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▶<u>M6</u> COMMISSION REGULATION (EU) No 605/2010

of 2 July 2010

laying down animal and public health and veterinary certification conditions for the introduction into the European Union of raw milk, dairy products, colostrum and colostrum-based products intended for human consumption ◄

(Text with EEA relevance)

(OJ L 175, 10.7.2010, p. 1)

Amended by:

Official Journal

		No	page	date
► <u>M1</u>	Commission Implementing Regulation (EU) No 914/2011 of 13 September 2011	L 237	1	14.9.2011
► <u>M2</u>	Commission Implementing Regulation (EU) No 957/2012 of 17 October 2012	L 287	5	18.10.2012
► <u>M3</u>	Commission Implementing Regulation (EU) No 300/2013 of 27 March 2013	L 90	71	28.3.2013
► <u>M4</u>	Commission Regulation (EU) No 519/2013 of 21 February 2013	L 158	74	10.6.2013
► <u>M5</u>	Commission Implementing Regulation (EU) No 556/2013 of 14 June 2013	L 164	13	18.6.2013
► <u>M6</u>	Commission Implementing Regulation (EU) No 209/2014 of 5 March 2014	L 66	11	6.3.2014
► <u>M7</u>	Commission Implementing Regulation (EU) 2018/83 of 19 January 2018	L 16	6	20.1.2018
► <u>M8</u>	Commission Implementing Regulation (EU) 2018/1120 of 10 August 2018	L 204	31	13.8.2018
► <u>M9</u>	Commission Implementing Regulation (EU) 2019/366 of 5 March 2019	L 65	1	6.3.2019

Corrected by:

►<u>C1</u> Corrigendum, OJ L 234, 10.9.2011, p. 47 (605/2010)

▼<u>B</u> ▼M6

COMMISSION REGULATION (EU) No 605/2010

of 2 July 2010

laying down animal and public health and veterinary certification conditions for the introduction into the European Union of raw milk, dairy products, colostrum and colostrum-based products intended for human consumption

▼<u>B</u>

(Text with EEA relevance)

Article 1

Subject matter and scope

This Regulation lays down:

▼M6

 (a) the public and animal health conditions and certification requirements for the introduction into the European Union of consignments of raw milk, dairy products, colostrum and colostrum-based products;

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(b) the list of third countries from which the introduction into the European Union of such consignments is authorised.

▼<u>M1</u>

This Regulation shall apply without prejudice to any specific certification requirements laid down in other Union acts or in agreements concluded by the Union with third countries.

▼<u>M6</u>

Article 2

Importation of raw milk, dairy products, colostrum and colostrum-based products from third countries or parts thereof listed in column A of Annex I

Member States shall authorise the importation of consignments of raw milk, dairy products, colostrum and colostrum-based products from the third countries or parts thereof listed in column A of Annex I.

▼<u>B</u>

Article 3

Imports of certain dairy products from third countries or parts thereof listed in column B of Annex I

Member States shall authorise the importation of consignments of dairy products derived from raw milk of cows, ewes, goats or buffaloes from the third countries or parts thereof not at risk from foot-and-mouth disease listed in column B of Annex I, provided that such dairy products have undergone, or been produced from raw milk which has undergone a pasteurisation treatment involving a single heat treatment:

(a) with a heating effect at least equivalent to that achieved by a pasteurisation process of at least 72 °C for 15 seconds;

(b) where applicable, sufficient to ensure a negative reaction to an alkaline phosphatase test applied immediately after the heat treatment.

Article 4

Imports of certain dairy products from third countries or parts thereof listed in column C of Annex I

1. \blacktriangleright M3 Member States shall authorise the importation of consignments of dairy products derived from raw milk of cows, ewes, goats, buffaloes or, where specifically authorised in Annex I, from camels of the species *Camelus dromedarius* from the third countries or parts thereof at risk of foot-and-mouth disease listed in column C of Annex I, provided that such dairy products have undergone, or been produced from raw milk which has undergone, a heat treatment involving: \blacktriangleleft

- (a) a sterilisation process, to achieve an F_0 value equal to or greater than three;
- (b) an ultra high temperature (UHT) treatment at not less than 135 °C in combination with a suitable holding time;
- (c) (i) a high temperature short time pasteurisation treatment (HTST) at 72 °C for 15 seconds applied twice to milk with a pH equal to or greater than 7.0 achieving, where applicable, a negative reaction to a alkaline phosphatase test, applied immediately after the heat treatment; or
 - (ii) a treatment with an equivalent pasteurisation effect to point (i) achieving, where applicable, a negative reaction to an alkaline phosphatase test, applied immediately after the heat treatment;
- (d) a HTST treatment of milk with a pH below 7.0; or
- (e) a HTST treatment combined with another physical treatment by either:
 - (i) lowering the pH below 6 for one hour, or
 - (ii) additional heating equal to or greater than 72 °C, combined with desiccation.

2. Member States shall authorise the importation of consignments of dairy products derived from raw milk of animals other than those referred to in paragraph 1, from the third countries or parts thereof at risk of foot-and-mouth disease listed in column C of Annex I, provided that such dairy products have undergone, or been produced from raw milk which has undergone a treatment involving:

(a) a sterilisation process, to achieve an F_0 value equal to or greater than three; or

(b) an ultra high temperature (UHT) treatment at not less than 135 °C in combination with a suitable holding time.

Article 5

Certificates

Consignments authorised for importation in accordance with Articles 2, 3 and 4 shall be accompanied by a health certificate drawn up in accordance with the appropriate model set out in Part 2 of Annex II for the commodity concerned and completed in accordance with the explanatory notes set out in Part 1 of that Annex.

However, the requirements laid down in this Article shall not preclude the use of electronic certification or of other agreed systems, harmonised at European Union level.

▼<u>M6</u>

Article 6

Transit and storage conditions

The introduction into the European Union of consignments of raw milk, dairy products, colostrum and colostrum-based products not intended for importation into the European Union but destined for a third country either by immediate transit or after storage in the Union, in accordance with Articles 11, 12 or 13 of Directive 97/78/EC, shall only be authorised if the consignments comply with the following conditions:

- (a) they come from a third country or part thereof listed in Annex I for the introduction into the European Union of consignments of raw milk, dairy products, colostrum or colostrum-based products and comply with the appropriate treatment conditions for such consignments, as provided for in Articles 2, 3 and 4;
- (b) they comply with the specific animal health conditions for importation into the European Union of the raw milk, dairy products, colostrum or colostrum-based products concerned, as laid down in the animal health attestation in point II.1 of the relevant model health certificate set out in Part 2 of Annex II;
- (c) they are accompanied by a health certificate drawn up in accordance with the appropriate model set out in Part 3 of Annex II for the consignment concerned and completed in accordance with the explanatory notes set out in Part 1 of that Annex;
- (d) they are certified as acceptable for transit, including for storage as appropriate, on the Common Veterinary Entry Document referred to in Article 2(1) of Regulation (EC) No 136/2004, signed by the official veterinarian of the border inspection post of introduction into the Union.

Article 7

Derogation concerning transit and storage conditions

1. By way of derogation from Article 6, the transit by road or by rail through the European Union, between designated border inspection posts in Latvia, Lithuania and Poland listed in Commission Decision 2009/821/EC (¹), of consignments coming from and destined to Russia directly or via another third country shall be authorised provided that the following conditions are complied with:

- (a) the consignment is sealed with a serially numbered seal at the border inspection post of introduction into the European Union by the veterinary services of the competent authority;
- (b) the documents accompanying the consignment and referred to in Article 7 of Directive 97/78/EC are stamped 'ONLY FOR TRANSIT TO RUSSIA VIA THE EU' on each page by the official veterinarian of the competent authority responsible for the border inspection post of introduction into the European Union;
- (c) the procedural requirements provided for in Article 11 of Directive 97/78/EC are complied with;
- (d) the consignment is certified as acceptable for transit on the common veterinary entry document by the official veterinarian of the border inspection post of introduction into the European Union.

2. Unloading or storage, as defined in Article 12(4) or in Article 13 of Directive 97/78/EC, of such consignments on European Union territory shall not be allowed.

3. Regular audits shall be made by the competent authority to ensure that the number of consignments and the quantities of products leaving the European Union territory matches the number and quantities entering the European Union.

▼<u>M5</u>

Article 7a

Derogation for transit through Croatia of consignments coming from Bosnia and Herzegovina and destined to third countries

1. By way of derogation from Article 6, the direct transit by road through the Union, between the border inspection post of Nova Sela and the border inspection post of Ploče, of consignments coming from Bosnia and Herzegovina and destined to third countries shall be authorised provided that the following conditions are complied with:

- (a) the consignment is sealed with a serially numbered seal by the official veterinarian at the border inspection post of entry;
- (b) the documents accompanying the consignment and referred to in Article 7 of Directive 97/78/EC are stamped 'ONLY FOR TRANSIT TO THIRD COUNTRIES VIA THE EU' on each page by the official veterinarian at the border inspection post of entry;

⁽¹⁾ OJ L 296, 12.11.2009, p. 1

- (c) the procedural requirements provided for in Article 11 of Directive 97/78/EC are complied with;
- (d) the consignment is certified as acceptable for transit on the Common Veterinary Entry Document referred to in Article 2(1) of Regulation (EC) No 136/2004 by the official veterinarian at the border inspection post of entry.

2. Unloading or storage, as defined in Article 12(4) or in Article 13 of Directive 97/78/EC, of such consignments on Union territory shall not be allowed.

3. Regular audits shall be made by the competent authority to ensure that the number of consignments and the quantities of products leaving the Union matches the number and quantities entering the Union.

▼<u>M6</u>

Article 8

Specific treatment

Consignments of dairy products and colostrum-based products authorised for introduction into the European Union in accordance with Articles 2, 3, 4,6 or 7 from third countries or parts thereof where an outbreak of foot-and-mouth disease has occurred within the period of 12 months preceding the date of signature of the health certificate, or which have carried out vaccination against that disease during that period, shall only be authorised for introduction into the European Union if such products have undergone one of the treatments listed in Article 4.

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Article 9

Repeal

Decision 2004/438/EC is repealed.

References to Decision 2004/438/EC shall be construed as references to this Regulation.

Article 10

Transitional provisions

For a transitional period until 30 November 2010, consignments of raw milk and milk-based products as defined in Decision 2004/438/EC in respect of which the relevant health certificates have been issued in accordance Decision 2004/438/EC may continue to be introduced into the European Union.

Article 11

Entry into force and applicability

This Regulation shall enter into force on the twentieth day following that of its publication in the *Official Journal of the European Union*.

It shall apply from 1 August 2010.

This Regulation shall be binding in its entirety and directly applicable in all Member States.

▼<u>M5</u>

ANNEX I

List of third countries or parts thereof authorised for the introduction into the European Union of consignments of raw milk, dairy products, colostrum (*) and colostrum-based products (*) and indicating the type of heat treatment required for such commodities

'+': third country is authorised

'0': third country is not authorised

	ISO code of third country	Third country or part thereof	Column A	Column B	Column C
▼ <u>M7</u>					
	AE	The Emirates of Abu Dhabi and Dubai of the United Arab Emirates $(^1)$	0	0	+ (2)
▼ <u>M6</u>					
	AD	Andorra	+	+	+
	AL	Albania	0	0	+
	AR	Argentina	0	0	+
	AU	Australia	+	+	+
	BR	Brazil	0	0	+
	BW	Botswana	0	0	+
	BY	Belarus	0	0	+
	BZ	Belize	0	0	+
▼ <u>M8</u>					
	BA	Bosnia and Herzegovina	+	+	+
▼ <u>M6</u>					
	CA	Canada	+	+	+
	СН	Switzerland (**)	+	+	+
	CL	Chile	0	+	+
	CN	China	0	0	+
	СО	Colombia	0	0	+
	CR	Costa Rica	0	0	+
	CU	Cuba	0	0	+
	DZ	Algeria	0	0	+
	ET	Ethiopia	0	0	+
	GL	Greenland	0	+	+
	GT	Guatemala	0	0	+
	НК	Hong Kong	0	0	+
	HN	Honduras	0	0	+
	IL	Israel	0	0	+
	IN	India	0	0	+

ISO code of third country	Third country or part thereof	Column A	Column B	Column C
IS	Iceland	+	+	+
JP	Japan	+	+	+
KE	Kenya	0	0	+
MA	Morocco	0	0	+
ME	Montenegro	+	+	+
MG	Madagascar	0	0	+
MK (***)	former Yugoslav Republic of Macedonia	0	+	+
MR	Mauritania	0	0	+
MU	Mauritius	0	0	+
MX	Mexico	0	0	+
NA	Namibia	0	0	+
NI	Nicaragua	0	0	+
NZ	New Zealand	+	+	+
РА	Panama	0	0	+
РҮ	Paraguay	0	0	+
RS (****)	Serbia	0	+	+
RU	Russia	0	0	+
SG	Singapore	0	0	+
SV	El Salvador	0	0	+
SZ	Swaziland	0	0	+
TH	Thailand	0	0	+
TN	Tunisia	0	0	+
TR	Turkey	0	0	+
UA	Ukraine	0	0	+
US	United States	+	+	+
UY	Uruguay	0	0	+
ZA	South Africa	0	0	+
ZW	Zimbabwe	0	0	+

(*) The colostrum and colostrum-based products can only be introduced into the European Union from countries authorised in column A.

column A.
(**) Certificates in accordance with the Agreement between the European Community and the Swiss Confederation on trade in agricultural products (OJ L 114, 30.4.2002, p. 132).
(***) The former Yugoslav Republic of Macedonia; the definitive nomenclature for this country will be agreed following the conclusion of the negotiations currently taking place on this subject at UN level.
(***) Not including Kosovo which is at present under international administration pursuant to United Nations Security Council Resolution 1244 of 10 June 1999.
(I) Only dairy product from complex of the preside of the preside Currently device.

(1) Only dairy products from camels of the species Camelus dromedarius.

(2) Dairy products from camels of the species Camelus dromedarius are authorised.

▼M6

ANNEX II

▼<u>M6</u>

PART 1

Models of health certificates

- 'Milk-RM': Health certificate for raw milk from third countries or parts thereof authorised in column A of Annex I intended for further processing in the European Union before being used for human consumption.
- 'Milk-RMP': Health certificate for dairy products derived from raw milk for human consumption, from third countries or parts thereof authorised in column A of Annex I intended for importation into the European Union.
- 'Milk-HTB': Health certificate for dairy products derived from milk of cows, ewes, goats and buffaloes for human consumption from third countries or parts thereof authorised in column B of Annex I intended for importation into the European Union.
- 'Milk-HTC': Health certificate for dairy products for human consumption from third countries or parts thereof authorised in column C of Annex I intended for importation into the European Union.
- 'Colostrum-C/CPB': Health certificate for colostrum of cows, ewes, goats and buffaloes and colostrum-based products derived from colostrum of the same species from third countries or parts thereof listed in column A of Annex I for human consumption intended for importation into the European Union.
- 'Milk/ Colostrum-T/S': Animal health certificate for raw milk, colostrum, dairy products or colostrum-based products for human consumption, intended for transit through or storage in the European Union.

Explanatory notes

- (a) The health certificates shall be issued by the competent authorities of the third country of origin, in accordance with the appropriate model set out in Part 2 of this Annex, according to the layout of the model that corresponds to the raw milk, colostrum, dairy products or colostrum-based products concerned. They shall contain, in the numbered order that appears in the model, the attestations that are required for any third country and, as the case may be, those supplementary guarantees that are required for the exporting third country concerned.
- (b) The original of the health certificate shall consist of a single sheet printed on either pages or, where more text is required, such that all the sheets form a whole and cannot be separated.
- (c) A separate, single health certificate must be presented for each consignment of the commodity concerned, exported to the same destination from a third country listed in the table in Annex I and transported in the same railway wagon, road vehicle, aircraft or ship.

- (d) The original of the health certificate and the labels referred to in the model certificate shall be drawn up in at least one official language of the Member State where border inspection takes place and of the Member State of destination. However, those Member States may allow it to be drawn up in another official language of the European Union instead of their own, accompanied, if necessary, by an official translation.
- (e) Where additional sheets are attached to the health certificate for the purpose of identifying the commodities making up the consignment, such additional sheets shall also be considered to form part of the original certificate, provided the signature and stamp of the certifying official veterinarian appear on each page.
- (f) Where the health certificate comprises more than one page, each page shall be numbered '-x(page number) of y(total number of pages)-' on the bottom of the page and shall bear the certificate reference number allocated by the competent authority on the top of the page.
- (g) The original of the health certificate must be completed and signed by a representative of the competent authority responsible for verifying and certifying that the raw milk, colostrum, dairy products or colostrum-based products meet the health conditions laid down in Section IX of Chapter I of Annex III to Regulation (EC) No 853/2004 and in Directive 2002/99/EC.
- (h) The competent authorities of the exporting third country shall ensure that principles of certification equivalent to those laid down in Council Directive 96/93/EC (¹) are complied with.
- (i) The colour of the signature of the official veterinarian shall be different from that of the printing on the health certificate. That requirement shall also apply to stamps other than embossed stamps or watermarks.
- (j) The original of the health certificate must accompany the consignment until it reaches the border inspection post of introduction into the European Union.
- (k) Where the model certificate states that certain statements shall be kept as appropriate, statements which are not relevant, may be crossed out and initialled and stamped by the certifying officer, or completely deleted from the certificate.

PART 2

Model Milk-RM

Health Certificate for raw milk from third countries or parts thereof authorised in column A of Annex I to Regulation (EU) No 605/2010 intended for further processing in the European Union before being used for human consumption

cou	NTRY	1	Veterinary certificate to EL		
	l.1.	Consignor	I.2. Certificate reference No I.2.a.		
		Name	I.3. Central competent authority		
		Address Tel.			
ment			I.4. Local competent authority		
sign	1.5.	Consignee Name	1.6.		
con		Address			
hed		Postcode			
Part I: Details of dispatched consignment		Tel.			
of dis	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10.		
ails o			destination		
Deta	1.11.	Place of origin	1.12.		
art I:		Name Approval number			
à		Address			
	112	Place of loading	I.14. Date of departure		
	1.13.	Frace of loading			
	l.15.	Means of transport	I.16. Entry BIP in EU		
		Aeroplane 🗌 Ship 🗌 Railway wagon 🗌	1		
		Road vehicle Other	1.17.		
		Identification Documentation references	1.17.		
	1.18.	Description of commodity	I.19. Commodity code (HS code)		
			I.20. Quantity		
	1.21.	Temperature of product	I.22. Number of packages		
		Ambient Chilled	Frozen		
	1.23.	Seal/Container No	I.24. Type of packaging		
	1.25.	Commodities certified for:			
		Further process			
	1.26.		I.27. For import or admission into EU		
	1.28.	Identification of the commodities			
		Manufacturing plant Number of packages	Species Net weight Batch number (Scientific name)		

▼<u>M1</u>

Model Milk-RM

	cou	OUNTRY	Raw milk
	II.	I. Health information II.a. Certificate reference number II.b.	
	11.1.	I.1. Animal Health Attestation	
		I, the undersigned official veterinarian, declare that I am aware of the relevant provisions of Directive 2002/99/EC and of F 853/2004 and hereby certify that the raw milk described above has been obtained from animals:	tegulation (EC) No
ation		(a) under the control of the official veterinary service,	
Part II: Certification		(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 prior to the date of this certificate, and where vaccination against foot-and-mouth disease has not been carried out	
art II:		(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 conditions laid down in Chapte Annex III to Regulation (EC) No 853/2004 and in Directive 2002/99/EC;	r I of Section IX of
	11.2.	I.2. Public Health attestation	
		I, the undersigned official inspector, declare that I am aware of the relevant provisions of Regulations (EC) No 178/2002, (EC) No 853/2004 and (EC) No 854/2004 and hereby certify that the raw milk described above was produced in acco provisions, in particular that:	
		 (a) it comes from holdings registered in accordance with Regulation (EC) No 852/2004 and checked in accordance Regulation (EC) No 854/2004, 	with Annex IV to
		(b) it was produced, collected, cooled, stored and transported in accordance with the hygiene conditions laid down in Cha of Annex III to Regulation (EC) No 853/2004,	pter I of Section IX
		(c) it 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,
		(d) the guarantees on the residues status of raw milk provided by the monitoring plans for the detection of residues or sub in accordance with Council Directive 96/23/EC, and in particular, Article 29 thereof, are fulfilled;	ostances submitted
		(e) pursuant to testing for residues of antibacterial drugs carried out by the food business operator in accordance with th Annex III, Section IX, Chapter I, Part III, point 4 of Regulation (EC) No 853/2004, it complies with the maximum residue of antibacterial veterinary medicinal products laid down in the Annex to Regulation (EU) No 37/2010;	
		(f) it has been produced under conditions guaranteeing compliance with the maximum residue levels for pesticides laid (EC) No 396/2005, and maximum levels for contaminants laid down in Regulation (EC) No 1881/2006.	lown in Regulation
	Note	Notes	
		This certificate is intended for raw milk from third countries or parts thereof authorised in column A of Annex I to Regulation ntended for further processing in the European Union before being used for human consumption.	(EU) No 605/2010
	Part	Part I:	
	— е	- Box reference I.7: Provide name and ISO code of the country or part thereof as appearing in Annex I to Regulation (EU) No 605/2010.
	— E	- Box reference I.11: Name, address and approval number of the establishment of dispatch.	
		— Box reference I.15: Registration number (railway wagons or container and road vehicle), flight number (aircraft) or name unloading and reloading, the consignor must inform the border inspection post of introduction into the European Union.	(ship). In case of
	— E	- Box reference I.19: Use the appropriate Harmonised System (HS) code under the following headings: 04.01; 04.02 or 04	.03.
	— E	- Box reference I.20: Indicate total gross weight and total net weight.	
	— E	- Box reference I.23: For containers or boxes, the container number and the seal number (if applicable) should be included	ł.
		 Box reference I.28: Manufacturing plant: introduce the approval number of the production holding(s), collection centre or star approved for exportation to the European Union. 	ndardization centre
	ı		

COUNTRY		<i>Model Milk-RM</i> Raw milk
II. Health information	II.a. Certificate reference number	II.b.
Part II: — The colour of the signature shall be different to the	at of the printing. The same rule applies to stam	ps other than those embossed or watermark.
Official veterinarian		
Name (in capital letters):	Qualifica	ion and title:
Date:	Signature	
Stamp:		

Model Milk-RMP

Health Certificate for dairy products derived from raw milk for human consumption from third countries or parts thereof authorised in column A of Annex I to Regulation (EU) No 605/2010 intended for importation into the European Union

COUN	NTRY	:			Veterinary certificate to EU
	1.1.	Consignor		I.2. Certificate reference No	I.2.a.
		Name			
		Address		I.3. Central competent authority	
Part I: Details of dispatched consignment	Tel.		I.4. Local competent authority		
lgnn			1.6.		
suo		Name Address			
pg c		Address			
tche		Postcode			
ispa		Tel.	1		
ofd	1.7.	Country of origin ISO code	I.8. Region of origin Code	I.9. Country of ISO code destination	l.10.
slis				destination	
Deta	1.11.	Place of origin		l.12.	
÷		Name	Approval number		
Pal		Address			_
	1.13.	Place of loading		I.14. Date of departure	
	l.15.	Means of transport		I.16. Entry BIP in EU	
		Aeroplane 🗌 Ship	Railway wagon		
		Road vehicle D Other			
		Identification		1.17.	
		Documentation references			
	l.18.	Description of commodity		I.19. Commodity co	de (HS code)
					I.20. Quantity
	1.21.	Temperature of product			I.22. Number of packages
		Ambient 🔲	Chilled 🔲	Frozen 🗖	
	1.23.	Seal/Container No			I.24. Type of packaging
	1.25.	Commodities certified for:			
		Human consumption			
				1	
	1.26.			I.27. For import or admission into I	
	_				
	1.28.	Identification of the commodities	3		
		Manufacturing start	Number of sectors	Out of the Allahout Allahout	usindat Datah mumban
		Manufacturing plant	Number of packages	Species Net w (Scientific name)	veight Batch number
				· · · · · · · · · · · · · · · · · · ·	

V MII	▼	M1
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	cou	NTRY	Dairy products deriv	Model Milk-RMP ved from raw milk for human consumption			
	II.	Health information	II.a. Certificate reference number	II.b.			
	II.1.	Animal Health Attestation					
		I, the undersigned official veterinarian, declare that I 853/2004 and hereby certify that the dairy products					
cation	(a) under the control of the official veterinary service,						
Part II: Certification		(b) which were in a country or part thereof that has I prior to the date of this certificate, and where values					
Part		(c) belonging to holdings which were not under res	trictions due to foot-and-mouth disease or	rinderpest, and			
		(d) subject to regular veterinary inspections to ensu Annex III to Regulation (EC) No 853/2004 and i		itions laid down in Chapter I of Section IX of			
	II.2.	Public Health attestation					
		I, the undersigned official inspector, declare that I an (EC) No 853/2004 and (EC) No 854/2004 and here accordance with those provisions, in particular that:	by certify that the dairy product made with				
		(a) it was manufactured from raw milk:					
	 (i) which comes from holdings registered in accordance with Regulation (EC) No 852/2004 and checked in accordance with Regulation (EC) No 854/2004, 						
		(ii) which was produced, collected, cooled, store Section IX of Annex III to Regulation (EC) N		hygiene conditions laid down in Chapter I of			
		(iii) which meets the plate and somatic cell co 853/2004,	unt criteria laid down in Chapter I of Sec	ction IX of Annex III to Regulation (EC) No			
		(iv) which complies with the guarantees on the re or substances submitted in accordance with					
			ter I, Part III, point 4 of Regulation (EC) I	d business operator in accordance with the No 853/2004, it complies with the maximum the Annex to Regulation (EU) No 37/2010;			
		 (vi) which has been produced under conditions Regulation (EC) No 396/2005, and maximur 					
		 (b) it comes from an establishment implementing No 852/2004, 	a programme based on the HACCP prir	nciples in accordance with Regulation (EC)			
		(c) it has been obtained from raw milk that has manufacturing process,	not undergone any heat treatment or any	/ physical or chemical treatment during the			
		 (d) it has been wrapped, packaged and labeled in No 853/2004, 	n accordance with Chapters III and IV of	Section IX of Annex III to Regulation (EC)			
		(e) it meets the relevant microbiological criteria laid	I down in Regulation (EC) No 2073/2005	on microbiological criteria for foodstuffs, and			
		(f) the guarantees covering live animals and proc 96/23/EC, and in particular Article 29 thereof, and		ans submitted in accordance with Directive			

COUNTRY	Dairy products	Dairy products derived from raw milk for human consumpt		
II. Health information	II.a. Certificate reference number	II.b.		
Notes				
	s derived from raw milk for human consumption, 605/2010 intended for importation into the Europ			
Part I:				
- Box reference I.7: Provide name and IS	O code of the country or part thereof as appea	ring in Annex I to Regulation (EU) No 605/2010		
- Box reference I.11: Name, address and	approval number of the establishment of dispatch).		
— Box reference I.15: Registration number (railway wagons or container and road vehicles), 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 the 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 under the following headings: 04.01; 04.02; 04.03; 04.04; 04.05; 04.05; 04.02; 21.05; 22.02; 35.01; 35.02 or 35.04.				
- Box reference I.20: Indicate total gross v	veight and total net weight.			
- Box reference I.23: For containers or bo	xes, the container number and the seal number (if applicable) should be included.		
 Box reference I.28: Manufacturing plant: i approved for exportation to the European 	ntroduce the approval number of the production ho n Union.	olding(s), collection centre or standardization centr		
Part II:				
- The colour of the signature shall be differ	ent to that of the printing. The same rule applies to	o stamps other than those embossed or watermark		
Official veterinarian				
Name (in capital letters):	Qu	alification and title:		
Date:	Sig	inature:		
Stamp:	Stamp:			

Model Milk-HTB

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

cour	ITRY			Veterinary certificate to EU	
	l.1.	Consignor	I.2. Certificate reference No I.2	2.a.	
		Name Address	I.3. Central competent authority		
ient		Tel.	I.4. Local competent authority		
Part I: Details of dispatched consignment	1.5.	Consignee Name Address	1.6.		
spatche	Postcode Tel.				
ails of dis	1.7.	Country of ISO code I.8. Region of origin Code origin	I.9. Country of ISO code I.1 destination	10.	
I: Det	l.11.	Place of origin	l.12.		
Part	Name Approval number Address				
	l.13.	Place of loading	I.14. Date of departure		
	l.15.	Means of transport	I.16. Entry BIP in EU		
	Aeroplane D Ship Railway wagon D				
		Road vehicle Other I Identification Documentation references	1.17.		
	l.18.	Description of commodity	I.19. Commodity code (H	HS code)	
			1.20.	. Quantity	
	l.21.	Temperature of product Ambient Chilled	I.22.	. Number of packages	
	1.23.	Seal/Container No	1.24.	. Type of packaging	
	1.25.	Commodities certified for:			
		Human consumption 🗌			
	1.26.		I.27. For import or admission into EU		
	1.28.	Identification of the commodities	1		
		Manufacturing plant Number of packages	Species Net weight (Scientific name)	t Batch number	

соι	UNTRY	Dairy products derived from milk of co consumption from third countries author	<i>Model Milk-H</i> ws, ewes, goats and buffaloes for huma prised in column B
Π.	Health information	II.a. Certificate reference number	II.b.
11.1	. Animal Health Attestation		
_	I, the undersigned official veterinarian, declare to 853/2004 and hereby certify that the dairy proc		ective 2002/99/EC and of Regulation (EC) N
	(a) has been obtained from animals:		
	(i) under the control of the official veterinal	ry service,	
	 (ii) which were in a country or part thereof months prior to the date of this certificate period, 	and of rinderpest for a period of at least 1 n disease has not been carried out during th	
	(iii) belonging to holdings which were not u	nder restrictions due to foot-and-mouth disease	e or rinderpest, and,
	(iv) subject to regular veterinary inspections Annex III to Regulation (EC) No 853/20	to ensure that they satisfy the animal health cor 04 and in Directive 2002/99/EC,	ditions laid down in Chapter I of Section IX
		milk which has been submitted to a pasteurisati to that achieved by a pasteurisation process of eaction to an alkaline phosphatase test applied	of at least 72°C for 15 seconds and whe
11.2	2. Public Health attestation		
	I, the undersigned official veterinarian, declar No 852/2004, (EC) No 853/2004 and (EC) N accordance with those provisions, in particular	o 854/2004 and hereby certify that the dairy	
	(a) it was manufactured from raw milk:		
	 (i) which comes from holdings registered in Regulation (EC) No 854/2004, 	accordance with Regulation (EC) No 852/2004	and checked in accordance with Annex IV
	(ii) which was produced, collected, cooled, Section IX of Annex III to Regulation (E	stored and transported in accordance with the C) No 853/2004,	hygiene conditions laid down in Chapter I
	(iii) which meets the plate and somatic ce 853/2004,	Il count criteria laid down in Chapter I of Se	ction IX of Annex III to Regulation (EC) N
		he residues status of raw milk provided by the with Council Directive 96/23/EC, and in partici	
	requirements of Annex III, Section IX, C	of antibacterial drugs carried out by the foo Chapter I, Part III, point 4 of Regulation (EC) ial veterinary medicinal products laid down in	No 853/2004, it complies with the maximu
		ions guaranteeing compliance with the maximu imum levels for contaminants laid down in Reg	
	(b) it comes from an establishment implemen No 852/2004,	ting a programme based on the HACCP pri	nciples in accordance with Regulation (EC
	(c) it has been processed, stored, wrapped, p Annex II to Regulation (EC) No 852/2004 a	ackaged and transported in accordance with nd Chapter II of Section IX of Annex III to Reg	
	(d) it meets the relevant criteria laid down in microbiological criteria laid down in Regulat	Chapter II of Section IX of Annex III to Reg ion (EC) No 2073/2005 on microbiological crite	

(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.

COUNTRY			<i>Model Milk-H</i> f cows, ewes, goats and buffaloes for huma uthorised in column B
II. Health information	II.a. Certificate r	eference number	II.b.
Notes			
This certificate is intended for dairy pr Regulation (EU) No 605/2010 intended			arts thereof authorised in column B of Annex I o
Part I:			
- Box reference I.7: Provide name a	and ISO code of the country or pa	art thereof as appeari	ng in Annex I to Regulation (EU) No 605/2010
- Box reference I.11: Name, address	and approval number of the estab	lishment of dispatch.	
transport in containers, the total nu	mber of containers and their registra	ation number and whe	ht number (aircraft) or name (ship). In the case a re there is a serial number of the seal it must b ne border inspection post of introduction into th
[∞] Box reference I.19: Use the appr 04.06; 15.17; 17.02; 21.05; 22.0	opriate Harmonised System (HS) c 2; 28.35; 35.01; 35.02 or 35.04. ◀	code under the follows	ing headings: 04.01; 04.02; 04.03; 04.04; 04.05
- Box reference I.20: Indicate total g	ross weight and total net weight.		
- Box reference I.23: For containers	or boxes, the container number an	d the seal number (if	applicable) should be included.
 Box reference I.28: Manufacturing export to the European Union. 	plant: introduce the approval num	ber of the treatment a	and/or processing establishment(s) approved for
Part II:			
- The colour of the signature shall be	different to that of the printing. The	same rule applies to s	stamps other than those embossed or watermar
Official veterinarian			
Name (in capital letters):		Qual	lification and title:
Date:		Sign	ature:
Stamp:			

Model Milk-HTC

Health certificate for dairy products for human consumption from third countries or parts thereof authorised in column C of Annex I to Regulation (EU) No 605/2010 intended for importation into the European Union

cou	NTR	(Veterinary certificate to EU			
	l.1.	Consignor Name	I.2. Certificate reference No I.2.a.			
		Address	I.3. Central competent authority			
nent		Tel.	I.4. Local competent authority			
Part I: Details of dispatched consignment	1.5.	Consignee Name Address	1.6.			
lispatche		Postcode Tel.				
etails of c	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10.			
ä	l.11.	Place of origin	I.12.			
Part		Name Approval number Address				
	l.13.	Place of loading	I.14. Date of departure			
	l.15.	Means of transport	I.16. Entry BIP in EU			
		Aeroplane 🗌 Ship 🗌 Railway wagon 🗌				
		Road vehicle Other				
		Identification Documentary references	I.17. I.19. Commodity code (HS code)			
	l.18.	Description of commodity				
		'				
			I.20. Quantity			
	I.21.	Temperature of product	I.22. Number of packages			
		Ambient Chilled	Frozen			
	1.23.	Seal/Container No	I.24. Type of packaging			
	1.25.	Commodities certified for:				
		Human consumption 🔲				
	1.26.		I.27. For import or admission into EU			
	1.28.	Identification of the commodities				
		Species Manufacturing plant Number of (scientific name)	packages Net weight Batch number			

▼<u>M3</u>

▼<u>M3</u>

	COUNTR	'RY		<i>Model Milk-HTC</i> Dairy products from third countries authorised in column C							
	11.	Health	n information	a. Certificate reference number	II.b.						
	II.1.	Anima	al Health Attestation								
		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:									
tion		(a) has been obtained from animals:									
Part II: Certification		(i) under the control of the official veterinary service;									
l: Ce		(ii) belonging to holdings which were not under restrictions due to foot-and-mouth disease or rinderpest; and									
Part	(iii) subject to regular veterinary inspections to ensure that they satisfy the animal health conditions laid down in Chap IX of Annex III to Regulation (EC) No 853/2004 and in Directive 2002/99/EC;										
	either	fo	ne dairy product was made from raw milk sourced from co botnote (2) of Annex I to Regulation (EC) No 605/2010, from - b import into the territory of the European Union:								
	(¹) eithe	r	[(i) a sterilisation process, to achieve an F_0 value equal to σ	or greater than three;]							
	(¹) or	[(ii) an ultra-high temperature (UHT) treatment at not less the	an 135 °C in combination with a s	uitable holding time;]						
	(¹) or	[(iii) a high temperature-short time pasteurisation treatment (H or greater than 7,0 achieving, where applicable, a negati the heat treatment;]								
	(¹) or	[(a treatment with an equivalent pasteurisation effect to po phosphatase test, applied immediately after the heat treatment 		e, a negative reaction to an alkaline						
	(¹) or	[(v) a HTST treatment of milk with a pH below 7,0;]									
(1) or [(vi) a HTST treatment combined with another physical treatment by											
	(1)	either	[(1) lowering the pH below 6 for one hour;]								
	(1)	(¹) or [(2) additional heating equal to or greater than 72 °C, combined with desiccation;]]									
	(¹) or	or [(b) the dairy product was made from raw milk sourced from animals other than cows, ewes, goats, buffaloes or camels of the species Camelus dromedarius, and has undergone, prior to import into the territory of the European Union:									
	(¹) either	· [(i) a sterilisation process, to achieve an F_0 value equal to c	or greater than three;]							
	(¹) or	[0	(ii) an ultra-high temperature (UHT) treatment at not less that	an 135 °C in combination with a si	uitable holding time;]]						
	II.2.	Public	e Health attestation								
		No 85	undersigned official inspector, declare that I am aware of s2/2004, (EC) No 853/2004 and (EC) No 854/2004 and her dance with those provisions, and in particular that:								
		(a) it	was manufactured from raw milk:								
			 which comes from holdings registered in accordance with IV to Regulation (EC) No 854/2004; 	Regulation (EC) No 852/2004 and	checked in accordance with Annex						
		(which was produced, collected, cooled, stored and transp of Section IX of Annex III to Regulation (EC) No 853/200		ne conditions laid down in Chapter I						
		(1	iii) which meets the plate and somatic cell count criteria la No 853/2004;	aid down in Chapter I of Section	IX of Annex III to Regulation (EC)						
		(i	 which complies with the guarantees on the residues sta residues or substances submitted in accordance with Co 								

COUNT	RY	Dairy products from third	Model Milk-HT
II.	Health information	II.a. Certificate reference number	II.b.
	 (v) which, pursuant to testing for residues of antibacteri requirements of Annex III, Section IX, Chapter I, Part residue limits for residues of antibacterial veterinary 	III, point 4 to Regulation (EC) No 853/2	2004, it complies with the maximun
	(vi) which has been produced under conditions guarantee Regulation (EC) No 396/2005, and maximum levels		
	(b) it comes from an establishment implementing a program No 852/2004;	nme based on the HACCP principles in	n accordance with Regulation (EC
	(c) it has been processed, stored, wrapped, packaged and t Annex II to Regulation (EC) No 852/2004 and Chapter II		
	(d) it meets the relevant criteria laid down in Chapter II of microbiological criteria laid down in Regulation (EC) No		
	(e) the guarantees covering live animals and products ther 96/23/EC, and in particular Article 29 thereof, are fulfilled		mitted in accordance with Directive
Notes			
	artificate is intended for dairy products for human consumption fr animal species only, in column C of Annex I to Regulation (El		
Part I:			
— Box	reference I.7: provide name and ISO code of the country o	r part thereof as appearing in Annex I	I to Regulation (EU) No 605/2010
— Вох	reference I.11: name, address and approval number of the es	tablishment of dispatch.	
In tł it m	reference I.15: registration number (railway wagons or containe he case of transport in containers, the total number of containers ust be indicated in box I.23. In the case of unloading and reload European Union.	and their registration number and where	there is a serial number of the sea
	reference I.19: use the appropriate Harmonised System (HS) o 17; 17.02; 19.01; 21.05; 21.06; 22.02; 28.35; 35.01; 35.02 or 3		; 04.02; 04.03; 04.04; 04.05; 04.06
— Box	reference I.20: indicate total gross weight and total net weight		
— Вох	reference I.23: for containers or boxes, the container number	and the seal number (if applicable) sho	uld be included.
	reference I.28: manufacturing plant: introduce the approval num he European Union.	ber of the treatment and/or processing e	establishment(s) approved for expor
Part II:	:		
(¹) Kee	ap as appropriate.		
— The	colour of the signature shall be different to that of the printing.	The same rule applies to stamps other t	than those embossed or watermark
Official	veterinarian		
Nar	ne (in capital letters):	Qualifica	ation and title:
Dat	e:	Signatur	e:
Star	mp:		

▼<u>M3</u>

Model Colostrum/Colostrum-based products- C/CBP

Health Certificate for colostrum of cows, ewes, goats and buffaloes and colostrum- based products derived from colostrum of the same species from third countries or parts thereof listed in column A of Annex I for human consumption intended for importation into the European Union

coul	NTRY	,	Veterinary certificate to E		
	l.1.	Consignor	I.2. Certificate reference No I.2.a.		
		Name			
		Address	I.3. Central competent authority		
		Tel.			
nent			I.4. Local competent authority		
ignn	1.5.	Consignee	1.6.		
üö		Name			
ed c		Address			
atch		Postcode			
disp		Tel.			
of	1.7.	Country of origin ISO code I.8.	I.9. Country of destination ISO code I.10.		
tails					
Part I: Details of dispatched consignment	1.11.	Place of origin	1.12.		
art		Name Approval number			
-		Address			
		Address			
	1 1 2	Place of loading	I.14. Date of departure		
	1.13.	Frace of roading			
	I.15.	Means of transport	I.16. Entry BIP in EU		
		Aeroplane 🗌 Ship 🗌 Railway wagon 🗌			
		Road vehicle Other O			
			1.17.		
		Identification Documentary references			
	1.18.	Description of commodity	I.19. Commodity code (HS code)		
			I.20. Quantity		
	1.21.	Temperature of product	I.22. Number of packages		
		Ambient Chilled	Frozen		
	1.23.	Seal/Container No	I.24. Type of packaging		
	1.25.	Commodities certified for:	I		
		Human consumption			
	1.26.		I.27. For import or admission into EU		
	1.28.	Identification of the commodities			
		Species Manufacturing plant Num (Scientific name)	ber of packages Net weight Batch number		
		· · · ·			

		Colostrum and colostrum b or parts thereof listed	Colostrum –Based Products C/CBF ased products from third countries in column A of Annex I for humar	
	JNTRY Health information	II.a. Certificate reference number	II.b.	
".			11.0.	
.1	Animal Health Attestation			
	I, the undersigned official veterinarian, declare that I am aware of t 853/2004 and hereby certify that the colostrum /colostrum-based p		02/99/EC and of Regulation (EC) No	
cation	have been obtained or manufactured from colostrum obtained from	n animals:		
	(i) under the control of the official veterinary service;			
Lau II:	 (ii) which were in a third country or part thereof that has been f months prior to the date of this certificate, and where vaccir period; 			
	(iii) belonging to holdings which were not under restrictions du	e to foot-and-mouth disease or rinder	rpest, and	
	(iv) subject to regular veterinary inspections to ensure that they Annex III to Regulation (EC) No 853/2004 and in Directive		aid down in Chapter I of Section IX of	
11.2	Public Health Attestation			
I, the undersigned official inspector, declare that I am aware of the relevant provisions of Regulations (EC) No (EC) No 853/2004 and (EC) No 854/2004 and hereby certify that the colostrum/colostrum-based products in described in Part I were produced in accordance with those provisions, and in particular that:				
(a) they were manufactured from colostrum:				
	 which comes from holdings registered in accordance with F Regulation (EC) No 854/2004; 	Regulation (EC) No 852/2004 and che	ecked in accordance with Annex IV to	
	 (ii) which was produced, collected, cooled, stored and transp- Section IX of Annex III to Regulation (EC) No 853/2004; 	orted in accordance with the hygiene	conditions laid down in Chapter I of	
	(iii) which complies with the guarantees on the residues status or substances submitted in accordance with Directive 96/2			
	 (iv) which, pursuant to testing for residues of antibacterial d requirements of point 4 in Part III of Chapter I of Secti maximum residue limits for residues of antibacterial veter 37/2010; 	on IX of Annex III to Regulation (E	C) No 853/2004, complies with the	
	 (v) which has been produced under conditions guaranteeing Regulation (EC) No 396/2005, and maximum levels for co 			
	(b) they come from an establishment implementing a programme 852/2004;	based on the HACCP principles in a	accordance with Regulation (EC) No	
	 (c) they have been processed, stored, wrapped, packaged and la Regulation (EC) No 853/2004; 	beled in accordance with Chapters II	I and IV of Section IX of Annex III to	
	(d) they meet the relevant requirements laid down in Chapter II, microbiological criteria laid down in Regulation (EC) No 2073/2			

(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.

Model Colostrum/Colostrum –Based Products C/CBP Colostrum and colostrum based products from third countries or parts thereof listed in column A of Annex I for human consumption intended for importation

COUN	ITRY	cons	sumption intended for importation						
П.;	Health information	II.a. Certificate reference number	II.b.						
Notes	Notes								
	This certificate is intended for colostrum or colostrum-based products from third countries or parts thereof authorised in column A of Annex I to Regulation (EU) No 605/2010.								
Part	1:								
60	— Box reference I.7: Provide name and ISO code of the country or part thereof as appearing in Annex I to Commission Regulation (EU) No 605/2010 of 2 July 2010 laying down animal and public health and veterinary certification conditions for the introduction into the European Union of raw milk and dairy products intended for human consumption (OJ L 175, 10.7.2010, p. 1).								
— в	ox reference I.11: Name, address and approval number of the esta	blishment of dispatch.							
	ox reference I.15: Registration number (railway wagons or contain nloading and reloading, the consignor must inform the border inspe								
	ox reference I.19: Use the appropriate Harmonised System (HS) cod 4.10; 15.17; 17.02; 19.01; 21.05; 21.06; 22.02; 28.35; 30.01; 35.01;		; 04.02; 04.03; 04.04; 04.05; 04.06;						
— в	ox reference I.20: Indicate total gross weight and total net weight.								
— в	ox reference I.23: For containers or boxes, the container number a	nd the seal number (if applicable) sho	ould be included.						
	ox reference I.28: Manufacturing plant: introduce the approval numb oproved for exportation to the European Union.	er of the production holding(s), collect	ion centre or standardization centre						
Part	И:								
(¹) K	eep as appropriate.								
— т	he colour of the signature shall be different to that of the printing. Th	ne same rule applies to stamps other t	han those embossed or watermark.						
Offici	al veterinarian								
N	Name (in capital letters): Qualification and title:								
D	Date: Signature:								
S	Stamp:								

PART 3

Model Milk/Colostrum-T/S

Animal Health Certificate for raw milk, dairy products, colostrum and colostrum-based products for human consumption intended for transit through or storage in the European Union

cou	INTRY	,							Veterinary certificate to EL
	1.1.	Consignor				1.2.	Certificate reference No	1.2	2.a.
	Name							_	
		Address				1.3.	I.3. Central competent authority		
dispatched consignment		Tel.			1.4.	I.4. Local competent authority			
l ū	1.5.	. Consignee				1.6.	Person responsible for the lo	ad in E	U
Suc		Name					Name		
S S		Address					Address		
hec		Postcode				Postal code			
patc		Tel.				Tel. No			
Part I: Details of dis	1.7.	Country of origin	ISO code	I.8. Region of origin	Code	1.9.	Country of ISO destination	code	1.10.
etail	1.11.	Place of origin				I.12.	Place of destination		e
0 ::		Name		Approval number			Customs warehouse 🗌		Ship supplier 🔲
art		Address					Name	App	proval number
•							Address		
							Postal code		
	1.13.	Place of loading				1.14.	Date of departure		
	1.45	-				1.40			
	1.15.	I.15. Means of transport			1.16.	Entry BIP in EU			
		Aeroplane	Ship [on 🗖				
		Road vehicle 🗌	Other			1.17.			
		Identification							
		Documentary references							
	1.18.	18. Description of commodity				I.19. Commodity co	ode (HS	S code)	
								1.20.	Quantity
	1.21.	Temperature of pro	oduct					1.22.	Number of packages
		Ambient 🔲		Chilled 🔲		Frozen 🗌			
	1.23.	Seal/Container No						1.24.	Type of packaging
	1.25.	Commodities certif	ied for:						
		Human consumption 🔲							
	1.26.	For transit through	EU to 3rd 0	Country 🔲		1.27.			
		3rd country		ISO code					
	1.28.	Identification of the	commoditie	95					
		Species (Scientific name)		Manufacturing plant	Nu	ımber	of packages Net w	eight	Batch number

▼	Μ	6

С	OUNTRY		ts, colostrum and colostrum-base consumption for transit or storag
I	I. Health information	II.a. Certificate reference number	II.b.
I	I.1 Animal Health Attestation		
	I, the undersigned official veterinarian, hereby certify that the [transit] / [storage] $(^2)$ in the European Union described in P] / [colostrum-based products] $(^{1})$ $(^{2})$ for
	(a) come from a country or part thereof authorised for importance based products as laid down in Annex I to Regulation (I		dairy products, colostrum or colostrum
	(b) comply with the relevant animal health conditions fo II.1 of the model health certificates [Milk-RM] / [Milk-F Regulation (EU) No 605/2010;	r the products concerned as laid dow RMP] / [Milk-HTB] / [Milk-HTC]/[Colostru	n in the animal health attestation in Par m-C/CBP](2) in Part 2 of Annex II to
	(c) was/were produced on	(³) or between	
r	Votes		
F	Part I:		
_	 Box reference I.7: Provide name and ISO code of the cc No 605/2010 of 2 July 2010 laying down animal and public h Union of raw milk and dairy products intended for human con 	ealth and veterinary certification condition	
_	 Box reference I.11: Name, address and approval number of t same as the country of export. 	the establishment of dispatch. Name of t	he country of origin which must be the
-	 Box reference I.15: Registration number (railway wagons or cor of transport in containers, the total number of containers a be indicated in box I.23. In case of unloading and reloading European Union. 	nd their registration number and where the	ere is a serial number of the seal it mus
_	 Box reference I.19: Use the appropriate Harmonised Sys 04.05; 04.06; 15.17; 17.02; 19.01; 21.05; 21.06; 22.02; 		
-	- Box reference I.20: Indicate total gross weight and total net w	veight.	
_	- Box reference I.23: For containers or boxes, the co	ntainer number and the seal numb	er (if applicable) should be included
-	 Box reference I.28: Manufacturing plant: introduce th standardization centre approved for exportation to the E 		tion holding(s), collection centre o
F	Part II:		
(Raw milk, dairy products, colostrum and colostrum -bass products for human consumption in transit or storage in ac December 1997 laying down the principles governing the or countries (OJ L 24, 30.1.1998, p. 9). 	cordance with Article 12(4) or Article 13	3 of Council Directive 97/78/EC of 18
(²) Keep as appropriate.		
(³) Date or dates of production. Imports of raw milk, dairy prod obtained either prior to the date of authorisation for export I.7 and I.8, or during a period where restrictive measures dairy products, colostrum and colostrum-based products	rtation to the European Union of the third have been adopted by the European	country or part thereof mentioned under

COUNTRY	Model Milk/Colostrum-7/S Raw milk, dairy products, colostrum and colostrum-based products for human consumption for transit or storage				
II. Health information	II.a. Certificate reference II.b. number				
Official veterinarian					
Name (in capital letters):	Qualification and title:				
Date:	Signature:				
Stamp:					