## Official certificate to the EU

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II. Health information

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

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 Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004:
  - (iii) which comes from animals belonging to herds free or officially free of brucellosis and tuberculosis;
  - (iv) which 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;
  - (v) which, pursuant to testing for residues of antibacterial drugs carried out by the food business operator in accordance with the requirements of point 4 in Part III of Chapter I of Section IX 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;
  - (vi) which 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;
- (b) they come 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) they have been processed, stored, wrapped, packaged and labelled in accordance with Chapters III and IV of Section IX of Annex III to Regulation (EC) No 853/2004;
- (d) they meet the relevant criteria laid down in Chapter II of Section IX of Annex III to Regulation (EC) No 853/2004 and the relevant microbiological criteria laid down in Commission Regulation (EC) No 2073/2005 :
- (e) the products described in Part I have been produced under conditions guaranteeing compliance with the maximum residue levels for pesticides laid down in Regulation (EC) No 396/2005, and the maximum levels for contaminants laid down in Regulation (EC) No 1881/2006.

The colostrum-based products(2) described in Part I:

II.2.1. originate from 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-based products 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 vaccination against these diseases has not been carried out during the same period;

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	II. Health info	rmation							
	II.2.2.	have been	processed f	rom colostrum obtained	<u> </u>				
		(1)	o either [in the zone referred to in point II.2.1.;]						
ication		(1)	o or	[in the zone/s with cod- certificate was/were lis	-				
		(1)	$\circ$ or	[in a Member State;]					
Part II : Certification	II.2.3.	has been obtained from animals of the species $\square$ [Bos Taurus,] (1) $\square$ [Ovis aries,] (1) $\square$ [Capra hircus,] (1) $\square$ [Bubalus bubalis,] (1) $\square$ [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							
III	II.2.4.	have been processed from colostrum obtained from animals kept in establishments:							
Pa		•	(a)	country or territory an	er the control of the competer d have a system in place to m icle 8 of Commission Delegate	aintain and to keep records			
		•	(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 Delegated Regulation (EU) 2020/692 and emerging diseases;					
			(c)	reasons, including the	t to national restriction meast relevant listed diseases refer EU) 2020/692 and emerging d n.	red to in Annex I to			
	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-based products, including when the Union is not the final destination of such products.								
	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 Part 1 of Annex XVII to Implementing Regulation (EU) 2021/404.							
	Box reference I.11:	Name, address and approval number of the establishment of dispatch.							
	Box reference I.15:	(vessel) is t there is a se	o be provid erial numbe	niner and road vehicles), fligh ort in containers their registr ndicated in box I.19. In the ca ler control post of entry into t	ation number and where use of unloading and				
	Box reference I.19:	For contain	ners or boxe	r and the seal number (if appl	licable) should be included.				
		narian or Offici	al inspector						
	Name (in cap Date of signa Stamp				Qualification and title Signature				
	•								