

Part I : Details of consignment	I.1. Consignor		I.2. IMSOC Reference	
	Name		I.2.a. Local Reference	
	Address			
	Country	ISO Code		
	I.5. Consignee		I.3. Central competent authority	
	Name		I.4. Local competent authority	
	Address			
	Country	ISO Code		
	I.7. Country of origin	ISO Code	I.9. Country of destination	ISO Code
	I.8. Region of origin	Code	<del>I.10. Region of destination</del>	
	I.11. Place of Dispatch		I.12. Place of destination	
	Name		Name	
	Address		Address	
	Approval Number		Approval Number	
Country	ISO Code	Country	ISO Code	
I.13. Place of Loading		I.14. Date and time of departure		
Name				
Address				
Approval Number				
Country	ISO Code			
I.15. Means of Transport		I.16 Entry Point		
Mode	International transport document	Identification		
I.18. Transport conditions		I.17. Accompanying documents		
Frozen <input type="checkbox"/>	Chilled <input type="checkbox"/>	Ambient <input type="checkbox"/>	Controlled temperature <input type="checkbox"/>	
		Commercial document reference	Date of issue	
		Country	Place of issue	
I.19. Container No / Seal No				
I.20. Certified as				
Human consumption <input type="checkbox"/>				
I.21. For transit through a third country <input type="checkbox"/>		I.22. For transit through Member State(s) <input type="checkbox"/>		
Country	ISO Code			
EU Exit Authority	BCP code			
EU Entry Authority	BCP code	Country	ISO Code	
I.23. Total number of packages	I.25. Total net weight	I.25. Total gross weight		
I.28. Description of consignment				
<b>1. 21 MISCELLANEOUS EDIBLE PREPARATIONS</b>				
<b>2106 Food preparations not elsewhere specified or included</b>				
Commodity	Species	Manufacturing plant	Package count	
			Net weight	

<b>Part II: Certification</b>	II. Health information			
	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:		
		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 of this certificate, and where vaccination against foot-and-mouth disease has not been carried out during that period.	
		c)	belonging to holdings which were not under restrictions due to foot-and-mouth disease or rinderpest, and	
		d)	subject to regular veterinary inspections to ensure that they satisfy the animal 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)	which, pursuant to testing for residues of antibacterial drugs carried out by the food business operator in accordance with the requirements of the EU, complies with the maximum residue limits for residues of antibacterial veterinary medicinal products laid down in the Annex to Commission Regulation (EU) No 37/2010		
	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.	They have been processed, stored, wrapped, packaged and transported in accordance with the relevant hygiene requirements of the EU.			
5.	They meet the relevant microbiological criteria of the EU Regulation.			
<b>Notes:</b>				
<b>Part I:</b>				
- 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				
<b>Signature and stamp must be different color that in the printed certificate.</b>				
Certifying Officer				
Name (in capital letters)		Qualification and title		
Date of signature		Signature		
Stamp				