

Country

Official certificate to the EU

Part I : Details of consignment

I.1. Consignor Name Address Country ISO Code		I.2. Certificate reference I.3. Central competent authority I.4. Local competent authority		I.2.a. IMSOC reference Specimen not to be used for imports into the EU	
I.5. Consignee Name Address Country ISO Code		I.6. Responsible for the consignment in EU Name Address Country ISO Code			
I.7. Country of origin ISO Code	I.8. Region of origin Code	I.9. Country of destination ISO Code	I.10. Region of destination Code		
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 Ambient <input type="checkbox"/> Chilled <input type="checkbox"/> Frozen <input type="checkbox"/>		I.17. Accompanying documents Type Number			
I.19. Container No / Seal No					
I.20. Certified as Relaying area/ purification centre <input type="checkbox"/> Products for human consumption <input type="checkbox"/> Other <input type="checkbox"/> Fattening <input type="checkbox"/> Traditional samples <input type="checkbox"/> Slaughter <input type="checkbox"/> Artificial reproduction <input type="checkbox"/> Further processing <input type="checkbox"/> Pets <input type="checkbox"/> Approved Bodies <input type="checkbox"/> Quarantine establishment <input type="checkbox"/> Feed stuff <input type="checkbox"/> Manufacture of petfood <input type="checkbox"/> Registered equine animal <input type="checkbox"/> Traveling circus/animal acts <input type="checkbox"/> Breeding/production <input type="checkbox"/> Pharmaceutical use <input type="checkbox"/> Further keeping <input type="checkbox"/> Production of petfood <input type="checkbox"/> Canning industry <input type="checkbox"/> Petfood <input type="checkbox"/> Technical Use <input type="checkbox"/> Game restocking <input type="checkbox"/>					
I.21. For transit <input type="checkbox"/> Non-EU ISO Code		I.22. For internal market <input type="checkbox"/> I.23. For re-entry <input type="checkbox"/>			
I.24. Total number of packages	I.25. Total quantity	I.26. Total net weight	I.26. Total gross weight		
I.27. Description of consignment 1. 04 DAIRY PRODUCE; BIRDS' EGGS; NATURAL HONEY; EDIBLE PRODUCTS OF ANIMAL ORIGIN, NOT ELSEWHERE SPECIFIED OR INCLUDED 0401 Milk and cream, not concentrated nor containing added sugar or other sweetening matter					
Commodity	Species	Cold store	Identification mark	Package count	
Net weight	Treatment type	Nature of commodity	Batch number	Final consumer	
Date of collection/production		Manufacturing plant		Plant / Establishment / Centre	

II. Health information

SPECIMEN

II. Health information

II.1. Public health attestation ○ [to delete when the Union is not the final destination of the colostrum]

I, the undersigned, declare that I am aware of the relevant requirements of Regulation (EC) No 178/2002 of the European Parliament and of the Council, Regulation (EC) No 852/2004 of the European Parliament and of the Council, Regulation (EC) No 853/2004 of the European Parliament and of the Council, Regulation (EU) 2017/625 of the European Parliament and of the Council and Commission Implementing Regulation (EU) 2019/627 and hereby certify that the colostrum(2) described in Part I was produced in accordance with these requirements, and in particular that:

(a) colostrum:

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

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

(iii) comes from animals belonging to herds free or officially free of brucellosis and tuberculosis;

(iv) pursuant to testing for residues of antibacterial drugs carried out by the food business operator in accordance with the requirements of point 4 in Section IX, Chapter I, Part III, 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 Commission Regulation (EU) No 37/2010 ;

(b) it comes from (an) establishment(s) applying general hygiene requirements and implementing a programme based on the hazard analysis and critical control points (HACCP) principles in accordance with Article 5 of Regulation (EC) No 852/2004, regularly audited by the competent authorities, and being listed as an EU approved establishment;

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

(d) it meets the relevant criteria laid down in Section IX, Chapter II, of Annex III to Regulation (EC) No 853/2004 and the relevant microbiological criteria laid down in Commission Regulation (EC) No 2073/2005 ;

(e) it 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 Article 29 of Council Directive 96/23/EC, and milk is listed in Commission Decision 2011/163/EU for the concerned country of origin;

Part II : Certification

II. Health information

(f) it has been produced under conditions guaranteeing compliance with the maximum residue levels for pesticides laid down in Regulation (EC) No 396/2005 of the European Parliament and of the Council, and the maximum levels for contaminants laid down in Commission Regulation (EC) No 1881/2006.

II.2. Animal health attestation
 ○ [to delete when the colostrum is derived from solipeds, leporidae or other wild land mammals others than ungulates]

The colostrum(2) described in Part I:

II.2.1. has been obtained in the zone/s with code/s: (3) which, at the date of issue of this certificate is/are authorised for entry into the Union of colostrum and listed in Part 1 of Annex XVII to Commission Implementing Regulation (EU) 2021/404, and in which foot and mouth disease and infection with rinderpest virus have not been reported for a 12 month period before the date of obtaining the colostrum, and during the same period vaccination against these diseases has not been carried out;

II.2.2. has been obtained from animals of the species ☐ [Bos Taurus,] (1) ☐ [Ovis aries,] (1) ☐ [Capra hircus,] (1) ☐ [Bubalus bubalis,] (1) ☐ [Camelus dromedarius] (1) that have remained in the zone/s referred to under point II.2.1 since birth, or for at least 3 months before the date of obtaining the colostrum;

II.2.3. has been obtained from animals coming from establishments:

- (a) registered by and under the control of the competent authority of the third country or territory and have a system in place to maintain and to keep records in accordance with Article 8 of Commission Delegated Regulation (EU) 2020/692 ;

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II. Health information

- (b) which receive regular animal health visits from a veterinarian for the purpose of the detection of, and information on, signs indicative of the occurrence of diseases, including the relevant listed diseases referred to in Annex I to Commission Delegated Regulation (EU) 2020/692 and emerging diseases;
- (c) which were not subject to national restriction measures for animal health reasons, including the relevant listed diseases referred to in Annex I to Delegated Regulation (EU) 2020/692 and emerging diseases, at the time of obtaining the colostrum.

Notes

In accordance with the Agreement on the withdrawal of the United Kingdom of Great Britain and Northern Ireland from the European Union and the European Atomic Energy Community, and in particular Article 5(4) of the Protocol on Ireland / Northern Ireland in conjunction with Annex 2 to that Protocol, references to European Union in this certificate include the United Kingdom in respect of Northern Ireland.

This certificate is intended for entry into the Union of colostrum, including when the Union is not the final destination of such colostrum.

This animal health/official certificate shall be completed according to the notes for the completion of certificates provided for in Chapter 4 of Annex I to Implementing Regulation (EU) 2020/2235.

Part I:

Box reference Provide the code of the zone as appearing in column 2 of the table in Part 1 of Annex XVII to Implementing Regulation (EU) 2021/404.

I.8:

Part II:

- (1) Keep as appropriate.
- (2) Colostrum as defined in Section IX, Point 1, of Annex III to Regulation (EC) No 853/2004.
- (3) Code of the zone in accordance with column 2 of the table in Part 1 of Annex XVII to Implementing Regulation (EU) 2021/404.
- (4) to be signed by:
- an official veterinarian when part II.2 Animal health attestation is not deleted
 - a certifying officer or an official veterinarian when part II.2 Animal health attestation is deleted

Part II : Certification

II. Health information			
Official veterinarian or Official inspector			
Name (in capital letters)		Qualification and title	
Date of signature		Signature	
Stamp			
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