	I.1. Consignor				I.2. IMSOC Reference		
	Name						
	Address				I.2.a. Local Reference		
	Country		ISO Code				
	I.5. Consignee				I.3. Central co	mpetent authority	
님	Name				I.4. Local com	petent authority	
ne	Address						
E	Country		ISO Code				
ŝ	I.7. Country of orig	gin	IS	SO Code	I.9. Country of	fdestination	ISO Code
8	1.7. country of ong	5				destination	
Part I : Details of consignment	I.8. Region of origi	in	Co	ode	I.10. Region of	destination	
S	L11 Dlaga of Diam				I.12. Place of d		
ail	I.11. Place of Dispa	aten				lesunation	
<u></u>	Name Address				Name Address		
귀	Approval Number	r			Approval Nui	mher	
뒨	Country	-	ISO Code		Country		SO Code
B					,		
	I.13. Place of Load	ing			I.14. Date and time of departure		
	Name						
	Address						
	Approval Number	r					
	Country		ISO Code				
	I.15. Means of Tra	nsport			I.16 Entry Poin	nt	
	Mode	International	Identification				
		transport document					
	I.18. Transport co	nditions				nying documents	
	Frozen 🗆	Controlled	Ambient 🛛 🛛 Cl	hilled \Box	Accompanyin reference	g document	
		temperature 🛛			Date of issue		
					Country		
					Place of issue		
	I.19. Container No	/ Seal No					
	I.20. Certified as						
	Production of petf	Food []	Animal Feedingstuff	Technical use			
	Production of peti	.000 🗀	Animai Feedingstun		rechnical use		
	I.21. For transit th	rough a third coun	try		I.22. For transit through Member State(s)		
	Country	-	ISO Code		Country ISO Code		
	Country EU Exit		BCP code				
	Authority						
	EU Entry		BCP code				
	Authority		88				
	I.25. Total net weig	-			I.25. Total gross weight		
	I.28. Description o	f consignment					
	1. 23 RESIDUES AI	ND WASTE FROM 7	THE FOOD INDUSTRIE	ES; PREPARED AN	NIMAL FODDER	R	
	2301 Flours me	eals and nellets of i	meat or meat offal of	fish or of crusta	reans molluscs	s or other aquatic invertebrat	es unfit for human
	consumption; g	reaves				-	
		-	s, of fish or of crustace		-		
			ellets, of fish or of cru				Detah susuah su
	#1. Commodity	Nature	of commodity	Manufacturing p	llant	Net weight	Batch number
	Species						

				-		-		0 1		
	II. Health informa	ation								
	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 and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011, and in particular Section 1 of Chapter II of Annex X, and Chapter I of Annex XIV thereto and certify that:									
				rotein or pr n consumpt		ibed above contains	exclusive	ely process	ed animal protein	
ificati	(a					d in an establishment or plant approved and supervised by the dance with Article 24 of Regulation (EC) No 1069/2009, and				
Cer	(b)	has been prepared exclusively with the following animal by-products:							
Part II: Certification	(1)		parts of ani	mals killed, ed EU law, k	animals slaughtered and which are fit fo out are not intended	r human	consumpti	ion in accordance	
	(1)		been slaugh human con following p	ntered in a s sumption fo	owing parts originati laughterhouse and v ollowing an ante-mon nals from game killeo ned EU law:	vere cons rtem insp	idered fit i ection or h	for slaughter for oodies and the	
					for human	bodies and parts of consumption in acco not show any signs of	ordance v	vith retain	ed EU law, but	
				(ii)	heads of po	oultry;				
					feet, includ	kins, including trimi ing the phalanges an metatarsus bones;				
				(iv)	pig bristles	• ,				
				(v)	feathers;]					
	(1)		through blo slaughtered	ood to huma l in a slaugh consumptio	ch did not show any ins or animals, obtain iterhouse after havir in following an ante-	ned from 1g been c	animals th onsidered :	nat have been fit for slaughter	
	(1)	□ and/or	human con	sumption, i	rising from the prod ncluding degreased l milk processing;]				
	(1)		which are r or due to p	no longer in coblems of r	rigin, or foodstuffs co tended for human co nanufacturing or pac or animal health ar	onsumpti ckaging c	on for com	mercial reasons	
	(1)		from live a	nimals that	l, feathers, hair, horr did not show signs o s or animals;]				
	(1)		-		parts of such anima liseases communical	-			
	(1)	□ and/or			rom aquatic animals products for human			stablishments or	
(1)										
				(i)	shells from	shellfish with soft ti	ssue or fl	esh:		
				(ii)		ng originating from t		l animals:		
					-	hatchery by-produc	ts,			

(GB) Processed animal protein, other than those derived from farmed insects, not intended for human consumption including mixtures and products other than perfood containing such protein GBHC078E.

BO (C) No 1069/2009:] and (c) has been subjected to the following processing standard: (1) • either [heating to a core temperature of more than 133°C for at least 30 bars produced by saturated steam, with a particle size prior to processing of not more than 50 millimetres;] (1) • or [in the case of non-mammalian protein other than fishmeal, the processing method 1-2-3-4-5-7 (1) • or [in the case of fishmeal the processing method 1-2-3-4-5-7 (1) • or [in the case of procise blood, the processing method 1-2-3-4-5-7 (1) • or [in the case of procise blood, the processing method 1-2-3-4-5-7 (1) • or [in the case of procise blood, the processing method 1-2-3-4-5-7 (10) • or [in the case of procise blood, the processing method 1-2-3-4-5-7 (11) • or [in the case of procise blood, the processing method 1-2-3-4-5-7 (11) • or [in the case of procise blood, the processing method 1-2-3-4-5-7 (12) (11) • or [in the case of procise blood, the processing method 1-2-3-4-5-7 (11) • or [in the case of procise blood, the processing method 1-2-3-4-5-7 (12) (12) [in the case of procise blood, the processing theatore processing sthot the case of procise blo	EUROPEAN UNION products other than petfood containing such protein GBHC078								
Image: Constraint of the second s	II. Health info	rmation							
Image: Constraint of the second s									
Upper Difference (ii) day-old chicks killed for commercial reasons.] (1) □ and/or I - squatic and terrestrial invertebrates other than species pathogenic to human or animals and other than insects.] (1) □ and/or I - anguatic and terrestrial invertebrates other than species pathogenic to human or animals and other than insects.] (1) □ and/or I - animals and parts thereof of the zoological orders of Rodentia and Lagomorpha, except Category 1 material as referred to in Article %(a)(ii), (iv) and (v) and (v) and (a Category 2 material as referred to in Article %(a)(iii), (iv) and (v) and (a Category 2 material as referred to a tharticle %(a)(iii), (iv) and (v) and (or a terrestricle %(a)(iii), (iv) and (iii) (iiii) (iii) (iiii) (iii) (iiii) (iii) (iii) (iiii) (iii) (iii) (iii									
Image: Second									
or animals and other than insects;] or animals and parts thereof of the zoological orders of Rodentia and Lagomorpha, except Category 1 material as referred to in Article 9(a)(ii), (iv) and (v) and (c) and (category 2 material as referred to in Article 9(a) to (g) of Regulatio (EC) No 1069/2009;] and (c) has been subjected to the following processing standard: (1) ○ either (hearing to a core temperature of more than 133°C for at least 20 minutes without interruption at a pressure (absolute) of at least 3 hars produced by saturated steam, with a particle size prior to processing of not more than 50 millimetres;] (1) ○ or fin the case of non-mammalian protein other than fishmeal, the processing method 1×2:3+45-7 (f) Corr fin the case of porcine blood, the processing method 1×2:3-45-7 (fin) corr fin the case of porcine blood, the processing method 1×2:3-45-7 (fin) the case of porcine blood, the processing method 1×2:3-45-7 (fin) corr fin the case of porcine blood, the processing method 1×2:3-45-7 (fin) corr fin the case of porcine blood, the processing method 1×2:3-45-7 (fin) the case of porcine blood, the processing method 1×2:3-45-7 (fin) corr fin the case of porcine blood, the processing method 1×2:3-45-7 (fin) the following standards (2): Salmonella: Absence in 25 g n = 5, c = 0, m = 0, M = 0 Enterobacteriacee: n = 5, c - 2, m = 10, M = 300 in 1g; 13. the product has undergone all precautions to avoid recontamination with pathogenic agents after treatment; (1) or (was transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected before use.									
 (1) ○ either Ineating to a core temperature of more than 133°C or at least 20 minutes without interruption at a pressure (absolute) of at least 3 bars produced by saturated steam, with a particle size prior to processing of not more than 50 millimetres.] (1) ○ or [in the case of non-mammalian protein other than fishmeal, the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011.] (1) ○ or [in the case of fishmeal the processing method 1-2-3-4-5-6-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011.] (1) ○ or [in the case of porcine blood, the processing method 1-2-3-4-5-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011.] (1) ○ or [in the case of porcine blood, the processing method 1-2-3-4-5-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011.) where in case of method 7 a heat treatment of at least 80 °C has been applied throughout its substance] II.2. the competent authority examined a random sample immediately prior to dispatch and found it to comply with the following standards (2): Salmonella: Absence in 25 g: n = 5, c = 0, m = 0, M = 0 Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1g; II.3. the product: II.4. the end product: II.4. the end product: II.5. the end product: II.5. the end product was stored in new or sterilised bags.] (1) ○ either [was packed in new or sterilised bags.] (1) ○ either [originates from a country or region, which is classified as posing a negligible BSE risk in accordance with Decision 2007/453/C, and in which there has been no indigenous BSE case, and the animaby products of runniant origin and: (1) ○ either [is derived from acountry or region classified as posing a negligible BSE risk in accordance with	10	(1)	⊔ and/or						
 (1) • either Ineating to a core temperature of more than 133°C or at least 20 minutes without interruption at a pressure (absolute) of at least 3 bars produced by saturated steam, with a particle size prior to processing of not more than 50 millimetres;] (1) • or [in the case of non-mammalian protein other than fishmeal, the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011.] (1) • or [in the case of fishmeal the processing method 1-2-3-4-5-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011.] (1) • or [in the case of porcine blood, the processing method 1-2-3-4-5-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011.] (1) • or [in the case of porcine blood, the processing method 1-2-3-4-5-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011.) (1) • or [in the case of porcine blood, the processing method 1-2-3-4-5-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011.) (1) • or [in the case of porcine blood, the processing method 1-2-3-4-5-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011.) (1) • or [in the case of porcine blood, the processing method 1-2-3-4-5-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011.) (1) • or [in the case of procine blood, the processing method 1-2-3-4-5-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011.] (1) • or [in the case of procine blood, the processing method 1-2-3-4-5-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011.] (1) • or [in the competent authority examined a random sample immed	· Certuicau	(1)	□ and/or	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					
 (1) ○ either Ineating to a core temperature of more than 133°C or at least 20 minutes without interruption at a pressure (absolute) of at least 3 bars produced by saturated steam, with a particle size prior to processing of not more than 50 millimetres.] (1) ○ or [in the case of non-mammalian protein other than fishmeal, the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011.] (1) ○ or [in the case of fishmeal the processing method 1-2-3-4-5-6-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011.] (1) ○ or [in the case of porcine blood, the processing method 1-2-3-4-5-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011.] (1) ○ or [in the case of porcine blood, the processing method 1-2-3-4-5-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011.) where in case of method 7 a heat treatment of at least 80 °C has been applied throughout its substance] II.2. the competent authority examined a random sample immediately prior to dispatch and found it to comply with the following standards (2): Salmonella: Absence in 25 g: n = 5, c = 0, m = 0, M = 0 Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1g; II.3. the product: II.4. the end product: II.4. the end product: II.5. the end product: II.5. the end product was stored in new or sterilised bags.] (1) ○ either [was packed in new or sterilised bags.] (1) ○ either [originates from a country or region, which is classified as posing a negligible BSE risk in accordance with Decision 2007/453/C, and in which there has been no indigenous BSE case, and the animaby products of runniant origin and: (1) ○ either [is derived from acountry or region classified as posing a negligible BSE risk in accordance with		and							
 withou[*] Interruption ar a pressure (absolute) of at least 3 bars produced by saturated steam, with a particle size prior to processing of not more than 50 millimetres;] (1) • or [in the case of non-mammalian protein other than fishmeal, the processing method 1-2-3-4-5-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011;] (1) • or [in the case of fishmeal the processing method 1-2-3-4-5-67 (indicate the processing method) no 142/2011;] (1) • or [in the case of porcine blood, the processing method 1-2-3-4-5-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011;] (1) • or [in the case of porcine blood, the processing method 1-2-3-4-5-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011;] (1) • or [in the case of porcine blood, the processing method 1-2-3-4-5-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011; where in case of method 7 a heat treatment of at least 80 °C has been applied throughout its substance;] II.2. the competent authority examined a random sample immediately prior to dispatch and found it to comply with the following standards (2): Salmonella: Absence in 25 gr. n = 5, c = 0, m = 0, M = 0 Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1g; II.3. the product has undergone all precautions to avoid recontamination with pathogenic agents after treatment; II.4. the end product: (1) • or [was transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected before use,] which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'; II.5. the end product was stored in enclosed storage; (1) □ or [triginates from a country or region, which is classified as posing a negligible BSE risk in accordance with Dec	Fai	(c) has been subjected to the following processing standard:							
method 1-2-3-4-5-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011;] (1) • or [in the case of fishmeal the processing method 1-2-3-4-5-6-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011;] (1) (1) • or [in the case of porcine blood, the processing method 1-2-3-4-5-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011, where in case of method 7 a heat treatment of at least 80 °C has been applied throughout its substance;] II.2. the competent authority examined a random sample immediately prior to dispatch and found it to comply with the following standards (2): Salmonella: Absence in 25 g; n = 5, c = 0, m = 0, M = 0 Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1g; II.3. the product has undergone all precautions to avoid recontamination with pathogenic agents after treatment; II.4 the end product: (1) • either [was packed in new or sterilised bags.] (1) • either [was stored in enclosed storage; (1) • either [originates from a country or region, which is classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC, and in which there has been no indigenous BSE case and] (1) • either [originates from a country or re		(1)	∘ either	without interruption at a pressure (absolute) of at least 3 bars produced by saturated steam, with a particle size prior to processing of not more than 50					
 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011;] (1) • or [in the case of porcine blood, the processing method 1-2-3-4-5-7 (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011, where in case of method 7 a heat treatment of at least 80 °C has been applied throughout its substance;] II.2. the competent authority examined a random sample immediately prior to dispatch and found it to comply with the following standards (2): Salmonella: Absence in 25 g; n = 5, c = 0, m = 0, M = 0 Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1g; II.3. the product has undergone all precautions to avoid recontamination with pathogenic agents after treatment; II.4. the end product: (1) • either [was packed in new or sterilised bags.] (1) • or [was transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected before use.] which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'; II.5. the end product was stored in enclosed storage; (1) • either [originates from a country or region, which is classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC, and in which there has been no indigenous BSE cas and]] (1) • or [originates from a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC, and in which there has been no indigenous BSE cas and]] (1) • oither [is derived from other runniants with meat-and-bone meal and greaves derived from runniants, as defined in the OIE Terrestrial Animal Health Code, has been effectively enforced in that country or region, and] (1) • either [is derived from other runniants than bovine, ovine or caprine animals.] (1) • or [is derived from bovine, ovine or caprine animals and does not contain and is not derived 		(1)	\circ or	method 1-2-3-4-5-7 (indicate the processing method) as set out in					
(indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011, where in case of method 7 a heat treatment of at least 80 °C has been applied throughout its substance;] II.2. the competent authority examined a random sample immediately prior to dispatch and found it to comply with the following standards (2): Salmonella: Absence in 25 g: n = 5, c = 0, m = 0, M = 0 Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1g; II.3. the product has undergone all precautions to avoid recontamination with pathogenic agents after treatment; II.4. the end product: (1) • either (was transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected before use,] which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'; II.5. the end product was stored in enclosed storage; (1) • either [I1.6. the processed animal protein or product described above contains or is derived from animal-by products of ruminant origin and: (1) • either [originates from a country or region, which is classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC, and in which there has been no indigenous BSE case, andI] (1) • 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 anima by-product or derived product were derived from animals born after the date from which the ban on the feeding of ruminan		(1)	◦ or	(indicate the processing method) as set out in Chapter III of Annex IV to					
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 II.3. the product has undergone all precautions to avoid recontamination with pathogenic agents after treatment; II.4. the end product: (1) o either [was packed in new or sterilised bags,] (1) o or [was transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected before use,] which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'; II.5. the end product was stored in enclosed storage; (1) □ [II.6. the processed animal protein or product described above contains or is derived from animal-by products of ruminant origin and: (1) o either [originates from a country or region, which is classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC, and in which there has been no indigenous BSE case, and]] (1) o 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 anima by-product or derived product were derived from animals born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants, as defined in the OIE Terrestrial Animal Health Code, has been effectively enforced in that country or region, and] (1) o either [is derived from other ruminants than bovine, ovine or caprine animals.] (1) o or 									
II.4. the end product: (1) • either [was packed in new or sterilised bags,] (1) • or [was transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected before use,] which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'; II.5. the end product was stored in enclosed storage; (1) □ [II.6. (1) • either [originates from a country or region, which is classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC, and in which there has been no indigenous BSE case and]] (1) • or (originates from a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC, and in which there has been no indigenous BSE case and]] (1) • or (1) • 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 anima by-product or derived product were derived from animals born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants, as defined in the OIE Terrestrial Animal Health Code, has been effectively enforced in that country or region, and] (1) • either [is derived from other ruminants than bovine, ovine or caprine animals.] (1) • or	Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1g;								
(1) ○ either [was packed in new or sterilised bags,] (1) ○ or [was transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected before use,] which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'; II.5. the end product was stored in enclosed storage; (1) □ [II.6. the processed animal protein or product described above contains or is derived from animal-by products of ruminant origin and: (1) □ either [originates from a country or region, which is classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC, and in which there has been no indigenous BSE case and]] (1) ○ 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 anima by-product or derived product were derived from animals born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants, as defined in the OIE Terrestrial Animal Health Code, has been effectively enforced in that country or region, and] (1) ○ either [is derived from other ruminants than bovine, ovine or caprine animals.] (1) ○ or [is derived from other, ovine or caprine animals and does not contain and is not derived	II.3.	-							
 (1) ○ or [was transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected before use,] which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'; II.5. the end product was stored in enclosed storage; (1) □ [II.6. the processed animal protein or product described above contains or is derived from animal-by products of ruminant origin and: (1) ○ either [originates from a country or region, which is classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC, and in which there has been no indigenous BSE case and]] (1) ○ 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 anima by-product or derived product were derived from animals born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants, as defined in the OIE Terrestrial Animal Health Code, has been effectively enforced in that country or region, and] (1) ○ either [is derived from other ruminants than bovine, ovine or caprine animals.] (1) ○ or [is derived from bovine, ovine or caprine animals and does not contain and is not derived 									
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 II.5. the end product was stored in enclosed storage; (1) □ [II.6. the processed animal protein or product described above contains or is derived from animal-by products of ruminant origin and: (1) ○ either [originates from a country or region, which is classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC, and in which there has been no indigenous BSE case and]] (1) ○ 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 anima by-product or derived product were derived from animals born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants, as defined in the OIE Terrestrial Animal Health Code, has been effectively enforced in that country or region, and] (1) ○ either [is derived from other ruminants than bovine, ovine or caprine animals.] (1) ○ or [is derived from bovine, ovine or caprine animals and does not contain and is not derived 	(1) • or [was transported in								
 (1) □ [II.6. the processed animal protein or product described above contains or is derived from animal-by products of ruminant origin and: (1) • either [originates from a country or region, which is classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC, and in which there has been no indigenous BSE case and]] (1) • 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 anima by-product or derived product were derived from animals born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants, as defined in the OIE Terrestrial Animal Health Code, has been effectively enforced in that country or region, and] (1) • either [is derived from other ruminants than bovine, ovine or caprine animals.] (1) • or [is derived from bovine, ovine or caprine animals and does not contain and is not derived 			which bear labels indicating 'NOT FOR HUMAN CONSUMPTION';						
 products of ruminant origin and: (1) • either [originates from a country or region, which is classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC, and in which there has been no indigenous BSE case and]] (1) • 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 anima by-product or derived product were derived from animals born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants, as defined in the OIE Terrestrial Animal Health Code, has been effectively enforced in that country or region, and] (1) • either [is derived from other ruminants than bovine, ovine or caprine animals.] (1) • or [is derived from bovine, ovine or caprine animals and does not contain and is not derived 	II.5.	the end pr	oduct was s	stored in enclosed storage;					
 accordance with Decision 2007/453/EC, and in which there has been no indigenous BSE case and]] o 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 anima by-product or derived product were derived from animals born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants, as defined in the OIE Terrestrial Animal Health Code, has been effectively enforced in that country or region, and] o either [is derived from other ruminants than bovine, ovine or caprine animals.] o or [is derived from bovine, ovine or caprine animals and does not contain and is not derived 	(1) 🗆 [II.6.								
 with Decision 2007/453/EC in which there has been an indigenous BSE case, and the animal by-product or derived product were derived from animals born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants, as defined in the OIE Terrestrial Animal Health Code, has been effectively enforced in that country or region, and] (1) • either [is derived from other ruminants than bovine, ovine or caprine animals.] (1) • or [is derived from bovine, ovine or caprine animals and does not contain and is not derived 	(1)	○ either	accordance with Decision 2007/453/EC, and in which there has been no indigenous BSE case						
(1) • or [is derived from bovine, ovine or caprine animals and does not contain and is not derived	(1)	∘ or	ruminants, as defined in the OIE Terrestrial Animal Health Code, has been effectively						
	(1)	\circ either	[is derived	l from other ruminants than bovine, ovine or caprine animals.]					
nom.	(1) • or [is derived from bovine, ovine or caprine animals and does not contain and is no from:								

(GB) 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 GBHC078E

	JROPEAN	UNION			products other than petfood containing such protein GBHC078E				
	II. Health info	ormation							
		(1)	\circ either	[bovine, ovine and caprine materials other than those derived from animals born, continuously reared and slaughtered in a country or region classified a posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]					
		(1)	o 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;				
Part II: Certification				[(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, in which there has been no indigenous BSE case,				
Dar				(c)	animal by-product or derived product obtained from bovine, ovine or caprine animals which have been killed, after stunning, by laceration of the central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, or by means of gas injected into the cranial cavity, except for those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk in accordance with Decision 2007/453/EC.]]]				
	II.7.	the proces	sed animal	protein or p	roduct described above:				
	(1)	\circ either		t contain milk or milk products of ovine or caprine animal origin or is not intended for farmed animals, other than fur animals.]					
	(1) • 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)		d from ovine and caprine animals which have been kept continuously in a country where the following conditions are fulfilled:				
				(i)	classical scrapie is compulsorily notifiable;				
				(ii)	an awareness, surveillance and monitoring system is in place for classical scrapie;				
				(iii)	official restrictions apply to holdings of ovine or caprine animals in the case of a suspicion of TSE or the confirmation of classical scrapie;				
				(iv)	ovine and caprine animals affected with classical scrapie are killed and destroyed;				
(v) the feeding to ovine and caprine animals of greaves, as defined in the Terrestrial Anima World Organisation for Animal Health (OII been banned and effectively enforced in the				the feeding to ovine and caprine animals of meat-and-bone meal or greaves, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health (OIE), of ruminant origin has been banned and effectively enforced in the whole country for a period of at least the preceding seven years;					
			(b) originate from holdings where no official restrictions are imposed due to a suspicion of TSE;						
			(c)	originate from holdings where no case of classical scrapie has been diagnose during a period of at least the preceding seven years or, following the confirmation of a case of classical scrapie:					
			(1)	 either [all ovine and caprine animals on the holding have been kildestroyed or slaughtered, except for breeding rams of the A genotype, breeding ewes carrying at least one ARR allele an allele and other ovine animals carrying at least one ARR all 					
			(1)	° 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				

EU	JROPEAN U	NION	nimal protein, other than t ded for human consumptic her than petfood containir	hose derived from farmed on including mixtures and ig such protein GBHC078E				
II. Health information								
			•	(EC) No 999/2001, of all of the e age of 18 months, except ov	-			
_			-	animals which have been sla consumption; and	ughtered for human			
Part II: Certification			-	animals which have died or l but which were not killed in eradication campaign.]]	5			
rt II: Cer	II.8.	-	sed animal protein or product descr f non-ruminant origin and is, accore		-			
Pa	(1)	\circ either	[not intended for the production o	f feed for farmed animals, oth	er than fur animals.]			
	(1)(3)	∘ or	[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 Control Post of entry will be provided with the results of the analyses carried out in accordance with the methods set out in Annex VI to Commission Regulation (EC) No 152/2009.]					
	Notes							
	(*) Those co		oject to the transitional import arran zerland and the Faroe Islands.	ngements include: an EU men	nber State; Liechtenstein;			
	EU legislation which has ndrawal) Act 2018).							
	References	to Great Br	itain in this certificate include Char	nel Islands and Isle of Man.				
	Part I:							
	-	Box reference I.6:	Person responsible for the consignment in Great Britain, Channel Islands or Isle of Man: this box is required to be filled in only if it is a certificate for a commodity to be transited through Great Britain, Channel Islands or Isle of Man; it may be filled in if the certificate is for a commodity that is to be imported into Great Britain, Channel Islands or Isle of Man.					
	-	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.					
	-	Box reference I.16:	Do not use this box until the end of	f the transitional staging perio	od.			
	-	Box reference I.19:	use the appropriate HS code: 05 05	; 05 06: 05.07; 05.11: 23 01 or 3	23 09.			
	-	Box reference I.25:	technical use: any use other than f production or manufacturing of pe		ner than fur animals, and the			
	-	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 or Suidae, Pesca, Molla Crustacea. In the case of farmed fis	usca, Crustacea, invertebrates	other than Mollusca and			
	Part II:							

EU	ROPEAN U	INION products of	her than petfood containir	ng such protein GBHC078E							
	II. Health infor	rmation									
	(1)	Delete as appropriate.									
	(2)	Where:									
		n = number of samples to be tested;									
no		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;									
Part II: Certification		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									
I: Cert		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									
Part I		The Person responsible for the load referred to in Box 1.6 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 a border control post of the point of entry into Great Britain, Channel Islands or Isle of Man.									
	-	The signature and the stamp must be in a diffe	erent colour to that of the prir	iting.							
	-	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 Great Britain, Channel Islands or Isl This certificate is only for veterinary purposes and must accompany the consignment until									
		the border control post.	and must accompany the con	isignment until it reaches							
	Certifying Offic Name (in capit Date of signatu Stamp	tal letters)	Qualification and title Signature								