

Part I: Details of dispatched consignment	I.1. Consignor Name Address Tel.		I.2. Certificate reference No		I.2.a	
			I.3. Central competent authority			
			I.4. Local competent authority			
	I.5. Consignee Name Address Postal code Tel.		I.6. Person responsible for the load in EU Name Address Postal code Tel.			
	I.7. Country of origin ISO code I.8. Region of origin Code		I.9. Country of destination ISO code		I.10. Region of destination Code	
	I.11. Place of origin Name Address Approval number		I.12. Place of destination Name Address Postal code Approval number Custom warehouse <input type="checkbox"/>			
	I.13. Place of loading		I.14. Date of departure			
	I.15. Means of transport <input type="checkbox"/> Aeroplane <input type="checkbox"/> Ship <input type="checkbox"/> Railway wagon <input type="checkbox"/> Road vehicle <input type="checkbox"/> Other Identification Documentation references		I.16. Entry BIP in EU I.17.			
	I.18. Description of commodity			I.19. Commodity code (HS code)		
				I.20. Quantity		
I.21. Temperature of product <input type="checkbox"/> Ambient <input type="checkbox"/> Chilled <input type="checkbox"/> Frozen			I.22. Number of packages			
I.23. Seal/Container No			I.24. Type of packaging			
I.25. Commodities certified for: Technical use <input type="checkbox"/>						
I.26. For transit through EU to third country <input type="checkbox"/> Third country ISO code			I.27. For import or admission into EU <input type="checkbox"/>			
I.28. Identification of the commodities Species (scientific name) Approval number of establishments Manufacturing plant Number of packages Net weight Batch number						

COUNTRY: UNITED KINGDOM

II.a Certificate reference No

II. Health information

I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009 of the European Parliament and of the Council ^(1a), and in particular Articles 8, 9 and 10 thereof, and Commission Regulation (EU) No 142/2011 ^(1b), and in particular Chapter II of Annex XIV thereto, and certify that the rendered fats described above:

II.1. consist of rendered fats not intended for human consumption that satisfy the health requirements below;

II.2. have been prepared exclusively with the following animal by-products:

(²)II.2.1. in the case of materials destined for the production of renewable fuels referred to in Chapter IV, Section 2, point L, of Annex IV to Regulation (EU) No 142/2011, biodiesel or oleochemical products, animal by-products referred to in Articles 8, 9 and 10 of Regulation (EC) No 1069/2009;]

(²)II.2.2. in the case of materials destined for the production of renewable fuels referred to in Chapter IV, Section 2, point J, of Annex IV to Regulation (EU) No 142/2011, the materials have been prepared exclusively from animal by-products referred to in Articles 9 and 10 of Regulation (EC) No 1069/2009;]

(²)II.2.3. in the case of materials destined for purposes other than cosmetics, pharmaceuticals or medical devices, the materials have been prepared exclusively from:

(²)either [- animal by-products containing residues of authorised substances or contaminants exceeding the permitted levels referred to in Article 15(3) of Council Directive 96/23/EC^(2a)];]

(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;]

(²)and/or [- carcasses 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 [- carcasses 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) carcasses 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 or animals;

(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;]

(²)and/or [- products of animal origin, or foodstuffs containing products of animal origin, which are no longer intended for human consumption for commercial reasons or due to problems of manufacturing or packaging defects or other defects from which no risk to public or animal health arises;]

(²)and/or [- petfood and feeding stuffs of animal origin, or feeding stuffs containing animal by-products or derived products, which are no longer intended for feeding for commercial reasons or due to problems of manufacturing or packaging defects or other defects from which no risk to public or animal health arises;]

(²)and/or [- blood, placenta, wool, feathers, hair, horns, hoof cuts and raw milk originating from live animals that did not show signs of any disease communicable through that product to humans or animals;]

(²)and/or [- aquatic animals, and parts of such animals, except sea mammals, which did not show any signs of diseases communicable to humans or animals;]

(²)and/or [- animal by-products from aquatic animals originating from plants or establishments manufacturing products for human consumption;]

(²)and/or [- the following material originating from animals which did not show any signs of disease communicable through that material to humans or animals:

(i) shells from shellfish with soft tissue or flesh;

(ii) the following originating from terrestrial animals:

- hatchery by-products,

- eggs,

- egg by-products, including egg shells,

(iii) day-old chicks killed for commercial reasons;]

(²)and/or [- aquatic and terrestrial invertebrates other than species pathogenic to humans or animals;]

(²)and/or [- animals and parts thereof of the zoological orders of Rodentia and Lagomorpha, except Category 1 material as referred to in Article 8, point (a)(iii), (iv) and (v), of Regulation (EC) No 1069/2009 and Category 2 material as referred to in Article 9, points (a) to (g), of that Regulation;]

Part II: Certification

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

COUNTRY: UNITED KINGDOM

II.a Certificate reference No

	<p>(²)and/or</p> <p>(²)and/or</p> <p>(²)II.2.4.</p> <p>(²)either</p> <p>(²)and/or</p> <p>(²)and/or</p> <p>(²)and/or</p> <p>II.3.</p> <p>(a)</p> <p>(²)II.4.</p> <p>(²)either</p> <p>(²)or</p> <p>(²) either</p> <p>(²)or</p> <p>Notes</p> <p>Part I:</p>	<p>[- hides and skins, hooves, feathers, wool, horns, hair and fur originating from dead animals that did not show any signs of disease communicable through that product to humans or animals;]</p> <p>[- adipose tissue from animals which did not show any signs of disease communicable through that material to humans or animals, which were slaughtered in a slaughterhouse and which were considered fit for slaughter for human consumption following an ante-mortem inspection in accordance with Union legislation;]]</p> <p>in the case of materials destined for purposes other than the production of organic fertilisers or soil improvers, cosmetics, pharmaceutical or medical devices :</p> <p>[- specified risk material as defined in Article 3(1), point (g), of Regulation (EC) No 999/2001 of the European Parliament and of the Council^(2b);]</p> <p>[- entire bodies or parts of dead animals containing specified risk material as defined in Article 3(1), point (g), of Regulation (EC) No 999/2001 at the time of disposal;]</p> <p>[- animal by-products which have been derived from animals which have been submitted to illegal treatment as defined in Article 1(2), point (d), of Council Directive 96/22/EC^(2c) or Article 2, point (b), of Directive 96/23/EC;]</p> <p>[- 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;]]</p> <p>the rendered fats:</p> <p>have been subjected to processing in accordance with method (indicate the processing method) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011, in order to kill pathogenic agents,</p> <p>(²) of Category 1 and 2 materials have been marked before dispatch to the European Union with glyceroltriheptanoate (GTH), so that a homogenous minimum concentration of at least 250 mg GTH per kilogramme fat is achieved,]</p> <p>in the case of rendered fats of ruminant origin, insoluble impurities in excess of 0,15 % in weight have been removed,</p> <p>have been transported under conditions which prevent their contamination, and</p> <p>bear labels on the packaging or container indicating "NOT FOR HUMAN OR ANIMAL CONSUMPTION";</p> <p>in the case of materials destined for organic fertilisers, cosmetics, pharmaceuticals, medical devices or soil improvers the rendered fats described above</p> <p>[are derived from other ruminants than bovine, ovine or caprine animals.]</p> <p>[are derived from bovine, ovine or caprine animals and does not contain and is not derived from:</p> <p>[bovine, ovine and caprine materials other than those derived from animals born, continuously reared and slaughtered in a country or region classified as posing a negligible bovine spongiform encephalopathy (BSE) risk in accordance with Commission Decision 2007/453/EC⁽³⁾.]</p> <p>[(a) specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001;</p> <p>(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 Decision 2007/453/EC, in which there has been no indigenous BSE case,</p> <p>(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.]]</p>	
	<p>- 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.</p> <p>- Box reference I.11: Approval number: the registration number of the establishment or plant, which has been issued by the competent authority.</p> <p>- Box reference I.12:</p> <p>- approval number: the registration number of the establishment or plant, which has been issued by the competent authority;</p> <p>- place of destination: this box is to be filled in only if it is a certificate for a transit commodity. Products in transit may only be stored in free zones, free warehouses and custom warehouses.</p> <p>- Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to be provided. In the case of unloading and reloading in the European Union, the consignor must inform the border inspection post of the point of entry into the European Union.</p> <p>- Box I.19: use the appropriate Harmonized System (HS) code under the following headings: 04.05; 15.01, 15.02; 15.03; 15.04; 15.05; 15.06; 15.16 or 15.18.</p> <p>- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) must be included.</p> <p>- Box reference I.25: technical use: any use other than feeding of farmed animals, other than fur animals or pet animals, and the production or manufacturing of petfood.</p> <p>- Box reference I.26 and I.27: fill in according to whether it is a transit or an import certificate.</p> <p>- Box reference I.28:</p> <p>- species: select from the following: Ruminantia, other than Ruminantia;</p> <p>- manufacturing plant: provide the registration number of the treatment/processing establishment.</p>		

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

COUNTRY: UNITED KINGDOM

II.a Certificate reference No

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. 10.

(2b) OJ L 147, 31.5.2001, p. 1.

(2c) OJ L 125, 23.5.1996, p. 3.

(3) 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 of the point of entry into the European Union.

Official veterinarian/Official inspector

Name (in capital letters):

Date:

Qualification and title:

Stamp:

Signature: