**Export Health Certificate** 

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	I.1. Consignor				I.2. IMSOC Reference					
	Name				I.2.a. Local Reference					
	Address									
			***							
	Country		ISO C	ode						
	LF Consigned				L2 Control competent soth ority					
	I.S. Consignee				I.3. Central competent authority					
Ħ	Name				I.4. Local competent authority					
er	Address									
띩	Country		ISO C	ode						
consignment										
Si	I.7. Country of origin ISO Code				I.9. Country of destination ISO C		ISO Code	9		
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oĮ	I.8. Region of origin	n		Code	I.10. Region of destination	<del>on</del>				
$\sim$	I.11. Place of Dispa	itch			I.12. Place of destination					
a;	Name				Name					
et										
	Address				Address					
:	Approval Number	ſ			Approval Number					
£	Country ISO Code				Country ISO Code					
Part I: Details of	,									
4	I.13. Place of Loadi	ing			I.14. Date and time of de	parture				
	Name									
	Address									
	Approval Number	ſ								
	Country		IS	O Code						
	_									
	I.15. Means of Tran	nsport			I.16 Entry Point					
	Mode	Internation	nal Identifica	ition						
		transport								
		document			-					
					<del></del>					
	I.18. Transport con				I.17. Accompanying doc	uments				
	Frozen $\square$	Controlled	Ambient	☐ Chilled ☐	Commercial	_		Commercial		
	temperature $\square$				document Date of issue					
		temperatu	ie 🗀			Date of	f issue			
		temperatu	ге ш		reference					
		temperatu	те 🗀			Place o				
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	I.19. Container No		le Li		reference	Place o				
	I.19. Container No I.20. Certified as	/ Seal No	le Li		reference	Place o				
	I.19. Container No	/ Seal No	le Li		reference	Place o				
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	I.19. Container No I.20. Certified as Human consumpti I.21. For transit thi	/ Seal No	rd country		reference	Place o issue				
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1/3

## **EUROPEAN UNION**

EUROPEAN UNION consumption					on from the EU to Georgia		
	II. Health info	ormation					
	I, the unde	rsigned Off	icial veterinarian hereby certify that	:			
	1.	The dairy	product described above, which is ex	xported to Georgia, has been	obtained from animals:		
Part II: Certification		a)	under the control of the official vet	erinary service,			
		b)	which were in a country or part thereof that has been free of foot-and-mouth disease and rinderpest for a period of at least 12 months prior to the date of this certificate, and wher vaccination against foot-and-mouth disease has not been carried out during that period,				
		c)	belonging to holdings which were rinderpest	not under restrictions due to	foot-and-mouth disease or		
		d)	subject to regular veterinary inspect requirements of the EU, and	ctions to ensure that they sati	isfy the animal health		
		e)	which were fed with feed produced genetically modified food and feed.		ion (EC) No 1829/2003 on		
	2.	pasteurisa	ergone pasteurisation or been produ tion treatment fulfilling the provisio quirements for heat treatment.				
	3.	It was mar	nufactured from raw milk:				
		a)	which comes from holdings registe checked in accordance with Article	_			
		b)	which was produced, collected, coo hygiene conditions laid down in Ar				
		c)	which meets the plate and somatic (EC) No 853/2004,	cell count criteria laid down	in Annex III to Regulation		
		d)	which, pursuant to testing for reside business operator in accordance with the maximum resimedicinal products laid down in Co	ith the requirements of the R due limits for residues of ant	egulation (EC) No 853/2004, ibacterial veterinary		
		e)	which has been produced under coresidue levels for pesticides in accoresidue.				
		f)	which has been produced under co Directive 96/22/EC concerning the p substances having a hormonal or the	prohibition on the use in stoc	k farming of certain		
	4.		om an establishment implementing e with Regulation (EC) No 852/2004.	a programme based on the H	ACCP principles in		
	5.		has been processed, stored, wrapped, packaged and transported in accordance with the relevant ygiene requirements of the EU.				
	6. It meets the relevant microbiological crite		e relevant microbiological criteria o	f Commission Regulation (EC	) No 2073/2005.		
		ntees covering live animals and proc e with the requirements of Regulatio					
	Notes:						
	Part I:						
	Box I.11: The approval number of the entity should be indicated, if applicable.						
	Box I.12: The approval number of the entity should be indicated, if applicable.						
	Box I.19: Either seal- or container number or both are to be indicated in this box.						
	Box I.28: "CN code": use the appropriate Harmonized System (HS) code of the World Customs Organisation: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 35.04.						
	Signature	and stamp r	nust be different color that in the pri	nted certificate.			
	The certifi						
Certifying Officer							

**2** / 3

## **EUROPEAN UNION**

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	II. Health information		
	Name (in capital letters)	Qualification and title	
	Date of signature	Signature	
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Part II: Certification			
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**3** / 3