Part I : Details of dispatched consignment

I.1. Consignor
Name
Address

Tel

I.2. Certificate reference number

I.2.a.

I.3. Central Competent Authority
DEPARTMENT FOR ENVIRONMENT, FOOD AND RURAL AFFAIRS

I.4. Local Competent Authority
ANIMAL & PLANT HEALTH AGENCY

I.5. Consignee
Name
Address

Postal Code
Tel

I.6. Person responsible for the load in the EU
Address

Postal Code
Phone

I.7. Country of origin, ISO code
UNITED KINGDOM GB

I.8. Region of origin, Code

I.9. Country of destination

ISO code

I.10. Region of destination

Code

I.11. Place of origin
Name

Approval number

Address

I.12. Place of destination

Custom warehouse ☐

Name
Address

Postal Code
Approval number

I.13. Place of loading

I.14. Date of departure

I.15 Means of transport

Aeroplane ☐ Ship ☐ Railway wagon ☐
Road vehicle ☐ Other ☐

Identification:
Documentary references:

I.16. Entry BIP in EU

I.17. No.(s) of CITES

I.18. Description of commodity

I.19. Commodity code (HS code)

I.21. Temperature of products

Ambient ☐ Chilled ☐ Frozen ☐

I.20. Quantity

I.22. Number of packages

I.23. Seal/Container No.

I.24. Type of packaging

I.25. Commodity certified for:

Animal feedingstuff ☐ Further process ☐ Production of petfood ☐ Technical use ☐

I.26. For transit to 3rd country by EU ☐

3rd country

ISO code

I.27. For import or admission into EU ☐

I.28. Identification of the commodities

Approval number of establishments

Species (Scientific name) | Manufacturing plant | Net weight | Batch number

specimen

COUNTRY: UNITED KINGDOM

Part II: Certification

II. Health information	II.a. Certificate reference no	II.b.

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 ([1a]), and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011 ([1b]), 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 ([4]), 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 during 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 at 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 inspection post of the European Union,]

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 of 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 colostrum/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.

Specimen

Notes

Part I:

— Box reference I.6: Person responsible for the load in the European Union: this box is required to be filled in only if it is a certificate for a commodity to be transited through the European Union; it may be filled in if the certificate is for a commodity to be imported into the European Union.

II. Health information	II.a. Certificate reference no	II.b.

— 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 the European Union, the consignor must inform the border inspection post of the European Union.

— 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 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: 'Manufacturing plant': provide the registration number of the treatment or processing establishment.

Part II:

([1a]) OJ L 300, 14.11.2009, p. 1.

([1b]) OJ L 54, 26.2.2011, p. 1.

([2]) Delete as appropriate.

([3]) For completion if the authorisation for introduction into the European Union is restricted to certain regions of the third country concerned.

([4]) OJ L 175, 10.7.2010, p. 1.

([5]) This condition applies only to third countries authorised in column 'A' of Annex I to Commission Regulation (EU) No 605/2010 (OJ L 175, 10.7.2010, p. 1).

([6]) Officially tuberculosis-free and brucellosis-free herd as laid down in Annex A to Council Directive 64/432/EEC (OJ 121, 29.7.1964, p. 1977/64) 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 in 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 inspection post of the European Union.

Official veterinarian/Official inspector

Name (in capital letters): Qualification and title:

Date: Signature:

Stamp:

Specimen

Specimen