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
Ambient ☐ Chilled ☐ Controlled temperature ☐ Frozen ☐

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 ☐ Pharmaceutical use ☐ Fattening ☐ Breeding and production ☐
Other ☐ Production of petfood ☐ Human consumption ☐ Slaughter ☐
Breeding ☐ Relaying ☐ Animal Feedingstuff ☐ Artificial reproduction ☐
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.23. Total number of packages	I.24. Total quantity	I.25. Total net weight	I.25. Total gross weight

I.28. Description of consignment

1. 05 PRODUCTS OF ANIMAL ORIGIN, NOT ELSEWHERE SPECIFIED OR INCLUDED
0511 Animal products not elsewhere specified or included; dead animals of Chapter|1|or 3, unfit for human consumption
051199 Other

Commodity	Species	Quantity	Net weight	Package count

Identification number	Identification system

Part II: Certification

II. Health information

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 Commission Regulation (EU) No 142/2011 and certify that the blood products described above:

II.1. consist of blood products that satisfy the health requirements below;

II.2. consist exclusively of blood products not intended for human consumption;

II.3. have been prepared and stored in a plant, approved and supervised by the competent authority in accordance with article 24 of Regulation (EC) No 1069/2009;

II.4. have been prepared exclusively with the following animal by-products:

(2) ☐ either [blood of slaughtered animals, which is fit for human consumption in accordance with retained EU law, but which is not intended for human consumption for commercial reasons;]

(2) ☐ and/or [blood of slaughtered animals, which has been rejected as unfit for human consumption in accordance with retained EU law, but which did not show any signs of diseases communicable to humans or animals, which has been derived from carcases that have been slaughtered in a slaughterhouse and which were considered fit for human consumption following an ante-mortem inspection in accordance with retained EU law;]

II.5. in order to inactivate pathogenic agents, have been submitted

(2) ○ either [to processing in accordance with processing method (3) as set out in Chapter III of Annex IV to Regulation (EU) No 142/2011;]

(2) ○ or [to a method and parameters which ensure that the product complies with the microbiological standards set out in Chapter I of Annex X to Regulation (EU) No 142/2011;]

(2) ○ or [in the case of blood products, including spray dried blood and blood plasma, of porcine origin intended for the feeding of porcine animals, to a heat treatment at a temperature of at least 80°C throughout the substance and the dry blood and blood plasma does not contain more than 8% w/w moisture with a water activity (Aw) of less than 0,60.]

II.6. the end product was:

(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 disinfectant approved by the competent authority before use,]

and which bear labels indicating ‘NOT FOR HUMAN CONSUMPTION’;

II.7. the end product was stored in enclosed storage;

II.8. the product has undergone all precautions to avoid contamination with pathogenic agents after treatment;

(2) and ☐ [in the case of blood products, including spray dried blood and blood plasma of porcine origin intended for the feeding of porcine animals, has been stored in dry warehouse conditions under room temperature for a period of at least 6 weeks.]

II.9. have been examined prior to dispatch under the responsibility of the competent authority by taking a random sample during or on removal from storage which was found to comply with the following standards (4):

Salmonella:	absence in 25g: n = 5, c = 0, m = 0, M = 0,
Enterobacteriaceae	n=5, c=2, m=10, M = 300 in 1 gram

(2) ☐ [II.10. the blood products described above

(2) ○ either [is derived from other ruminants than bovine, ovine or caprine animals.]]

Part II: Certification

II. Health information

(2) ○ or [is derived from bovine, ovine or caprine animals and does not contain and is not derived from:

(2) ○ 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.]]

(2) ○ 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.11. the blood products described above:

(2) ○ either [do 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.]

(2) ○ or [contain milk or milk products of ovine or caprine animal origin and is intended for feed for farmed animals, other than fur animals, which:

(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 Organization 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 the period of at least the preceding seven years or, following the confirmation of a case of classical scrapie:

Part II: Certification

II. Health information

(2)	○ 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;]
(2)	○ 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 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.12. the blood products described above contain or are derived from animal-by products of non-ruminant origin, and are, according to the statement of the Consignor referred to in Box I.1,

(2)	○ either	[not intended for the production of feed for farmed animals, other than fur animals.]
(2)(7)	○ 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.]

Part II: Certification

II. Health information

Notes

(*) Those countries subject to the transitional import arrangements include: an EU member State; Liechtenstein; Norway; Iceland and Switzerland.

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 that is 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 a 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 case of unloading and reloading in Great Britain, Channel Islands or Isle of Man.
- Box reference I.16: Box I.16: do not use this box until the end of the transitional staging period.
- Box reference I.19: use the appropriate HS code: 05.11.91, 05.11.99, 35.02 or 35.04
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) should be included.
- 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, Reptilia.

Part II

(2) Delete as appropriate.

(3) Insert method 1 to 5 or method 7 as applicable.

(4) 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.

Part II: Certification

II. Health information

(7) The person responsible for the load referred to in Box I.6 must ensure that, if the blood products described in this health certificate are 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 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 of the point of entry into Great Britain, Channel Islands or Isle of Man.

Certifying Officer

Name (in capital letters)	Qualification and title
Date of signature	Signature
Stamp	