Part I : Details of consignment

I.1. Consignor
Name
Address
Country ISO Code

I.2. IMSOC Reference
I.2.a. Local Reference

I.5. Consignee
Name
Address
Country ISO Code

I.3. Central competent authority
I.4. Local competent authority

I.7. Country of origin ISO Code

I.9. Country of destination ISO Code

I.8. Region of origin Code

~~I.10. Region of destination~~

I.11. Place of Dispatch
Name
Address
Approval Number
Country ISO Code

I.12. Place of destination
Name
Address
Approval Number
Country ISO Code

I.13. Place of Loading
Name
Address
Approval Number
Country ISO Code

I.14. Date and time of departure

I.15. Means of Transport

Mode	International transport document	Identification

I.16 Entry Point

I.18. Transport conditions
Frozen ☐ Controlled temperature ☐ Ambient ☐ Chilled ☐

I.17. Accompanying documents
Accompanying document reference
Date of issue
Country
Place of issue

I.19. Container No / Seal No

I.20. Certified as
Production of petfood ☐ Animal Feedingstuff ☐ Technical use ☐

I.21. For transit through a third country ☐
Country ISO Code
EU Exit Authority BCP code
EU Entry Authority BCP code

I.22. For transit through Member State(s) ☐
Country ISO Code

I.25. Total net weight

I.25. Total gross weight

I.28. Description of consignment

1. 23 RESIDUES AND WASTE FROM THE FOOD INDUSTRIES; PREPARED ANIMAL FODDER

2309 Preparations of a kind used in animal feeding

#1. Commodity	Nature of commodity	Manufacturing plant	Net weight	Batch number
Species				

Part II: Certification

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 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:

II.1. the processed animal protein or product described above contains exclusively processed animal protein not intended for human consumption that:

(a) has been prepared and stored in an establishment or plant approved and supervised by the competent authority in accordance with Article 24 of Regulation (EC) No 1069/2009, and

(b) has been prepared exclusively with the following animal by-products:

(1) ☐ 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 retained EU law, but are not intended for human consumption for commercial reasons;]

(1) ☐ 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 retained EU law:

(i) carcases or bodies and parts of animals which are rejected as unfit for human consumption in accordance with retained EU law, 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;]

(1) ☐ 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 retained EU law;]

(1) ☐ 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;]

(1) ☐ 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 arise;]

(1) ☐ 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;]

(1) ☐ 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;]

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

(1) ☐ 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,

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II. Health information

- eggs,
- egg by-products, including egg shells;

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

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

(1) ☐ and/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) and Category 2 material as referred to in Article 9(a) to (g) of Regulation (EC) 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 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 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) ○ 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 = 1O, 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) ☐ [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 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 from:

Part II: Certification

II. Health information

(1) ○ either [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 BSE risk in accordance with Decision 2007/453/EC.]]

(1) ○ 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;

[(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,

(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 processed animal protein or product described above:

(1) ○ either [does not contain milk or milk products of ovine or caprine animal origin or is not intended for feed 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) are derived from ovine and caprine animals which have been kept continuously since birth 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 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 diagnosed 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 killed and destroyed or slaughtered, except for breeding rams of the ARR/ARR genotype, breeding ewes carrying at least one ARR allele and no VRQ allele and other ovine animals carrying at least one ARR allele;]

(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

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II. Health information

Regulation (EC) No 999/2001, of all of the following animals which are over the age of 18 months, except ovine animals of the ARR/ARR genotype:

- animals which have been slaughtered for human consumption; and
- animals which have died or been killed on the holding but which were not killed in the framework of a disease eradication campaign.]]

II.8. the processed animal protein or product described above contains or is derived from animal-by products of non-ruminant origin and is, according to the statement of the Consignor referred to in Box I.1,

(1) ○ either [not intended for the production of feed for farmed animals, other 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 countries subject to the transitional import arrangements include: an EU member State; Liechtenstein; Norway; Iceland, Switzerland and the Faroe Islands.

References to European Union legislation within this certificate are references to direct EU legislation which has been retained in Great Britain (retained EU law as defined in the European Union (Withdrawal) Act 2018).

References to Great Britain in this certificate include Channel 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 the transitional staging period.
- Box reference I.19: use the appropriate HS code: 05 05; 05 06: 05.07; 05.11: 23 01 or 23 09.
- 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. In the case of farmed fish, specify the scientific name of the fish.

Part II:

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II. Health information

(1) Delete as appropriate.

(2) 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

(3) 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 different colour to that of the printing.
- Note for the person responsible for the consignment in Great Britain, Channel Islands or Isle of Man: This certificate is only for veterinary purposes and must accompany the consignment until it reaches the border control post.

Certifying Officer

Name (in capital letters) | Qualification and title

Date of signature | Signature

Stamp