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
Chilled ☐ Ambient ☐ Frozen ☐ Controlled temperature ☐

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

I.19. Container No / Seal No

I.20. Certified as
Relaying ☐ Technical use ☐ Pharmaceutical use ☐ Slaughter ☐
Breeding and production ☐ Fattening ☐ Breeding ☐ Production ☐
Animal Feedingstuff ☐ Human consumption ☐ Artificial reproduction ☐ Other ☐
Production of petfood ☐

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

3504 Peptones and their derivatives; other protein substances and their derivatives, not elsewhere specified or included; hide powder, whether or not chromed

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

Part II: Certification

II. Health information

II.1. Animal Health Attestation

I, the undersigned official veterinarian, declare that I am aware of the relevant provisions of Directive 2002/99/EC and of Regulation (EC) No 853/2004 and hereby certify that the ☐ colostrum/ ☐ colostrum-based products (1) described in Part I:

have been obtained or manufactured from colostrum obtained from animals:

(i) under the control of the official veterinary service;

(ii) which were in a third country or part thereof that has been free of foot-and-mouth disease and of rinderpest for a period of at least 12 months prior to the date of this certificate, and where vaccination against foot-and-mouth disease has not been carried out during that period;

(iii) belonging to holdings which were not under restrictions due to foot-and-mouth disease or rinderpest, and

(iv) subject to regular veterinary inspections to ensure that they satisfy the animal health conditions laid down in Chapter I of Section IX of Annex 3 to Regulation (EC) No 853/2004 and in Directive 2002/99/EC.

II.2. Public Health Attestation

I, the undersigned official inspector, declare that I am aware of the relevant provisions of Regulations (EC) No 178/2002, (EC) No 852/2004, (EC) No 853/2004 and (EU) 2019/627 and hereby certify that the ☐ colostrum/ ☐ colostrum-based products made with colostrum(1) described in Part I were produced in accordance with those provisions, and in particular that:

(a) they were manufactured from colostrum:

(i) which comes from holdings registered in accordance with Regulation (EC) No 852/2004 and checked in accordance with Articles 49 and 50 of 2019/627;

(ii) which was produced, collected, cooled, stored and transported in accordance with the hygiene conditions laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004;

(iii) which complies with the guarantees on the residues status of colostrum provided by the monitoring plans for the detection of residues or substances submitted in accordance with Directive 96/23/EC, and in particular, Article 29 thereof;

(iv) which, pursuant to testing for residues of antibacterial drugs carried out by the food business operator in accordance with the requirements of point 4 in Part III of Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004, complies with the maximum residue limits for residues of antibacterial veterinary medicinal products laid down in the Annex to Regulation (EU) No 37/2010;

(v) which has been produced under conditions guaranteeing compliance with the maximum residue levels for pesticides laid down in Regulation (EC) No 396/2005, and maximum levels for contaminants laid down in Regulation (EC) No 1881/2006;

(b) they come from an establishment implementing a programme based on the HACCP principles in accordance with Regulation (EC) No 852/2004;

(c) they have been processed, stored, wrapped, packaged and labelled in accordance with Chapters III and IV of Section IX of Annex III to Regulation (EC) No 853/2004;

(d) they meet the relevant requirements laid down in Chapter II, Section IX of Annex III to Regulation (EC) NO 853/2004 and the relevant microbiological criteria laid down in Regulation (EC) No 2073/2005 on microbiological criteria for foodstuffs, and

(e) the guarantees covering live animals and products thereof provided by the residue plans submitted in accordance with Directive 96/23/EC, and in particular Article 29 thereof are fulfilled.

Notes

Part II: Certification

II. Health information

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) and can be viewed on the UK legislation website (legislation.gov.uk).

References to Great Britain in this certificate include Channel Islands and Isle of Man.

This certificate is intended for colostrum or colostrum-based products from third countries or parts thereof authorised in column A as set out in a document relating to 'milk and milk products' published on gov.uk in accordance with Regulation (EU) No 605/2010. (2)

Part I:

Box reference I.7:	Provide name and ISO code of the country or part thereof as appearing in a document relating to 'milk and milk products' published on gov.uk in accordance with Commission Regulation (EU) No 605/2010 . (2)

Box reference I.11: Name, address and approval number of the establishment of dispatch.

Box reference I.15: Registration number (railway wagons or container and road vehicle), flight number (aircraft) or name (ship). In case of unloading and reloading, the consignor must inform the border control post of introduction into Great Britain.

Box reference I.19:	For containers or boxes, the container number and the seal number (if applicable) should be included.
Box reference I.25:	Indicate total gross weight and total net weight.
Box reference I.28:	Use the appropriate Harmonised System (HS) code under the following headings: 04.01; 04.02; 04.03; 04.04; 04.05; 04.06; 04.10; 15.17; 17.02; 19.01; 21.05; 21.06; 22.02; 28.35; 30.01; 35.01; 35.02 or 35.04.
Box reference I.28:	Manufacturing plant: introduce the approval number of the production holding(s), collection centre or standardization centre approved for exportation to Great Britain.

Part II:

(1)	Keep as appropriate.
(2)	A document relating to 'milk and milk products' for EU and EFTA states published by the Secretary of State, with the consent of the Scottish and Welsh Ministers, may be found here:

EU and EFTA states approved to export animals and animal products to Great Britain - data.gov.uk

The colour of the signature shall be different to that of the printing. The same rule applies to stamps other than those embossed or watermark.

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