CHAPTER 3(E)

Health certificate

For flavouring innards for use in the manufacture of petfood, intended for dispatch to $\frac{1}{2}$ the European Union

COUNTRY: UNITED STATES Veterinary certificate to EU

	I.1. Consignor	I.2. Certificate reference No I.2.a.				
	Name Address					
	7.44.650	I.3. Central competent authority				
	Tel.	APHIS-VS I.4. Local competent authority				
		10.5				
	I.5. Consignee Name Address	I.6. Person responsible for the load in EU Name Address				
	Postal code Tel.	Postal code Tel.				
ment						
sign	17 Country of 100 and 10 Parism of Oak					
uoo pe	I.7. Country of ISO code origin I.8. Region of Code origin	I.9. Country of ISO I.10. Region of Code destination code destination				
atche	I.11. Place of origin	I.12. Place of destination				
Part I : Details of dispatched consignment	Name Approval number Address	Custom warehouse Name Approval number Address				
Detai		Postal code				
1 =						
Par						
	Name Approval number Address					
	Name Approval number					
	Address					
	I.13. Place of loading	I.14. Date of departure				
	I.15. Means of transport	I.16. Entry BIP in EU				
	Aeroplane □ Ship □ Railway wagon □					
	Road vehicle Other	1.17.				
	Identification					
	Documentation references					
	I.18. Description of commodity	I.19. Commodity code (HS code)				
		I.20. Quantity				
	I.21. Temperature of product	I.22. Number of packages				
	Ambient ☐ Chilled ☐ Frozen ☐					

COUNTRY: UNITED STATES Flavouring innards for use in the manufacture of petfood I.2. Certificate reference No I.2.a. I.23. Seal/Container No I.24. Type of packaging I.25. Commodities certified for: Petfood Technical use \Box I.26. For transit through EU to third country I.27. For import or admission into EU Third country ISO code I.28. Identification of the commodities Species Nature of commodity Approval number of establishments Net weight Batch number (Scientific name) Manufacturing plant

	COUNTRY: UNITED STATES		3	Flavouring limarus for use in the manufacture of petiood				
	II.	Health information	on	II.a. Certificate reference No	II.b.			
	-	I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council(1a), and in particular Article 8 and 10 thereof, and Commission Regulation (EU) No 142/2011(1b), and in particular Chapter III of Annex XIII and Chapter II of Annex XIV thereto, and certify that the flavouring innards products described above:						
	II.1.	consist of anima	al by-products that satisf	fy the animal health requirements below;				
o								
Part II: Certification		(²)either [-	carcases and parts of parts of animals killed	animals slaughtered or, in the case of gan , and which are fit for human consumption ir n, but are not intended for human con	ne, bodies or naccordance			
=		(2)and/or [**	wing parts originating either from animals th	at have been			
Par		(²)and/or [-	slaughtered in a slaugl	wing parts originating sittler from attilinate th hterhouse and were considered fit for slaugh _I an ante-mortem inspection or bodies and	ter for human			
				game killed for human consumption in acc				
	human consum			lies and parts of animals which are rejecte ption in accordance with Union legislation, l gns of disease communicable to humans or	out which did			
			(ii) heads of poultry	<u></u>				
			(iii) hides and skins,	, including trimmings and splitting thereof, horns and feet,				
			including the ph and metatarsus	nalanges and the carpus and metacarpus bones, tarsus				
			(iv) pig bristles;					
			(v) feathers;]					
		(²)and/or [-	through blood to hum slaughtered in a slaug	ch did not show any signs of disease or nans or animals, obtained from animals the phterhouse after having been considered fit n following an ante mortem inspection in acc	at have been for slaughter			
		(²)and/or [-		ising from the production of products intende g degreased bone, greaves and centrifuge essing;]				
		(²)and/or [- products of animal or which are no longer ir or due to problems of r		igin, or foodstuffs containing products of a ntended for human consumption for comme manufacturing or packaging defects or other or animal health arise;]	rcial reasons			
		(²)and/or [- petfood and feedingstuffs of animal origin, or feedingstuffs containing ani by-products or derived products, which are no longer intended for feeding commercial reasons or due to problems of manufacturing or packaging deformation or other defects from which no risk to public or animal health arise;]			or feeding for aging defects			
		(²) and/or [- blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk orig from live animals that did not show signs of any disease communicable that product to humans or animals;]						
(²)and/or [- aquatic animals, and p			aquatic animals, and p	varts of such animals, except sea mammals, eases communicable to humans or animals;	which did not]			
		(²)and/or [from aquatic animals originating from acturing products for human consumption;	n plants or			
		(²)and/or [-	disease communicable	originating from animals which did not show any signs of through that material to humans or animals:				
			` '	lfish with soft tissue or flesh; ginating from terrestrial animals:				

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II.	Health inf	formation		II.a.	a. Ceri	ificate ref	erence N	0		II.b.
			- hatchery	by-prod	ducts					/
	eggs,									
	egg by products, including egg shells;									
	(iii) day old chicks killed for commercial reasons;									
	(a) animal by products from aquatic or terrestrial invertebrates other than species									
	()4.74.07	pathogenic to humans or animals;]								
	(²)and/or	nd/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	and/or [material from animals which have been treated with certain substances which are prohibited by Council Directive 96/22/EC(^{2a}), the import of the material being permitted in accordance with Article 35(a)(ii) of Regulation (EC) No 1069/2009;]								
II.3.		e been subjected to processing in accordance with Chapter III of Annex XIII to Regulation) No 142/2011, in order to kill pathogenic agents;								
II.4.	during or	after storage	indom sampling e at the process	ng plar	int and	complie	s with th			
	Salmone		absence in 25	-						
			n = 5, c = 2, m	= 10, 1	M = 30	00 in 1 g	ramme;			
II.5.		product was:								
	(²)either		new or sterilise							
	(²)or [transported in bulk in containers or other means of transport that were thoroug cleaned and disinfected with a disinfectant approved by the competent authobefore use.]									
	and which bear labels indicating 'NOT FOR HUMAN CONSUMPTION';									
II.6.	the end p	he end product was stored in enclosed storage;								
II.7.	the produ treatmen		rgone all preca	utions to	to avoi	d contar	nination	with pat	hogenic a	gents after
$(^{2})[II.8.$	the flavor	uring innards	products descr	bed ab	bove					
	(²)either	: [is derived f	rom other rumin	ants th	han bo	vine, ovir	ne or ca	orine anii	mals.]]	
	(²) <i>or</i>	[is derived derived from	from bovine, ov n:	ine or	r caprii	ne anima	als and	does no	t contain	and is not
		(²) either	[bovine, ovine animals born, classified as 2007/453/EC:	continu cosing	uously	reared a	and slau	ghtered i	in a countr	y or region
		(²)or	Regul		EC) No					nnex V to nent and of
			ovine born, c classif Comm	or capr continudied as	orine a uously s posin Decis	nimals, e reared a g a neg ion 2007	except fr nd slaug lligible E	om thos phtered in BSE risk	e animals n a countr in accord	of bovine, that were y or region dance with has been
			or cap lacera rod-sh means	orine ar tion of t aped in s of gas	animals the ce instrum as inje	which ntral ner nent intro cted into	have be ous tiss oduced the cra	en killed ue by me into the inial cav	d, after st eans of ar cranial ca ity, excep	vine, ovine unning, by elongated wity, or by t for those htered in a

COUNTRY: UNITED STATES

II. Health information

II.a. Certificate reference No

II.b.

country or region classified as posing a negligible 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.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.19: use the appropriate HS code: 05.04; 05.06, 05.11 or 23.09.
- 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 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
 - 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.
- (^{2a}) OJ L 125, 23.5.1996, p. 3.
- (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.
- (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 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.

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Flavouring innards for use in the manufacture of petfood

II. Health information	I	I.a. Certificate reference No	II.b.
Official veterinarian/Official inspector	•		
Name (in capital letters):	Qualific	cation and title:	
Date:	Signati	ure:	
	Stamp		