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 health 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 or in the establishment of collection, 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 carcases 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 derived from the production of products intended for human consumption;]

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

(2) ☐ and/or [- animal by-products 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 level laid down in retained EU law or, in the absence thereof, in national legislation;]

II.4. the blood, that such products were manufactured from, was 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 obtained from animals belonging to the taxa Artiodactyla, Perissodactyla and Proboscidea, including crossbreds between species of those taxa, the blood was collected in a country or region where no case of rinderpest, peste des petits ruminants and Rift Valley fever has been recorded for a period of at least the preceding 12 months and in which vaccination has not been carried out against those diseases for a period of at least the preceding 12 months, and;

(2) ○ either [in third countries, territories or parts thereof (insert ISO country code in the case of a country, or codes (3) in the case of territories or parts thereof) where no case of foot-and-mouth disease has been recorded for a period of at least the preceding 12 months and in which vaccination has not been carried out against this disease for a period of at least the preceding 12 months, and]

(2) ○ or [in third countries, territories or parts thereof (insert ISO country code in the case of a country or codes (3) for territories or parts thereof) where no case of foot-and-mouth disease has been recorded for a period of at least the preceding 12 months and in which vaccination programmes against foot-and-mouth disease are being officially carried out and controlled in domestic ruminant animals for a period of at least the preceding 12 months (4), and]]

(2) ☐ [II.5.1. in the case of animals other than Suidae and Tayassuidae, in third countries or regions in which:

(2) ○ either [no case of vesicular stomatitis and bluetongue (2) (including the presence of

Part II: Certification

II. Health information

seropositive animals) has been recorded for a period of at least the preceding 2 months and in which vaccination has not been carried out against those diseases for a period of at least the preceding 12 months;]

(2) ○ or [vesicular stomatitis and bluetongue (2) seropositive animals are present(4);]]]]

(2) ☐ [II.5.2. in the case of Suidae and Tayassuidae, in third countries or regions in which no case of swine vesicular disease, classical swine fever and African swine fever has been recorded for a period of at least the preceding 12 months and vaccination has not been carried out against those diseases for a period of at least the preceding 12 months in the susceptible species and:

(2) ○ either [no case of vesicular stomatitis (including the presence of seropositive animals) has been recorded for a period of at least the preceding 12 months and in which vaccination has not been carried out against this disease for a period of at least the preceding 12 months;]]

(2) ○ or [vesicular stomatitis seropositive animals are present(4);]]]

(2) ☐ [II.6. in the case of blood products derived from poultry or other avian species the animals and the products come from the territory of the country or region with code (5)which has been free from Newcastle disease and highly pathogenic avian influenza as defined in the Terrestrial Animal Health Code of the OIE, which for a period of at least the preceding 12 months has not carried out vaccination against avian influenza, where the animals from which the products are derived, have not been vaccinated against Newcastle disease with vaccines prepared from a Newcastle disease master strain showing a higher pathogenicity than lentogenic virus strains;]

II.7. 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,]

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

II.8. the products were stored in enclosed storage;

II.9. all precautions were taken to avoid contamination of the products with pathogenic agents during transport;

(2) ☐ [II.10. the untreated 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, 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

Part II: Certification

II. Health information		

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 l:

-	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.11 and I.12:	Approval number: the registration number of the establishment or plant, which has been issued by the competent authority.
-	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 border control post of the point 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 or 35.02.
-	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 Species:	select from the following: Aves, Ruminantia, Suidae, Mammalia, other than Ruminantia or Suidae, Pesca, Reptilian.

Part II:

(2)	Delete as appropriate.
(3)	Code of the territory as it appears in Part 1 of Annex II to Regulation (EU) No 206/2010
(4)	In this case following the veterinary checks provided for in Directive 97/78/EC, and in accordance with the conditions laid down in Article 8(4) of that Directive, the products must be transported directly to the establishment at the place of destination.
(5)	Code of the territory as it appears in Part 1 of Annex I to Commission Regulation (EC) No 798/2008

Part II: Certification

II. Health information		

- 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	