Export Health Certificate

	I.1. Consignor		I.2. IMSOC Reference											
	Name		I.2.a. Local Reference											
	Address													
	Country ISO Code													
	,													
	I.S. Consignee						I.3. Central competent authority							
nt	Name						I.4. Local competent authority							
ne	Address Country ISO Code													
ם	Country													
Sig	I.7. Country of orig	ISO Code	I.9. Country of destination ISO Code											
Part I: Details of consignment	I.8. Region of origin	n				Code	I.10. Region of destination							
S	I.11. Place of Dispa	Code	I.12. Place of d											
E	Name		Name											
et	Address						Address							
\Box	Approval Number	,					Approval Nui	nher						
ίI	Country			ISO	Code		Country	ilber			ISO Code			
ar														
"	I.13. Place of Loadi	ing					I.14. Date and	time of de	parture					
	Name													
	Address													
	Approval Number	•												
	Country			ISO	Code									
ŀ	I.15. Means of Trar	enort					I.16 Entry Poi	nt.						
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	Mode	Internation transport		Identification	on									
		document												
ŀ	I 10 Transport con	ditions					I 17 Assembe	nring door	ımanta					
	I.18. Transport con Ambient	Chilled \square		Frozen 🗆	Cor	ntrolled	I.17. Accompa	nymg doci	intents					
	Ambient ☐ Chilled ☐ Frozen ☐ Controlled temperature ☐						Commercial document			Date o	f issue			
							reference							
											Country Place of issue			
							Country			Place o	of			
	I.19. Container No	/ Seal No					Country			Place of issue	of			
	I.19. Container No	/ Seal No					Country			Place of issue	of			
	I.20. Certified as						Country			Place of issue	of			
							Country			Place (issue	of			
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	I.20. Certified as Human consumpti I.21. For transit thr Country EU Exit Authority EU Entry	on □ rough a thin		ISO Code BCP code		l net weight	I.22. For trans	it through	Member Sta	re(s)	□			
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UROPEAN UNION					(PE) Dairy products V.2						
	II. Health info	rmation									
	II.	Health info	ormation								
				by the undersigned office	cial votorinarian, cortifies tha	t·					
	II.1.	-	ity, represented by the undersigned official veterinarian, certifies that: oducts have been produced in an establishment that has been subject to health approval by the								
_	11.1.	competent	authority a	nd applies the Hazard A	nalysis and Critical Control P nd Sanitation Standard Opera	oints (HACCP) system, Good					
rart II: cermication	II.2.	•			n process specifications to ens of pathogens for animal heal						
3		The milk v	was subjected to one of the following treatments or equivalent:								
II. Ce	(either) (1)	○ [II.2.1.	-	perature Short-Time (HT less than 7]	ST) pasteurisation at at least	72° C for at least 15 seconds					
rari	(or) (1)	○ [II.2.2.		perature Short-Time (HT er than or equal to 7]	ST) pasteurisation on two (2)	consecutive occasions if the					
	(or) (1)	○ [II.2.3.	Slow paste	urisation at a temperatu	are of at least 63°C for at least	30 minutes]					
	(or) (1)	○ [II.2.4.		gh temperature (UHT) to olding time.]	reatment at not less than 135	C in combination with a					
	(or) (1)	○ [II.2.5.	A HTST treatment combined with another physical treatment by either: lowering the pH below 6 for one hour or additional heating equal to or greater than 72°C combined with desiccation]								
		o II.2.6.	Dairy prod	ucts derived from raw n	milk(2):						
			II.2.6.1. The milk used to make the product comes from herds free from Brucellosis and Tuberculosis.								
			II.2.6.2.	The product has under temperature of 2°C or a	gone a maturing process of a above.	t least 60 days at a					
	II.3.	They are f	it for human	consumption.							
	II.4.	Additiona	l animal hea	alth attestation: the prod	ucts comply with the require	ments mentioned below:					
		II.4.1.		from herds and primar rictions at the time of th	y production establishments e milk collection.	that were not subject to					
		II.4.2.	The primary production establishment and the area of at least 10 km surrounding it have not been under quarantine or subject to animal movement restrictions in the sixty (60) days prior to dispatch.								
		II.4.3.	The produ	ct was subject to an iden	atity check at the place of load	ing.					
		II.4.4.	its product	s with any micro-organi	een taken after treatment to a ism that is potentially pathogo diseases according to the OIE	enic to animals that cause					
	Notes										
	Part I										
	(1)		at does not a								
	(2)	Only appli		ble to countries that are free from foot-and-mouth disease.							
		-	The signature and the stamp must be in a different colour to that of the printing. The certificate must be issued in Spanish and in the language of the EU Members.								
	0 1101	-		-	panish and in the language of nd stamps of the issuing heal						
	Certifying Offi Name (in cap Date of signa	ital letters)	Qualification and title Signature								

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