EUROPEAN UNION

	I.1. Consignor					I.2. IMSOC Reference							
	Name		I.2.a. Local Reference										
	Address												
	Country												
	I.5. Consignee		I.3. Central competent authority										
	Name		I.4. Local competent authority										
en	Address												
nu	Country ISO Code												
Part I : Details of consignment	I.7. Country of orig	ISO Code	I.9. Country of destination ISO Code										
f C	I.8. Region of origi	n			Code	I.10. Region of	doctinatio	n					
SO	I.11. Place of Dispa				coue	I.12. Place of d		11					
ail	Name		Name										
let	Address					Address							
Ξ		Approval Number						Approval Number					
뒨	Country							Country ISO Code					
Pai	I.13. Place of Loadi	ing				I.14. Date and time of departure							
	Name					ו.ו.ד. שמופ מווע נוווד טו עבףמו עודפ							
	Address												
	Approval Number	•											
	Country		ISO	Code									
	I.15. Means of Trai	anont				140 Estima Desint							
	Mode	International	Idontificati			I.16 Entry Point							
	Mode	transport	l Identification										
		document											
	I.18. Transport cor Chilled 🗌	ditions Controlled	Frozen 🛛	٨٣	ıbient 🗆	I.17. Accompanying documents							
	temperature					Commercial document Date of issue							
				reference Place of									
						Country issue							
	I.19. Container No	/ Seal No											
	I.20. Certified as												
	Human consumpti	Human consumption											
	I.21. For transit thi		I.22. For transit through Member State(s)										
	Country		ISO Code										
	EU Exit Authority		BCP code			Country ISO Code							
	EU Entry												
	<u>Authority</u> I.23. Total number	ofpackages	Der coue	I 25 Tota	l net weight	I.25. Total gross weight							
				1.25. 1000	i net weight								
	I.28. Description of												
	1.35 ALBUMINOI												
	matter), albumins	nates and other a	trates of two lbumin deriv	or more w atives	ontaining by w	eight more	e than 80 % whey p	oroteins, calcula	ted on the dry				
	Commodity Species Manufacturin						Package c		Net weight				
							0						

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	II. Health info	rmation								
	I, the undersigned Official veterinarian hereby certify that:									
	1. The colostrum/colostrum-based products described above which are exported to the Republic of									
		Moldova, has been obtained from animals:								
Part II: Certification		a) under the control of the official veterinary service.								
		b) which were in a country or part thereof that has been free of foot-and-mouth disease and of rinderpest for a period of at least 12 months prior to the date this certificate, and where vaccination against foot-and-mouth disease has no been carried out during that period.								
	 belonging to holdings which were not under restrictions due to foot-and-disease or rinderpest, and d) subject to regular veterinary inspections to ensure that they satisfy the a health requirements of the EU. 									
									2.	They were manufactured from colostrum:
		a) which come from holdings registered in accordance with Regulation (EC) No 852/2004 and checked in accordance with article 49 and article 50 of Regulation (EU) 2019/627.								
			b) which was produced, collected, cooled, stored and transported in accordance with the hygiene conditions laid down in Annex III to Regulation (EC) No 853/2004.							
		c)	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 Regulation (EU) 2017/625.							
		d)	food business operator with the maximum res	ing for residues of antibacter in accordance with the requ idue limits for residues of an d down in the Annex to Comr	irements of the EU, complies tibacterial veterinary					
		e) which has been produced under conditions guaranteeing compliance with the maximum residue levels for pesticides in accordance with the requirements of the EU.								
	3.	They come from an establishment implementing a programme based on the HACCP principles in accordance with Regulation (EC) No 852/2004.								
	4.									
	5.	They meet the relevant microbiological criteria of the EU Regulation.								
	Notes:									
	Part I:									
- Box I.19: Indicate total gross weight and total net weight										
	-	Box I.21: Either seal- or container number or both is to be indicated in this box.								
	Box I.25: Custom code and title: Use the appropriate Harmonized System (HS) code under the following headings: 04.01; 04.02; 04.03; 04.04; 04.05; 04.06; 04.10; 15.17; 17.02; 19.01; 21.05; 21.06; 22.02; 28.35; 30.01; 35.01; 35.02; 35.04									
			different color that in the pr	inted certificate.						
Certifying Officer Name (in capital letters) Qualification and title										
	Name (in cap Date of signa Stamp			Qualification and title Signature						