

Heat-Treated Milk Products Intended for Human Consumption

With the exception of certificates for shipments destined to Belgium, the United Kingdom, and Ireland, the Agricultural Marketing Service (AMS) endorses both the animal health and public health parts of the EU Certificates for dairy products for human consumption.

Veterinary Services (VS) does not endorse certificates for dairy products exported to other EU Member States for human consumption. (Exception: Shipments only transiting the EU. For information on transit certificates, please go back to the previous page on your browser and select the pertinent link.)

Even though they are not members of the EU, Iceland, Norway, Liechtenstein, and Switzerland also require the same certification for dairy products for human consumption. AMS endorses both sections of the below certificate for certificates for shipment to these countries.

Special requirements for shipments to the United Kingdom, Belgium, and Ireland:

Both AMS and VS endorsement is required for certificates for shipments of dairy products (intended for human consumption) to the United Kingdom, Belgium, and Ireland. When exporting to these EU Countries, the exporter should first take the certificate to AMS for endorsement, and then take the certificate to Veterinary Services (VS) for endorsement. The certificate is not transferred to VS Security Paper. VS countersigns and dates the certificate below the AMS signature, and makes a photocopy of the final version for VS records. Standard user fees apply for the VS countersignature. **VS may only endorse the certificate after AMS endorsement.**

Health Certificate for dairy products derived from milk of cows, ewes, goats, and buffaloes for human consumption from third countries authorised in column B of Annex I to Regulation (EU) No 605/2010 intended for importation to the European Union

COUNTRY

Veterinary certificate to EU

Part I. Details of dispatched consignment	I.1 Consignor Name			I.2 Certificate reference number		I.2.a	
	Address			I.3 Central Competent Authority			
	Tel.N°			I.4 Local competent Authority			
	I.5 Consignee Name			I.6			
	Address						
	Tel.N°						
	I.7 Country of Origin	ISO code	I.8 Region of Origin	Code	I.9 Country of Destination	ISO Code	I.10
	I.11 Place of origin Name			I.12			
	Approval Number						
	Address						
	I.13 Place of loading			I.14 Date of departure			
	I.15 Means of transport Aeroplane <input type="checkbox"/> Ship <input type="checkbox"/> Railway wagon <input type="checkbox"/> Road vehicle <input type="checkbox"/> Other <input type="checkbox"/>			I.16 Entry BIP in EU			
	Identification: Documentation reference:			I.17			
I.18 Description of commodity				I.19 Commodity code (HS code)			
				I.20 Quantity			
I.21 Temperature of product Ambient <input type="checkbox"/> Chilled <input type="checkbox"/> Frozen <input type="checkbox"/>			I.22 Number of packages				
I.23 Identification of container/Seal number			I.24 Type of packaging				
I.25 Commodities certified for: Human consumption <input type="checkbox"/>							
I.26			I.27 For import or admission into EU <input type="checkbox"/>				
I.28 Identification of the commodities							
Species (Scientific name)		Approval number of establishment Manufacturing plant		Number of packages	Net weight	Batch number	

II. Health information	II.a Certificate reference number	II.b
II.1 Animal Health Attestation		
<p>I, the undersigned official veterinarian, declare that I am aware of the relevant provisions of Directive 2002/99/EC and of Regulation (EC) No 853/2004 and hereby certify that the dairy product described above:</p>		
<p>(a) has been obtained from animals:</p> <ul style="list-style-type: none"> (i) under the control of the official veterinary service, (ii) 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, (iii) belonging to holdings which were not under restrictions due to foot-and-mouth disease or rinderpest, and, (iv) subject to regular veterinary inspections to ensure that they satisfy the animal health conditions laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004 and in Directive 2002/99/EC. <p>(b) Has undergone or been produced from raw milk which has been submitted to a pasteurization treatment involving a single heat treatment with a heating effect at least equivalent to that achieved by a pasteurization process of at least 72°C for at 15 seconds and where applicable, sufficient to ensure a negative reaction to an alkaline phosphatase test applied immediately after the heat treatment.</p>		
II.2 Public Health Attestation		
<p>I, the undersigned official inspector, declare that I am aware of the relevant provisions of Regulations (EC) No 178/2002, (EC) No 852/2004, (EC) No 853/2004 and (EC) No 854/2004 and hereby certify that the dairy product described above was produced in accordance with those provisions, in particular that:</p>		
<p>(a) it was manufactured from raw milk:</p> <ul style="list-style-type: none"> (i) which comes from holdings registered in accordance with Regulation (EC) No 852/2004 and checked in accordance with Annex IV to Regulation (EC) No 854/2004. (ii) which was produced, collected, cooled, stored and transported in accordance with the hygiene conditions laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004. (iii) which meets the plate and somatic cell count criteria laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004, and (iv) which does not contain antibiotic residues exceeding the limits authorised under the Annex to Commission Regulation (EC) No 853/2004; (v) which does not contain pesticide residues exceeding the limits authorized by Regulation (EC) No 396/2005, and (vi) which does not contain contaminants exceeding the maximum tolerances laid down by Regulation (EC) No 1881/2006. <p>(b) it comes from an establishment implementing a programme based on the HACCP principles in accordance with Regulation (EC) No 852/2004;</p> <p>(c) it has been processed, stored, wrapped, packaged and transported in accordance with the relevant hygiene conditions laid down in Annex II to Regulation (EC) No 852/2004 and Chapter II of Section IX of Annex III to Regulation (EC) No 853/2004;</p> <p>(d) it meets the relevant criteria laid down in Chapter II of Section IX of Annex III to Regulation (EC) No 853/2004 and the relevant microbiological criteria laid down in Commission Regulation (EC) No 2073/2005 on microbiological criteria in foodstuffs;</p> <p>(e) the guarantees covering live animals and products thereof provided by the residue plans submitted in accordance with Directive 96/23/EC, and in particular Article 29 thereof, are fulfilled.</p>		

II. Health Information	II.a Certificate reference number	II.b						
<p>Notes</p> <p>This certificate is intended for dairy products for human consumption from third countries or parts thereof authorized in column B of Annex I of Regulation (EU) No 605/2010 intended for importation into the European Union.</p> <p>Part I:</p> <ul style="list-style-type: none"> • Box reference I.7: Provide the country and ISO code of the country or part thereof as appearing in Annex I to Regulation (EU) No 605/2010. • Box reference I.11: Name, address and approval number of establishment of dispatch. • Box reference I.15: Registration number (railway wagons or container and road vehicle), flight number (aircraft) or name (ship). In the case of transport in containers, the total number of containers and their registration number and where there is a serial number of the seal it must be indicated in box I.23. In case of unloading and reloading, the consignor must inform the border inspection post of introduction into the European Union. • Box reference I.19: Use the appropriate Harmonised System (HS) code of the World Customs Organisation: 04.01; 04.02; 04.03; 04.04; 04.05; 04.06 or 21.05. • Box reference I.20: Indicate total gross weight and total net weight. • Box reference I.23: for containers or boxes, the container number and the seal number (if applicable) should be included. • Box reference I.28: manufacturing plant: introduce the approval number of the treatment and/or processing establishment(s) approved for export to the European Union. <p>Part II:</p> <ul style="list-style-type: none"> • The colour of the signature shall be different to that of the printing. The same rule applies to stamps other than those embossed or watermark. 								
<p>Official veterinarian</p> <table border="0" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 50%; border-top: 1px solid black; text-align: center; padding-top: 5px;">Name (in capital letters)</td> <td style="width: 50%; border-top: 1px solid black; text-align: center; padding-top: 5px;">Qualification and title</td> </tr> <tr> <td style="border-top: 1px solid black; text-align: center; padding-top: 5px;">Date</td> <td></td> </tr> <tr> <td style="text-align: center; padding-top: 5px;">Stamp</td> <td style="border-top: 1px solid black; text-align: center; padding-top: 5px;">Signature</td> </tr> </table>			Name (in capital letters)	Qualification and title	Date		Stamp	Signature
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