

Part I : Details of consignment	I.1. Consignor Name Address Country ISO Code		I.2. IMSOC Reference I.2.a. Local Reference																
	I.5. Consignee Name Address Country ISO Code		I.3. Central competent authority I.4. Local competent authority																
	I.7. Country of origin ISO Code		I.9. Country of destination ISO Code																
	I.8. Region of origin Code		I.10. Region of destination																
	I.11. Place of Dispatch Name Address Approval Number Country ISO Code		I.12. Place of destination Name Address Approval Number Country ISO Code																
	I.13. Place of Loading Name Address Approval Number Country ISO Code		I.14. Date and time of departure																
	I.15. Means of Transport		I.16 Entry Point																
	<table border="1" style="width:100%; border-collapse: collapse;"> <thead> <tr> <th style="width:20%;">Mode</th> <th style="width:20%;">International transport document</th> <th style="width:60%;">Identification</th> </tr> </thead> <tbody> <tr><td> </td><td> </td><td> </td></tr> <tr><td> </td><td> </td><td> </td></tr> <tr><td> </td><td> </td><td> </td></tr> <tr><td> </td><td> </td><td> </td></tr> </tbody> </table>		Mode	International transport document	Identification														
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I.18. Transport conditions Frozen <input type="checkbox"/> Chilled <input type="checkbox"/> Ambient <input type="checkbox"/> Controlled temperature <input type="checkbox"/>		I.17. Accompanying documents 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"/> Country ISO Code EU Exit Authority BCP code EU Entry Authority BCP code		I.22. For transit through Member State(s) <input type="checkbox"/> Country ISO Code																	
I.23. Total number of packages		I.25. Total net weight	I.25. Total gross weight																
I.28. Description of consignment 1. 22 BEVERAGES, SPIRITS AND VINEGAR 2202 Waters, including mineral waters and aerated waters, containing added sugar or other sweetening matter or flavoured, and other non-alcoholic beverages, not including fruit or vegetable juices of heading 2009																			
Commodity	Species	Manufacturing plant	Package count	Net weight															
Batch number																			

Part II: Certification	II. Health information			
	I, the undersigned Official veterinarian hereby certify that:			
	1.	The dairy product described above, which is 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.	It has undergone pasteurisation or been produced from raw milk which has been submitted to a pasteurisation treatment fulfilling the provisions of Regulation (EC) No 853/2004 laying down specific rules or requirements for heat treatment.		
	3.	It was manufactured from raw milk:		
		a)	which comes 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 meets the plate and somatic cell count criteria laid down in Annex III to Regulation (EC) No 853/2004,		
	d)	which, pursuant to testing for residues of antibacterial drugs carried out by the food business operator in accordance with the requirements of the Regulation (EC) No 853/2004, it complies with the maximum residue limits for residues of antibacterial veterinary medicinal products laid down in 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.		
4.	It comes from an establishment implementing a programme based on the HACCP principles in accordance with Regulation (EC) No 852/2004.			
5.	It has been processed, stored, wrapped, packaged and transported in accordance with the relevant hygiene requirements of the EU.			
6.	It meets the relevant microbiological criteria of Commission Regulation (EC) No 2073/2005.			
7.	The guarantees covering live animals and products thereof provided by the residue plans submitted in accordance with the requirements of Regulation (EU) 2017/625 are fulfilled.			
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; 21.05			
Signature and stamp must be different color that in the printed certificate.				
Certifying Officer				
Name (in capital letters)		Qualification and title		
Date of signature		Signature		
Stamp				