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Animal health/Official certificate to the EU

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

<b>I.27 Description of consignment</b>							
<b>1</b>	CN Code	Species	Cold store	Identification mark	Type of packaging	Net weight	Treatment type
	Nature of commodity	Number of packages	Batch No	Date of collection /production	Manufacturing plant	Approval or registration number of plant /establishment /centre	Final consumer <input type="checkbox"/>
<b>2</b>	CN Code	Species	Cold store	Identification mark	Type of packaging	Net weight	Treatment type
	Nature of commodity	Number of packages	Batch No	Date of collection /production	Manufacturing plant	Approval or registration number of plant /establishment /centre	Final consumer <input type="checkbox"/>
<b>3</b>	CN Code	Species	Cold store	Identification mark	Type of packaging	Net weight	Treatment type
	Nature of commodity	Number of packages	Batch No	Date of collection /production	Manufacturing plant	Approval or registration number of plant /establishment /centre	Final consumer <input type="checkbox"/>
<b>4</b>	CN Code	Species	Cold store	Identification mark	Type of packaging	Net weight	Treatment type
	Nature of commodity	Number of packages	Batch No	Date of collection /production	Manufacturing plant	Approval or registration number of plant /establishment /centre	Final consumer <input type="checkbox"/>
<b>5</b>	CN Code	Species	Cold store	Identification mark	Type of packaging	Net weight	Treatment type
	Nature of commodity	Number of packages	Batch No	Date of collection /production	Manufacturing plant	Approval or registration number of plant /establishment /centre	Final consumer <input type="checkbox"/>

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**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<sup>A</sup>, Regulation (EC) No 852/2004 of the European Parliament and of the Council<sup>B</sup>, 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<sup>C</sup> and hereby certify that the colostrum<sup>(2)</sup> 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<sup>D</sup>;

(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<sup>E</sup>;

(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<sup>F</sup>, and milk is listed in Commission Decision 2011/163/EU<sup>G</sup> for the concerned country of origin;

(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<sup>H</sup>, and the maximum levels for contaminants laid down in Commission Regulation (EC) No 1881/2006<sup>I</sup>.

**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<sup>(2)</sup> described in Part I:

II.2.1. has been obtained in the zone/s with code/s: .....<sup>(3)</sup> 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<sup>J</sup>, 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*.]<sup>(1)</sup> [*Ovis aries*.]<sup>(1)</sup> [*Capra hircus*.]<sup>(1)</sup> [*Bubalus bubalis*.]<sup>(1)</sup> [*Camelus dromedarius*.]<sup>(1)</sup> 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;

<sup>A</sup> Regulation (EC) No 178/2002 of the European Parliament and of the Council of 28 January 2002 laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety (OJ L 31, 1.2.2002, p. 1).

<sup>B</sup> Regulation (EC) No 852/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs (OJ L 139, 30.4.2004, p. 1).

<sup>C</sup> Commission Implementing Regulation (EU) 2019/627 of 15 March 2019 laying down uniform practical arrangements for the performance of official controls on products of animal origin intended for human consumption in accordance with Regulation (EU) 2017/625 of the European Parliament and of the Council and amending Commission Regulation (EC) No 2074/2005 as regards official controls (OJ L 131, 17.5.2019, p. 51).

<sup>D</sup> Commission Regulation (EU) No 37/2010 of 22 December 2009 on pharmacologically active substances and their classification regarding maximum residue limits in foodstuffs of animal origin (OJ L 15, 20.1.2010, p. 1).

<sup>E</sup> Commission Regulation (EC) No 2073/2005 of 15 November 2005 on microbiological criteria for foodstuffs (OJ L 338, 22.12.2005, p. 1).

<sup>F</sup> Council Directive 96/23/EC of 29 April 1996 on measures to monitor certain substances and residues thereof in live animals and animal products and repealing Directives 85/358/EEC and 86/469/EEC and Decisions 89/187/EEC and 91/664/EEC (OJ L 125, 23.5.1996, p. 10).

<sup>G</sup> Commission Decision 2011/163/EU of 16 March 2011 on the approval of plans submitted by third countries in accordance with Article 29 of Council Directive 96/23/EC (OJ L 70, 17.3.2011, p. 40).

<sup>H</sup> Regulation (EC) No 396/2005 of the European Parliament and of the Council of 23 February 2005 on maximum residue levels of pesticides in or on food and feed of plant and animal origin and amending Council Directive 91/414/EEC (OJ L 70, 16.3.2005, p. 1).

<sup>I</sup> Commission Regulation (EC) No 1881/2006 of 19 December 2006 setting maximum levels for certain contaminants in foodstuffs (OJ L 364, 20.12.2006, p. 5).

<sup>J</sup> Commission Implementing Regulation (EU) 2021/404 of 24 March 2021 laying down the lists of third countries, territories or zones thereof from which the entry into the Union of animals, germinal products and products of animal origin is permitted in accordance with Regulation (EU) 2016/429 of the European Parliament and of the Council (OJ L 114, 31.3.2021, p. 1).

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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<sup>K</sup>;
- (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<sup>L</sup> 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 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:**

- (<sup>1</sup>) Keep as appropriate.
- (<sup>2</sup>) Colostrum as defined in Section IX, Point 1, of Annex III to Regulation (EC) No 853/2004.
- (<sup>3</sup>) Code of the zone in accordance with column 2 of the table in Part 1 of Annex XVII to Implementing Regulation (EU) 2021/404.
- (<sup>4</sup>) 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

[Official veterinarian]<sup>(1)(4)</sup>/[Certifying officer]<sup>(1)(4)</sup>

Name (in capital letters)

Date

Qualification and title

Stamp

Signature

<sup>K</sup> Commission Delegated Regulation (EU) 2020/692 of 30 January 2020 supplementing Regulation (EU) 2016/429 of the European Parliament and of the Council as regards rules for entry into the Union, and the movement and handling after entry of consignments of certain animals, germinal products and products of animal origin (OJ L 174, 3.6.2020, p. 379)

<sup>L</sup> Commission Delegated Regulation (EU) 2020/692 of 30 January 2020 supplementing Regulation (EU) 2016/429 of the European Parliament and of the Council as regards rules for entry into the Union, and the movement and handling after entry of consignments of certain animals, germinal products and products of animal origin (OJ L 174, 3.6.2020, p. 379)