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												
ဌျ	I.8. Region of origin	Code	I.10. Region of destination									
S	I.11. Place of Dispa	Couc	I.12. Place of d									
<u> </u>	Name		Name									
et	Address						Address					
\Box	Approval Number						Approval Nui	mher				
τI	Country			ISO	Code		Country				ISO Code	
ar												
	I.13. Place of Loadi	ng					I.14. Date and	time of de	parture			
	Name											
	Address											
	Approval Number	•										
	Country			ISO	Code							
ŀ	I.15. Means of Tran	enort					I.16 Entry Poi	nt				
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	Mode	Internation transport	nai	Identificati	on							
		document										
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ŀ	I 10 Transport con	ditiono					I 17 Assamns	nring door	ımanta			
	I.18. Transport con Chilled □	Ambient [٦	Controlled	Fre	ozen 🗆	I.17. Accompa	nymg doci	unems			
	Crimeu 🗀	Allibleili L	_	temperatur	re 🗆 - 110	ozen 🗀	Commercial document			Date o	f issue	
							reference					
								Country Place of issue				
							Country				J1	
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	I.19. Container No	/ Seal No					Country					
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							Country					
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	I.20. Certified as Human consumpti I.21. For transit thr Country	on 🗆	d coun	ISO Code				it through	Member Sta	issue		
	I.20. Certified as Human consumpti I.21. For transit thr	on 🗆	rd coun	-				it through	Member Sta	issue		
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	I.20. Certified as Human consumpti I.21. For transit thr Country EU Exit Authority EU Entry Authority	on □ rough a thir		ISO Code BCP code		I net weight	I.22. For trans	it through		te(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.	-	y, represented by the undersigned official veterinarian, certifies that: ducts 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	vas 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 Tuberculosis.	the product comes from here	ls free from Brucellosis and					
			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 fit 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		ple 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 printin The certificate must be issued in Spanish and in the language of the EU Membe								
	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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