

UNITED KINGDOM

Animal health/Official certificate to the EU

Part I: Description of consignment	I.1 Consignor/Exporter		I.2 Certificate reference		I.2a	
	Name					
	Address		I.3 Central Competent Authority			
			DEPARTMENT FOR ENVIRONMENT, FOOD & RURAL AFFAIRS			
	Country		ISO country code		I.4 Local Competent Authority	
					ANIMAL AND PLANT HEALTH AGENCY	
	I.5 Consignee/Importer		I.6 Operator responsible for the consignment			
	Name		Name			
	Address		Address			
	Country		ISO country code		Country	
					ISO country code	
	I.7 Country of origin		ISO country code		I.9 Country of destination	
					ISO country code	
	I.8 Region of origin		Code		I.10 Region of destination	
					Code	
	I.11 Place of dispatch		Registration/Approval No		I.12 Place of destination	
					Registration/Approval No	
	Name		Name			
	Address		Address			
	Country		ISO country code		Country	
					ISO country code	
	I.13 Place of loading		I.14 Date and time of departure			
	I.15 Means of transport		I.16 Entry Border Control Post			
	☐ Aircraft ☐ Vessel					
	☐ Railway ☐ Road vehicle					
	Identification		I.17 Accompanying documents			
			Type			
			Code			
			Country			
			ISO country code			
			Commercial document reference			
	I.18 Transport conditions		☐ Ambient ☐ Chilled ☐ Frozen			
	I.19 Container number/Seal number					
	Container No			Seal No		
	I.20 Certified as or for					
	☐ Products for human consumption					
	I.21			I.22		
	☐ For transit			☐ For internal market		
	Third country			ISO country code		
				I.23		
	I.24 Total number of packages		I.25		I.26 Total net weight/gross weight (kg)	

UNITED KINGDOM

1.27 Description of consignment							
1	CN Code	Species	Cold store	Type of packaging	Net weight	Treatment type	Final consumer ☐
	Nature of commodity	Number of packages	Batch No	Date of collection/production	Manufacturing plant		
2	CN Code	Species	Cold store	Type of packaging	Net weight	Treatment type	Final consumer ☐
	Nature of commodity	Number of packages	Batch No	Date of collection/production	Manufacturing plant		
3	CN Code	Species	Cold store	Type of packaging	Net weight	Treatment type	Final consumer ☐
	Nature of commodity	Number of packages	Batch No	Date of collection/production	Manufacturing plant		
4	CN Code	Species	Cold store	Type of packaging	Net weight	Treatment type	Final consumer ☐
	Nature of commodity	Number of packages	Batch No	Date of collection/production	Manufacturing plant		
5	CN Code	Species	Cold store	Type of packaging	Net weight	Treatment type	Final consumer ☐
	Nature of commodity	Number of packages	Batch No	Date of collection/production	Manufacturing plant		

UNITED KINGDOM

II.a Certificate reference

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II. Health information⁽¹⁾ **[II.1. Public health attestation** (*Delete when the Union is not the final destination of the colostrum-based products*)

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 and 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-based products ⁽²⁾ described in Part I were produced in accordance with these requirements, and in particular that:

- (a) they were produced 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 Implementing Regulation (EU) 2019/627;
 - (ii) which 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) which comes from animals belonging to herds officially free of tuberculosis and free or officially free of brucellosis;
 - (iv) which complies with the guarantees covering animals and products thereof provided by the control plan submitted in accordance with Article 6(2) of Commission Delegated Regulation (EU) 2022/2292 and the third country or region thereof of its origin is listed in Annex -I to Commission Implementing Regulation (EU) 2021/405 and marked with an "X" for the category "milk";
 - (v) which, pursuant to testing for antibiotic residues carried out by the food business operator in accordance with the requirements of Section IX, Chapter I, Part III, point 4, 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) they come from establishments 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 listed as Union approved establishments;
- (c) they have been processed, stored, wrapped, packaged and labelled in accordance with Section IX, Chapters III and IV, of Annex III to Regulation (EC) No 853/2004;
- (d) they meet 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.]

⁽¹⁾⁽⁷⁾ **[II.1a. Attestation as regards Commission Delegated Regulation (EU) 2023/905** (*Delete when the Union is not the final destination of the colostrum-based products*)

I, the undersigned, declare that I am aware of the relevant requirements of Regulation (EU) 2019/6 of the European Parliament and of the Council and Delegated Regulation (EU) 2023/905 and hereby certify that the colostrum-based products described in Part I were produced in accordance with these requirements, and in particular that the animals from which the colostrum-based products have been obtained have not been administered antimicrobial medicinal products for growth promotion or yield increase or antimicrobial medicinal products containing an antimicrobial that is included in the list of antimicrobials reserved for the treatment of certain infections in humans laid down in Commission Implementing Regulation (EU) 2022/1255 as set out in Article 3 of Delegated Regulation (EU) 2023/905 and originate from a third country or region thereof listed in the Annex to Commission Implementing Regulation (EU) 2024/2598.]

⁽¹⁾ **[II.2. Animal health attestation** (*Delete when the colostrum-based products are derived from solipeds, leporidae or wild land mammals other than ungulates*)The colostrum-based products ⁽²⁾ described in Part I:

- ⁽¹⁾ *either* [II.2.1. originate from the **zone(s)** with code(s) _____ ⁽³⁾ which, at the date of issue of this animal health/official certificate is/are authorised for the entry into the Union of colostrum-based products and is/are 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 the last 12 months before the date of obtaining the colostrum, and vaccination against these diseases has not been carried out during that period;]
- ⁽¹⁾⁽⁴⁾ *or* [II.2.1. originate from the **zone** with code _____ ⁽⁵⁾ which, at the date of issue of this animal health/official certificate is authorised for the transit through the Union of colostrum-based products intended for a destination outside the Union and is listed in Part 1 of Annex XXII to Commission Implementing Regulation (EU) 2021/404;]
- II.2.2. have been processed from **colostrum** obtained in
- ⁽¹⁾ *either* [the zone referred to in point II.2.1;]
- ⁽¹⁾ *or* [the zone(s) with code(s) _____ ⁽³⁾ which, at the date of issue of this animal health/official certificate is/are listed in Part 1 of Annex XVII to Implementing Regulation (EU) 2021/404 for entry into the Union of milk, colostrum and colostrum-based products;]
- ⁽¹⁾ *or* [Member States;]
- II.2.3. have been processed from colostrum 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.2.4. have been processed from colostrum obtained from animals kept in **establishments**:
- (a) which are registered by and are under the control of the competent authority of the third country or territory and have a

UNITED KINGDOM

II.a Certificate reference

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system in place to maintain and to keep records in accordance with Article 8 of Commission Delegated Regulation (EU) 2020/692;

- (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 listed diseases referred to in Annex I to Delegated Regulation (EU) 2020/692 relevant for the species and emerging diseases;
- (c) which were not subject to national restriction measures for animal health reasons, including the listed diseases referred to in Annex I to Delegated Regulation (EU) 2020/692 relevant for the species and emerging diseases, at the date 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 Windsor Framework (see Joint Declaration No 1/2023 of the Union and the United Kingdom in the Joint Committee established by 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 of 24 March 2023, OJ L 102, 17.4.2023, p. 87) in conjunction with Annex 2 to that Framework, references to the Union in this animal health/official certificate include the United Kingdom in respect of Northern Ireland.

This animal health/official certificate is intended for the entry into the Union of colostrum-based products, including when the Union is not the final destination of such products.

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

Part I:

Box reference I.8: 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.

Part II:

- (1) Delete if not applicable.
- (2) “Colostrum-based products” as defined in Section IX, point 2, 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) Only applicable to consignments transiting through the Union and intended for a destination outside the Union, originating from third country or territory, or zone thereof listed in Part 1 of Annex XXII to Implementing Regulation (EU) 2021/404 as authorised for transit through the Union of colostrum-based products accompanied by an animal health certificate corresponding to the present model certificate in accordance with column 5 of the table in Part 1 of Annex XXII to Implementing Regulation (EU) 2021/404.
- (5) Code of the zone in accordance with column 2 of the table in Part 1 of Annex XXII to Implementing Regulation (EU) 2021/404.
- (6) To be signed by:
 - (a) an official veterinarian when Part II.2 Animal health attestation is not deleted,
 - (b) a certifying officer or an official veterinarian when Part II.2 Animal health attestation is deleted.
- (7) Applicable to consignments entering the Union as from 3 September 2026.

[Official veterinarian]⁽¹⁾⁽⁶⁾/[Certifying officer]⁽¹⁾⁽⁶⁾

Name (in capital letters)

Date

Qualification and title

Stamp

Signature