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

ANNEX XV

MODEL HEALTH CERTIFICATES

CHAPTER 3(E)

Health certificate

For flavouring innards for use in the manufacture of petfood, intended for dispatch to or for transit through $\binom{2}{2}$ the European Union

cou	COUNTRY Veterinary certificate to							ate to EU		
	l.1.	Consignor			1.2.	Certificat	e reference	No	1.2.a.	
		Name					competent s	uthority		
		Address				I.3. Central competent authority				
						I.4. Local competent authority				
	1.5.	Tel.				_				
eut	1.5.	Consignee			1.6.		esponsible	for the load	in EU	
틸		Name			Name					
l sig		Address				Address				
8		Postcode			Postcode					
Je		Tel.				Tel.	,			
atc					_				I	
g	1.7.	Country of origin ISO code I.8. Region of	origin	Code	1.9.	Country		ISO code	I.10. Region of destination	Code
5		1	- 1			destination			destination	ı
is is	144	Place of origin			1.40	D . (
Part I: Details of dispatched consignment	1.11.	Place of origin			1.12.	Place of	destination			
=		Name Approval numb	er			Name			Custom warehouse	
at		Address				Address			Approval number	
"		Name Approval numb	er							
						Postcode	9			
		Name Approval numb	er							
	I.13.	Place of loading			1.14.	Date of o	departure			
	l.15.	Means of transport			I.16.	Entry BIF	o in EU			
		Aeroplane ☐ Ship ☐ Railway wagon ☐				-				
		Road vehicle Other O			1.17.					
		Identification			". " / .					
		Documentation references			_					
	I.18.	Description of commodity				I.19. Commodity code (HS code)				
						I.20. Quantity				
	1.21.	Temperature of product				I.22. Number of packages				
		Ambient Chilled Chilled			Froze	n \square		1.22. 1	uniber of packages	
	1.23	Seal/Container No						124 T	no of pookoging	
		- Court Contain Contai						1.24. 1	pe of packaging	
	1.25.	Commodities certified for:								
			hnical use	П						
		Animal reedingston	iiiicai use							
	1.26.	6. For transit through EU to third country			1.27.	For impo	rt or admiss	sion into EU		
		Third country ISO code								
		-								
	1.28.	Identification of the commodities								
		Species Nature of commodit	y A	pproval nu				Net v	weight Batch r	number
		(Scientific name)	-	Mar	ufactu	ring plant			-	

со	UNTRY				Flavouring innards for u	se in the manufacture of petfood		
	II.	Health inf	orn	nation	II.a. Certificate reference No	II.b.		
		and of the	C	ouncil (1a) and in particular Article 8 and 10 there	and understood Regulation (EC) No 1069/2009 of the European Parliament cof, and Commission Regulation (EU) No 142/2011 (^{1b}), and in particular certify that the flavouring innards products described above:			
tion	II.1. consist of animal by-products that satisfy the animal health requirements below;							
ertifica	II.2.	have been	pre	epared including the following animal by-products	which are exclusively:			
Part II: Certification		(2) either [- carcases and parts of animals slaughtered or, in the case of game, bodies or parts of animals killed, and which are human consumption in accordance with Union legislation, but are not intended for human consumption for commeasons;]						
		(2) and/or [- carcases and the following parts originating either from animals that have been slaughtered in a slaughterhouse and wer considered fit for slaughter for human consumption following an ante-mortem inspection or bodies and the following parts of animals from game killed for human consumption in accordance with Union legislation:						
	(i) carcases or bodies and parts of animals which are rejected as unfit for human consumption in accordance was legislation, but which did not show any signs of disease communicable to humans or animals;							
				(ii) heads of poultry;				
	(iii) hides and skins, including trimmings and splitting thereof, horns and feet, including the phalanges and the of metacarpus bones, tarsus and metatarsus bones, of animals other than ruminants;							
	(iv) pig bristles;							
(v) feathers;]								
	(²) and/or [- blood of animals which did not show any signs of disease communicable through blood to humans or animals, o animals other than ruminants that have been slaughtered in a slaughterhouse after having been considered fit for human consumption following an ante-mortem inspection in accordance with Union legislation;]				ng been considered fit for slaughter			
	(²) and/or [- animal by-products arising from the production of products intended for human consumption, including deg greaves and centrifuge or separator sludge from milk processing;]				mption, including degreased bone,			
	(²) and/or [- products of animal origin, or foodstuffs containing products of animal origin, which are no longer inten- consumption for commercial reasons or due to problems of manufacturing or packaging defects or othe 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 product longer intended for feeding for commercial reasons or due to problems of manufacturing or packaging defects from which no risk to public or animal health arises;]								
		(2) and/or	[-	blood, placenta, wool, feathers, hair, horns, hoof any disease communicable through that product		animals that did not show signs of		
	(2) and/or [- aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of diseases conto humans or animals;] (2) and/or [- animal by-products from aquatic animals originating from plants or establishments manufacturing products consumption;]							
(²) and/or [- the following material originating from animals which did not show any signs of disease communicable through to humans or animals:						communicable through that material		
	(i) shells from shellfish with soft tissue or flesh;							

COUNTRY

Flavouring innards for use in the manufacture of petfood

II.a. Certificate reference No	COUNT	RY		Flavouring innards for use in the manufacture of petfood		
	II.	Health inf	ormation	II.a. Certificate reference No	II.b.	
- eggs, - egg by-products, including egg shells; - (iii) day-old chicks killed for commercial reasons.] - (c) and/or [- animal by-products from aquatic or terrestrial invertebrates other than species pathogenic to humans or animals.] - (d) and/or [- animal by-products from aquatic or terrestrial invertebrates other than species pathogenic to humans or animals.] - (e) and/or [- animal by-products from aquatic or terrestrial invertebrates other than species pathogenic to humans or animals.] - (e) and/or [- animal by-products from aquatic or terrestrial invertebrates other than species pathogenic to humans or animals.] - (e) and/or [- animal by-products from aquatic or terrestrial invertebrates other than species pathogenic to humans or animals.] - (e) and/or [- animal by-products with Annex XIII, Chapter III of Regulation (EU) No 142/2011, in order to kill pathogenic agents; - (e) animals of the examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply with the following standards (e): - Salmonella: absence in 25g: n = 5, c = 0, m = 0, M = 0, Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram; - (e) either [packed in new or sterilised bags.] - (f) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectar approved by the competent authority before use.] - and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'; - (e) either [packed in new or sterilised bags.] - (f) or [transported in bulk in enclosed storage; - (g) either [packed in the competent authority before use.] - (g) either [packed in the contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) N 599/2001 of the European Parliament and of the Council (f) or mechanically separated meat obtained from bowers or caprine animals; and the animals from which the special care special and selection of contain and is not derived from bowine, ovine or caprine materials other than those deri			(ii) the following originating from terrestrial anima	als:		
- eggs, - egg by-products, including egg shells; - (iii) day-old chicks killed for commercial reasons.] - (c) and/or [- animal by-products from aquatic or terrestrial invertebrates other than species pathogenic to humans or animals.] - (d) and/or [- animal by-products from aquatic or terrestrial invertebrates other than species pathogenic to humans or animals.] - (e) and/or [- animal by-products from aquatic or terrestrial invertebrates other than species pathogenic to humans or animals.] - (e) and/or [- animal by-products from aquatic or terrestrial invertebrates other than species pathogenic to humans or animals.] - (e) and/or [- animal by-products from aquatic or terrestrial invertebrates other than species pathogenic to humans or animals.] - (e) and/or [- animal by-products with Annex XIII, Chapter III of Regulation (EU) No 142/2011, in order to kill pathogenic agents; - (e) animals of the examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply with the following standards (e): - Salmonella: absence in 25g: n = 5, c = 0, m = 0, M = 0, Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram; - (e) either [packed in new or sterilised bags.] - (f) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectar approved by the competent authority before use.] - and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'; - (e) either [packed in new or sterilised bags.] - (f) or [transported in bulk in enclosed storage; - (g) either [packed in the competent authority before use.] - (g) either [packed in the contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) N 599/2001 of the European Parliament and of the Council (f) or mechanically separated meat obtained from bowers or caprine animals; and the animals from which the special care special and selection of contain and is not derived from bowine, ovine or caprine materials other than those deri			— batchery by-products			
- egg by-products, including egg shells; (iii) day-old chicks killed for commercial reasons:] (?) and/or [- animal by-products from aquatic or terrestrial invertebrates other than species pathogenic to humans or animals:] (?) and/or [- animal by-products from aquatic or terrestrial invertebrates other than species pathogenic to humans or animals:] (?) and/or [- animal by-products from aquatic or terrestrial invertebrates other than species pathogenic to humans or animals:] (?) and/or [- animal by-products from aquatic or terrestrial products of the material being permitted in accordance with Aniel So(a)(ii) of Regulation (EC) No 1089/2009.] (8) have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply with the following standards (?): Salmonella: absence in 25g; n = 5, c = 0, m = 0, M = 0. Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram; (8) either [packed in new or sterilleed bags.] (9) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectar approved by the competent authority before use.] and which bear labels indicating "NOT FOR HUMAN CONSUMPTION: (9) either [the product was stored in enclosed storage: (1) either [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) N 999/2001 of the European Parliament and of the Council (?) or mechanically separated meat obtained from bone or bone of subjustered after stunning became of gas injected into the cranial cavity or killed by the same method or staughtered by loceration of central nervous test by means of an elongated of shaped instrument introduced into the carallal cavity or killed by the same method or staughtered by loceration of central nervous test by means of an elongated of shaped instrument introduced into the carallal cavity or killed by the same method or staughtered by loceration of central nervous test by means of an elongated						
(iii) day-old chicks killed for commercial reasons.] (c) and/or [- animal by-products from aquatic or terrestrial invertebrates other than species pathogenic to humans or animals.] (d) and/or [- material from animals which have been treated with certain substances which are prohibited pursuant to Directive 96/22/EC the import of the material being permitted in accordance with Article 33(a)(iii) of Regulation (EC) No 1059/2009.] II.3. have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply with the following standards (e): Salmonella: absence in 25g: n = 5, c = 0, m = 0, M = 0, Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram; II.5. the end product was: (e) either [packed in new or sterilised bags.] (f) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectar approved by the competent authority before use.] and which bear labels indicating "NOT FOR HUMAN CONSUMPTION"; II.6. the end product was stored in enclosed storage: II.7. the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) N 959/2001 of the European Parliament and of the Council (f) or mechanically separated meat obtained from bones of boving oving or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning be means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration or central nervous tissus by means of an elongated rod-shaped instrument introduced into the cranial cavity.] (e) or [the product does not contain and is not derived from boning or caprine materials other than those derived from animals born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk by a decision is accordance with Article 5(2) of Regulation (EC) No 999/2001) II.9. in addition as regards TS. or cranial cavity or c						
(c) and/or [- animal by-products from aquatic or terrestrial invertebrates other than species pathogenic to humans or animals.] (d) and/or [- material from animals which have been treated with certain substances which are prohibited pursuant to Directive 96/22/EC the import of the material being permitted in accordance with Article 36(a)(ii) of Regulation (EC) No 1069/2009.] II.3. have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply with the following standards (e): Salmonella: absence in 25g: n = 5, c = 0, m = 0, M = 0, Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram; II.5. the end product was: (e) either [packed in new or sterilised bags.] (f) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectar approved by the competent authority before use.] and which bear labels indicating "NOT FOR HUMAN CONSUMPTION"; II.6. the end product was stored in enclosed storage; II.7. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment; II.8. (f) either [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) N 959/2001 of the European Parliament and of the Council (f) or mechanically separated meat obtained from bones of boring owners of gas injected into the cranial cavity or killed by the same method or stauphreted by laceration of central nervous tissu by means of gas injected into the cranial cavity or killed by the same method or stauphreted by laceration of central nervous tissus by means of gas injected into the cranial cavity or region classified as posing a negligible BSE risk by a decision is accordance with Article (EC) No 999/2001.) II.9. in addition as regards Tatile: (f) either [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin the ovine and caprine animals from				1		
(?) and/or [- material from animals which have been treated with certain substances which are prohibited pursuant to Directive 98/22/EC the import of the material being permitted in accordance with Article 35(3)(ii) of Regulation (EC) No 1069/2009.] II.3. have been subjected to processing in accordance with Annex XIII, Chapter III of Regulation (EU) No 142/2011, in order to kill pathogeni agents: II.4. have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply with the following standards (?): Salmonella: absence in 25g: n = 5, c = 0, m = 0, M = 0, Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram; II.5. the end product was: (?) either [packed in new or sterilised bags.] (?) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectar approved by the competent authority before use.] and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'; II.6. the end product was stored in enclosed storage; II.7. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment; II.8. (?) either [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) N 999/2001 of the European Parliament and of the Council (?) or mechanically separated meat obtained from bones of bovin orine or caprine animals; and the animals from which this product is derived have not been slaughtered after sturning by means of an elongated rod-shaped instrument introduced into the oranial cavity.] (?) or [the product does not contain and is not derived from bovine, ovine or caprine materials other than those derived from accordance with Article 5(2) of Regulation (EC) No 999/2001; II.9. in addition as regards TSE: (?) either [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin the ovine and caprine animals from which these prod		(2) and (a)			sia ta humana ay animalad	
the import of the material being permitted in accordance with Article 35(a)(ii) of Regulation (EC) No 1069/2009.] II.3. have been subjected to processing in accordance with Annex XIII, Chapter III of Regulation (EU) No 142/2011, in order to kill pathogeni agents: II.4. have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply with the following standards (?): Salmonella: absence in 25g: n = 5, c = 0, m = 0, M = 0, Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram; II.5. the end product was: (*) either [packed in new or sterilised bags.] (*) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectar approved by the competent authority before use.] and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'; II.6. the end product was stored in enclosed storage; II.7. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment; II.8. (*) either [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) N 999/2001 of the European Parliament and of the Council (*) or mechanically separated meat obtained from bones of boving ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning by means of an elongated rod-shaped instrument introduced into the cranial cavily.] (*) or [the product does not contain and is not derived from bovine, ovine or caprine materials other than those derived from accordance with Article 5(2) of Regulation (EC) No 999/2001; II.9. in addition as regards TSE: (*) either [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last three years on a holding where no official novement restrictio					•	
agents; II.4. have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply with the following standards (*): Salmonella: absence in 25g: n = 5, c = 0, m = 0, M = 0, Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram; II.5. the end product was: (*) either [packed in new or sterilised bags.] (*) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectar approved by the competent authority before use.] and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'; II.6. the end product was stored in enclosed storage; II.7. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment; II.8. (*) either [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) N 998/2001 of the European Parliament and of the Council (*) or mechanically separated meat obtained from bones of boving ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning be means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissu by means of an elongated rod-shaped instrument introduced into the cranial cavity.] (*) or [the product does not contain and is not derived from bovine, ovine or caprine materials other than those derived from animal born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk by a decision is accordance with Article 5(2) of Regulation (EC) No 999/2001.] III.9. in addition as regards TSE: (*) either [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origing the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last three years: (i) It has been subject to regular of		(-) and/or				
following standards (*): Salmonella: absence in 25g: n = 5, c = 0, m = 0, M = 0, Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram; II.5. the end product was: (*) either [packed in new or sterilised bags.] (*) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectar approved by the competent authority before use.] and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'; II.6. the end product was stored in enclosed storage; II.7. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment; II.8. (*) either [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) N 999/2001 of the European Parliament and of the Council (*) or mechanically separated meat obtained from bones of boving ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning be means of gas injected into the oranial cavity or killed by the same method or slaughtered by laceration of central nervous tissus by means of an elongated cod-shaped instrument introduced into the oranial cavity.] (*) or [the product does not contain and is not derived from bovine, ovine or caprine materials other than those derived from animal born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk by a decision is accordance with Article 5(2) of Regulation (EC) No 999/2001:] II.9. in addition as regards TSE: (*) either [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last three years: (*) It has been subject to regular official weterinary checks; (*) no classical scrapic case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or f	II.3.		subjected to processing in accordance with Annex X	(III, Chapter III of Regulation (EU) No 1	42/2011, in order to kill pathogenic	
Enterobacteriaceae: n = 5, c = 2, m = 10, M = 300 in 1 gram; II.5. the end product was: (*) either [packed in new or sterilised bags.] (*) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectar approved by the competent authority before use.] and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'; II.6. the end product was stored in enclosed storage; II.7. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment; II.8. (*) either [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) N 999/2001 of the European Parliament and of the Council (*) or mechanically separated meat obtained from bones of bovinin ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning b means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissu by means of an elongated rod-shaped instrument introduced into the cranial cavity) (*) or [the product does not contain and is not derived from bovine, ovine or caprine materials other than those derived from animal born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk by a decision in accordance with Article 5(2) of Regulation (EC) No 999/2001;] II.9. in addition as regards TSE: (*) either [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origing the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the lat three years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfie the following requirements for the last three years: (i) it has been subject to regular official veterinary checks; (ii) no classical scrapie case, as defined	II.4.	have been examined by the competent authority taking a random sample immediately prior to dispatch and found it to comply with the following standards (3):				
II.5. the end product was: (*) either [packed in new or sterilised bags.] (*) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectar approved by the competent authority before use.] and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'; II.6. the end product was stored in enclosed storage; II.7. the end product has undergone all precautions to avoid contamination with pathogenic agents after treatment; II.8. (*) either [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) N 999/2001 of the European Parliament and of the Council (*) or mechanically separated mean of basinghiered after stunning breason of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissu by means of an elongated rod-shaped instrument introduced into the cranial cavity.] (*) or [the product does not contain and is not derived from bovine, ovine or caprine materials other than those derived from all above, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk by a decision is accordance with Article 5(2) of Regulation (EC) No 999/2001;] II.9. in addition as regards TSE: (*) either [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origing the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the lat three years. (*) it has been subject to regular official veterinary checks; (i) in classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed of following the confirmation of a classical scrapie was confirmed have been killed and destroyed, and — all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARI		Salmonella	a: absence in 25g: $n = 5$, $c = 0$, $m = 0$,	M = 0,		
(²) either [packed in new or sterilised bags.] (²) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectar approved by the competent authority before use.] and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'; II.6. the end product was stored in enclosed storage; II.7. the end product has undergone all precautions to avoid contamination with pathogenic agents after treatment; II.8. (²) either [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) N 999/2001 of the European Parliament and of the Council (²) or mechanically separated mean at obtained from bones of boving over parliaments and the animals from which this product is derived have not been slaughtered after stunning be means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissu by means of an elongated rod-shaped instrument introduced into the cranial cavity.] (²) or [the product does not contain and is not derived from bovine, ovine or caprine materials other than those derived from animal born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk by a decision is accordance with Article 5(2) of Regulation (EC) No 999/2001;] II.9. in addition as regards TSE: (²) either [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the lat three years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfie the following requirements for the last three years: (i) it has been subject to regular official evterinary checks; (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed of following th		Enterobact	teriaceae: n = 5, c = 2, m = 10, M = 300 in 1 g	ıram;		
(a) or [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectar approved by the competent authority before use.] and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'; II.6. II.7. II.8. (b) either [the product was stored in enclosed storage; the product has undergone all precautions to avoid contamination with pathogenic agents after treatment; II.8. (c) either [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) N 999/2001 of the European Parliament and of the Council (†) or mechanically separated meat obtained from bones of bovine ovine or caprine animals; and the animals from which this product is derived any not been slaughtered after stunning be means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissus by means of an elongated rod-shaped instrument introduced into the cranial cavity.] (d) or [the product does not contain and is not derived from bovine, ovine or caprine materials other than those derived from animal born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk by a decision is accordance with Article 5(2) of Regulation (EC) No 999/2001;] II.9. in addition as regards TSE: (e) either [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the lat three years on a holding where no official novement restriction is imposed due to a suspicion of TSE and which has satisfie the following requirements for the last three years: (i) it has been subject to regular official veterinary checks; (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed of following the confirmation of a classical	II.5.	the end pr	oduct was:			
and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION'; II.6. the end product was stored in enclosed storage; II.7. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment; II.8. (?) either [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) N 999/2001 of the European Parliament and of the Council (*) or mechanically separated meat obtained from bones of boving ovine or caprine animals; and the animals from which this product is devided have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissus by means of an elongated rod-shaped instrument introduced into the cranial cavity.] (?) or [the product does not contain and is not derived from bovine, ovine or caprine materials other than those derived from animal born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk by a decision is accordance with Article 5(2) of Regulation (EC) No 999/2001:] II.9. in addition as regards TSE: (?) either [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origing the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last three years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfied the following requirements for the last three years: (i) It has been subject to regular official veterinary checks; (ii) no classical scrapic case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or following the confirmation of a classical scrapic case: — all animals in which classical scrapie was confirmed have been killed and destroyed, and — all goats and sheep on the holding have been killed and destroyed, except for breeding		(2) either	[packed in new or sterilised bags,]			
II.6. the end product was stored in enclosed storage; II.7. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment; II.8. (2) either [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) N 999/2001 of the European Parliament and of the Council (*) or mechanically separated meat obtained from bones of boving ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning be means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissus by means of an elongated rod-shaped instrument introduced into the cranial cavity.] (2) or [the product does not contain and is not derived from bovine, ovine or caprine materials other than those derived from animal born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk by a decision is accordance with Article 5(2) of Regulation (EC) No 999/2001;] III.9. in addition as regards TSE: (2) either [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origing the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last three years: (i) it has been subject to regular official oveterinary checks; (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed or following the confirmation of a classical scrapie case: — all animals in which classical scrapie was confirmed have been killed and destroyed, and — all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARI		(²) or		transport that were thoroughly cleaned	and disinfected with a disinfectant	
II.7. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment; II.8. (2) either		and which	bear labels indicating 'NOT FOR HUMAN CONSUM	IPTION';		
(2) either (2) either (3) either (4) either (4) either (5) either (6)	II.6.	the end pr	oduct was stored in enclosed storage;			
(²) either [the product does not contain and is not derived from specified risk material as defined in Annex V to Regulation (EC) N 999/2001 of the European Parliament and of the Council (⁴) or mechanically separated meat obtained from bones of boving ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning be means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissu by means of an elongated rod-shaped instrument introduced into the cranial cavity.] (²) or [the product does not contain and is not derived from bovine, ovine or caprine materials other than those derived from animal born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk by a decision is accordance with Article 5(2) of Regulation (EC) No 999/2001;] II.9. in addition as regards TSE: (²) either [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origing the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last three years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfies the following requirements for the last three years: (i) it has been subject to regular official veterinary checks; (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed of following the confirmation of a classical scrapie case: — all animals in which classical scrapie was confirmed have been killed and destroyed, and — all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARI	II.7.	the produc	et has undergone all precautions to avoid contaminati	on with pathogenic agents after treatm	nent;	
999/2001 of the European Parliament and of the Council (*) or mechanically separated meat obtained from bones of boving ovine or caprine animals; and the animals from which this product is derived have not been slaughtered after stunning be means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration of central nervous tissus by means of an elongated rod-shaped instrument introduced into the cranial cavity;] (²) or [the product does not contain and is not derived from bovine, ovine or caprine materials other than those derived from animal born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk by a decision is accordance with Article 5(2) of Regulation (EC) No 999/2001;] II.9. in addition as regards TSE: (²) either [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last three years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfie the following requirements for the last three years: (i) it has been subject to regular official veterinary checks; (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed of following the confirmation of a classical scrapie case: — all animals in which classical scrapie was confirmed have been killed and destroyed, and — all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARI	II.8.					
born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk by a decision is accordance with Article 5(2) of Regulation (EC) No 999/2001;] II.9. in addition as regards TSE: (2) either [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origing the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the last three years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfied the following requirements for the last three years: (i) it has been subject to regular official veterinary checks; (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed of following the confirmation of a classical scrapie case: — all animals in which classical scrapie was confirmed have been killed and destroyed, and — all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARI		(²) either	999/2001 of the European Parliament and of the Co ovine or caprine animals; and the animals from wh means of gas injected into the cranial cavity or killed	ouncil (4) or mechanically separated m nich this product is derived have not by the same method or slaughtered by	eat obtained from bones of bovine, been slaughtered after stunning by	
(2) either [in case of animal by-products intended for feeding ruminants and containing milk or milk products of ovine or caprine origin the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the lat three years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfie the following requirements for the last three years: (i) it has been subject to regular official veterinary checks; (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed of following the confirmation of a classical scrapie case: — all animals in which classical scrapie was confirmed have been killed and destroyed, and — all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARI		(²) or	born, continuously reared and slaughtered in a cou	ntry or region classified as posing a r		
the ovine and caprine animals from which these products are derived have been kept continuously since birth or for the lat three years on a holding where no official movement restriction is imposed due to a suspicion of TSE and which has satisfie the following requirements for the last three years: (i) it has been subject to regular official veterinary checks; (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed of following the confirmation of a classical scrapie case: — all animals in which classical scrapie was confirmed have been killed and destroyed, and — all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARI	II.9.	in addition	as regards TSE:			
 (ii) no classical scrapie case, as defined in point 2(g) of Annex I to Regulation (EC) No 999/2001, has been diagnosed of following the confirmation of a classical scrapie case: all animals in which classical scrapie was confirmed have been killed and destroyed, and all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARI 		(²) either	the ovine and caprine animals from which these pro three years on a holding where no official movemen	oducts are derived have been kept co	ntinuously since birth or for the last	
following the confirmation of a classical scrapie case: — all animals in which classical scrapie was confirmed have been killed and destroyed, and — all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARI			(i) it has been subject to regular official veterinary	checks;		
 — all goats and sheep on the holding have been killed and destroyed, except for breeding rams of the ARR/ARI 					999/2001, has been diagnosed or,	
			- all animals in which classical scrapie was co	onfirmed have been killed and destroy	ed, and	

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Flavouring innards for use in the manufacture of petfood

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

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

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

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

Notes

Part I:

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

Part II:

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

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Flavouring innards for use in the manufacture of petfood

II. Health information	II.a. Certificate reference No	II.b.					
(⁴) OJ L 147, 31.5.2001, p. 1.							
(⁵) OJ L 94, 11.4.2006, p. 28.							
— The signature and the stamp must be in a different colour to that of the printing.							
 Note for the person responsible for the consignment in the European Union: This certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border inspection post. 							
Official veterinarian/Official inspector							
Name (in capital letters): Qualification and title:							
Date: Signature:							
Stamp:							