entry BIP in EU	
	1.15.	means of transport	1.10.	Lifty bir iii Lo	
		Aeroplane ☐ Ship ☐ Railway wagon ☐			
		Road vehicle Other O	1.17.		
		Identification			
		Documentation references			
	I.18.	Description of commodity		I.19. Com	modity 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

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

1.25.	Commodities certi	fied for:				
	Manufacture of pe	tfood 🗆	Further pro	ocess 🗆	Technical use	
1.26.	For transit through	EU to third country		I.27. For import or	admission into EU	
	Third country	ISO cod	е			
1.28.	Identification of the	e commodities	Approval number	of establishments		
(Sci	Species entific name)	Nature of commodity	Manufacturing plant	Number of packages	Net weight	Batch number

II.b.

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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 Commission Regulation (EU) No 142/2011 (¹b), and in

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

**Health information** 

### COUNTRY

II.

## Animal by-products for the manufacture of petfood

		particular	Chap	ter II of	Annex XIV thereto, and certify that the animal by-products described above:
	II.1.1.	consist of	anim	al by-pro	oducts that satisfy the animal health requirements below;
tion	II.1.2.	have beer	obta	ined in t	the territory of: (1c) from animals:
Part II: Certification		(²) either	[(a)		ave remained in this territory since birth or for a period of at least three months preceding te of slaughter or production;]
± ∷		(²) or	[(b)	killed ir	n the wild in this territory ( <sup>1d</sup> );]
Pa		(²) or	[(c)	derive	d from rodents, lagomorphs, aquatic animals or terrestrial or aquatic invertebrates;]
	II.1.3.	have beer	n obta	ined fro	m or produced by animals:
		(²) either	[(a)	coming	g from holdings:
				(i)	where, for the following diseases for which the animals are susceptible, there has been no case/outbreak of rinderpest, swine vesicular disease, Newcastle disease or highly pathogenic avian influenza during the period of the preceding 30 days, nor of classical or African swine fever during the period of the preceding 40 days; nor in the holdings situated in their vicinity within a 10 km radius, during the period of the preceding 30 days; and
				(ii)	where there has been no case/outbreak of foot-and-mouth disease during the period of the preceding 60 days, nor in the holdings situated in their vicinity within a 25 km radius, during the period of the preceding 30 days; and
			(b)	which:	
				(i)	were not killed to eradicate any epizootic disease;
				(ii)	have remained in their holdings of origin for a period of at least 40 days before the date of departure and which have been transported directly to the slaughterhouse without any contact with other animals which did not comply with the same health conditions;
				(iii)	at the slaughterhouse, have passed the ante-mortem health inspection during the period of 24 hours preceding the time of slaughter and have shown no evidence of the diseases referred to above for which the animals are susceptible; and
				(iv)	have been handled in the slaughterhouse before and at the time of slaughter or killing in accordance with the relevant provisions of Union legislation and have met requirements at least equivalent to those laid down in Chapters II and III of Council Regulation (EC) No 1099/2009 (4)]
		(²) or	[(a)	capture	ed and killed in the wild in an area:
				(i)	in which within a 25 km radius there has been no case/outbreak of any of the following diseases for which the animals are susceptible: foot-and-mouth disease, rinderpest, Newcastle disease or highly pathogenic avian influenza during the period of the preceding 30 days, nor of classical or African swine fever during the period of the preceding 40 days; and
				(ii)	situated at a distance of at least 20 km from any country or part of the territory of a country not authorised for export to the European Union of poultry material during the preceding 30 days or of porcine material during the preceding 40 days; and
			(b)	either	after killing were transported within a period of 12 hours following the killing for chilling to a collection centre and immediately afterwards to a game handling establishment, or

directly to a game handling establishment;]

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

### COUNTRY

### Animal by-products for the manufacture

				of petfood
II.	Health inform	ation	II.a. Certificate reference No	II.b.
II.1.4.	of the diseases 30 days or, in Union has bee	s referred to in point the event of a case an authorised only a	ment around which, within a radius of 10 II.1.3 for which the animals are susceptile of disease, the preparation of raw matafter the removal of all meat, and the tan official veterinarian;	ble during the period of the preceding terial for exportation to the European
II.1.5.			ed without contact with any other mate as been handled so as to avoid contamin	
II.1.6.	indicating 'RAV		ng preventing any leakage and in officiall FOR THE MANUFACTURE OF PET FO European Union;	
II.1.7.	consist only of	the following animal	by-products:	
	(²) either [-	killed which were	s of animals slaughtered or, in the case deemed fit for human consumption in a dad as animal by-products for commercial	ccordance with Union legislation until
	(²) and/or [-	slaughterhouse an mortem inspection	following parts originating either from ani id were considered fit for slaughter for hi n or bodies and the following parts of a cordance with Union legislation:	uman consumption following an ante-
		consumption	or bodies and parts of animals which on in accordance with Union legislation, ommunicable to humans or animals;	
		(ii) heads of po	oultry;	
			skins, including trimmings and splitting and the carpus and metacarpus bones, t	
		(iv) pig bristles	S	
		(v) feathers;]		
	(²) and/or [-		s arising from the production of produced bone, greaves and centrifuge or separa	
	(²) and/or [-	intended for humar	origin, or foodstuffs containing products n consumption for commercial reasons or or other defects from which no risk to pul	r due to problems of manufacturing or
	(²) and/or [-		nd parts of such animals, except sea ma unicable to humans or animals;]	mmals, which did not show any signs
	(²) and/or [-	animal by-products products for humar	s from aquatic animals originating from pl n consumption;]	ants or establishments manufacturing
	(²) and/or [-		erial originating from animals which dough that material to humans or animals:	lid not show any signs of disease
		(i) shells from	shellfish with soft tissue or flesh;	
		(ii) the following	ng originating from terrestrial animals:	
		— hat	chery by-products,	
		— egg	gs,	

egg by-products, including egg shells;

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

### COUNTRY

## Animal by-products for the manufacture of petfood

					of petrood		
II.	Health in	formation	II.a.	Certificate reference No	II.b.		
		(iii) day-old cl	icks kil	led 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	Category 1 mater	al as re		of Rodentia and Lagomorpha, except and (v) of Regulation (EC) No 1069/2009 g) of that Regulation;]		
	(²) and/or [- material from animals which have been treated with certain Council Directive 96/22/EC (⁴a), the import of the material by Article 35(a)(ii) of Regulation (EC) No 1069/2009;]						
II.1.8.	legislation	have been deep-frozen at the plant of origin or have been preserved in accordance with European Unic legislation in such a way that they will not spoil between dispatch and delivery to the plant of destination in the European Union or during the transit through the European Union;					
II.1.9.	Directive 9				ed with certain substances prohibited by mitted in accordance with Article 35(a)(ii)		
	lique trans of de of ea	(a) it has been marked in the third country before entry into the territory of the European Union by a cros liquefied charcoal or activated carbon on each outer side of each frozen block, or, when the raw materi transported in pallets which are not divided into separate consignments during transport to the petfood of destination in the European Union or during the transit through the European Union, on each outer of each pallet, in a way that the marking covers at least 70 % of the diagonal length of the frozen block be of at least 10 cm width;					
	entr	in the case of material which is not frozen, the raw material has been marked in the third country bet entry into the territory of the European Union by spraying it with liquefied charcoal or by applying charco powder in such a way that the charcoal is clearly visible on the material; and					
	othe	where the animal by-products are made up of raw material which has been treated as referred to above an other non-treated raw material, all the raw materials have been marked as referred to in point (a) and (a) above.					
(²) ( <sup>5</sup> ) [II.2.	Specific requirements						
(²) ( <sup>6</sup> ) [II.2.1.	(II.1.2), w	The by-products in this consignment come from animals that have been kept in the territory referred to in poir (II.1.2), where vaccination programmes against foot-and-mouth disease are being regularly carried out an officially controlled in domestic bovine animals.]					
(²) ( <sup>7</sup> ) [II.2.2.	ruminants hours, or	The by-products in this consignment consist only of animal by-products derived from trimmed offal of domestic ruminants, which have maturated at an ambient temperature of more than + 2 °C for a period of at least three hours, or in the case of masseter muscles of bovine animals and deboned meat of domestic animals, for a period of at least 24 hours.]]					
(²) [II.3.		the animal by-products for the manufacture of petfood contains or is derived from animal-by products of ruminan origin and:					
	(²) either		-	egion, which is classified as poon and in which there has been no in	sing a negligible BSE risk in accordance ndigenous BSE case, and]]		
	(²) or	Decision 2007/453/EC or derived product wer ruminants with meat-a	in whice derive nd-bon	ch there has been an indigenous ed from animals born after the d e meal and greaves derived	negligible BSE risk in accordance with us BSE case, and the animal by-product late from which the ban on the feeding of from ruminants, as defined in the OIE I in that country or region, and]]		
	(²) either	[is derived from other r	uminan	ts than bovine, ovine or caprine	animals.]		

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

#### COUNTRY

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

II.	Health i	nformation		II.a.	Certificate reference No	II.b.
	(²) or	[is derived	from bovine	ovine (	or caprine animals and does n	ot contain and is not derived from:
		(²) either	continuou	sly rea		than those derived from animals born, ountry or region classified as posing a a 2007/453/EC.]]]
		(²) or			risk material as defined in 001 of the European Parliame	point 1 of Annex V to Regulation (EC) nt and of the Council (8);
			ar sla ac	imals, aughter cordan	except from those animals t ed in a country or region clas	I from bones of bovine, ovine or caprine that were born, continuously reared and saified as posing a negligible BSE risk in 2007/453/EC(9), in which there has been
			ar ne th th	imals v rvous t e crania ose ani	which have been killed, afte issue by means of an elongat al cavity, or by means of gas mals that were born, continuc classified as posing a negligit	obtained from bovine, ovine or caprine or stunning, by laceration of the central ted rod-shaped instrument introduced into injected into the cranial cavity, except for busly reared and slaughtered in a country ble BSE risk in accordance with Decision

#### Notes

#### Part I:

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

### Part II:

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

ANNEX II
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Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

### COUNTRY

## Animal by-products for the manufacture of petfood

II.	Health information	II.a.	Certificate reference No	II.b.		
(1c)	The name and ISO code number of the expo	orting c	ountry as laid down in:			
	— Part 1 of Annex II to Regulation (EU) No 206/2010;					
	<ul> <li>Part 1 of Annex I to Regulation (EC) N</li> </ul>	o 798/	2008, and			
	<ul> <li>Part 1 of Annex I to Regulation (EC) N</li> </ul>	o 119/	2009.			
	In addition the ISO code of regionalisation is concerned) must be included.	n the a	bovementioned Annexes (whe	ere applicable for the susceptible species		
( <sup>1d</sup> )	Only for countries from which game meat in importation into the European Union.	itended	d for human consumption of the	e same animal species is authorised for		
(2)	Delete as appropriate.					
(3)	Excluding raw blood, raw milk, hides and certificates in that Annex for the import of the			les and feathers (see relevant specific		
(4)	OJ L 303, 18.11.2009, p. 1.					
( <sup>4a</sup> )	OJ L 125, 23.5.1996, p. 3.					
(5)	Supplementary guarantees to be provided when the material of domestic ruminants originated in the territory of a South American or South African country or part thereof from where only maturated and deboned fresh meat of domestic ruminants for human consumption is permitted for exportation to the European Union. The whole masseter muscles of bovine animals, incised in accordance with Part B.1 of Chapter I of Section IV of Annex I to Regulation (EC) No 854/2004 of the European Parliament and of the Council (OJ L 139, 30.4.2004, p. 206), are also permitted.					
( <sup>6</sup> )	Only for certain South American countries.					
( <sup>7</sup> )	Only for certain South American and South	African	countries.			
(8)	OJ L 147, 31.5.2001, p. 1.					
(9)	OJ L 172, 30.6.2007, p. 84.					
_	The signature and the stamp must be in a di	fferent	colour to that of the printing.			
_	Note for the person responsible for the cons and must accompany the consignment until					
Offic	cial veterinarian/Official inspector					
	Name (in capital letters):		Qu	alification and title:		
	Date:		Sig	nature:		
	Stamp:					

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

CHAPTERealth certificateFor blood products not intended for human consumption 4(B) that could be used as feed material, intended for dispatch to or for transit through (2) the European Union

Road vehicle Other

Documentation references

I.18. Description of commodity

Identification

I.21. Temperature of product

Ambient ☐

I.23. Seal/Container No

I.19. Commodity code (HS code)

I.20. Quantity

I.22. Number of packages

I.24. Type of packaging

Veterinary certificate to EU

Document Generated: 2023-08-31

COUNTRY:

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

	l.1.	Consignor			1.2.	Certificate reference	ce No	I.2.a.	
		Name			1.3.	Central competent	authority		
		Address			1.4.	Local competent a	uthority		
		Tel.							
	1.5.	Consignee			1.6.	Person responsible	e for the load	d in EU	
i i		Name				Name			
Ĕ		Address				Address			
nsiç									
05		Postcode				Postcode			
chec		Tel.				Tel.			
Part I : Details of dispatched consignment	1.7.	Country ISO code	I.8. Region of	Code	1.9.	Country of	ISO	I.10. Region of	Code
ğ		of origin	origin			destination	code	destination	I
ils o									
Deta	I.11.	Place of origin			l.12.	Place of destinatio	n		
Ξ									_
Part		Name	Approval number					Custom warehouse	
-		Address				Name		Approval number	
		Name	Approval number			Address			
		Address							
		Name	Approval number			Postcode			
		Address							
	I.13.	Place of loading			I.14.	Date of departure			
	I.15.	Means of transport			I.16.	Entry BIP in EU			
		Aeroplane D Ship	Railway wag	gon 🗖					

I.17.

Frozen

Chilled

1.25.	Commodities certified for:					
	Animal feedingstuff □		Manufactu	re of petfood $\square$	Technical	use 🗆
1.26.	For transit through EU to third	d country		I.27. For import or admissio	n into EU	
	Third country	ISO code				
1.28.	Identification of the commodit	ties				
		Approv	al number	of establishments		
	Species (Scientific name)	Nature of comm	odity	Manufacturing plant		Batch number

### COUNTRY

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

			courd be used as reed materi						
	II.	Health inforn	nation	II.a.	Certificate reference No	II.b.			
		the European	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 bloc	onsist of blood products that satisfy the health requirements below;						
ıtion	II.2.	consist exclus	ively of blood products no	t intend	ded for human consumption;				
Part II: Certification	II.3.		epared and stored in a pla egulation (EC) No 1069/2		proved and supervised by the compete	ent authority in accordance with			
art II:	II.4.	have been pre	epared exclusively with the	follow	ing animal by-products:				
Δ.		(²) either			s, which is fit for human consumpti inded for human consumption for comm				
		(²) and/or	accordance with Union humans or animals, who was a second to the contract of t	egislat nich h ich we	s, which has been rejected as unf ion, but which did not show any signs as been derived from carcases that re considered fit for human consump Jnion legislation;]	of diseases communicable to have been slaughtered in a			
	II.5.	in order to ina	activate pathogenic agents, have been submitted						
		(²) either	[to processing in accordance with processing method						
		(²) or	[to a method and parameters which ensure that the product complies with the microbiologic standards set out in Chapter I of Annex X to Regulation (EU) No 142/2011;]						
		(²) or	intended for the feeding	of po	s, including spray dried blood and b rcine animals, to a heat treatment at the dry blood and blood plasma does kw) of less than 0,60.]	a temperature of at least 80°C			
	II.6.	the end produ	ct was:						
		(²) either	[packed in new or sterilis	ed bag	gs;]				
		(²) or			ers or other means of transport that approved by the competent authority be				
		and which bea	ar labels indicating 'NOT F	OR HI	JMAN CONSUMPTION';				
	II.7.	the end produ	ct was stored in enclosed	storag	e;				
	II.8.	the product ha	ct has undergone all precautions to avoid contamination with pathogenic agents after treatment;						
		(²) and	[in the case of blood products, including spray dried blood and blood plasma of porcine origin intended for the feeding of porcine animals, has been stored in dry warehouse conditions under room temperature for a period of at least 6 weeks.]						
	II.9.				r the responsibility of the competent ch was found to comply with the followi				
		Salmonella:	absence in 2	5g: n =	5, c = 0, m = 0, M = 0,				
		Enterobacteria	aceae: n = 5, c = 2,	m = 10	, M = 300 in 1 gram;				

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

(c)

### COUNTRY

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

				could be used as feed material
II.	Health infor	mation		II.a. Certificate reference No II.b.
(²) [II.10.	the blood pro	oducts descri	bed above	
	(²) either	[is derived	d from other	ruminants than bovine, ovine or caprine animals.]]
	(²) or	[is derived	d from bovin	e, ovine or caprine animals and does not contain and is not derived from:
		(²) either	continuous	ovine and caprine materials other than those derived from animals born, sly reared and slaughtered in a country or region classified as posing a BSE risk in accordance with Decision 2007/453/EC.]]
		(²) or	[(a)	specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001 of the European Parliament and of the Council ( $^5$ );
			(b)	mechanically separated meat obtained from bones of bovine, ovine or caprine animals, except from those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Commission Decision 2007/453/EC ( $^6$ ), in which there has been no indigenous BSE case,
			(c)	animal by-product or derived product obtained from bovine, ovine or caprine animals which have been killed, after stunning, by laceration of the central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, or by means of gas injected into the cranial cavity, except for those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]]
II.11.	the blood pro	oducts descri	bed above:	
	(²) either			or milk products of ovine or caprine animal origin or is not intended for feed for than fur animals.]
	(²) or			products of ovine or caprine animal origin and is intended for feed for farmed r animals, which:
		(a)		d from ovine and caprine animals which have been kept continuously since country where the following conditions are fulfilled:
			(i)	classical scrapie is compulsorily notifiable;
			(ii)	an awareness, surveillance and monitoring system is in place for classical scrapie; $ \\$
			(iii)	official restrictions apply to holdings of ovine or caprine animals in the case of a suspicion of TSE or the confirmation of classical scrapie;
			(iv)	ovine and caprine animals affected with classical scrapie are killed and destroyed;
			(v)	the feeding to ovine and caprine animals of meat-and-bone meal or greaves, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health (OIE), of ruminant origin has been banned and effectively enforced in the whole country for a period of at least the preceding seven years;
				Sevell years,

originate from holdings where no case of classical scrapie has been diagnosed during the period of at least the preceding seven years or, following the confirmation of a case of classical scrapie:

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

#### COUNTRY

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

II.	Health inform	nation		II.a.	Certificate reference No	II.b.
		(²) either	or :	slaug es ca	e and caprine animals on the holding hitered, except for breeding rams of the arrying at least one ARR allele and no carrying at least one ARR allele;]	ARR/ARR genotype, breeding
		(²) or	two inte pre poi the	stroye yea ensific esence int 3.2 follo	hals in which classical scrapie was co bed, and the holding has been subjects since the date of confirmation of the bed TSE monitoring, including testing e of TSE in accordance with the late of Chapter C of Annex X to Regulation wing animals which are over the age of the ARR/ARR genotype:	cted for a period of at least elast classical scrapie case to with negative results for the boratory methods set out in on (EC) No 999/2001, of all of
			_	an	imals which have been slaughtered for	human consumption; and
			-		imals which have died or been killed o t killed in the framework of a disease en	
II.12.		ducts described above one statement of the Cor			r are derived from animal-by products of ferred to in Box I.1,	f non-ruminant origin, and are,
	(²) either	[not intended for the p	orod	uction	n of feed for farmed animals, other than	fur animals.]
	(²) ( <sup>7</sup> ) or	Consignor has undert	ake ses	n to e carri	feed for non-ruminant farmed animals, on sure that the border inspection post of ed out in accordance with the meth lo 152/2009 (8).]	entry will be provided with the

### Notes

# Part I:

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

ANNEX II
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Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

# COUNTRY

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

Part II:  (**) OJ L 300, 14.11.2009, p. 1.  (**) OJ L 54, 28.2.2011, p. 1.  (**) Delete as appropriate.  (**) Insert method 1 to 5 or method 7 as applicable.  (**) 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.  (**) OJ L 147, 31.5.2001, p. 1.  (**) OJ L 172, 30.6.2007, p. 84.  (**) The person responsible for the load referred to in Box 16 must ensure that, if the blood products described in this health certificate are intended to be used for the production of feed for non-ruminant farmed animals, other than fur animals, the consignment must be snallysed, in accordance with the methods set out in Annex VI to Regulation (EC) No 1522/200, in unsure that the state of the production of the order of minds of m	II.	Health information	II.a. Certificate reference No	II.D.		
(*) Delete as appropriate. (*) Insert method 1 to 5 or method 7 as applicable. (*) 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.  (*) OJ L 147, 31.5.2001, p. 1. (*) OJ L 172, 30.6.2007, p. 84.  (*) The person responsible for the load referred to in Box I.6 must ensure that, if the blood products described in this health certificate are intended to be used for the production of feed for non-ruminant farmed animals, other than fur animals, the consignment must be analysed, in accordance with the methods set out in Annex V I to Regulation (EC) No 1522009, in order to verify the absence of unauthorised consistents of animal origin. The information on the result of such analysis must be attached to this health certificate when presenting the consignment at a border inspection post of the European Union.  (*) OJ L 54, 26.2.2009, p. 1.  The signature and the stamp must be in a different colour to that of the printing.  Note for the person responsible for the consignment in the European Union: this certificate is only for veterinary purposes and must accompany the consignment until it reaches the border inspection post of the point of entry into the European Union.  Official veterinarian/Official inspector  Name (in capital letters):  Qualification and title:  Signature:	Par	II:				
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Date: Signature:	Offic	cial veterinarian/Official inspector				
		Name (in capital letters):	Qualificatio	n and title:		
Stamp:		Date:	Signature:			
		Stamp:				

CHAPTERealth certificateFor untreated blood products, excluding those of equidae, 4(C) 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

I.21. Temperature of product

Ambient  $\square$ 

I.23. Seal/Container No

I.22. Number of packages

I.24. Type of packaging

Document Generated: 2023-08-31

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

cou	JNTRY	<b>/</b> :								Vete	erinary certifica	te to EU
	l.1.	Consignor					1.2.	Certificate refere	nce No	1.2	2.a.	
		Name					1.3.	Central competer	nt authority			
		Address					1.4.	Local competent	authority			
		Tel.										
	1.5.	Consignee					1.6.	Person responsib	le for the loa	d in El	U	
ent		Name						Name				
gnm		Address						Address				
Part I : Details of dispatched consignment		Postcode						Postcode				
ped o		Tel.						Tel.				
atch	1.7.	Country	ISO code	1.8.	Region of	Code	1.9.	Country of	ISO	110	Region of	Code
disp	1.7.	of origin	130 code	1.0.	origin	Code	1.9.	destination	code	1.10.	destination	Code
s of												
tails	l.11.	Place of or	igin				I.12.	Place of destinati	on			
ă												
art		Name		Appro	val number					Cust	tom warehouse	
-		Address						Name		Appı	roval number	
		Name		Appro	val number			Address				
		Address										
		Name		Appro	val number			Postcode				
		Address										
	I.13.	Place of lo	ading				I.14.	Date of departure	•			
	I.15.	Means of to	ransport				I.16.	Entry BIP in EU				
		Aeroplane			Railway wa	agon 🗖						
		Road vehic	cle 🔲 Othe	er 🗆			I.17.					
		Identification	on									
		Documenta	ation reference	ces								
	I.18.	Description	of commodi	ty					I.19. Comm	nodity o	code (HS code)	
										Ι.		
										1.20	. Quantity	

Chilled

Frozen 🗖

1.25.	Commodities certified for:	
	Technical use □	
1.26.	For transit through EU to third country	1.27. For import or admission into EU
	Third country ISO code	
1.28.	Identification of the commodities	
	Approval number	of establishments
	Species (Scientific name) Manufactu	uring plant Batch number

# COUNTRY

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

	II.	Health inforn	nation	II.a. Certificate reference No II.b.				
		the European	Parliament and of the	n, declare that I have read and understood Regulation (EC) No 1069/2009 of council (1*), and in particular Article 8(c) and Article 8(d) and Article 10 thereof, a 142/2011 (1b), and in particular Chapter II of Annex XIV thereto, and certify				
_	II.1.	the blood prod	ducts described above	onsist of blood products that satisfy the health requirements below;				
catior	II.2.	they consist e	exclusively of blood prod	ucts not intended for human or animal consumption;				
Part II: Certification	II.3.		en prepared and store	in a plant supervised by the competent authority or in the establishment of an animal by-products:				
Part II		(²) either		red animals, which is fit for human consumption in accordance with Union of intended for human consumption for commercial reasons;]				
		(²) and/or	with Union legislate animals, derived	ed animals, which is rejected as unfit for human consumption in accordance on, but which did not show any signs of diseases communicable to humans or from carcases that have been slaughtered in a slaughterhouse and were numan consumption following an ante-mortem inspection in accordance with				
		(²) and/or	for [- blood of slaughtered animals, which did not show any signs of diseases communicable to humans or animals, obtained from animals that have been slaughtered in a slaughterhouse after having been considered fit for human consumption following an ante-mortem inspection in accordance with Union legislation;]					
		(²) and/or	[- blood and blood consumption;]	products derived from the production of products intended for human				
		(²) and/or		roducts originating from live animals that did not show signs of any disease hugh that product to humans or animals;]				
		(²) and/or		s derived from animals which have been submitted to illegal treatment as $1(2)(d)$ of Council Directive $96/22/EC$ ( $^{2a}$ ) or Article $2(b)$ of Council Directive				
		(²) and/or	listed in Group B(	s containing residues of other substances and environmental contaminants ) of Annex I to Directive 96/23/EC, if such residues exceed the permitted level legislation or, in the absence thereof, in national legislation;]				
	II.4.	with Union le	gislation, in slaughterh	nanufactured from, was collected in slaughterhouses approved in accordance buses approved and supervised by the competent authority of the country of illities approved and supervised by the competent authority of the country of				
	(²) [II.5.	Proboscidea, where no cas least the pred	including crossbreds be of rinderpest, peste of	ined from animals belonging to the taxa Artiodactyla, Perissodactyla and atween species of those taxa, the blood was collected in a country or region es petits ruminants and Rift Valley fever has been recorded for a period of at in which vaccination has not been carried out against those diseases for a norths, and;				
		(²) either	country, or codes (3) disease has been rec	tories or parts thereof				
		(²) or	country or codes (3) been recorded for programmes against	tories or parts thereof				

# COUNTRY

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

							the fe	ed chain fo	or farmed animals
II.	Health inform	ation		II.a.	Certificate ref	ference No		II.b.	
(²) [II.5.1.	in the case of a	animals othe	er than Suidae	e and Ta	ayassuidae, in	third countries	or region	s in which :	
	(²) either	has been r	ecorded for a	period (		preceding 12 n	nonths an	id in which v	ropositive animals) vaccination has not nonths;]
	(2) or	[vesicular	stomatitis and	blueton	igue (²) seropo	sitive animals	are prese	nt (4);]]	
(²) [II.5.2.	classical swine	e fever and n has not b	African swine een carried or	fever h	nas been recor	rded for a peri	od of at I	east the pre	vesicular disease, eceding 12 months eding 12 months in
	(²) either	for a perio	d of at least t	he prec		hs and in which	ch vaccin	ation has no	has been recorded ot been carried out
	(²) or	[vesicular	stomatitis sero	positive	e animals are p	resent (4);]]]			
(²) [II.6.	in the case of the territory of					vian species th	ne animal	s and the p	roducts come from
	which has bee			lisease	and highly pat	thogenic avian	influenza	a as define	d in the Terrestrial
	which for a per	riod of at lea	st the preced	ing 12 n	nonths has not	carried out va	ccination	against avia	an influenza,
									castle disease with an lentogenic virus
II.7.	the products w	ere:							
	(²) either	[packed in	new or sterilis	sed bags	s or bottles,]				
	(²) or				ers or other moproved by the				ughly cleaned and
	the outer pack	aging or cor	ntainers bear	abels in	dicating 'NOT	FOR HUMAN	OR ANIM	IAL CONSU	IMPTION';
II.8.	the products w	ere stored i	n enclosed st	orage;					
II.9.	all precautions	were taken	to avoid cont	aminatio	on of the produ	icts with patho	genic age	ents during t	ransport;
(²) [II.10.	the untreated b	olood produ	cts described	above					
	(²) either	[is derived	from other ru	minants	than bovine, o	vine or caprine	e animals	.]]	
	(²) or	[is derived	from bovine,	ovine or	caprine anima	als and does no	ot contain	and is not	derived from:
		(²) either	continuously	reared		ered in a cou	ntry or r	egion class	om animals born, sified as posing a
		(²) or			k material as of the Europea				Regulation (EC)
			anima slaug accor	als, exc htered i	ept from thos in a country or	e animals tha r region classi	it were b	orn, continuosing a neg	e, ovine or caprine uously reared and gligible BSE risk in ich there has been

no indigenous BSE case,

Health information

Document Generated: 2023-08-31

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

#### COUNTRY

II.

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

(c)	animal by-product or derived product obtained from bovine, ovine or caprine animals which have been killed, after stunning, by laceration of the central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, or by means of gas injected into the cranial cavity, except for those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSF risk in accordance with Decision

2007/453/EC.]]]

Certificate reference No

#### Notes

#### Part I:

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

# Part II:

- (1a) OJ L 300, 14.11.2009, p. 1.
- (1b) OJ L 54, 26.2.2011, p. 1.
- (2) Delete as appropriate.
- (2a) OJ L 125, 23.5.1996, p. 3.
- (2b) OJ L 125, 23.5.1996, p. 10.
- (3) Code of the territory as it appears in Part 1 of Annex II to Regulation (EU) No 206/2010 (OJ L 73, 20.3.2010, p. 1).
- (4) In this case following the veterinary checks provided for in Directive 97/78/EC (OJ L 24, 30.1.1998, p. 9), and in accordance with the conditions laid down in Article 8(4) of that Directive, the products must be transported directly to the establishment at the place of destination.

ANNEX II
Document Generated: 2023-08-31

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

### COUNTRY

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

II.	Health information	II.a.	Certificate reference N	lo	II.b.
(5)	Code of the territory as it appears in Part 1 of A p. 1).	nnex	to Commission Regula	tion (EC) No 7	98/2008 (OJ L 226, 23.8.2008,
( <sup>6</sup> )	OJ L 147, 31.5.2001, p. 1.				
( <sup>7</sup> )	OJ L 172, 30.6.2007, p. 84.				
_	The signature and the stamp must be in a differ	ent co	lour to that of the printir	ng.	
_	Note for the person responsible for the consignand must accompany the consignment until it Union.				
Offic	cial veterinarian/Official inspector				
	Name (in capital letters):			Qualification a	and title:
	Date:			Signature:	
	Stamp:				

CHAPTERealth certificateFor treated blood products, excluding those of equidae, for 4(D) 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

I.21. Temperature of product

Ambient  $\square$ 

I.23. Seal/Container No

I.22. Number of packages

I.24. Type of packaging

Document Generated: 2023-08-31

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

cou	JNTRY	<b>/</b> :								Vete	erinary certificat	te to EU
	l.1.	Consignor					1.2.	Certificate refere	nce No	1.2	2.a.	
		Name					1.3.	Central competer	nt authority			
		Address					1.4.	Local competent	authority			
									,			
		Tel.										
	1.5.	Consignee	•				1.6.	Person responsib	ole for the loa	id in El	U	
ent		Name						Name				
E E		Address						Address				
nsić												
00 p		Postcode						Postcode				
tche		Tel.						Tel.				
spat	1.7.	Country of origin	ISO code	1.8.	Region of origin	Code	1.9.	Country of destination	ISO code	I.10.	Region of destination	Code
of di		or origin	I	l	origin			destination			destination	
ails	111	Place of or	igin :				112	Place of destinat	ion			
Det		1 1000 01 01	·g···				1.12.	ridos or dosundo				
Part I : Details of dispatched consignment		Name		Appro	val number					Cust	tom warehouse	
Ра		Address						Name			roval number	
		Name		Appro	val number			Address				
		Address										
		Name		Appro	val number			Postcode				
		Address										
	I.13.	Place of lo	ading				1.14.	Date of departure	•			
	115	Means of t	ransport				116	Entry BIP in EU				
	1.10.	mound of t	гапорот				1.10.	Linay 511 111 20				
		Aeroplane	☐ Ship		Railway wa	agon 🗖						
		Road vehic			•	•	1.17.					
		Identification	on									
		Documenta	ation reference	ces								
	I.18.	Description	n of commodi	ty					I.19. Comm	nodity o	code (HS code)	
										1.20	. Quantity	

Chilled

Frozen 🗖

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	
I.28.	Identification of the commodities	
	Approval number	of establishments
	Species (Scientific name) Manufacto	uring plant Batch number

check.]]

# COUNTRY

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

	II.	Health informat	ion	II.a.	Certificate reference No	II.b.		
		the European Pa	rliament and of the Cou	uncil (18	that I have read and understood Reg ), and in particular Article 8(c) and Arti 1 (1b), and in particular Chapter II of A	cle 8(d) and Article 10 thereof,		
_	II.1.	the blood produc	ts described above con	sist of I	blood products that satisfy the requirem	nents below;		
icatio	II.2.	they consist excl	usively of blood product	ts not ir	ntended for human or animal consumpt	ion;		
Part II: Certification	II.3.	they have been panimal by-produc		a plant	supervised by the competent authority	, exclusively with the following		
Part		(²) either [-			als, which is fit for human consumpti ed for human consumption for commerc			
		(²) and/or [-	with Union legislation animals, derived from	, but w m card	ls, which is rejected as unfit for huma hich did not show any signs of disease asses that have been slaughtered in onsumption following an ante-mortem	s communicable to humans or a slaughterhouse and were		
		(²) and/or [-	humans or animals, on having been consider	blood of slaughtered animals, which did not show any signs of diseases communicable to humans or animals, obtained from animals that have been slaughtered in a slaughterhouse after having been considered fit for human consumption following an ante-mortem inspection in accordance with Union legislation;]				
		(²) and/or [-			originating from live animals that did r ugh these products to humans or anima			
		(²) and/or [-	blood and blood pr consumption;]	roducts	derived from the production of	roducts intended for human		
		(²) and/or [-		in Artic	ave been derived from animals which lole 1(2)(d) of Council Directive 96/22/E0			
		(²) and/or [-	listed in Group B(3)	of Anr	ing residues of other substances and nex I to Directive 96/23/EC, if such regislation or, in the absence thereof, in r	esidues exceed the permitted		
	II.4.	accordance with	Union legislation, in slation or from live anima	aughtei	ctured from was been collected in rhouses approved and supervised by t acilities approved and supervised by the	he competent authority of the		
	(²) [II.5.	crossbreeds, oth guaranteeing the	er than Suidae and Ta	ayassui of foot	om Artiodactyla, Perissodactyla and dae, the products have undergone on -and-mouth disease, vesicular stomatit	e of the following treatments,		
		(²) either	[heat treatment at a check;]	temper	rature of 65 °C for at least three hours	, followed by an effectiveness		
		(²) and/or	[irradiation at 25 kGy	by gan	nma rays, followed by an effectiveness	check;]		
		(²) and/or	[change in pH to pH 5	5 for tw	o hours, followed by an effectiveness c	heck;]		
		(²) and/or	[heat treatment of a	at least	t 80 °C throughout their substance,	followed by an effectiveness		

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

# COUNTRY

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

II.	Health informati	ion	II.a. Certificate reference No II.b.		
(²) [II.6.	undergone one of and-mouth disea	of the following transe, vesicular st	red from Suidae, Tayassuidae, poultry and other avian species, the products have eatments guaranteeing the absence of pathogens of the following diseases: footomatitis, swine vesicular disease, classical swine fever, African swine fever logenic avian influenza, as appropriate to the species:		
	(²) either	[heat treatment check;]	at a temperature of 65 °C for at least three hours, followed by an effectiveness		
	(²) and/or	[irradiation at 2	5 kGy by gamma rays, followed by an effectiveness check;]		
	(²) and/or		of at least 80 °C for Suidae/Tayassuidae (²) and at least 70°C for poultry and cies (²) throughout the substance of the product, followed by an effectiveness		
(²) [II.7.			ived from species other than those listed in point II.5 or II.6, the products have the theoretical point (please specify):		
II.8.	The products were	re:			
	(²) either	[packed in new	or sterilised bags or bottles,]		
	(²) or		bulk in containers or other means of transport that were thoroughly cleaned and a disinfectant approved by the competent authority before use;] and		
	the outer packaging or containers bear labels indicating 'NOT FOR HUMAN OR ANIMAL CONSUMPTION';				
II.9.	the products were	e stored in enclos	ed storage;		
II.10.	all precautions w	ere taken to avoi	the contamination of the products with pathogenic agents after treatment;		
(²) [II.11.	The treated blood	d products descri	ped above		
	(²) either	[is derived from	other ruminants than bovine, ovine or caprine animals.]]		
	(²) or	[is derived from	bovine, ovine or caprine animals and does not contain and is not derived from:		
		(²) either	[bovine, ovine and caprine materials other than those derived from animals born continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]		
		(²) or	(a) specified risk material as defined in point 1 of Annex V to Regulation (EC No 999/2001 of the European Parliament and of the Council (3);		
			(b) mechanically separated meat obtained from bones of bovine, ovine o caprine animals, except from those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Commission Decision 2007/453/EC (4), in which there has been no indigenous BSE case,		
			(c) animal by-product or derived product obtained from bovine, ovine of caprine animals which have been killed, after stunning, by laceration of the central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, or by means of gas injected into the cranial cavity, except for those animals that were born continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]]		

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

# COUNTRY

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

		I		outoide the it	To ramica animais
II.	Health information	II.a.	Certificate reference No	)	II.b.
Not	es				
Par	t I:				
-	Box reference I.6: Person responsible for the it is a certificate for a commodity to be trans commodity to be imported into the European	ited th	rough the European Uni		
-	Box reference I.11 and I.12: Approval numbissued by the competent authority.	er: the	e registration number of	the establishm	nent or plant, which has been
-	Box reference I.12: Place of destination: this in transit may only be stored in free zones, free				a transit commodity. Products
_	Box reference I.15: Registration number (rail is to be provided. In the case of unloading a entry into the European Union.				
-	Box I.19: use the appropriate Harmonized Sy	stem (	HS) code under the follow	wing headings:	05.11, 30.02, 35.02 or 35.04.
_	Box reference I.23: for bulk containers, the co	ntaine	er number and the seal nu	ımber (if applic	able) must be included.
-	Box reference I.25: technical use: any use production or manufacturing of pet food.	othe	r than feeding of farme	d animals, oth	er than fur animals, and the
_	Box reference I.26 and I.27: fill in according to	whet	her it is a transit or an im	port certificate.	
-	Box reference I.28 in case of Species: se Ruminantia or Suidae, Pesca, Reptilian.	lect fr	om the following: Aves,	Ruminantia, S	Suidae, Mammalia other than
Par	t II:				
( <sup>1a</sup> )	OJ L 300, 14.11.2009, p. 1.				
(1b)	OJ L 54, 26.2.2011, p. 1.				
(2)	Delete as appropriate.				
( <sup>2a</sup> )	OJ L 125, 23.5.1996, p. 3.				
( <sup>2b</sup> )	OJ L 125, 23.5.1996, p. 10.				
(3)	OJ L 147, 31.5.2001, p. 1.				
(4)	OJ L 172, 30.6.2007, p. 84.				
_	The signature and the stamp must be in a diff	erent	colour to that of the printi	ng.	
-	Note for the person responsible for the consignand must accompany the consignment until it				
Offic	cial veterinarian/Official inspector				
	Name (in capital letters):			Qualification a	and title:
	Date:			Signature:	
	Stamp:				

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

CHAPTERealth certificateFor game trophies or other preparations of birds and 6(B) ungulates consisting of entire parts which have not been treated, intended for dispatch to or for transit through (2) the European Union

cou	JNTRY	<b>′</b> :				Veterinary certificate to EU						
	l.1.	Consignor	1.2.	Certificate refere	ence No	I.2.a.						
		Name	1.3.	Central compete	nt authority							
		Address	1.4.	Local competent	authority							
		Tel.										
	1.5.	Consignee	1.6.	Person responsi	ble for the loa	ad in EU						
ent		Name		Name								
gu		Address		Address								
nsi												
D D		Postcode		Postcode								
tche		Tel.		Tel.								
ispa	1.7.	Country ISO code I.8. Region of Code of origin	1.9.	Country of destination	ISO code	I.10. Region of Code destination						
of d				accuration								
Part I : Details of dispatched consignment	l.11.	Place of origin	1.12.	Place of destination	tion							
Det		· ·										
Į.		Name Approval number				Custom warehouse						
2		Address		Name		Approval number						
		Name Approval number		Address								
		Address										
		Name Approval number		Postcode								
		Address										
	I.13.	Place of loading	1.14.	Date of departur	е							
	115	Means of transport	116	Entry BIP in EU								
	1.10.	Means of transport	1.10.	Lifty bir iii Lo								
		Aeroplane ☐ Ship ☐ Railway wagon ☐										
		Road vehicle Other O	1.17.	Number(s) of Cl	TES							
		Identification		(,,								
		Documentation references										
	I.18.	Description of commodity			I.19. Comm	nodity code (HS code)						
						I.20. Quantity						
	I.21.					I.22. Number of packages						
	1.23.	Seal/Container No				I.24. Type of packaging						

Commission	Regulation	(EU)	2019/319	of 6	February	2019	amending	Annex I	IX to .	Regulation	(EC)
4NNFYII											

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ANNEX II

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Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

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	
I.28.	Identification of the commodities	
	Species (Scientific name)	Number of packages

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

# COUNTRY

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

	II.	Health inf	ormatio	n	II.a.	Certificate reference	No	II.b.
		the Europ	ean Pa	liament and of the	Counc		on Regulation (E	ulation (EC) No 1069/2009 of U) No 142/2011 ( <sup>1b</sup> ), and in d above:
	(²) either	[II.1.	with re	spect to game trophic	es or o	ther preparations of clo	oven-hoofed anim	als, excluding swine:
Part II: Certification			(a)		eding 1	2 months, and during		uth disease and rinderpest for ccination against any of those
ü			(b)	the game trophies o	r other	preparations describe	d above:	
Pa				authorised for susceptible of there have b	r the e domest een no	xportation to the Euro ic species and where	pean Union of fre , during the perion	itory of that region, which is sh meat of the corresponding od of the preceding 60 days, eaks of diseases to which the
				of another th	ird cou		country not author	least 20 km from the borders ised to export untreated game uropean Union;]
	(²) or	[II.1.	with re	spect to game trophic	es or o	ther preparations of wi	ld swine:	
			(a)	classical swine feve porcine enteroviral	r, Afric enceph	an swine fever, swine	vesicular disease isease) and no va	ng 12 months, was free from t, foot-and-mouth disease and accinations have been carried
			(b)	the game trophies o	r other	preparations describe	d above:	
				exportation domestic sp	to the ecies a mal he	European Union of and where, during the	fresh meat of the property	ry, which is authorised for the le corresponding susceptible eceding 60 days, there have eases to which the swine are
				of another th	ird cou		country not author	least 20 km from the borders ised to export untreated game
	(²) or	[II.1.	describ					trophies or other preparations the territory of the exporting
	(²) or	[II.1.	with re	spect to game trophic	es or o	ther preparations of ga	ime birds:	
			(a)	disease; and	(r	egion) is free from hig	ghly pathogenic a	vian influenza and Newcastle
			(b)	that were killed in th	at reg	on and where during	the period of the	btained from wild game birds preceding 30 days there have to which the wild birds are
	II.2.	products of	of anima					out being in contact with other osed packages so as to avoid

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

#### COUNTRY

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

II.	Health inf	formation		II.a.	Certificate reference No	II.b.				
(²) [II.3.	The game	trophies or ot	her preparatior	s desc	cribed above					
	(²) either	[are derived	from other rum	inants	than bovine, ovine or caprine animals.]]					
	(²) or	[are derived	are derived from bovine, ovine or caprine animals and does not contain and is not derived from:							
		(²) either	continuously	reare	d caprine materials other than those ed and slaughtered in a country or re in accordance with Decision 2007/453/E	egion classified as posing a				
		(²) or			isk material as defined in point 1 of a 01 of the European Parliament and of the					
			anim slau acco	als, e ghtered rdance	lly separated meat obtained from bone xcept from those animals that were be d in a country or region classified as po e with Commission Decision 2007/453/Edus BSE case,	orn, continuously reared and osing a negligible BSE risk in				
			anim nerv the o thos or re	als word with the court of the	product or derived product obtained fr hich have been killed, after stunning, sue by means of an elongated rod-shap cavity, or by means of gas injected into lals that were born, continuously reared lassified as posing a negligible BSE risk EC.]]]	by laceration of the central ed instrument introduced into the cranial cavity, except for and slaughtered in a country				

### Notes

# Part I:

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

ANNEX II
Document Generated: 2023-08-31

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

# COUNTRY

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

Health information	II.a.	Certificate reference No		II.b.				
II:								
OJ L 300, 14.11.2009, p. 1.								
OJ L 54, 26.2.2011, p. 1.								
Delete as appropriate.								
OJ L 147, 31.5.2001, p. 1.								
OJ L 172, 30.6.2007, p. 84.								
The signature and the stamp must be in a diff	ferent	colour to that of the printir	ıg.					
ial veterinarian/Official inspector								
Name (in capital letters):			Qualification ar	nd title:				
Date:			Signature:					
Stamp:								
	II:  OJ L 300, 14.11.2009, p. 1.  OJ L 54, 26.2.2011, p. 1.  Delete as appropriate.  OJ L 147, 31.5.2001, p. 1.  OJ L 172, 30.6.2007, p. 84.  The signature and the stamp must be in a diff Note for the person responsible for the consignand must accompany the consignment until Union.  ial veterinarian/Official inspector  Name (in capital letters):  Date:	II:  OJ L 300, 14.11.2009, p. 1.  OJ L 54, 26.2.2011, p. 1.  Delete as appropriate.  OJ L 147, 31.5.2001, p. 1.  OJ L 172, 30.6.2007, p. 84.  The signature and the stamp must be in a different Note for the person responsible for the consignment and must accompany the consignment until it reac Union.  ial veterinarian/Official inspector  Name (in capital letters):  Date:	II:  OJ L 300, 14.11.2009, p. 1.  OJ L 54, 26.2.2011, p. 1.  Delete as appropriate.  OJ L 147, 31.5.2001, p. 1.  OJ L 172, 30.6.2007, p. 84.  The signature and the stamp must be in a different colour to that of the printing signature and the stamp must be in a different colour to that of the printing signature and must accompany the consignment until it reaches the border inspection Union.  ital veterinarian/Official inspector  Name (in capital letters):  Date:	II:  OJ L 300, 14.11.2009, p. 1.  OJ L 54, 26.2.2011, p. 1.  Delete as appropriate.  OJ L 147, 31.5.2001, p. 1.  OJ L 172, 30.6.2007, p. 84.  The signature and the stamp must be in a different colour to that of the printing.  Note for the person responsible for the consignment in the European Union: this certificate is and must accompany the consignment until it reaches the border inspection post of the poi Union.  ial veterinarian/Official inspector  Name (in capital letters):  Qualification and Date:  Signature:				

(4) Chapter 8 is replaced by the following:

CHAPTERealth certificateFor 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

Ambient  $\square$ 

I.23. Seal/Container No

Document Generated: 2023-08-31

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

col	JNTRY	<b>'</b> :					Veterinary certifica	te to EU
	I.1.	Consignor		1.2.	Certificate refere	nce No	I.2.a.	
		Name		1.3.	Central compete	nt authority		
		Address		1.4.	Local competent	authority		
		Tel.						
	1.5.	Consignee		1.6.	Person responsil	ble for the loa	d in EU	
ent		Name			Name			
gnm		Address			Address			
onsi								
o pa		Postcode			Postcode			
atch		Tel.	D-1		Tel.	100	140 8-1	0 1
disp	1.7.	Country ISO code I.8. of origin	Region of Code origin	1.9.	Country of destination	ISO code	I.10. Region of destination	Code
o de								
Part I : Details of dispatched consignment	l.11.	Place of origin	'	I.12.	Place of destinat	ion		
ă								
art		Name Appr	oval number				Custom warehouse	
п.		Address			Name		Approval number	
		Name Appr	oval number		Address			
		Address						
			oval number		Postcode			
		Address						
	l.13.	Place of loading		1.14.	Date of departure	е		
	I.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. Comm	nodity code (HS code)	
					L		I.20. Quantity	
	121	Temperature of product					I.22. Number of page	ckages
	1.21.	remperature or product					1.22. Number of par	ckayes

Chilled

Frozen

I.24. Type of packaging

1.25.	Commodities certi	fied for:				
	Technical use					
1.26.	For transit through	EU to third country	у 🗆	I.27. For import or a	admission into EU	
	Third country	ISO co	de			
1.28.	Identification of the	e commodities				
			Approval number	of establishments		
	Species entific name)	Nature of commodity	Manufacturing plant	Number of packages	Net weight	Batch number

# COUNTRY

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

									the f	feed cha	in or for	trade sa	mples	( <sup>2</sup> )
	II.	Health in	forma	tion		II.a.	Certificate	referenc	ce No		II.b.			
		of the Eu	ropear	Parliamer	veterinarian, nt and of the nex XIV there	Council	l ( <sup>1a</sup> ), and (	Commiss	ion Regu	lation (E	U) No 14	2/2011 (1		
ion		(²) either	referre	ed to in the	es which con definition of el 'TRADE S	trade s	samples in	point 39	of Annex	κ I to Reg	gulation (			
Part II: Certification		(²) or	[satisf	y the anim	al health requ	uiremen	ts set out	in point II	l.1.];					
Ë	II.1.	The anim	al by p	roducts de	scribed abov	re								
Pa	II.1.1.	have bee	n											
		(²) either	[(a)	obtained thereof: .	from ma	terials	imported (³) aut		a thi o export f			,		art ;]
		(²) and/or	r [(b)	obtained i animals th	n the exporti	ng third	country, to	erritory or	r part ther	reof:			( <sup>3</sup> ) fro	om
				either:										
				(i)	have remai meat to the three month	e Europ	ean Unior	since b	irth or fo	r a perio				
				(ii)	were killed	in the w	vild in that	third cour	ntry, territ	tory or pa	rt thereo	of (4);]		
		(²) and/or	r [(c)	derived fr	rom eggs, m ites;]	ilk, rod	ents, lago	morphs,	or aquat	tic anima	ls or te	rrestrial o	r aqua	atic
	(²) [II.1.2.				other than more or aquatic in									
		(²) either	[(a)	coming fro	om holdings:									
				(i)	where, for not been a disease or 30 days, not 40 days; not the period of	any cas highly or of cla or in the	e/outbreal pathogeniassical or / holdings	k of rinde c avian i African sv situated	erpest, s influenza wine feve in their v	wine ves during the r during	sicular d ne period the perio	isease, No of the pool of the p	lewcas recedi recedi	ing ing
				(ii)	where there period of th a 25 km rac	e prece	eding 60 d	ays, nor i	in the hol	ldings sit	uated in			
			(b)	which:										
				(i)	were not ki	lled to e	radicate a	ny epizod	otic disea	se;				
				(ii)	remained of of departur contact with	e and	which wer	e transp	orted dire	ectly to t	he slaug	hterhous	e with	out
				(iii)	at the slaug of 24 hours referred to	before	the time	of slaugh	ter and s	howed n	o eviden			
				(iv)	were handl accordance requiremen Regulation	with the	he relevai ast equival	nt provisi ent to the	ions of U	Inion leg	islation a	and comp	olied w	/ith
	I													ı

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

# COUNTRY

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

									the ree	eu chai	11 01 10	i traue s	samples (
II.	Health inf	orma	ition		II.a.	Certific	ate refe	rence 1	No		II.b.		
	(²) or	[(a)	captured a	nd killed in t	the wil	d in an a	rea:						
			(i)	where with following di rinderpest, period of th period of th	isease Newo he pre	es for whi castle dis ceding 3	ch the a sease of 0 days r	nimals r highly nor of c	are sus	ceptible genic a	e: foot-a avian ir	and-mou nfluenza	ith diseas during th
			(ii)	that is situ another ter dates for th	ritory (	of a third	country	or par	t thereof	f, which	is not	authoris	
		(b)	which after centre and establishm	d immediat									
(²) [II.1.3.	obtained in diseases r 30 days of exportation	n an referr r, in n to t	materials oth establishme ed to in poi the event o he Europea of the estab	ent around int II.1.2 for of a case/ou in Union wa	which, r which utbreak as auth	, within a h the an k of one norised o	radius imals ar of those nly after	of 10 e susce disea the re	km, there eptible ases, the moval of	re has during e prepa of all m	been n a perio aration	od case/od of the	outbreak o precedin material fo
II.1.4.				nd prepared without contact with other material which does not comply with the re, and it has been handled so as to avoid contamination with pathogenic agents;									
II.1.5.	disinfected sealed un PRODUCT	l befo ider ΓS ΟΙ	ore use and the respons NLY FOR Th	n new packaging which prevents any leakage or in packaging which has been cleaned and use and, in the case of consignments shipped other than via parcel post, in containers responsibility of the competent authority, bearing the label indicating 'ANIMAL BYFOR THE MANUFACTURE OF DERIVED PRODUCTS FOR USES OUTSIDE THE FEED me and address of the establishment of destination in the European Union;									
II.1.6.	consist onl	ly of t	he following	animal by-	produc	cts:							
	(²) either	[-	killed which	and parts of h were deer declared a	med fit	for huma	an consu	imption	in acco	rdance	with U		
	(²) and/or	[-	slaughterhoante	and the followouse and we minspection insumption in	vere con or l	onsidere bodies a	d fit for nd the f	slaugh followir	iter for l	numan	consur	mption fo	ollowing a
			(i)	carcases of consumption signs of dis	on in a	accordan	ce with	Union	legislati	ion, bu			
			(ii)	heads of po	oultry;								
			(iii)	hides and s the phalan bones;	skins, i iges a	including and the d	trimmin carpus a	gs and and me	splitting tacarpu	thereo	f, horns s, tars	and fee	et, includin metatarsu
			(iv)	pig bristles	Ģ								
			(v)	feathers;]									
	(²) and/or	[-	Article 1(3	products from the products from (b) of Reg	gulatio	n (EC)	No 853/	2004	of the E	uropea	an Parl	liament	and of th
	(²) and/or	[-	humans or after havin	nimals which r animals, ong been conspection in a	btaine nsider	ed from a red fit for	nimals t slaugh	hat ha	ve been human	slaugh	itered i	n a slau	ghterhous

**Health information** 

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Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

# COUNTRY

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

Certificate reference No

	(²) and/or	[-	animal by-products arising from the production of products intended for human consumption, including degreased bone, greaves and centrifuge or separator sludge from milk processing;]
	(²) and/or	[-	products of animal origin, or foodstuffs containing products of animal origin, which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects from which no risk to public or animal health arises;]
	(²) and/or	[-	petfood and 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 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:
			<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 invertebrates, other than species pathogenic to humans or animals;]
	(²) and/or	[-	animals and parts thereof of the zoological orders of Rodentia and Lagomorpha, except Category 1 material as referred to in Article 8(a)(iii), (iv) and (v) of Regulation (EC) No 1069/2009 and Category 2 material as referred to in Article 9(a) to (g) of that Regulation;]
	(²) and/or	[-	furs originating from dead animals that did not show clinical signs of any disease communicable through that product to humans or animals;]
II.1.7.		in sı	ep-frozen at the plant of origin or have been preserved in accordance with European Union uch a way that they will not spoil between the time of dispatch and the time of delivery to the ation.
(²) ( <sup>6</sup> ) [II.1.8.			
(²) ( <sup>7</sup> )			

either [II.1.8.1. The animal by-products in this consignment come from animals that have been obtained in the country,

territory or part thereof referred to in point II.1.1, where vaccination programmes against foot-and-mouth disease are regularly carried out and officially controlled in domestic bovine animals.]]

# COUNTRY

# Animal by-products to be used for purposes outside

				the feed ch	ain or for trade samples (2)				
II.	Health info	ormation		II.a. Certificate reference No	II.b.				
(²) ( <sup>8</sup> )									
and/or [II.1.8.2.	The anima meat.]]	the animal by-products in this consignment consist of animal by-products derived from offal or deboned neat.]]							
(²) [II.1.9.	the animal	by-product	s described above	е					
	(²) either	[are derive	ed from other rum	ninants than bovine, ovine or caprine anima	ls.]]				
	(²) or	[are derive	ed from bovine, o	vine or caprine animals and does not conta	in and is not derived from:				
		(²) either	continuously	e and caprine materials other than those reared and slaughtered in a country or re E risk in accordance with Decision 2007/450	egion classified as posing a				
		(²) or		d risk material as defined in point 1 of A 2001 of the European Parliament and of the					
			(b) mechanically separated meat obtained from bones of bovine, animals, except from those animals that were born, continuo slaughtered in a country or region classified as posing a negliq accordance with Commission Decision 2007/453/EC ( <sup>10</sup> ), in the been no indigenous BSE case,						
			animals nervous into the for those country	by-product or derived product obtained from which have been killed, after stunning, this tissue by means of an elongated rod-shoranial cavity, or by means of gas injected if a enimals that were born, continuously not region classified as posing a negligible in 2007/453/EC.]]]	by laceration of the central naped instrument introduced into the cranial cavity, except eared and slaughtered in a				
II.1.10	the animal	by-product	s described above	e:					
	(²) either		ntain milk or milk imals, other than	products of ovine or caprine animal origin of fur animals.]	or is not intended for feed for				
	(²) or			cts of ovine or caprine animal origin and is mals, and the milk or milk products:	intended for feed for farmed				
				e and caprine animals which have been ke llowing conditions are fulfilled:	pt continuously since birth in				
		(i)	classical so	crapie is compulsorily notifiable;					
		(ii)	an awarene	ess, surveillance and monitoring system is i	n place for classical scrapie;				
		(iii)		rictions apply to holdings of ovine or capri of TSE or the confirmation of classical scrap					
		(iv)	ovine and o	caprine animals affected with classical scrap	oie are killed and destroyed;				
		(v)	defined in t Health (OIE	to ovine and caprine animals of meat-an the Terrestrial Animal Health Code of the W E), of ruminant origin has been banned an try for a period of at least the preceding se	orld Organisation for Animal deflectively enforced in the				
		(b) origin	ate from holdings	s where no official restrictions are imposed	due to a suspicion of TSE;				
			d of the precedi	s where no case of classical scrapie has ng seven years or, following the confirma					

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

#### COUNTRY

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

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

# Notes

#### Part I:

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

killed in the framework of a disease eradication campaign. 11.

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

# COUNTRY

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

II.	Health information	II.a.	Certificate reference No		II.b.					
Part	t 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.									
( <sup>2a</sup> )	OJ L 139, 30.4.2004, p. 55.									
(3)	The name and ISO code number of the exporting	ig cour	itry as laid down in:							
_	Part 1 of Annex II to Commission Regulation (E	U) No :	206/2010 (OJ L 73, 20.3.:	2010, p. 1);						
_	Annex I to Commission Regulation (EC) No 798	3/2008	(OJ L 226, 23.8.2008, p.	1), and						
_	Annex I to Commission Regulation (EC) No 119	/2009	(OJ L 39, 10.2.2009, p. 1	2).						
	In addition the ISO code of territories and parts thereof referred to in the Annexes to Regulations (EU) No 206/2010, (EC) No 798/2008 and (EC) No 119/2009 referred to in this note (where applicable for the susceptible species concerned) must be included where applicable.									
(4)	Only for countries from where the game meat intended for human consumption of the same animal species is authorised for importation into the European Union.									
( <sup>5</sup> )	OJ L 303, 18.11.2009, p. 1.									
( <sup>6</sup> )	Supplementary guarantees to be provided where the material of domestic ruminants originated in the territory of a South American or South African country or part thereof from where only maturated and deboned fresh meat of domestic ruminants for human consumption is authorised for exportation to the European Union. The whole masseter muscles of bovine animals, incised in accordance with the requirements of Part B.1 of Chapter I of Section IV of Annex Ito Regulation (EC) No 854/2004 of the European Parliament and of the Council (OJ L 139 30.4.2004, p. 206), are also permitted.									
( <sup>7</sup> )	Only for certain South American countries.									
(8)	Only for certain South American and South Africa	can cou	untries.							
( <sup>9</sup> )	OJ L 147, 31.5.2001, p. 1.									
( <sup>10</sup> )	OJ L 172, 30.6.2007, p. 84.									
-	The signature and the stamp must be in a difference	ent col	our to that of the printing.							
_	Note for the person responsible for the consignand must accompany the consignment until it r Union.									
Offic	cial veterinarian/Official inspector									
	Name (in capital letters):		Q	ualification and	d title:					
	Date:		S	ignature:						
	Stamp:									

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

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

CHAPTERealth certificateFor rendered fats not intended for human consumption to 10(A) be used as feed material, intended for dispatch to or for transit through (2) the European Union

COL	JNTRY	<b>/</b> :			Veterinary certificate to EU		
	l.1.	Consignor	1.2.	Certificate reference No	I.2.a.		
		Name	1.3.	I.3. Central competent authority			
		Address	1.4.	Local competent authority			
		Tel.					
	1.5.	Consignee	1.6.	Person responsible for the I	oad in EU		
nent		Name		Name			
ignn		Address		Address			
sons		Postcode		Postcode			
ped o		Tel.		Tel.			
atch	1.7.	Country ISO code I.8. Region of Code	1.9.	Country of ISO	I.10. Region of Code		
disp	1.7.	of origin origin	1.0.	destination code	destination		
s of							
Part I : Details of dispatched consignment	l.11.	Place of origin	I.12.	Place of destination			
<u>.</u>							
art		Name Approval number			Custom warehouse		
-		Address		Name	Approval number		
		Name Approval number		Address			
		Address					
		Name Approval number		Postcode			
		Address					
	I.13.	Place of loading	I.14.	Date of departure			
	I.15.	Means of transport	I.16.	Entry BIP in EU			
		Aeroplane ☐ Ship ☐ Railway wagon ☐					
		Road vehicle  Other	I.17.				
		Identification					
		Documentation references					
	I.18.	Description of commodity		I.19. Con	nmodity 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 cert	ified for:						
	Animal feedingstu	ıff 🗆	Manufactu	re of petfood $\square$	Technical use	Technical use □		
1.26.	For transit through	h EU to third country	, 🗆	I.27. For import or	admission into EU			
	Third country	ISO co	de					
1.28.	28. Identification of the commodities  Approval number of establishments							
(Sci	Species entific name)	Nature of commodity	Manufacturing plant	Number of packages	Net weight	Batch number		

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

# COUNTRY

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

							used as feed material				
	II.	Health informati	ion		II.a.	Certificate reference No	II.b.				
	I, the undersigned official veterinarian, declare that I have read and understood Regulation (the European Parliament and of the Council (1a), and in particular Article 10 thereof, and Co (EU) No 142/2011 (1b), and in particular Chapter II of Annex XIV thereto, and certify that the reabove:										
_	II.1.	<ol> <li>consist of rendered fats that satisfy the health requirements below;</li> </ol>									
icatio	II.2.	consist of rendered fats not intended for human consumption;									
Part II: Certification	II.3.	have been prepared and stored in a plant approved and supervised by the competent authority in accordance with Article 24 of Regulation (EC) No 1069/2009 or in accordance with Article 4(2) of Regulation (EC) No 853/2004 of the European Parliament and of the Council (3), in order to kill pathogenic agents;									
å	II.4.	have been prepa	red ex	clusively with th	ne follow	ring animal by-products:					
		(²) either	[-	animals kille	d, and	of animals slaughtered or, in the case which are fit for human consumption t intended for human consumption for c	on in accordance with Union				
		(²) and/or	[-	slaughtered consumption	in a sla followin	following parts originating either from aughterhouse and were considered g an ante-mortem inspection or bodi alled for human consumption in accorda	fit for slaughter for human es and the following parts of				
				co	onsumpti	or bodies and parts of animals which a on in accordance with Union legislatio sease communicable to humans or ani	n, but which did not show any				
				(ii) he	heads of poultry;						
				in	cluding 1	I skins, including trimmings and split the phalanges and the carpus and m s bones;					
				(iv) pi	pig bristles;						
				(v) fe	athers;]						
		(²) and/or	[-	humans or ar after having	nimals, o been co	ch did not show any signs of disease c btained from animals that have been sl nsidered fit for slaughter for human c accordance with Union legislation;]	aughtered in a slaughterhouse				
		(²) and/or	[-		includin	arising from the production of pr ig degreased bone, greaves and centr					
	longer intende		led for h	igin, or foodstuffs containing products numan consumption for commercial re kaging defects or other defects from w	easons or due to problems of						
		(²) and/or	[-	or derived pr	oducts, ms of m	tuffs of animal origin, or feedingstuffs which are no longer intended for feed anufacturing or packaging defects or or alth arises;]	ing for commercial reasons or				
		(²) and/or	[-		did not	I, feathers, hair, horns, hoof cuts and show signs of any disease communi					

# COUNTRY

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

			used as feed material
II.	Health informati	on	II.a. Certificate reference No II.b.
	(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:
			<ul> <li>hatchery by-products,</li> </ul>
			<ul><li>eggs,</li></ul>
			<ul> <li>egg by-products, including egg shells;</li> </ul>
			(iii) day-old chicks killed for commercial reasons;]
1.5.	(²) either	[-	in the case of material of porcine origin, come from a country or part of the territory of a country free from foot-and-mouth disease for the period of the preceding 24 months and free from classical swine fever and African swine fever for the period of the preceding 12 months;]
	(²) and/or	[-	in the case of material of poultry origin, come from a country or part of a territory of a country free from Newcastle disease and avian influenza for a period of the preceding 6 months;]
	(²) and/or	[-	in the case of material of ruminant origin, come from a country or part of a territory of a country free from foot-and-mouth disease for the period of the preceding 24 months and free from rinderpest for the period of the preceding 12 months;]
	(²) and/or	[-	where there has been an outbreak of one of the diseases referred to in point II.5. during the relevant period referred to in point II.5, and where the rendered fats 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.			t animals, were purified in such way that the maximum levels of remaining total insoluble $0.15\%$ in weight;
1.7.	the rendered fats:	:	
		(a)	have been subjected to processing in accordance with the requirements of Section 3 of Chapter II of Annex X to Regulation (EU) No 142/2011, or a 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 have been 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 la	abels ir	dicating 'NOT FOR HUMAN CONSUMPTION';

(c)

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

# COUNTRY

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

II.	Health information			II.a.	Certificate reference No	II.b.			
(²) [II.8.	the rendere	d fats descr	ibed above						
	(²) either	[is derived from other ruminants than bovine, ovine or caprine animals.]]							
	(²) or	[is derived	[is derived from bovine, ovine or caprine animals and does not contain and is not derived from:						
		(²) either	continuously	reared	caprine materials other than those and slaughtered in a country or region ce with Decision 2007/453/EC.]]				
		(²) or			risk material as defined in point 1 c 001 of the European Parliament and of				
			ar sla ac	nimals, aughter ccordan	cally separated meat obtained from boto except from those animals that were ed in a country or region classified as ce with Commission Decision 2007/4 ndigenous BSE case,	born, continuously reared and posing a negligible BSE risk in			
			ar ne in fo	animal by-product or derived product obtained from bovine, ovine or caprine animals which have been killed, after stunning, by laceration of the central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, or by means of gas injected into the cranial cavity, except for those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]]					
II.9.	the rendere	d fats descr	ibed above:						
	(²) either		contain milk or imals, other tha		roducts of ovine or caprine animal orig nimals.]	n or is not intended for feed for			
	(²) or				of ovine or caprine animal origin and and the milk or milk products:	is intended for feed for farmed			
		(a)			ne and caprine animals which have be le following conditions are fulfilled:	en kept continuously since birth			
			(i) cla	assical	scrapie is compulsorily notifiable;				
				aware rapie;	eness, surveillance and monitoring s	ystem is in place for classical			
					strictions apply to holdings of ovine or of TSE or the confirmation of classical				
				ovine and caprine animals affected with classical scrapie are killed a destroyed;					
			de Ar er	efined in nimal H	ng to ovine and caprine animals of mean n the Terrestrial Animal Health Code lealth (OIE), of ruminant origin has in the whole country for a period of	of the World Organisation for been banned and effectively			
		(b)	originate fron TSE;	n holdir	ngs where no official restrictions are	imposed due to a suspicion of			

originate from holdings where no case of classical scrapie has been diagnosed during the preceding seven years or, following the confirmation of a case of classical scrapie:

# COUNTRY

Health information

II.

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

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

Certificate reference No

(2) either [all ovine and caprine animals on the holding have been killed and destroyed or

#### Notes

#### Part I:

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

#### Part II:

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

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

COL	INI	

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

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

CHAPTERealth certificateFor rendered fats not intended for human consumption to 10(B) 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

		'•				votormary continuat	0 10 20
	l.1.	Consignor	1.2.	Certificate reference No		I.2.a.	
		Name	I.3. Central competent authority				
		Address	1.4.	I.4. Local competent authority			
		Tel.					
	1.5.	Consignee	1.6.	I.6. Person responsible for the load in EU			
ent		Name		Name			
gnm		Address		Address			
onsi							
Ö		Postcode		Postcode			
che		Tel.		Tel.			
Part I: Details of dispatched consignment	1.7.	Country ISO code I.8. Region of Code of origin	1.9.	Country of destination ISO code		I.10. Region of destination	Code
s of							
stails	l.11.	Place of origin	I.12.	Place of destination			
ă							
art		Name Approval number				Custom warehouse	
Δ.		Address		Name		Approval number	
		Name Approval number		Address			
		Address					
		Name Approval number		Postcode			
		Address					
	I.13.	Place of loading	I.14.	Date of departure			
	I.15.	Means of transport	I.16.	Entry BIP in EU			
				,			
		Aeroplane ☐ Ship ☐ Railway wagon ☐					
		Road vehicle  Other	1.17.				
		Identification					
		Documentation references					
	I.18.	Description of commodity		I.19. C	ommo	dity code (HS code)	
						I.20. Quantity	
	I.21.	Temperature of product				I.22. Number of pac	kages
		Ambient ☐ Chilled ☐		Frozen 🗆			
	1.23.	Seal/Container No				I.24. Type of packag	ging

Commission Regulation	on (EU) 2019/319	of 6 February 20	19 amending Annex	IX to Regulation (EC)
ANNEYII				

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ANNEX II

Document Generated: 2023-08-31

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

1.25.	Commodities certifie	ed for:				
	Technical use					
1.26.	For transit through E	EU to third country		I.27. For import of	or admission into EU	
	Third country	ISO code				
1.28.	Identification of the					
		Appro	oval number	of establishments		
(8	Species Scientific name)	Manufacturing plant	Number of	· packages	Net weight	Batch number

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

# COUNTRY

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

					certain purposes outside the feed chain					
	II.	Health inform	ation		II.a. Certificate reference No	II.b.				
		I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council (1a), and in particular Articles 8, 9 and 10 thereof, and Commission Regulation (EU) No 142/2011 (1b), and in particular Chapter II of Annex XIV thereto, and certify that the rendered fadescribed above:								
_	II.1.	consist of rendered fats not intended for human consumption that satisfy the health requirements below;								
catior	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 point L of Section 2 of Chapter IV of Annex IV to Regulation (EU) No 142/2011, biodiesel or oleochemical products, animal by-products referred to in Articles 8, 9 and 10 of Regulation (EC) No 1069/2009;]								
Pa	(²) [II.2.2.	in the case of materials destined for the production of renewable fuels referred to in point J of Section 2 of Chapter IV of Annex IV to Regulation (EU) No 142/2011, the materials have been prepared exclusively from animal by-products referred to in Articles 9 and 10 of Regulation (EC) No 1069/2009;]								
	(²) [II.2.3.	<ol> <li>in the case of materials destined for purposes other than cosmetics, pharmaceuticals or medical devices, the materials have been prepared exclusively from:</li> </ol>								
		(²) either	[-		oducts containing residues of authorised permitted levels referred to in Article 15(3) of					
		(²) and/or	[-		nimal origin which have been declared unfit for preign bodies in those products;]	r human consumption due to the				
		(²) and/or	[-	(EC) No 1069	parts of animals, other than those referred to in /2009, that died other than being slaughtered nals killed for disease control purposes;]					
		(²) and/or	[-	animals kille	d parts of animals slaughtered or, in the ca d, and which are fit for human consumpt t are not intended for human consumption for o	ion in accordance with Union				
		(²) and/or	[-	carcasses and the following parts originating either from animals that have been sla in a slaughterhouse and were considered fit for slaughter for human consumption an ante-mortem inspection or bodies and the following parts of animals from game human consumption in accordance with Union legislation:						
				consu	ses or bodies and parts of animals which a nption in accordance with Union legislation, b ase communicable to humans or animals;					
				(ii) heads	of poultry;					
					and skins, including trimmings and splitting thalanges and the carpus and metacarpus bones					
				(iv) pig bris	stles;					
				(v) feather	rs;]					
(²) and/or [- blood of animals which did not show any signs humans or animals obtained from animals that after having been considered fit for slaughter to mortem inspection in accordance with Union legical contents.					imals obtained from animals that have been been considered fit for slaughter for human	slaughtered in a slaughterhouse				
		(²) and/or	[-		oducts arising from the production of pincluding degreased bone, greaves and center in the contract of the con					

milk processing;]

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

# COUNTRY

# Rendered fats not intended for human consumption for

			certain purposes outside the feed chain
II.	Health informat	tion	II.a. Certificate reference No II.b.
	(²) and/or	[-	products of animal origin, or foodstuffs containing products of animal origin, which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects from which no risk to public or animal health arises;]
	(²) and/or	[-	petfood and feeding stuffs of animal origin, or feeding stuffs containing animal by-products or derived products, which are no longer intended for feeding for commercial reasons or due to problems of manufacturing or packaging defects or other defects from which no risk to public or animal health arises;]
	(²) and/or	[-	blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk originating from live animals that did not show signs of any disease communicable through that product to humans or animals;]
	(²) and/or	[-	aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of diseases communicable to humans or animals;]
	(²) and/or	[-	animal by-products from aquatic animals originating from plants or establishments manufacturing products for human consumption;]
	(²) and/or	[-	the following material originating from animals which did not show any signs of disease communicable through that material to humans or animals:
			(i) shells from shellfish with soft tissue or flesh;
			(ii) the following originating from terrestrial animals:
			<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	[-	aquatic and terrestrial invertebrates other than species pathogenic to humans or animals;]
	(²) and/or	[-	animals and parts thereof of the zoological orders of Rodentia and Lagomorpha, except Category 1 material as referred to in Article 8(a)(iii), (iv) and (v) of Regulation (EC) No 1069/2009and Category 2 material as referred to in Article 9(a) to (g) of that Regulation;]
	(²) and/or	[-	hides and skins, hooves, feathers, wool, horns, hair and fur originating from dead animals that did not show any signs of disease communicable through that product to humans or animals;]
	(²) and/or	[-	adipose tissue from animals which did not show any signs of disease communicable through that material to humans or animals, which were slaughtered in a slaughterhouse and which were considered fit for slaughter for human consumption following an ante-mortem inspection in accordance with Union legislation;]]
(²) [II.2.4.			Is destined for purposes other than the production of organic fertilisers or soil improvers, ical or medical devices :
	(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 $\binom{2b}{2}$ ;
	(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 Council Directive 96/22/EC ( <sup>2c</sup> ) or Article 2(b) of Council Directive 96/23/EC;]

#### COUNTRY

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

II.	Health	information		II.a.	Certificate reference No	II.b.				
	(²) and	/or [-	contaminants the permitted	products containing residues of other substances and environmens listed in Group B(3) of Annex I to Directive 96/23/EC, if such residues exced levels laid down by Union legislation or, in the absence thereof, by legislation or state of importation;]]						
II.3.	the ren	dered fats:								
	(a)				accordance with methodAnnex IV to Regulation (EU) No 142/20					
	(b)				t to the European Union with glycerolt of at least 250 mg GTH per kilogramme f					
	(c)	in the case of r removed,	endered fats of	rumin	ant origin, insoluble impurities in excess	of 0,15% in weight have been				
	(d)	have been trans	sported under co	onditio	ns which prevent their contamination, and	i				
	(e)	bear labels on t	he packaging or	conta	niner indicating "NOT FOR HUMAN OR A	NIMAL CONSUMPTION";				
(²) [II.4.		case of materials dered fats descr		ganic f	fertilisers, cosmetics, pharmaceuticals, m	edical devices or soil improvers				
	(²) eith	er [are derive	ed from other ru	minan	ts than bovine, ovine or caprine animals.]					
	(²) or	[are derive	ed from bovine,	ne, ovine or caprine animals and does not contain and is not derived from:						
		(²) either	continuously r	eared	d caprine materials other than those and slaughtered in a country or region once with Decision 2007/453/EC.]					
		(²) or			k material as defined in point 1 of of the European Parliament and of the C					
	anima slaug accor		animal slaugh accord	s, exc ered ance v	separated meat obtained from bones cept from those animals that were be in a country or region classified as powith Commission Decision 2007/453/EC SE case,	orn, continuously reared and osing a negligible BSE risk in				
	(c) anim: which mear by m born,		which I means by mea born, c	nave by of an ans of ontinu	product or derived product obtained from bovine, ovine or caprine a e been killed, after stunning, by laceration of the central nervous tis an elongated rod-shaped instrument introduced into the cranial car of gas injected into the cranial cavity, except for those animals tha inuously reared and slaughtered in a country or region classified as e BSE risk in accordance with Decision 2007/453/EC.]]]					

# Notes

## Part I:

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

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

cou	UNTRY		Rendered fats		for human consumption for poses outside the feed chain
II.	Health information	II.a.	Certificate reference No		II.b.
-	Box reference I.12: Place of destination: this transit may only be stored in free zones, free				transit commodity. Products in
-	Box reference I.15: Registration number (rail- to be provided. In the case of unloading a inspection post of the point of entry into the E	nd rela	oading in the European U		
-	Box I.19: use the appropriate Harmonized S 15.04; 15.05; 15.06; 15.16 or 15.18.	System	n (HS) code under the follo	owing heading	s: 04.05; 15.01, 15.02; 15.03;
_	Box reference I.23: for bulk containers, the co	ontaine	er number and the seal nun	nber (if applica	ble) must be included.
-	Box reference I.25: technical use: any use of the production or manufacturing of pet food.	her tha	an feeding of farmed anima	ils, other than t	fur animals or pet animals, and
_	Box reference I.26 and I.27: fill in according to	o whet	her it is a transit or an impo	ort certificate.	
_	Box reference I.28:				
	Species: select from the following: Ruminanti	a, othe	er than Ruminantia		
	Manufacturing plant: provide the registration	numbe	er of the treatment/processi	ng establishme	ent.
Part	II:				
( <sup>1a</sup> )	OJ L 300, 14.11.2009, p. 1.				
(1b)	OJ L 54, 26.2.2011, p. 1.				
(²)	Delete as appropriate.				
( <sup>2a</sup> )	OJ L 125, 23.5.1996, p. 10.				
( <sup>2b</sup> )	OJ L 147, 31.5.2001, p. 1.				
( <sup>2c</sup> )	OJ L 125, 23.5.1996, p. 3.				
(3)	OJ L 147, 31.5.2001, p. 1.				
(4)	OJ L 172, 30.6.2007, p. 84.				
_	The signature and the stamp must be in a diff	ferent	colour to that of the printing	<b>]</b> .	
-	Note for the person responsible for the cons and must accompany the consignment until Union.				
Offic	cial veterinarian/Official inspector				
	Name (in capital letters):			Qualification ar	nd title:
	Date:		;	Signature:	
	Stamn:				

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

#### COUNTRY:

# Veterinary certificate to EU

		•			,				
	l.1.	Consignor	1.2.	Certificate reference No	I.2.a.				
		Name	1.3.	Central competent authorit	у				
		Address	1.4.	Local competent authority					
		Tel.							
	1.5.	Consignee	1.6.	Person responsible for the	load in EU				
ent		Name		Name					
gnm		Address		Address					
onsi									
ğ		Postcode		Postcode					
tche		Tel.		Tel.					
ispa	1.7.	Country ISO code I.8. Region of Code of origin origin	1.9.	Country of ISO destination code	I.10. Region of Code destination				
of d									
tails	l.11.	Place of origin	1.12.	Place of destination					
: De		•							
Part I : Details of dispatched consignment		Name Approval number			Custom warehouse				
ď		Address		Name	Approval number				
		Name Approval number		Address					
		Address							
		Name Approval number							
		Address							
	I.13.	Place of loading	1.14.	Date of departure					
	L15	Means of transport	1.16	Entry BIP in EU					
				2, 5 20					
		Aeroplane ☐ Ship ☐ Railway wagon ☐							
		Road vehicle  Other	1.17.						
		Identification							
		Documentation references							
	I.18.	Description of commodity		I.19. Co	mmodity 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				

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

1.25.	Commodities certifie	ed for:								
	Animal feedingstuff	Manufactu	re of petfood	☐ Technical	Technical use □					
1.26.	For transit through E		I.27. For ir	mport or admission into EU						
	Third country	ISO code								
1.28.	dentification of the commodities  Approval number of establishments									
(8	Species Manufacturing plant (Scientific name)		Number of	packages	Net weight	Batch number				

#### COUNTRY

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

							chain			
	II.	Health information			II.a.	Certificate reference No	II.b.			
		the European Parl	liament a	and of the	Council (	e that I have read and understood Regi <sup>1a</sup> ), and in particular Article 10 thereof, oter I of Annex XIV thereto, and certify	and Commission Regulation			
	II.1.	consists of gelatine	e/collage	n (²) that s	satisfy the	health requirements below;				
ation	II.2.	consist exclusively	of gelati	ine/collage	en (²) not i	ntended for human consumption;				
Part II: Certification	II.3.					oved and supervised by the competent order to kill pathogenic agents;	t authority in accordance with			
art	II.4.	has been prepared	d exclusiv	vely with the	he followin	ng animal by-products:				
а.		(²) either	ar	nimals kill	ed, and	of animals slaughtered or, in the case which are fit for human consumption to intended for human consumption for co	n in accordance with Union			
		(²) and/or	sla	aughtered onsumption	in a sla n followin	ollowing parts originating either froi aughterhouse and were considered g an ante-mortem inspection or bodie led for human consumption in accordan	fit for slaughter for human es and the following parts of			
			(i)	cons	ases or bodies and parts of animals which are rejected as unfit for human sumption in accordance with Union legislation, but which did not show any s of disease communicable to humans or animals;					
		(ii) heads of poultry;								
			(iii		es and skins, including trimmings and splitting thereof, horns and feet, including phalanges and the carpus and metacarpus bones, tarsus and metatarsus nes;					
			(iv	/) pig b	pristles;					
			(v	) feath	ners;]					
		(²) and/or	cc		n, includin	arising from the production of prog g degreased bone, greaves and centrif				
		(²) and/or	lo	nger inter	ided for h ng or pacl	igin, or foodstuffs containing products of uman consumption for commercial re- kaging defects or other defects from wh	asons or due to problems of			
		(²) and/or	or du	derived purchase desired to	products, volems of ma	tuffs of animal origin, or feedingstuffs on which are no longer intended for feeding anufacturing or packaging defects or oth alth arises;]	ng for commercial reasons or			
		(²) and/or			mals, and parts of such animals, except sea mammals, which did not show any eases communicable to humans or animals;]					
		(²) and/or			-products from aquatic animals originating from plants or establishments ring products for human consumption;]					
	II.5.	the gelatine/collag	en (²):							
			ar	nd in part	icular wra	ged, stored and transported under sa apping and packaging took place in ad under Union legislation were used.				

#### COUNTRY

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

									chain	
II.	Health info	rmation		II.a.	Certificate reference No		II.b.			
					packages containing gela GEN(2) SUITABLE FOR ANIM			the	words	
	(²) either	[(b)	Category 3 more rinse	materia s, involv followe	latine, was produced by a p I was subjected to a treatmen ving pH adjustment, extraction d by purification by means of	nt with acid on by heat	or alkali, followe	d by o	one or nes in	
	(²) or	[(b)	Category 3	materia ali follo	lagen, was produced by a p I was subjected to a treatment wed by one or more rinses,	involving v	vashing, pH adjus	stment	using	
(²) [II.6.	in the case	e of gelatine/o	ollagen (²) fro	om mate	erials other than hides and skin	s				
	(²) either	[is derived fi	om other rur	ninants t	than bovine, ovine or caprine a	nimals.]]				
	(²) or	[is derived fi	rom bovine, o	ovine or	caprine animals and does not o	contain and	is not derived fro	m:		
		(²) either	continuousl	y reared	d caprine materials other th I and slaughtered in a country once with Decision 2007/453/E0	or region cla				
		(²) or			sk material as defined in po 1 of the European Parliament a			ulation	(EC)	
			anin slau acc	nals, ex ghtered ordance	y separated meat obtained from those animals that in a country or region classi with Commission Decision 20 us BSE case,	at were bo ified as pos	rn, continuously sing a negligible	reared BSE i	d and risk in	
			anin tissu cavi that clas	nals whi ue by mo ty, or by were	product or derived product or ch have been killed, after stuni- eans of an elongated rod-shap means of gas injected into the born, continuously reared an as posing a negligible BSI C.]]]	ning, by lac ed instrume e cranial ca nd slaughte	eration of the cer ent introduced int wity, except for the ered in a countr	ntral ne o the d ose ar y or i	ervous cranial nimals region	
II.7.	in the case	e of gelatine/c	ollagen (2) fro	om mate	erials other than hides and skin	s described	above:			
	(²) either		ontain milk on nals, other tha		roducts of ovine or caprine ani nimals.]	imal origin	or is not intended	for fe	ed for	
	(²) or				of ovine or caprine animal ori and the milk or milk products:	igin and is	intended for feed	d for fa	armed	
				from ovine and caprine animals which were kept continuously since birth in a country llowing conditions are fulfilled:						
		(i)	clas	sical scr	rapie is compulsorily notifiable;					
		(ii)	an a	warene	ss, surveillance and monitoring	system is	in place for classi	cal scr	apie;	
		(iii)			ictions apply to holdings of ov TSE or the confirmation of clas			e cas	e of a	

ANNEX II

Document Generated: 2023-08-31

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

#### COUNTRY

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

							CHAIN				
II.	Health information	l		II.a.	Certificate reference No		II.b.				
		(iv)	vine	and capr	ine animals affected with classical	scra	pie are killed and destroyed;				
		de H	the feeding to ovine and caprine animals of meat-and-bone meal or greaves, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health (OIE), of ruminant origin has been banned and effectively enforced in the whole country for a period of at least the preceding seven years;								
	(b)	originate from h	n holdings where no official restrictions are imposed due to a suspicion of TSE;								
	(c)		holdings where no case of classical scrapie has been diagnosed during the perioding seven years or, following the confirmation of a case of classical scrapie:								
		sl	augh arryir	htered, e ng at lea	caprine animals on the holding had accept for breeding rams of the AF ast one ARR allele and no VRC st one ARR allele;]	RR/A	RR genotype, breeding ewes				
		do si m aa A	estro ince ionito ccoro nnex	oyed, and the date oring, ind dance w x X to Re	n which classical scrapie was of the holding has been subjected of confirmation of the last classic cluding testing with negative resith the laboratory methods set of gulation (EC) No 999/2001, of all f 18 months, except ovine animals	for a al so ults out in of th	a period of at least two years crapie case to intensified TSE for the presence of TSE in n point 3.2 of Chapter C of the following animals which are				
		_	- 4	animals v	which have been slaughtered for hi	ımar	n consumption; and				
		_			which have died or been killed on the framework of a disease eradica						

# Notes

#### Part I:

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

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

#### COUNTRY

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

II.	Health information	II.a.	Certificate reference No	II.b.
Part	: II:			
( <sup>1a</sup> )	OJ L 300, 14.11.2009, p. 1.			
(1b)	OJ L 54, 26.2.2011, p. 1.			
(²)	Delete as appropriate.			
( <sup>3</sup> )	OJ L 147, 31.5.2001, p. 1.			
(4)	OJ L 172, 30.6.2007, p. 84.			
_	The signature and the stamp must be in a d	ifferent co	olour to that of the printing.	
-	Note for the person responsible for the consand must accompany the consignment until			e is only for veterinary purposes
Offic	cial veterinarian/Official inspector			
	Name (in capital letters):		Qualification	and title:
	Date:		Signature:	
	Stamp:			

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

#### COUNTRY:

#### Veterinary certificate to EU

		•					rotormary continua				
	l.1.	Consignor		1.2.	Certificate referen	nce No	I.2.a.				
		Name		1.3.	Central competer	nt authority					
		Address		1.4.	Local competent	authority					
		Tel.									
	1.5.	Consignee		1.6.							
ent		Name			Name						
mug		Address			Address						
nsi											
ğ		Postcode			Postcode						
che		Tel.			Tel.						
spat	1.7.	Country ISO code I.8. Region of origin	Code	1.9.	Country of destination	ISO code	I.10. Region of destination	Code			
of d			I		destination						
ails	111	Place of origin		112	Place of destinati	on					
Det		- Naccon Singin			r idoo or doomida						
Part I : Details of dispatched consignment		Name Approval number					Custom warehouse				
ď		Address			Name		Approval number				
		Name Approval number			Address						
		Address									
		Name Approval number		Postcode							
		Address									
	I.13.	Place of loading		1.14.	Date of departure	)					
	115	Means of transport		116	Entry BIP in EU						
	1.10.	mound of dunisport		1.10.	Linay 511 111 20						
		Aeroplane ☐ Ship ☐ Railway w	agon 🗖								
		Road vehicle Other		1.17.							
		Identification									
		Documentation references									
	I.18.	Description of commodity				I.19. Comm	nodity code (HS code)				
							I.20. Quantity				
	I.21.	Temperature of product					I.22. Number of pa	ckages			
		Ambient ☐ Chilled [	]		Frozen $\Box$	<u> </u>					
	1.23.	Seal/Container No					I.24. Type of packa	iging			

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

1.25.	Commodities cert	ified for:						
	Animal feedingstu	ff 🗆	Manufactu	re of petfood $\square$	Technical us	Technical use □		
1.26.	. For transit through EU to third country			I.27. For import or a	27. For import or admission into EU			
	Third country	ISO co	de					
1.28.	Identification of th	e commodities	Approval number	of establishments				
(Sci	Species entific name)	Nature of commodity	Manufacturing plant	Number of packages	Net weight	Batch number		

#### COUNTRY

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

							useu	as ieeu i	iateriai or io	ı us	es outside the leed th	alli
	II.	Health inf	ormation			II.a. Cert	ificate refe	erence No		II.	b.	
		the Europe (EU) No	ean Parliament	t and of and in	f the Co n partic	uncil ( <sup>1a</sup> ), a ular Chapt	ind in part er I of A	ticular Arti Annex XIV	cle 10 thereo	of, a	ntion (EC) No 1069/2009 and Commission Regulat certify that the hydrolys	ion
ion	II.1.	consists o	f hydrolysed p	orotein/d	dicalciun	n phosphate	e/tricalciur	m phosph	ate (²) that s	satis	fy the health requireme	nts
ertificati	II.2.	consists e		nydrolys	ed prot	ein/dicalciui	m phosph	ate/tricalc	ium phospha	ite (	<sup>2</sup> ) not intended for hum	nan
Part II: Certification	II.3.		prepared and s of Regulation (I							nt a	uthority in accordance v	vith
ď	II.4.	has been p	prepared exclu	sively w	ith the f	ollowing ani	imal by-pr	oducts:				
		(²) either	[in the case of dicalcium phosphate derived from defatted bones, carcases and parts of anim slaughtered or, in the case of game, bodies or parts of animals killed, and which are fit for hur consumption in accordance with Union legislation, but are not intended for human consumption commercial reasons;]  [in the case of other materials:									
		(²) or	[in the case or	f other r	material	S:						
	of an					als killed, a egislation, l	and which	are fit fo	or human co	nsu	se of game, bodies or pa mption in accordance v ensumption for commerc	vith
			(²) and/or		slaughte consum	es and the following parts originating either from animals that have been tered in a slaughterhouse and were considered fit for slaughter for human aption following an ante-mortem inspection or bodies and the following parts hals from game killed for human consumption in accordance with Union ion:					nan arts	
					со	nsumption i	n accorda	nce with l		ion,	rejected as unfit for hum but which did not show a als;	
					(ii) he	ads of poult	try;					
					inc		phalanges				g thereof, horns and fe acarpus bones, tarsus a	
					(iv) pig	bristles;						
					(v) fea	athers;]]						
			(²) and/or		blood to slaughte	humans or erhouse af ption follov	animals o ter havin	obtained fi ig been	om animals to considered	that I	ase communicable throu have been slaughtered i for slaughter for hum n accordance with Un	n a nan
			(²) and/or	-	consum		ding deg	reased be			ducts intended for hum d centrifuge or separa	
			(²) and/or		are no problem	onger inten	ided for hacturing or	uman con r packagin	sumption for	con	ucts of animal origin, wh nmercial reasons or due r defects from which no r	to to

**Health information** 

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

#### COUNTRY

II.

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

	(²) and	d/or [-	petfood and feedingstuffs of animal origin, or feedingstuffs containing animal by- products or derived products, which are no longer intended for feeding for commercial reasons or due to problems of manufacturing or packaging defects or other defects from which no risk to public or animal health arises;]]
	(²) and	d/or [-	blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk originating from live animals that did not show signs of any disease communicable through that product to humans or animals;]]
	(²) and	d/or [-	aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of diseases communicable to humans or animals;]]
	(²) and	d/or [-	animal by-products from aquatic animals originating from plants or establishments manufacturing products for human consumption;]]
	(²) and	d/or [-	the following material originating from animals which did not show any signs of disease communicable through that material to humans or animals:
			(i) shells from shellfish with soft tissue or flesh;
			(ii) the following originating from terrestrial animals:
			<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;]]
II.5. the hy	drolysed pr	otein/dicalciur	n phosphate/tricalcium phosphate (²):
	(a)	CONSUMPT particular the	and and packaged in packaging which bear labels indicating 'NOT FOR HUMAN TON' and was stored and transported under satisfactory hygiene conditions, and in a wrapping and packaging took place in a dedicated room, and only preservatives der Union legislation were used; and
(²) eitl	her [(b)		of hydrolysed protein, was produced by a process involving appropriate measures to ntamination of raw Category 3 material.
		produced in	of hydrolysed proteins entirely or partly derived from ruminants hides and skins, was a processing plant dedicated only to hydrolysed proteins production, using a process preparation of the raw Category 3 material by brining, liming and intensive washing
		temp	exposure of the material to a pH of more than 11 for more than 3 hours at a perature of more than 80 °C and subsequently by heat treatment at a temperature of than 140 °C for 30 minutes at more than 3,6 bar; or
			exposure of the material to a pH of 1 to 2, followed by a pH of more than 11, followed heat treatment at a temperature of more than 140 °C for 30 minutes at 3 bar.]
(²) or	[(b)	in the case of	of dicalcium phosphate, was produced by a process that:
		and	ares that all Category 3 bone-material is finely crushed and degreased with hot water treated with dilute hydrochloric acid (at a minimum concentration of 4 % and a pH of than 1,5) over a period of at least two days,
			wed by a treatment of the obtained phosphoric liquor with lime, resulting in a ipitate of dicalcium phosphate at pH 4 to 7, and

Certificate reference No

#### COUNTRY

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

	Health information									uses outside the reed chain
II.	Health ini	rormation				II.a.	Certificate r	eference inc		II.b.
			(iii)				precipitate, v een 30 °C ar		temperature of	of 65 °C to 325 °C and an end
	(²) or	[(b)	in the c	case of tr	ricalciur	m pho	sphate, was	produced b	y a process er	suring:
			(i)				one-material ss than 14 m		ushed and deg	reased in counter-flow with hot
			(ii)	the conf	ntinuous	cook	ing with stea	m at 145 °C	during 30 mir	nutes at 4 bars,
			(iii)	the sep centrifug	•		ne protein bi	roth from th	ne hydroxyapa	atite (tricalcium phosphate) by
			(iv)	the grad 200 °C.		n of t	he tricalciun	n phosphate	e after drying	in a fluidised bed with air at
(²) [II.6.	the hydroly	ysed pro	otein/dic	alcium pl	hospha	ate/tric	alcium phos	phate (²) de	scribed above	
	(²) either	[is der	ived fror	m other r	ruminar	nts tha	ın bovine, ov	ine or capri	ne animals.]]	
	(²) or	[is der	ived fror	m bovine	e, ovine	or ca <sub>l</sub>	prine animals	s and does	not contain an	d is not derived from:
		(²) eith	ier	continuo	ously r	reared	and slaugl	htered in a		derived from animals born, region classified as posing a EC.]]
		(²) or		. , .					n point 1 of ent and of the	Annex V to Regulation (EC) Council (3);
				ar sla ac	inimals, laughtei iccordar	exce red in nce wi	pt from tho a country	se animals or region cl	that were b	s of bovine, ovine or caprine orn, continuously reared and osing a negligible BSE risk in C (4), in which there has been
				ar tis ca th cla	nimals of ssue by avity, or hat wer	which mear by m e bor as	have been has of an elor neans of gas on, continuo posing a	killed, after s ngated rod-s injected into usly reared	stunning, by la shaped instrun o the cranial c d and slaught	om bovine, ovine or caprine ceration of the central nervous nent introduced into the cranial avity, except for those animals ered in a country or region accordance with Decision
II.7.	the hydroly	ysed pro	otein/dic	alcium p	hospha	ate/tric	alcium phos	phate (²) de	scribed above	:
	(²) either			tain milk ls, other t				e or caprine	animal origin	or is not intended for feed for
	(²) or						ovine or cap the milk or			s intended for feed for farmed
		(a)					caprine anin		have been ke	ot continuously since birth in a
			(i)	classica	al scrap	oie is c	compulsorily	notifiable;		
			(ii)	an awar	reness,	, surve	eillance and i	monitoring s	system is in pla	ace for classical scrapie;
			(iii)				oply to holdir nation of clas			mals in the case of a suspicion

#### COUNTRY

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

II.	Health information	II.a. Certificate reference No II.b.
	(iv) ovine a	d caprine animals affected with classical scrapie are killed and destroyed;
	in the of rum	ng to ovine and caprine animals of meat-and-bone meal or greaves, as defined rrestrial Animal Health Code of the World Organisation for Animal Health (OIE), ant origin has been banned and effectively enforced in the whole country for a at least the preceding seven years;
	(b) originate from	ldings where no official restrictions are imposed due to a suspicion of TSE;
		oldings where no case of classical scrapie has been diagnosed during the period seven years or, following the confirmation of a case of classical scrapie:

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

## Notes

# Part I:

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

ANNEX II
Document Generated: 2023-08-31

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

#### COUNTRY

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

п.	Health information	II.a. Certificate reference No	II.D.						
	<ul> <li>Nature of commodity: specify if hydrolysed protein, dicalcium phosphate or tricalcium phosphate.</li> </ul>								
	<ul> <li>Manufacturing plant: provide the registration number of treatment/processing establishment.</li> </ul>								
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 147, 31.5.2001, p. 1.								
(4)	OJ L 94, 1.4.2006, p. 28.								
-	The signature and the stamp must be in a diff	erent 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 must accompany the consignment until it reaches the border inspection post of the point of entry into the European Union.</li> </ul>								
Offic	ial veterinarian/Official inspector								
	Name (in capital letters):	Qualification a	and title:						
	Date:	Signature:							
	Stamp:								
1									

(6) Chapter 18 is replaced by the following:

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

COU	INIT	DV.	
	ואוי	RI.	

#### Veterinary certificate to EU

		•					
	l.1.	Consignor	1.2.	Certificate refere	ence No	I.2.a.	
		Name	1.3.	Central compete	ent authority		
		Address	1.4.	Local competent	t authority		
		Tel.					
	1.5.	Consignee	1.6.	Person responsi	ible for the loa	d in EU	
ent		Name		Name			
gnm		Address		Address			
onsi							
ğ		Postcode		Postcode			
tch		Tel.	-	Tel.			
Part I : Details of dispatched consignment	1.7.	Country ISO code I.8. Region of Code of origin origin	1.9.	Country of destination	ISO code	I.10. Region of destination	Code
of							
tails	l.11.	Place of origin	I.12.	Place of destina	tion		
: De		·					
Ĭ,		Name Approval number				Custom warehouse	
č		Address		Name		Approval number	
		Name Approval number		Address			
		Address					
		Name Approval number		Postcode			
		Address					
	I.13.	Place of loading	I.14.	Date of departur	re		
_	I.15.	Means of transport	I.16.	Entry BIP in EU			
				,			
		Aeroplane ☐ Ship ☐ Railway wagon ☐					
		Road vehicle  Other	l.17.	Number(s) of CI	TES		
		Identification					
		Documentation references					
	I.18.	Description of commodity			I.19. Comm	odity code (HS code)	
						05.07	
						I.20. Quantity	
	I.21.	Temperature of product				I.22. Number of page	kages
		Ambient ☐ Chilled ☐		Frozen D	]		
	1.23.	Seal/Container No				I.24. Type of package	ging

1.25.	Commodities certified for:				
	Further process		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 commoditi		/al number	of establishments	
	Species (Scientific name)	Manufacturing	plant	Net weight	Batch number

#### 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 inf	ormation			II.a. Certificate reference No	II.b.				
		the Europ particular	ean Parliame Chapter II of	ent and of Annex XIV	the Co	eclare that I have read and understood Regul buncil ( <sup>1a</sup> ), and Commission Regulation (EU o, and certify that the horns and horn product meal ( <sup>2</sup> ) described above	No 142/2011	(1b), and in			
_	II.1.	originate from animals									
tificatio		(²) either	either [that were slaughtered in a slaughterhouse, after undergoing ante-mortem inspection, and were fit, as a result of such inspection, for slaughter for human consumption;]								
Part II: Certification		(²) or [that did not show clinical signs of any disease communicable through that product to humans of animals;]									
ď	II.2.	horns, horn products, hooves and hoof products must have undergone a heat treatment for one hour at a core temperature of at least 80 °C;									
	II.3.	horns mus	t have been r	removed wit	thout of	pening the cranial cavity;					
	II.4.	at any sta contamina		essing, stor	rage o	r transport every precaution must have be	en taken to a	avoid cross-			
	II.5.	the horns packed:	and horn pr	oducts, exc	cluding	horn meal, and hooves and hoof products,	excluding hoof	meal, were			
		(²) either	[in new pacl	kaging or co	ontaine	rs;]					
		(²) or	[in vehicles authority;]	or bulk con	ntainers	disinfected prior to loading using a product	approved by th	e competent			
			'NOT FOR H			ked so as to indicate the type of the animal by IAL CONSUMPTION' and the name and addi					
	(²)[II.6.	The horns above	and horn pro	oducts, excl	luding h	norn meal, and hooves and hoof products, exc	cluding hoof me	eal described			
		(²) either	[is derived f	rom other ru	uminan	ts than bovine, ovine or caprine animals.]]					
		(²) or	[is derived f	rom bovine,	ovine	or caprine animals and does not contain and is	not derived fro	m:			
			(²) either	continuous	sly rear	and caprine materials other than those do red and slaughtered in a country or region class redance with Decision 2007/453/EC.]					
			(²) or			risk material as defined in point 1 of An 001 of the European Parliament and of the Co		ulation (EC)			
				ani sla acc	imals, aughter cordan	ally separated meat obtained from bones of except from those animals that were borned in a country or region classified as positive with Commission Decision 2007/453/EC (nous BSE case,	i, continuously ng a negligible	reared and BSE risk in			
		(c) animal by-product or derived product obtained from bovine, ovine or ca animals which have been killed, after stunning, by laceration of the central ne tissue by means of an elongated rod-shaped instrument introduced into the cavity, or by means of gas injected into the cranial cavity, except for those an that were born, continuously reared and slaughtered in a country or classified as posing a negligible BSE risk in accordance with De 2007/453/EC.]]]									

#### 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.							
Not	es							
Par	Part I:							
_	<ul> <li>Box reference I.6: Person responsible for the consignment in the European Union: this box is required to be filled in only if it is a certificate for a commodity to be transited through the European Union; it may be filled in if the certificate is for a commodity to be imported into the European Union.</li> </ul>							
-	<ul> <li>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.</li> </ul>							
-	Box reference I.12: Place of destination: this be in transit must only be stored in free zones, free			transit commodity. Products				
-	Box reference I.15: Registration number (railwainformation is to be provided in the event of unl			per (aircraft) or name (ship);				
-	Box reference I.23: for bulk containers, the con	tainer number and the seal nu	mber (if applicabl	e) must be given.				
_	Box reference I.25: technical use: any use other	er than for animal consumption	l.					
_	Box reference I.26 and I.27: fill in according to	whether it is a transit or an imp	oort certificate.					
_	Box reference I.28: Nature of commodity.							
Par	t II:							
( <sup>1a</sup> )	OJ L 300, 14.11.2009, p. 1.							
(1b)	OJ L 54, 26.2.2011, p. 1.							
(2)	Delete as appropriate.							
(3)	Type of product: horns, horn products, hooves,	, hoof products.						
(4)	OJ L 147, 31.5.2001, p. 1.							
( <sup>5</sup> )	OJ L 172, 30.6.2007, p. 84.							
_	The signature and the stamp must be in a diffe	rent colour to that of the printing	ng.					
_	Note for the person responsible for the consignand must accompany the consignment until it Union.							
Offic	cial veterinarian/Official inspector							
	Name (in capital letters):		Qualification and	i title:				
	Date:		Signature:					
	Stamp:							

(7) Chapter 20 is replaced by the following:

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319. (See end of Document for details)

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

								-				
COL	JNTRY	<b>/</b> :								Vete	rinary certificat	te to EU
	l.1.	Consignor					1.2.	Certificate referen	nce No	1.2	2.a	
		Name					1.3.	Central competent authority				
		Address					1.4.	Local competent	authority			
		Tel.										
	1.5.	Consignee						Person responsib	le for the loa	ad in El	J	
ent	Name						Name					
ŭ L		Address						Address				
nsiç												
oo p		Postcode						Postcode				
che		Tel.						Tel.				
spat	1.7.		ISO code	1.8.	Region of	Code	1.9.	Country of destination	ISO code	I.10.	Region of destination	Code
of di		of origin		I	origin	I		destination		I	destination	I
sils	111	Place of origi	in				112	Place of destinati				
Deta	1.11.	Flace of ong	""				1.12.	riace of destinati	OII			
Part I : Details of dispatched consignment		Name		Annroy	val number					Cust	om warehouse	
Pal		Address		Дрио	varriamber			Name			roval number	_
		Name		Approx	val number			Address		, (pp.	oval nambol	
		Address		прріо	varriambor			Addiess				
		Name		Approv	val number			Postcode				
		Address										
	I.13.	Place of load	dina				1.14.	Date of departure	<u> </u>			
								<u> </u>				
	I.15.	Means of tra	nsport				I.16.	Entry BIP in EU				
		_	_	_		_						
		Aeroplane 🗆			Railway wa	agon 🏻						
		Road vehicle		rЦ			I.17.					
		Identification										
		Documentati										
	I.18.	Description of	of commodit	ty					I.19. Comn	nodity o	code (HS code)	
								L		1.00	Overette	
		_									Quantity	
	1.21.	Temperature	of product						ı	1.22	Number of page	ckages
		Ambient $\square$			Chilled C	]		Frozen				
	1.23.	Seal/Contain	ier No							1.24	Type of packa	ging

1.25.	Commodities certified for:						
	Technical use $\square$						
1.26.	For transit through EU to third	d country	I.27. For import or admission into EU				
	Third country	ISO code					
I.28.	B. Identification of the commodities  Approval number of establishments						
	Species (Scientific name)	Manufacturing plant	Net weight	Batch number			

#### COUNTRY

Part II: Certification

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

					vet	devices, in vitro diagnostics medi erinary purposes, laboratory reage			
II.	Health	n infor	matio	n	II.a.	Certificate reference No	II.b.		
DEC	CLARATION								
tran	I, the undersigned, declare that the intermediate product referred to above is intended to be imported by me into or to be transited through the European Union and satisfies the definition of an intermediate product provided for in point 35 of Annex I to Commission Regulation (EU) No 142/2011 ( <sup>1a</sup> ), and in particular that:								
(1)	it is intended	for the	e man	nufacture of:					
	(²) either	[-	medi	cinal products,]					
	(²) and/or	[-	veter	rinary medicinal products	,]				
	(²) and/or	[-	medi	cal devices for medical a	nd vet	erinary purposes,]			
	(²) and/or	[-	activ	e implantable medical de	vices,]				
	(²) and/or	[-	in vit	ro diagnostic medical dev	vices fo	or medical and veterinary purposes,]			
	(²) and/or	[-	labor	ratory reagents,]					
	(²) and/or	[-	cosm	netic products;]					
(2)	directly or as or transform into service active impla	s a cor ation s as a m ntable	npone uch a nedicir medi	ent of a product intended is mixing, coating, assem nal product, veterinary mi ical devices, an in vitro	for tha bling o edicina diagn	nave been sufficiently completed in it purpose, except for the fact that it in or packaging to make it suitable for pla al product, medical device for medical sostic medical device for medical ar- tion legislation (1b) applicable to those	equires further manufacturing acing on the market or putting I and veterinary purposes, an nd veterinary purposes or a		
(3)	it has been o	derived	from:	:					
	(²) either	[-				ed from animals submitted to an ille 6/22/EC ( <sup>2a</sup> ) or in Article 2(b) of Coun			
	(²) and/or	[-	and v		onsum	ghtered or, in the case of game, bod ption in accordance with Union legisla reasons;]			
	(²) and/or	[-	slaug morte	ghterhouse and were co	onsider s and	originating either from animals that red fit for slaughter for human con the following parts of animals fr ion legislation:	sumption following an ante-		
			(i)			of animals which are rejected as un tion, but which did not show any signs			
			(ii)	heads of poultry;					
			(iii)			trimmings and splitting thereof, ho nd metacarpus bones, tarsus and m			
			(iv)	pig bristles;					
			(v)	feathers;]					

COUNTRY

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

He	alth info	rmation II.a. Certificate reference No II.b.							
(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-morten inspection in accordance with Union legislation;]							
(²) and/or	[-	animal by-products arising from the production of products intended for human consumption, includin degreased bone, greaves and centrifuge or separator sludge from milk processing;]							
(²) and/oi	[-	products of animal origin, or foodstuffs containing products of animal origin, which are no longe intended for human consumption for commercial reasons or due to problems of manufacturing o 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 the 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:							
		<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 invertebrates other than species pathogenic to huma or animals;]							
(²) and/or	[-	animals and parts thereof of the zoological orders of Rodentia and Lagomorpha, except Category material as referred to in Article 8(a)(iii), (iv) and (v) and Category 2 material as referred to Article 9(a) to (g) of Regulation (EC) No 1069/2009;]							
(²) and/or	[-	products derived from or generated by:							
		<ul> <li>aquatic animals, and parts of such animals, except sea mammals, which did not show any sign of disease communicable to humans or animals,</li> </ul>							

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;]

#### COUNTRY

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

II.	Health Infori			n	II.a.	Certificate reference i	NO	II.D.				
	(²) and/or [- animals and parts of animals, other than those referred to in Article 8 or Article 10 of Regulatio No 1069/2009,											
			<ul> <li>that died other than by being slaughtered or killed for human consumption, including a killed for disease control purposes;</li> </ul>									
			(ii)	foetuses;								
			(iii)	oocytes, embryos and	semer	which are not destine	d for breeding purp	oses; and				
			(iv)	dead-in-shell poultry;]								
	(²) and/or	[-	anim	nal by-products other tha	n Cate	egory 1 material or Cate	egory 3 material;]					
(4)	its outer packaging is labelled 'FOR MEDICINAL PRODUCTS / VETERINARY MEDICINAL PRODUCTS / MEDICAL DEVICES FOR MEDICAL AND VETERINARY PURPOSES / ACTIVE IMPLANTABLE MEDICAL DEVICES / IN VITRO DIAGNOSTIC MEDICAL DEVICES FOR MEDICAL AND VETERINARY PURPOSES / LABORATORY REAGENTS / COSMETIC PRODUCTS ONLY' and it is not intended to be diverted at any stage within the European Union for any other use;											
(5)	the consignment will be transported directly to the place of destination in the European Union as indicated under point I.12 of this declaration, that is:											
	(²) either	[an establishment or plant for the production of medicinal products, veterinary medicinal products, medical devices for medical and veterinary purposes, active implantable medical devices, in vitro diagnostic medical devices for medical and veterinary purposes, laboratory reagents or cosmetic products, which has been registered in accordance with Article 23 of Regulation (EC) No 1069/2009],										
	(²) or	No	n establishment or plant which has been approved in accordance with Article 24(1)(i) of Regulation (EC) to 1069/2009, from where they may only be dispatched to an establishment or plant referred to in the receding indent of this point.]									
Note	Notes											
_	Box reference I.19: use appropriate Harmonised System (HS) code in accordance with Commission Decision 2007/275/EC of 17 April 2007 concerning lists of animals and products to be subject to controls at border inspection posts in accordance with Council Directives 91/496/EEC and 97/78/EC (OJ L 116, 4.5.2007, p.9)											
_	Box reference I.25: technical use: any use other than for animal consumption.											
( <sup>1a</sup> )	OJ L 54, 26.2.2011, p. 1.											
( <sup>1b</sup> )	Directive 2001/82/EC of the European Parliament and of the Council of 6 November 2001 on the Community code relating to veterinary medicinal products (OJ L 311, 28.11.2001, p. 1), Directive 2001/83/EC of the European Parliament and of the Council of 6 November 2001 on the Community code relating to medicinal products for human use (OJ L 311, 28.11.2001, p. 67), Council Directive 93/42/EEC of 14 June 1993 concerning medical devices (OJ L 169, 12.7.1993, p. 1) and Directive 98/79/EC of the European Parliament and the Council of 27 October 1998 on in vitro diagnostic medical devices (OJ L 331, 7.12.1998, p. 1), Regulation (EC) No 1223/2009 of the European Parliament and of the Council of 30 November 2009 on cosmetic products (OJ L 342, 22.12.2009, p. 59), as appropriate.											
( <sup>2</sup> )	Delete as appropriate.											
( <sup>2a</sup> )	OJ L 125, 23.5.1996, p. 3.											
( <sup>2b</sup> )	OJ L 125, 23.5.1996, p. 10.											
The importer												
	Name (in capi	tal le	tters):	:		Address:						
	Date:						Signature:					

- (1) OJ L 147, 31.5.2001, p. 1.
- (2) OJ L 300, 14.11.2009, p. 1.
- (3) Commission Decision 2007/453/EC of 29 June 2007 establishing the BSE status of Member States or third countries or regions thereof according to their BSE risk (OJ L 172, 30.6.2007, p. 84).
- (4) Commission Regulation (EU) 2016/1396 of 18 August 2016 amending certain Annexes to Regulation (EC) No 999/2001 of the European Parliament and of the Council laying down rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies (OJ L 225, 19.8.2016, p. 76).
- (5) Commission Regulation (EU) No 142/2011 of 25 February 2011 implementing Regulation (EC) No 1069/2009 of the European Parliament and of the Council laying down health rules as regards animal by-products and derived products not intended for human consumption and implementing Council Directive 97/78/EC as regards certain samples and items exempt from veterinary checks at the border under that Directive (OJ L 54, 26.2.2011, p. 1).
- (6) http://www.oie.int/international-standard-setting/terrestrial-code/access-online/
- (7) Council Regulation (EEC) No 2658/87 of 23 July 1987 on the tariff and statistical nomenclature and on the Common Customs Tariff (OJ L 256, 7.9.1987, p. 1).
- (8) Commission Regulation (EU) No 630/2013 of 28 June 2013 amending the Annexes to Regulation (EC) No 999/2001 of the European Parliament and of the Council laying down rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies (OJ L 179, 29.6.2013, p. 60).
- (9) Commission Decision 2007/275/EC of 17 April 2007 concerning lists of animals and products to be subject to controls at border inspection posts under Council Directives 91/496/EEC and 97/78/ EC (OJ L 116, 4.5.2007, p. 9).

# **Changes to legislation:**

There are currently no known outstanding effects for the Commission Regulation (EU) 2019/319.