**Export Health Certificate** 

	I.1. Consignor				I.2. IMSOC Ref	erence			
	Name				I.2.a. Local Reference				
	Address				1.2.a. Local Reference				
	Country ISO Code								
	I.S. Consigned				I.3. Central competent authority				
	I.5. Consignee Name								
en	Address				I.4. Local comp	petent authority			
틝	Country ISO Code								
.59	<u> </u>						700.0		
on S	I.7. Country of origin ISO Code				I.9. Country of	destination	ISO Code		
of consignment	I.8. Region of origin Code				I.10. Region of	destination			
	I.11. Place of Dispatch				I.12. Place of destination				
: Details					Name				
De	Name Address				Address				
$\mathbf{I}$	Approval Numbe	r			Approval Number				
Part I	Country		ISO Code		Country		ISO Code		
P	I.13. Place of Load	ing			I.14. Date and	time of departure			
	Name	o .				•			
	Address								
	Approval Numbe	r							
	Country		ISO Code						
	I.15. Means of Tra	nsport			I.16 Entry Poir	nt			
	Mode	International	Identification						
		transport document							
	I.18. Transport conditions				I.17. Accompa	nying documents			
	Frozen Controlled Ambient Chilled				Accompanying document reference				
		temperature $\square$			Date of issue Country Place of issue				
	I.19. Container No	/ Seal No							
	I.20. Certified as								
	Human consumpt	ion 🗆							
	-				122 For transit through Mambar State(s)				
		rough a third coun	,		I.22. For transit through Member State(s)				
	Country		ISO Code		Country ISO Code		ISO Code		
	EU Exit Authority BCP code  EU Entry BCP code				-				
	EU Entry						I.25. Total gross weight		
	Authority	r of packages	_	Total net weight	•	I 25 Total m	ross weight		
	Authority I.23. Total number		_	Total net weight		I.25. Total gr	ross weight		
	Authorify I.23. Total number I.28. Description o	f consignment	I.25.			I.25. Total gr	ross weight		
	Authorify 1.23. Total number 1.28. Description o 1. 35 ALBUMINOR	f consignment DAL SUBSTANCES;	I.25.	CHES; GLUES; ENZY					
	Authorify 1.23. Total number 1.28. Description o 1. 35 ALBUMINOR	f consignment DAL SUBSTANCES;	I.25.	CHES; GLUES; ENZY			ross weight whey proteins, calculated on the dry		
	Authorify 1.23. Total number 1.28. Description o 1. 35 ALBUMINOR	f consignment  DAL SUBSTANCES;  (including concent nates and other all	I.25.	CHES; GLUES; ENZY					
	Authorify 1.23. Total number 1.28. Description o 1. 35 ALBUMINOT 3502 Albumins matter), albumin	f consignment  DAL SUBSTANCES;  (including concent nates and other all	I.25.  MODIFIED STARC rates of two or mobumin derivatives	CHES; GLUES; ENZY		eight more than 80 % v	whey proteins, calculated on the dry		
	Authorify I.23. Total number I.28. Description o 1. 35 ALBUMINOI 3502 Albumins matter), albumi #1. Commodity	f consignment  DAL SUBSTANCES;  (including concent nates and other all	I.25.  MODIFIED STARC rates of two or mobumin derivatives	CHES; GLUES; ENZY		eight more than 80 % v	whey proteins, calculated on the dry		
	Authorify I.23. Total number I.28. Description o 1. 35 ALBUMINOI 3502 Albumins matter), albumi #1. Commodity	f consignment  DAL SUBSTANCES;  (including concent nates and other all	I.25.  MODIFIED STARC rates of two or mobumin derivatives	CHES; GLUES; ENZY		eight more than 80 % v	whey proteins, calculated on the dry		
	Authorify I.23. Total number I.28. Description o 1. 35 ALBUMINOI 3502 Albumins matter), albumi #1. Commodity	f consignment  DAL SUBSTANCES;  (including concent nates and other all	I.25.  MODIFIED STARC rates of two or mobumin derivatives	CHES; GLUES; ENZY		eight more than 80 % v	whey proteins, calculated on the dry		
	Authorify I.23. Total number I.28. Description o 1. 35 ALBUMINOI 3502 Albumins matter), albumi #1. Commodity	f consignment  DAL SUBSTANCES;  (including concent nates and other all	I.25.  MODIFIED STARC rates of two or mobumin derivatives	CHES; GLUES; ENZY		eight more than 80 % v	whey proteins, calculated on the dry		
	Authorify I.23. Total number I.28. Description o 1. 35 ALBUMINOI 3502 Albumins matter), albumi #1. Commodity	f consignment  DAL SUBSTANCES;  (including concent nates and other all	I.25.  MODIFIED STARC rates of two or mobumin derivatives	CHES; GLUES; ENZY		eight more than 80 % v	whey proteins, calculated on the dry		
	Authorify I.23. Total number I.28. Description o 1. 35 ALBUMINOI 3502 Albumins matter), albumi #1. Commodity	f consignment  DAL SUBSTANCES;  (including concent nates and other all	I.25.  MODIFIED STARC rates of two or mobumin derivatives	CHES; GLUES; ENZY		eight more than 80 % v	whey proteins, calculated on the dry		
	Authorify I.23. Total number I.28. Description o 1. 35 ALBUMINOI 3502 Albumins matter), albumi #1. Commodity	f consignment  DAL SUBSTANCES;  (including concent nates and other all	I.25.  MODIFIED STARC rates of two or mobumin derivatives	CHES; GLUES; ENZY		eight more than 80 % v	whey proteins, calculated on the dry		

n 1/4

# **EUROPEAN UNION**

					-			Tourities abileouth (vs.o)
	II. Health information							
	II.1.	Animal H	ealth Attest	ation				
			dersigned official veterinarian, declare that I am aware of the relevant provisions of Directive CC and of Regulation (EC) No 853/2004 and hereby certify that the dairy product described					
Ħ		(a)	has been	obtained fr	om animals:			
äţį			(i)	under the	e control of th	e official veterinary s	service;	
ertific			(ii)		g to holdings v r rinderpest: a		r restric	tions due to foot-and-mouth
Part II: Certification			(iii)	health co	nditions laid			hat they satisfy the animal of Annex 3 to Regulation (EC)
	(1)(2)either	○ [(b)	where au milk prod	the dairy product was made from raw milk sourced from cows, ewes, goats, buffaloes or, where authorised, countries with footnote (b) as set out in a document relating to 'milk and milk products' published on gov.uk, in accordance with Regulation (EC) No 605/2010, from camels of the species Camelus dromedarieus, and has undergone, prior to import into Great Britain				
	(1)	either	○ [(i)	a steriliza	ation process,	to achieve an F0 valu	ıe equal	to or greater than three;]
	(1)	or	○ [(ii)		-	ure (UHT) treatment table holding time;]	at not le	ess than 135 °C in
	(1)	or	o [(iii)	a high temperature-short time pasteurisation treatment (HTST) at 72 °C for 15 seconds applied twice to milk with a pH equal to or greater than 7,0 achieving, where applicable, a negative reaction to an alkaline phosphatase test, applied immediately after the heat treatment;]				
	(1)	or	○ [(iv)	where ap	plicable, a ne	-		t to point (iii) achieving, phosphatase test, applied
	(1)	or	∘ [(v)	a HTST tr	eatment of m	ilk with a pH below 7	7,0;]	
	(1)	or	○ [(vi)	a HTST tr	eatment com	bined with another p	hysical t	reatment by
	(1)		either	o [(1)	lowering th	ne pH below 6 for one	e hour;]	
	(1)		or	o [(2)	additional desiccation		reater th	an 72 °C, combined with
	(1)or	○ [(b)	goats, but	y product was made from raw milk sourced from an uffaloes or camels of the species Camelus dromedari nto Great Britain:				
	(1)	-			ation process,	to achieve an F0 valu	ıe equal	to or greater than three;]
	(1)	or	○ [(ii)			ure (UHT) treatment table holding time;]]		ess than 135 °C in
	II.2.	Public He	alth attestation					
	I, the undersigned official inspector, declare the (EC) No 178/2002, (EC) No 852/2004, (EC) No 85 dairy product described above was produced that:				04, (EC) No 85	3/2004 and (EU ) 2019	9/627 an	d hereby certify that the
		(a)	it was ma	s manufactured from raw milk:				
				and checked			e with Regulation (EC) No -50 of Regulation (EU)	
			(ii)	with the l	_	tions laid down in Ch		ransported in accordance of Section IX of Annex III to
			(iii) which meets the plate and somatic cell count criteria laid do Section IX of Annex III to Regulation (EC) No 853/2004;					

en 2 / 4

#### **EUROPEAN UNION**

_	TO HOME TO COMMENCE OF THE COMMENT OF THE COMMENT OF THE COMMENT OF THE COMMENT OF THE COMMENCE OF THE COMMENT OF THE						
	II. Health information						
Certification		(iv)	which complies with the guarantees on the residues status of raw milk provided by the monitoring plans for the detection of residues or substances submitted in accordance with Council Directive 96/23/EC, and in particular, Article 29 thereof;				
		(v)	which, pursuant to testing for residues of antibacterial drugs carried out by the food business operator in accordance with the requirements of Annex III, Section IX, Chapter I, Part III, point 4 to Regulation (EC) No 853/2004, it complies with the maximum residue limits for residues of antibacterial veterinary medicinal products laid down in the Annex to Regulation (EU) No 37/2010;				
Part II: Cert		(vi)	which has been produced under conditions guaranteeing compliance with the maximum residue levels for pesticides laid down in Regulation (EC) No 396/2005, and maximum levels for contaminants laid down in Regulation (EC) No 1881/2006;				
	(b)	it comes from an establishment implementing a programme based on the HACCP principling in accordance with Regulation (EC) No 852/2004;					
	(c)	it has been processed, stored, wrapped, packaged and transported in accordance with the relevant hygiene conditions laid down in Annex II to Regulation (EC) No 852/2004 and Chapter II of Section IX of Annex III to Regulation (EC) No 853/2004;					
	(d)	it meets 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 Regulation (EC) No 2073/2005 on microbiological criteria for foodstuffs;					
	(e)		antees covering live animals and products thereof provided by the residue plans ed in accordance with Directive 96/23/EC, and in particular Article 29 thereof, are				

#### Notes

References to European Union legislation within this certificate are references to direct EU legislation which has been retained in Great Britain (retained EU law as defined in the European Union (Withdrawal) Act 2018) and can be viewed on the UK legislation website (legislation.gov.uk).

References to Great Britain in this certificate include Channel Islands and Isle of Man.

This certificate is intended for dairy products for human consumption from third countries or parts thereof authorised, where applicable, for milk from certain animal species only, listed in column C in a document relating to 'milk and milk products' published on gov.uk, in accordance with Regulation (EU) No 605/2010 intended for importation into Great Britain.(2)

### Part I:

I.28:

35.04.

Box reference I.7:	to 'milk and milk products' published on gov.uk, in accordance with Regulation (EU) No 605/2010.(2)
Box refere	nce I.11: Name, address and approval number of the establishment of dispatch.
Box reference I.15:	registration number (railway wagons or container and road vehicles), flight number (aircraft) or name (ship) is to be provided. In the case of transport in containers, the total number of containers and their registration number and where there is a serial number of the seal it must be indicated in box I.19. In the case of unloading and reloading, the consignor must inform the border control post of introduction into Great Britain.
Box reference I.19:	for containers or boxes, the container number and the seal number (if applicable) should be included.
Box reference I.25:	indicate total gross weight and total net weight.
Box reference	use the appropriate Harmonised System (HS) code under the following headings: 04.01; 04.02; 04.03; 04.04; 04.05; 04.06;15.17; 17.02; 19.01; 21.05; 21.06; 22.02; 28.35; 35.01; 35.02 or

n 3 / 4

## **EUROPEAN UNION**

	TT TT. 3:3 1 C			·					
	II. Health info	rmation							
		Box reference I.28:	use the appropriate Harmonised S 04.02; 04.03; 04.04; 04.05; 04.06;15. 35.04.						
	Part II:								
ı	(1)	Keep as ap	appropriate.						
tior		(2)	A document relating to 'milk and milk products' for EU and EFTA states published by						
ica	Secretary of State, with the consent of the Scottish and Welsh Ministers, may be found here:								
rtif	EU and EFT	'A states ap	proved to export animals and anim	al products to Great Britain - o	data.gov.uk				
Cel	The colour	(1) Keep as appropriate.  (2) A document relating to 'milk and milk products' for EU and EFTA states published by the Secretary of State, with the consent of the Scottish and Welsh Ministers, may be found here:  EU and EFTA states approved to export animals and animal products to Great Britain - data.gov.uk  The colour of the signature shall be different to that of the printing. The same rule applies to stamps other than							
П:	those embo	ssed or wat	termark.	printing. The sume rule uppr	les to stamps other than				
Part II:	Certifying Offi								
Ъ	Name (in capit			Qualification and title					
	Date of signati	ıre		Signature					
	Stamp								

en 4/4