# COUNTRY: UNITED KINGDOM Part I: Details of dispatched consignment

## Veterinary certificate to EU

I.1. Consignor Name		I.2. Certificate reference number		I.2.a.	
Address		L3. Central Competent Authority			
		DEPARTMENT FOR ENVIRONMENT, FOOD AND RURAL AFFAIRS			
Tal		I.4. Local Competent Authority			
Tel.  I.5. Consignee		I.6. Person responsible for the load in the	EU		
Name		Name			
Address		Address			
Postcode Tel.		Postcode Tel.			
I.7. Country of origin, ISO code UNITED KINGDOM	I.8. Region of origin Code	I.9. Country of destination	ISO code	I.10. Region of destination	Code
I.11. Place of origin		I.12. Place of destination			
Name	Approval number	N.	Custom	warehouse	
Address		Name Address			
		Postcode			
L12 Diversity For		Approval number			
I.13. Place of loading		I.14. Date of departure			
I.15 Means of transport		I.16. Entry BIP in EU			
	hip Railway wagon				
Road vehicle Identification:	Other	I.17.			
Documentary references:		1.17.			
I.18. Description of commodity		I.19. Commodi	ty code (HS	S code)	
I.21. Temperature of products	11.20.	Quantity	I.22. N	Number of packages	
Ambient Chilled	_				
I.23. Seal/Container No.			І.24. Т	Type of packaging	
I.25. Commodity certified for:					
Technical use					
I.26. For transit to 3rd country by EU	1 27/	For import or admission into EU			
1.20. For dailor to 3rd country by 20					
3rd country	ISO code				
I.28. Identification of the commodities					
Species (Scientific name)	Approval number of establishr Manufacturing plant	nents  Number of packages	N	let weight Batch number	er
species (selentine name)		rumoti oi puolugos		Silver mans	
,					

II.	Health information	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 (<sup>1a</sup>) and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011 (<sup>1b</sup>), and in particular Chapter II of Annex XIV thereto, and certify that the fat derivatives described above:

- II.1. consist of fat derivatives that satisfy the health requirements below;
- II.2. consist of fat derivatives intended for purposes outside the feed chain, other than in cosmetics, pharmaceuticals and medical devices;
- II.3. have been prepared and stored in a plant approved, validated and supervised by the competent authority in accordance with Article 24 of Regulation (EC) No 1069/2009, in order to kill pathogenic agents;
- II.4. have been prepared from rendered fats exclusively produced from the following materials:
- II.4.1. in case the fat derivatives are intended for uses outside the feed chain, other than in organic fertilisers, soil improvers, cosmetics, pharmaceuticals and medical devices, the following Category 1 materials:
  - (2) either [- the following material:
    - (i) specified risk material;
    - (ii) entire bodies or parts of dead animals containing specified risk material at the time of disposal;]
  - (2) and/or [- animal by-products which have been derived from animals which have been submitted to illegal treatment as defined in Article 1(2)(d) of Directive 96/22/EC or Article 2(b) of Directive 96/23/EC;]
  - (2) and/or [- animal by-products containing residues of other substances and environmental contaminants listed in Group B (3) of Annex I to Directive 96/23/EC, if such residues exceed the permitted levels laid down by Union legislation or, in the absence thereof, by legislation of the Member State of importation;]
- II.4.2. in case the fat derivatives are intended for use in organic fertilisers or soil improvers or other uses outside the feed chain, other than in cosmetics, pharmaceuticals and medical devices, the following Category 2 materials:
  - (2) either [- animal by-products containing residues of authorised substances or contaminants exceeding the permitted levels referred to in Article 15(3) of Directive 96/23/EC;]
  - (²) and/or [- products of animal origin which have been declared unfit for human consumption due to the presence of foreign bodies in those products;]
  - (2) and/or [- animals and parts of animals, other than those referred to in Articles 8 and 10 of Regulation (EC) No 1069/2009, that died other than being slaughtered or killed for human consumption, including animals killed for disease control purposes;]
- II.4.3. the following Category 3 materials:
  - (2) either [- carcases and parts of animals slaughtered or, in the case of game, bodies or parts of animals killed, and which are fit for human consumption in accordance with Union legislation, but are not intended for human consumption for commercial reasons;]
  - (2) and/or [- carcases and the following parts originating either from animals that have been slaughtered in a slaughterhouse and were 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 with Union legislation, but which did not show any signs of disease communicable to humans;
    - (ii) heads of poultry;
    - (iii) hides and skins, including trimmings and splitting thereof, horns and feet, including the phalanges and the carpus and metacarpus bones, tarsus and metatarsus bones;
    - (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, obtained from animals that have been slaughtered in a slaughterhouse after having been considered fit for slaughter for human consumption following an ante-mortem inspection in accordance with Union legislation;]
  - (2) and/or [- animal by-products arising from the production of products intended for human consumption, including degreased bone, greaves and centrifuge or separator sludge from milk processing;]

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### Fat derivatives not intended for human onsumption to be used outside the feed chain

	consumption to be used outside the feed c			n to be used outside the feed chain	
II.	Health information		II.a. Certificate reference No	II.b.	
				or due to problems of manufacturing	gin, which are no longer intended for human g or packaging defects or other defects from
	(2) and/or [- petfood and feedingstuffs of animal origin, or feedingstuffs containing animal by-products or derived products, which are no longer intended for feeding for commercial reasons or due to problems of manufacturing or packaging defects of other defects from which no risk to public or animal health arises;]				
	( <sup>2</sup> ) and/or	[- blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk originating from live animals that did not show signs of any disease communicable through that product to humans or animals;]			
	(2) and/or [- aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of disease communicable to humans or animals;]				
	(2) and/or [- animal by-products from aquatic animals originating from plants or establishments manufacturing products for h consumption;]				lishments manufacturing products for human
	(²) and/or		ng material originating from humans or animals:	animals which did not show any	signs of disease communicable through that
(i) shells from shellfish with soft tissue or flesh;					
	(ii) the following originating from terrestrial animals:				
	<ul><li>— hatchery by-products,</li><li>— eggs,</li><li>— egg by-products, including egg shells;</li></ul>				
	(iii) day-old chicks killed for commercial reasons;]				
II.5.	5. in case of fat derivatives produced from animal by-products referred to in point II.4.1 and point II.4.2:				
	(a) have been	produced usin	g the following methods:		
	(²) either	[transesterification or hydrolysis at least 200 °C, under corresponding appropriate pressure, for 20 minutes (glycerol, fatty acids and esters)]			
	(²) or [saponification with NaOH 12M (glycerol and soap):				
		(²) either	[in a batch process at	95 °C for three hours;]	
		( <sup>2</sup> ) or	[in a continuous proces	s at 140 °C, 2 bars (2000 hPa) for	eight minutes;]]
	(²) or [hydrogenation at 160 °C at 12 bars (12000 hPa) pressure for 20 minutes;]				

- (b) are packaged in new containers or in containers that have been cleaned, and all precautions are taken to prevent its contamination which bear labels indicating "NOT FOR HUMAN OR ANIMAL CONSUMPTION";
- II.6. in case of fat derivatives produced from animal by-products referred to in point II.4.3, the fat derivatives have been produced in accordance with one of the processing methods [1]-[2]-[3]-[4]-[6]-[7] (2) referred to in Chapter III of Annex IV to Regulation (EU) No 142/2011.

### 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) is to be provided. In case of unloading and reloading, the consignor must inform the BIP of entry into the EU.
- Box I.19: use the appropriate Harmonized System (HS) code under the following headings: 15.16 or 15.08.

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# Fat derivatives not intended for human consumption to be used outside the feed chain

		II.a. Certificate reference No	II.b.		
II.	Health information				
— Вох	<ul> <li>Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.</li> </ul>				
— Вох	reference I.25: technical use: any use other than for animal cons	sumption.			
— Вох	reference I.26 and I.27: fill in according to whether it is a transit	or an import certificate.			
— Вох	reference I.28:				
Spe	ecies: select from the following: Ruminantia, Other;				
Ма	nufacturing plant: provide the registration number of treatment/pro-	cessing establishment.			
Part II	:				
( <sup>1a</sup> ) O	J L 300, 14.11.2009, p. 1.				
( <sup>1b</sup> ) O	J L 54, 26.2.2011, p. 1.				
(²) D	elete as appropriate.				
	e signature and the stamp must be in a different colour to that of	the printing.			
— Not	e for the person responsible for the consignment in the European U		ary purposes and has to accompany		
the	consignment until it reaches the border inspection post.	,	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,		
— 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				

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