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
Pharmaceutical use ☐ Other ☐ Slaughter ☐ Relaying ☐
Artificial reproduction ☐ Fattening ☐ Production ☐ Breeding ☐
Production of petfood ☐ Breeding and production ☐ Animal Feedingstuff ☐ Human consumption ☐
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. 35 ALBUMINOIDAL SUBSTANCES; MODIFIED STARCHES; GLUES; ENZYMES

3502 Albumins (including concentrates of two or more whey proteins, containing by weight more than 80|% whey proteins, calculated on the dry matter), albuminates and other albumin derivatives

#1.	Commodity	Quantity	Net weight	Package count
Species		Identification number	Identification system	

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 8 (c) and Article 8(d) and Article 10 thereof, and Commission Regulation (EU) No 142/2011, and in particular Chapter II of Annex XIV thereto, and certify that:

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

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

II.3. they have been prepared and stored in a plant supervised by the competent authority, 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 is not intended for human consumption for commercial reasons;]

(2) ☐ and/or [- blood of slaughtered animals, which is 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, derived from carcasses that have been slaughtered in a slaughterhouse and were considered fit for human consumption following an ante-mortem inspection in accordance with retained EU law;]

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

(2) ☐ and/or [- blood and blood products originating from live animals that did not show clinical signs of any disease communicable through these products to humans or animals;]

(2) ☐ and/or [- blood and blood products derived from the production of products intended for human consumption;]

(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 Council Directive 96/22/EC or Article 2(b) of Council 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 retained EU law or, in the absence thereof, in national legislation;]

II.4. the blood that these products were manufactured from has been collected in slaughterhouses approved in accordance with retained EU law, in slaughterhouses approved and supervised by the competent authority of the country of collection or from live animals in facilities approved and supervised by the competent authority of the country of collection.

(2) ☐ [II.5. In the case of blood products derived from Artiodactyla, Perissodactyla and Proboscidea including their crossbreeds, other than Suidae and Tayassuidae, the products have undergone one of the following treatments, guaranteeing the absence of pathogens of foot-and-mouth disease, vesicular stomatitis, rinderpest, peste des petits ruminants, Rift Valley fever and bluetongue:

(2) ☐ either [heat treatment at a temperature of 65°C for at least three hours, followed by an effectiveness check;]

(2) ☐ and/or [irradiation at 25 kGy by gamma rays, followed by an effectiveness check;]

(2) ☐ and/or [change in pH to pH 5 for two hours, followed by an effectiveness check;]

(2) ☐ and/or [heat treatment of at least 80 °C throughout their substance, followed by an effectiveness check]]

(2) ☐ [II.6. In the case of blood products derived from Suidae, Tayassuidae, poultry and other avian species, the products have undergone one of the following treatments guaranteeing the absence of pathogens of the following diseases: foot-and-mouth disease, vesicular stomatitis, swine vesicular disease, classical swine fever, African swine fever, Newcastle disease and highly pathogenic avian influenza, as appropriate to the species:

(2) ☐ either [heat treatment at a temperature of 65°C for at least three hours, followed by an effectiveness check;]

(2) ☐ and/or [irradiation at 25 kGy by gamma rays, followed by an effectiveness check;]

(2) ☐ and/or [heat treatment of at least 80°C for Suidae/Tayassuidae (2) and at least 70°C for poultry and

Part II: Certification

II. Health information

other avian species (2) throughout the substance of the product, followed by an effectiveness check]].

(2) ☐ [II.7. In the case of blood products derived from species other than those listed in point II.5 or II.6, the products have undergone the following treatment (please specify): .]

II.8. The products were:

(2) ○ either [packed in new or sterilised bags or bottles,]

(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 the outer packaging or containers bear labels indicating 'NOT FOR HUMAN OR ANIMAL CONSUMPTION';

II.9. the products were stored in enclosed storage;

II.10. all precautions were taken to avoid the contamination of the products with pathogenic agents after treatment;

(2) ☐ [II.11. The treated blood products described above

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

(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, 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.]]]

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 to be transited through Great Britain, Channel Islands or Isle of Man; it may be filled in if the certificate is for a commodity to be imported into Great Britain, Channel Islands or Isle of Man.
- Box reference Approval number; the registration number of the establishment or plant, which has been issued by the competent authority.

Part II: Certification

II. Health information

I.11 and I.12:

- 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) is to be provided. In the case of unloading and reloading in Great Britain, Channel Islands or Isle of Man, the consignor must inform the BCP of entry into Great Britain, Channel Islands or Isle of Man.
- Box reference I.16: Do not use this box until the end of the transitional staging period.
- Box reference I.19: use the appropriate Harmonized System (HS) code under the following headings: 05.11, 30.02, 35.02 or 35.04.
- Box reference I.23: for bulk containers, the container number and the seal number (if applicable) must 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 in case of Species: select from the following: Aves, Ruminantia, Suidae, Mammalia other than Ruminantia or Suidae, Pesca, Reptilian.

Part II:

(2) Delete as appropriate.

- 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 Great Britain, Channel Islands or Isle of Man.

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