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

2309 Preparations of a kind used in animal feeding

230990 other than 2309 10

#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 10 thereof, and Commission Regulation (EU) No 142/2011 , and in particular Section 4 of Chapter II of Annex X and Chapter I of Annex XIV thereto, and certify that the colostrum (2) or the colostrum products (2) referred to in box I.28 comply with the following conditions:

II.1. they were produced and derived in … (insert name of exporting country) (3), … (insert name of region) (3), which is listed in Annex I to Commission Regulation (EU) No 605/2010 , and which has been free from foot-and-mouth disease (FMD) and rinderpest for a period of 12 months immediately prior to export and has not practised vaccination against rinderpest uring that period;

II.2. they were produced from colostrum derived from animals which at the time of milking did not show clinical signs of any disease transmissible through colostrum to humans or animals, and which had been kept for a period of at least 30 days prior to the date of production on holdings that were not subject to official restrictions due to foot-and-mouth disease or rinderpest;

II.3. they are colostrum or colostrum products of bovine animals that have been subject to high temperature short time pasteurisation in 72°C for at least 15 seconds, or an equivalent pasteurisation achieving a negative reaction to a phosphatase test in bovine colostrum, in combination with:

(2)(5) ○ either [the condition that the colostrum or colostrum products have been produced during a period at least 21 days before the date of shipping and during this period no cases of FMD have been detected in the exporting country,]

(2)(5) ○ or [the condition that the colostrum or colostrum products have been produced on (insert the date), this date, in consideration of the foreseen voyage duration, being at least 21 days before the consignment is presented to a border control post of the point of entry into Great Britain, Channel Islands or Isle of Man Great Britain ,]

and have been obtained from animals subject to regular veterinary inspections to ensure that they come from holdings on which all bovine herds are:

(2)(5) ○ either [recognised as officially tuberculosis and brucellosis free (6),]

(2)(5) ○ or [not restricted under the national legislation of the third country of origin for the eradication of tuberculosis and brucellosis,]

and (2)(5) ○ either [recognised as official enzootic-bovine-leukosis-free (6),]

(2)(5) ○ or [included in an official system for the control or enzootic bovine leukosis and there has been no evidence as result of clinical and laboratory testing of this disease in the herd during the period of the preceding two years,]]

II.4. every precaution has been taken to avoid contamination of the colostrum/colostrum product after processing;

II.5. the colostrum or colostrum product was packed:

(2) ○ either [in new containers,]

(2) ○ or [in vehicles or bulk containers disinfected prior to loading using a product approved by the competent authority,]

and the containers are marked so as to indicate the nature of the olostrum/colostrum product and bear labels indicating that the product is Category 3 material and not intended for human consumption;

II.6. the colostrum or colostrum product does not contain milk or milk products of ovine or caprine animal origin.

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 II: Certification

II. Health information

Part I:

- Box reference I.6: Person responsible for the load 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 I.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity.
- 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 Harmonised System (HS) code of the World Customs Organisation: 04.04.90; 23.09.10, 23.09.90, 35.01, 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 or 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: 'Manufacturing plant': provide the registration number of the treatment or processing establishment.

Part II:

(2) Delete as appropriate.

(3) For completion if the authorisation for introduction into Great Britain, Channel Islands or Isle of Man is restricted to certain regions of the third country concerned.

(5) This condition applied only to third countries authorised in column 'A' of Annex I to Commission Regulation (EU) No 605/2010 .

(6) Officially tuberculosis-free and brucellosis-free herd as laid down in Annex A to Council Directive 64/432/EEC and officially enzootic-bovine-leukosis-free herd as laid down in Chapter I of Annex D to that Directive.

- The signature and the seal must be a different colour from that of the printing.
- Note for the importer: this certificate is only for veterinary purposes and must accompany the consignment until it reaches the border control post of 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	