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►<u>B</u>

▶<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
► <u>M10</u>	Commission Delegated Regulation (EU) 2019/2124 of 10 October 2019	L 321	73	12.12.2019

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:

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 (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;

▼<u>B</u>

(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.

▼<u>M10</u>

▼<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;

- (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.

▼M6

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	+	+	+
PA	Panama	0	0	+
PY	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 drive resoluted of the meaning Council of demonstration.

⁽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	OUNTRY: Veterinary certificate to EU						
	l.1.	Consignor			I.2. Certifica	te reference No	l.2.a.
		Name			I.3. Central	competent authority	
		Address					
nent		Tel.			I.4. Local co	ompetent authority	
signr	1.5.	Consignee			1.6.		
cons		Name Address					
hed		Postcode					
Part I: Details of dispatched consignment		Tel.					
f dis	1.7.	Country of origin ISO code	I.8. Region of origin	Code	I.9. Country		I.10.
ils o		I			destinat	lion	
Deta	l.11.	. Place of origin	LL		l.12.		
벁		Name	Approval numbe	er			
Pa		Address					
	1.40	Disco of incolling					
	1.13.	Place of loading			I.14. Date of	departure	
	l.15.	Means of transport			I.16. Entry B	IP in EU	
		Aeroplane 🗌 Ship	Railway wa	agon 🔲			
		Road vehicle Other			l.17.		
		Identification			1.17.		
		Documentation references					
	l.18.	. Description of commodity				I.19. Commodity co	de (HS code)
							I.20. Quantity
	1.01	.					
	1.21.	. Temperature of product Ambient □	Chilled 🔲			Frozen 🗖	I.22. Number of packages
	123	. Seal/Container No					I.24. Type of packaging
	1.25.	Commodities certified for:					
		Further process					
	1.26.				I.27. For imp	ort or admission into I	
	1.28.	Identification of the commodities	5				
		Manufacturing plant	Number of packages		Species (Scientific nam		veight Batch number

Model Milk-RM

	COUNTRY Raw milk				
	II.	Health information II.	a. Certificate reference number	II.b.	
	11.1.	. Animal Health Attestation			
		I, the undersigned official veterinarian, declare that I ar 853/2004 and hereby certify that the raw milk describ			
ation		(a) under the control of the official veterinary service,			
Part II: Certification		(b) which were in a country or part thereof that has be prior to the date of this certificate, and where vac			
art II		(c) belonging to holdings which were not under restrict	ctions due to foot-and-mouth disease or r	inderpest, and	
<u>c</u>		(d) subject to regular veterinary inspections to ensure Annex III to Regulation (EC) No 853/2004 and in		ions laid down in Chapter I of Section IX of	
	II.2.	. Public Health attestation			
		I, the undersigned official inspector, declare that I am a (EC) No 853/2004 and (EC) No 854/2004 and hereb provisions, in particular that:			
		 (a) it comes from holdings registered in accordance Regulation (EC) No 854/2004, 	with Regulation (EC) No 852/2004 and	checked in accordance with Annex IV to	
		(b) it was produced, collected, cooled, stored and trans of Annex III to Regulation (EC) No 853/2004,	sported in accordance with the hygiene co	nditions laid down in Chapter 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 p in accordance with Council Directive 96/23/EC, ar			
		(e) pursuant to testing for residues of antibacterial dru Annex III, Section IX, Chapter I, Part III, point 4 of F of antibacterial veterinary medicinal products laid	Regulation (EC) No 853/2004, it complies	with the maximum residue limits for residues	
		 (f) it has been produced under conditions guaranteeir (EC) No 396/2005, and maximum levels for conta 			
	Note	tes			
		s certificate is intended for raw milk from third countries ended for further processing in the European Union befo		of Annex I to Regulation (EU) No 605/2010	
	Part	rt I:			
	— E	Box reference I.7: Provide name and ISO code of the	e 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	er of the establishment of dispatch.		
		Box reference I.15: Registration number (railway wagor unloading and reloading, the consignor must inform the			
	— E	Box reference I.19: Use the appropriate Harmonised Sy	ystem (HS) code under the following head	dings: 04.01; 04.02 or 04.03.	
	— е	Box reference I.20: Indicate total gross weight and total	l net weight.		
	— E	Box reference I.23: For containers or boxes, the contain	ner number and the seal number (if appli	cable) should be included.	
		Box reference I.28: Manufacturing plant: introduce the ap approved for exportation to the European Union.	oproval number of the production holding(s), collection centre or standardization centre	

COUN.	TRY		<i>Model Milk-RM</i> Raw milk
II.	Health information	II.a. Certificate reference number	II.b.
Part I — Th	I: e colour of the signature shall be different to that o	f the printing. The same rule applies to stamp	s other than those embossed or watermark.
Officia	al veterinarian		
	Name (in capital letters):	Qualificati	on 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

COUNTRY: Veterinary certificate to EU					
	l.1.	0	I.2. Certificate reference No I.2.a.		
		Name Address	I.3. Central competent authority		
ţ			I.4. Local competent authority		
Part I: Details of dispatched consignment		Tel.			
nsigr	1.5.	Consignee Name	1.6.		
d co		Address			
Itche		Postcode			
dispa		Tel.			
s of e	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10. destination		
etail					
	1.11.	Place of origin Name Approval number	1.12.		
Parl		Address			
	l.13.	Place of loading	I.14. Date of departure		
	l.15.	Means of transport	I.16. Entry BIP in EU		
		Aeroplane			
		Road vehicle Other I Identification	1.17.		
		Documentation references			
	l.18.	Description of commodity	I.19. Commodity code (HS code)		
			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			
		Manufacturing plant Number of packages	Species Net weight Batch number (Scientific name)		

	Model Milk-R COUNTRY Dairy products derived from raw milk for human consumpt				
	11.	Health information	II.a. Certificate reference number	II.b.	
	II.1.	Animal Health Attestation	<u></u>		
		I, the undersigned official veterinarian, declare that 853/2004 and hereby certify that the dairy produc			
cation		(a) under the control of the official veterinary serv	ice,		
Part II: Certification		(b) which were in a country or part thereof that has prior to the date of this certificate, and where			
Part		(c) belonging to holdings which were not under re	estrictions due to foot-and-mouth disease or	rinderpest, and	
		(d) subject to regular veterinary inspections to ens Annex III to Regulation (EC) No 853/2004 and		itions laid 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 178/2002, (EC) No 852/2004 (EC) No 853/2004 and (EC) No 854/2004 and hereby certify that the dairy product made with raw milk described above was produced in accordance with those provisions, in particular that:				
		(a) it was manufactured from raw milk:			
		 (i) which comes from holdings registered in ac Regulation (EC) No 854/2004, 	cordance with Regulation (EC) No 852/2004	and checked in accordance with Annex IV to	
		(ii) which was produced, collected, cooled, sto Section IX of Annex III to Regulation (EC)		hygiene conditions laid down in Chapter I of	
		(iii) which meets the plate and somatic cell o 853/2004,	count 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 or substances submitted in accordance with 	residues status of raw milk provided by the in the Council Directive 96/23/EC, and in particular		
			apter 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 condition Regulation (EC) No 396/2005, and maximum	s guaranteeing compliance with the maximu um levels for contaminants laid down in Reg		
		 (b) it comes from an establishment implementing No 852/2004, 	g 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 No 853/2004,	in accordance with Chapters III and IV of	Section IX of Annex III to Regulation (EC)	
		(e) it meets the relevant microbiological criteria la	id down in Regulation (EC) No 2073/2005	on microbiological criteria for foodstuffs, and	
		(f) the guarantees covering live animals and pr 96/23/EC, and in particular Article 29 thereof,		ans submitted in accordance with Directive	

COUNTRY	Dairy products o	derived from raw milk for human consumpti			
II. Health information	II.a. Certificate reference number	II.b.			
Notes					
	his certificate is intended for dairy products derived from raw milk for human consumption, from third countries or parts thereof authorised in olumn A of Annex I to Regulation (EU) No 605/2010 intended for importation into the European Union.				
Part I:	art I:				
- Box reference I.7: Provide name and IS	O code of the country or part thereof as appeari	ng in Annex I to Regulation (EU) No 605/201			
- Box reference I.11: Name, address and a	approval number of the establishment of dispatch.				
transport in containers, the total number of	ailway wagons or container and road vehicles), flig of containers and their registration number and whe ading and reloading, the consignor must inform th	ere there is a serial number of the seal it must b			
 Box reference I.19: Use the appropriate Ha 17.02; 21.05; 22.02; 35.01; 35.02 or 35.0 	armonised System (HS) code under the following he 4.	eadings: 04.01; 04.02; 04.03; 04.04; 04.05; 04.0			
- Box reference I.20: Indicate total gross w	eight and total net weight.				
- Box reference I.23: For containers or box	es, the container number and the seal number (if	applicable) should be included.			
 Box reference I.28: Manufacturing plant: in approved for exportation to the European 	troduce the approval number of the production hole Union.	ding(s), collection centre or standardization cent			
Part II:					
- The colour of the signature shall be differe	ent to that of the printing. The same rule applies to s	stamps other than those embossed or waterma			
Official veterinarian					
Name (in capital letters):	Qual	lification and title:			
Date:	Sign	ature:			
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

cou	OUNTRY: Veterinary certificate to EU					
	l.1.	Consignor	I.2. Certificate reference No I.2.a.			
		Name Address	I.3. Central competent authority			
lent		Tel.	I.4. Local competent authority			
of dispatched consignment	1.5.	Consignee Name Address Postcode Tel.	1.6.			
ails of dis	l.7.	Country of ISO code I.8. Region of origin Code origin	I.9. Country of ISO code I.10.			
Part I: Details	l.11.	Place of origin Name Approval number Address	1.12.			
	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 Railway wagon Road vehicle Other C				
		Identification Documentation references	l.17.			
	l.18.	Description of commodity	I.19. Commodity code (HS code)			
			I.20. Quantity			
	l.21.	Temperature of product Ambient Chilled	I.22. Number of packages			
	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				
		Manufacturing plant Number of packages	Species Net weight Batch number (Scientific name)			

	сои	NTRY	Dairy products derived from milk of co consumption from third countries author	<i>Model Milk-HTE</i> ws, ewes, goats and buffaloes for human orised in column B	
	Π.	Health information	II.a. Certificate reference number	II.b.	
	II.1.	Animal Health Attestation			
	-	I, the undersigned official veterinarian, declare that 853/2004 and hereby certify that the dairy produc		ective 2002/99/EC and of Regulation (EC) No	
		(a) has been obtained from animals:			
Б		(i) under the control of the official veterinary a	service,		
Part II: Certification		 which were in a country or part thereof the months prior to the date of this certificate, a period, 		and of rinderpest for a period of at least 12 a disease has not been carried out during that	
Part II:		(iii) belonging to holdings which were not unde	er restrictions due to foot-and-mouth disease	e or rinderpest, and,	
		(iv) subject to regular veterinary inspections to Annex III to Regulation (EC) No 853/2004		nditions laid down in Chapter I of Section IX of	
	_	(b) has undergone or been produced from raw mill with a heating effect at least equivalent to the applicable, sufficient to ensure a negative read	nat achieved by a pasteurisation process o	of at least 72°C for 15 seconds and where	
	II.2.	Public Health attestation			
	I, the undersigned official veterinarian, declare that I am aware of the relevant provisions of Regulations (EC) No 178/2002, (E No 852/2004, (EC) No 853/2004 and (EC) No 854/2004 and hereby certify that the dairy product described above was produced accordance with those provisions, in particular that:				
		(a) it was manufactured from raw milk:			
		 (i) which comes from holdings registered in ac Regulation (EC) No 854/2004, 	cordance with Regulation (EC) No 852/2004	and checked in accordance with Annex IV to	
		(ii) which was produced, collected, cooled, sto Section IX of Annex III to Regulation (EC)		hygiene conditions laid down in Chapter I of	
		(iii) which meets the plate and somatic cell o 853/2004,	count criteria laid down in Chapter I of Se	ction IX of Annex III to Regulation (EC) No	
		(iv) which complies with the guarantees on the or substances submitted in accordance wi	residues status of raw milk provided by the th Council Directive 96/23/EC, and in partici		
			apter I, Part III, point 4 of Regulation (EC)	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 condition Regulation (EC) No 396/2005, and maximu	s guaranteeing compliance with the maximu um levels for contaminants laid down in Reg		
		(b) it comes from an establishment implementing a programme based on the HACCP principles in accordance with Regulation (E No 852/2004,			
		(c) it has been processed, stored, wrapped, pack Annex II to Regulation (EC) No 852/2004 and			
		(d) it meets the relevant criteria laid down in Cl microbiological criteria laid down in Regulation			
		(e) the guarantees covering live animals and pr 96/23/EC, and in particular Article 29 thereof,		ans submitted in accordance with Directive	

COUNTRY		from milk of cows, ewes d countries authorised in	<i>Model Milk-H</i> , goats and buffaloes for huma column B		
II. Health information	II.a. Certificate reference	e number II.b.			
Notes					
This certificate is intended for dairy products for human consumption from third countries or parts thereof authorised in column B of Annex I of Regulation (EU) No 605/2010 intended for importation into the European Union.					
Part I:					
- Box reference I.7: Provide name an	d ISO code of the country or part there	of as appearing in Annex I	to Regulation (EU) No 605/201		
- Box reference I.11: Name, address a	nd approval number of the establishmen	t of dispatch.			
transport in containers, the total num	er (railway wagons or container and road per of containers and their registration nu unloading and reloading, the consignor	mber and where there is a s	erial number of the seal it must b		
[∞] — Box reference I.19: Use the approp 04.06; 15.17; 17.02; 21.05; 22.02;	priate Harmonised System (HS) code und 28.35; 35.01; 35.02 or 35.04. ◀	der the following headings:	04.01; 04.02; 04.03; 04.04; 04.03		
- Box reference I.20: Indicate total gro	ss weight and total net weight.				
- Box reference I.23: For containers of	boxes, the container number and the se	eal number (if applicable) sh	ould be included.		
 Box reference I.28: Manufacturing p export to the European Union. 	ant: introduce the approval number of t	he treatment and/or proces	sing establishment(s) approved f		
Part II:					
- The colour of the signature shall be d	ifferent to that of the printing. The same r	ule applies to stamps other t	han those embossed or watermar		
Official veterinarian					
Name (in capital letters):	Name (in capital letters): Qualification and title:				
Date:		Signature:			
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

ου	NTR	(Veterinary certificate to EU					
	1.1.	Consignor Name	I.2. Certificate reference No I.2.a.					
		Address	I.3. Central competent authority					
uent		Tel.	I.4. Local competent authority					
Part I: Details of dispatched consignment	1.5.	Consignee Name Address	1.6.					
lispatched		Postcode Tel.						
etails of d	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of ISO code I.10.					
ă ≓	1.11.	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	1.17.					
	l.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:						
		Human consumption 🔲						
	1.26.		I.27. For import or admission into EU					
	1.28.	Identification of the commodities	1					
		Species Manufacturing plant Number of (scientific name)	packages Net weight Batch number					
l								

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▼<u>M3</u>

	COUN	TR۱	(D	airy p	prod	lucts	from	third	l cou	ntries	s au	thor			<i>lilk-HTC</i> Jumn C
	II.		Hea	th in	forma	ation										II.a.	Certi	ficate	refe	erenc	e num	nber	II.b).					
Part II: Certification	II.1.		l, the No t	e unc 353/2 has (i) (ii) (iii) :	dersig 2004 beer unde belor subje	and he n obtain r the c nging to	ficial v reby o ned fro ontrol o holdi	veterir certify om an of the ngs v veteri	that himals offic which	t the s: icial v n wer	dain veter re no	y pro rinaŋ ot un ons t	oduc y sei nder to en	rvice; restri	; ; ictior	ed ab ns du	ove: e to fc satisfy	oot-an	id-mo	outh nal he	diseas	se or	rinde	rpest;	and	ł			on (EC) Section
	either		[(b)	footr	note		nnex	to Re	egula	ation	(EC)) No	605	/2010															ce with e, prior
	(¹) <i>eit</i>	her		[(i)	a st	erilisat	on pro	ocess	, to a	achie	eve a	an F ₍	o val	lue e	qual	to or	great	er tha	an th	ree;]	I								
	(¹) or			[(ii)	an i	ultra-hi	gh ten	perat	ture ((UHT	T) tre	eatme	ent a	at no	t les	s thar	135	°C in	com	nbina	ation w	/ith a	suitab	ole ho	ldin	g tim	ne;]		
	(¹) or			[(iii)	or g		than 7	',0 ac																					qual to ly after
	(1) or			[(iv)		eatmer sphata													ving,	whe	əre ap	plicab	le, a	negat	ive	reac	tion to	o an a	alkaline
	(¹) or			[(v)	a H	TST tr	eatme	nt of I	milk	with	a pł	H be	elow	7,0;]															
	(¹) or			[(vi)	a H	TST tr	eatme	nt cor	mbine	ed w	vith a	anoth	ner p	ohysio	cal tr	eatm	ent by												
		(1)	eithe	ər	[(1)	lower	ng the	pH I	belov	w 6 f	for o	ne h	nour;]]															
		(1)	or		[(2)	additi	onal h	eating) equ	ual to	o or i	grea	iter t	han 1	72 °C	C, cor	nbinec	l with	des	siccat	tion;]]								
	(1) or		[(b)			produ drome																		faloes	or	cam	els of	the s	species
	(¹) eiti	her		[(i)	a st	erilisati	on pro	cess,	, to a	achie	eve a	an F _o	_o valı	ue eo	qual	to or	greate	er tha	n thr	ree;]									
	(1) or			[(ii)	an u	ıltra-hiç	h tem	perat	ure (UHT	r) tre	atme	ent a	at not	t less	s thar	135 °	°C in	com	binat	tion w	ith a s	suitab	le hol	lding	g tim	e;]]		
	II.2.		Pub	lic H	lealth	n attes	tation																						
			No	352/2	2004,		No 85	3/200)4 an	nd (E	EC) 1	No 8	354/2	2004															2, (EC) uced in
			(a)	it wa	as m	anufac	tured t	rom r	raw n	milk:																			
				(i)		h com Regu							n acc	corda	nce	with F	legula	tion (E	EC) I	No 8	352/20	04 an	d che	cked	in a	ccor	dance	e with	Annex
				(ii)		h was ection												acco	rdan	nce w	vith the	e hygi	ene c	onditi	ons	laid	down	in Cł	napter I
				(iii)		h mee 853/20		plate	e anc	d sor	matic	c cel	II co	ount d	criter	ia lai	dow	n in (Chap	pter	l of S	ectior	n IX c	of Anr	nex	III to	Reg	ulatio	n (EC)
				(iv)		ch com dues o																							ction of

COUNTR	Y	Dairy products from third	<i>Model Milk-HT</i> countries authorised in column (
11.	Health information	II.a. Certificate reference number	II.b.			
	(v) which, pursuant to testing for residues of antibacterial requirements of Annex III, Section IX, Chapter I, Part II residue limits for residues of antibacterial veterinary metabolic	I, point 4 to Regulation (EC) No 853/2	004, it complies with the maximum			
	 (vi) which has been produced under conditions guaranteein Regulation (EC) No 396/2005, and maximum levels fo 					
	 (b) it comes from an establishment implementing a programm No 852/2004; 	ne based on the HACCP principles in	n accordance with Regulation (EC			
	(c) it has been processed, stored, wrapped, packaged and tra Annex II to Regulation (EC) No 852/2004 and Chapter II o					
	(d) it meets the relevant criteria laid down in Chapter II of Si microbiological criteria laid down in Regulation (EC) No 20					
	(e) the guarantees covering live animals and products thereor 96/23/EC, and in particular Article 29 thereof, are fulfilled.	f provided by the residue plans subr	nitted in accordance with Directive			
Notes						
	tificate is intended for dairy products for human consumption fron nimal species only, in column C of Annex I to Regulation (EU)					
Part I:						
— 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.			
— Box	reference I.11: name, address and approval number of the esta	blishment of dispatch.				
In the it mu	— Box reference 1.15: registration number (railway wagons or container and road vehicles), flight number (aircraft) or name (ship) is to be provided. 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 1.23. In the case of unloading and reloading, the consignor must inform the border inspection post of introduction into the European Union.					
	reference I.19: use the appropriate Harmonised System (HS) coc 7; 17.02; 19.01; 21.05; 21.06; 22.02; 28.35; 35.01; 35.02 or 35.0		; 04.02; 04.03; 04.04; 04.05; 04.06;			
— Box	reference I.20: indicate total gross weight and total net weight.					
— Box	reference I.23: for containers or boxes, the container number ar	nd the seal number (if applicable) sho	uld be included.			
	reference I.28: manufacturing plant: introduce the approval numbe e European Union.	er of the treatment and/or processing e	stablishment(s) approved for expor			
Part II:						
(¹) Keep	as appropriate.					
— The	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			
Official v	reterinarian					
Nam	e (in capital letters):	Qualifica	tion and title:			
Date		Signature	ə:			
Stam	p:					

▼<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

cou	NTRY	,	Veterinary certificate to EU					
	l.1.	Consignor	I.2. Certificate reference No I.2.a.					
		Name						
		Address	I.3. Central competent authority					
nent		Tel.	I.4. Local competent authority					
signn	1.5.	Consignee	1.6.					
üö		Name						
p		Address						
che		Postcode						
ispat		Tel.						
of d	1.7.	Country of origin ISO code I.8.	I.9. Country of destination ISO code I.10.					
tails o								
Part I: Details of dispatched consignment	I.11.	Place of origin	1.12.					
art		Name Approval number						
-		Address						
		Address						
	1.13.	Place of loading	I.14. Date of departure					
	I.15.	Means of transport	I.16. Entry BIP in EU					
		Aeroplane 🗋 Ship 🗋 Railway wagon 🗋						
		Road vehicle Other	1.17.					
		Identification						
		Documentary references						
	I.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:						
		Human consumption						
	1.26.		I.27. For import or admission into EU					
	1.28.	Identification of the commodities	1					
		Species Manufacturing plant Num (Scientific name)	ber of packages Net weight Batch number					

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	coui	NTRY	Colostrum and colostrum b or parts thereof listed	Colostrum -Based Products C/CBP ased products from third countries in column A of Annex I for human nsumption intended for importation
	11.	Health information	II.a. Certificate reference number	II.b.
	11.1	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		002/99/EC and of Regulation (EC) No
cation		have been obtained or manufactured from colostrum obtained from	n animals:	
Part II: Certification		(i) under the control of the official veterinary service;		
Part 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 vaccin period; 		
		(iii) belonging to holdings which were not under restrictions due	e to foot-and-mouth disease or rinde	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 (EC) No 853/2004 and (EC) No 854/2004 and hereby certify th described in Part I were produced in accordance with those provis	at the colostrum/colostrum-based	
		(a) they were manufactured from colostrum:		
		 (i) 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 transpo Section IX of Annex III to Regulation (EC) No 853/2004; 	orted in accordance with the hygiene	e 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 dr requirements of point 4 in Part III of Chapter I of Sectir maximum residue limits for residues of antibacterial veteri 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 con- 		
		 (b) they come from an establishment implementing a programme 852/2004; 	based on the HACCP principles in	accordance with Regulation (EC) No
		(c) they have been processed, stored, wrapped, packaged and lat Regulation (EC) No 853/2004;	peled in accordance with Chapters I	II and IV of Section IX of Annex III to
		(d) they meet the relevant requirements laid down in Chapter II, 5 microbiological criteria laid down in Regulation (EC) No 2073/2		
		(e) the guarantees covering live animals and products thereof provid and in particular Article 29 thereof are fulfilled.	ded by the residue plans submitted ir	n accordance with Directive 96/23/EC,

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

COUNTRY	consumption intended for importation						
II. 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 I:							
— 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).							
- Box reference I.11: Name, address and approval number of the est	ablishment of dispatch.						
 Box reference I.15: Registration number (railway wagons or contail unloading and reloading, the consignor must inform the border inspectively) 							
 Box reference I.19: Use the appropriate Harmonised System (HS) co 04.10; 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;					
- Box reference I.20: Indicate total gross weight and total net weight.							
- Box reference I.23: For containers or boxes, the container number a	and the seal number (if applicable) she	buld be included.					
 Box reference I.28: Manufacturing plant: introduce the approval numl approved for exportation to the European Union. 	per of the production holding(s), collect	ion centre or standardization centre					
Part II:							
(¹) Keep as appropriate.							
- The colour of the signature shall be different to that of the printing. T	- 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.						
Official veterinarian	Official veterinarian						
Name (in capital letters):	Q	ualification and title:					
Date:	Si	gnature:					
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

COUNTRY

1.1. Consignor Name Address 1.2. Certificate reference No 1.2.a. Tel. 1.3. Central competent authority 1.4. Local competent authority 1.5. Consignee Name Address 1.4. Local competent authority 1.4. Local competent authority 1.5. Consignee Name Address 1.6. Person responsible for the load in EU Name Address Name Address Postoode Tel. 1.7. Country of origin Name 1.8. Region of origin Address Code 1.9. Country of destination ISO code 1.0. 1.11. Place of origin Address Name Approval number 1.12. Place of destination Customs warehouse in Ship supplier Name Ship supplier 1.13. Place of loading 1.14. Date of departure 1.16. Entry BIP in EU 1.17. I.18. Description of commodity 1.19. Commodity code (HS code) 1.20. Quantity 1.23. Seal/Container No Chilled in Frozen in Internet in Chilled in Frozen in Internet in In	
Address 1.3. Central competent authority Tel. 1.4. Local competent authority 1.5. Consignee Name Name Address Postcode Postal code Tel. 1.6. Person responsible for the load in EU Name Address Postcode Tel. No 1.7. Country of origin ISO code 1.8. Region of origin Code 1.11. Place of origin I.8. Region of origin Name Approval number Address Postal code 1.11. Place of loading I.12. Place of destination Name Approval number Address Postal code I.13. Place of loading I.14. Date of departure I.15. Means of transport I.16. Entry BIP in EU Acorplane Ship Road vehicle Other I.18. Description of commodity I.17. I.18. Description of commodity I.17. I.18. Description of commodity I.19. Commodity code (HS code) I.18. Description of commodity I.19. Commodity code (HS code) I.18. Description of commodity I.20. Quantity	
Tel: I.4. Local competent authors 1.5. Consignee I.4. Local competent authors Name Address Postode Postal code Tel. I.6. Person responsible for the load in EU Name Address Postode Tel. No I.7. Country of origin ISO code I.7. Country of origin ISO code I.1.1. Place of origin Approval number Address Approval number Address Postal code I.11. Place of loading I.12. Place of destination Customs warehouse Ship supplier Name Approval number Address Postal code I.13. Place of loading I.14. Date of departure I.15. Means of transport I.16. Entry BIP in EU Acroplane Ship Road vehicle Other I.18. Description of commodity I.19. Commodity code (HS code) I.19. Commodity code (HS code) I.20. Quantity I.21. Temperature of product I.22. Number of packad Ambient Chilled Frozen	
I.4. Local competent authority I.5. Consignee Name Address Postode Tel. I.7. Country of origin ISO code I.8. Region of origin Code I.7. Country of origin ISO code I.1. Place of origin Name Address Postal code Tel. I.1. Place of origin Name Address Postal code I.1.1. Place of loading I.1.3. Place of loading I.1.4. Date of departure I.1.5. Means of transport Active so I.1.6. Entry BIP in EU Active so I.1.8. Description of commodity I.1.9. Commodity code (HS code) I.20. Quantity I.21. Temperature of product </th <td></td>	
I.4. Local competent authority I.5. Consignee Name Address Postoode Postal code Tel. Tel. No I.7. Country of origin I.1. Place of origin Name Address Postoode Tel. No I.7. Country of origin I.1. Place of origin Name Approval number Address Postal code I.1.1. Place of origin Name Approval number Address Postal code I.1.1. Place of loading I.1.1. Place of loading I.1.3. Place of loading I.1.4. Date of departure I.1.5. Means of transport Actroplane Ship Name Approval number Actoress Postal code I.1.1. Place of loading I.1.2. Name Actoress Postal code I.1.1. Place of loading I.1.2. Name Acor	
Address Name Approval number Address Postal code 1.13. Place of loading 1.14. Date of departure 1.13. Place of loading 1.14. Date of departure 1.15. Means of transport 1.16. Entry BIP in EU Aeroplane Ship Railway wagon Road vehicle Other 1.17. Identification 1.17. Documentary references 1.19. Commodity code (HS code) 1.18. Description of commodity 1.19. Commodity code (HS code) 1.20. Quantity 1.20. Quantity	
Address Name Approval number Address Postal code 1.13. Place of loading 1.14. Date of departure 1.13. Place of loading 1.14. Date of departure 1.15. Means of transport 1.16. Entry BIP in EU Aeroplane Ship Railway wagon Road vehicle Other 1.17. Identification 1.17. Documentary references 1.19. Commodity code (HS code) 1.18. Description of commodity 1.19. Commodity code (HS code) 1.20. Quantity 1.20. Quantity	
Address Name Approval number Address Postal code 1.13. Place of loading 1.14. Date of departure 1.15. Means of transport 1.16. Entry BIP in EU Aeroplane Ship Road vehicle Other Identification 1.17. Identification 1.18. Description of commodity 1.18. Description of commodity 1.19. Commodity code (HS code) 1.20. Quantity 1.20. Quantity 1.21. Temperature of product Chilled Frozen	
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Ambient Chilled Frozen	
	iges
I.23. Seal/Container No I.24. Type of packagin	
	ng
I.25. Commodities certified for:	
Human consumption	
I.26. For transit through EU to 3rd Country	
3rd country ISO code	
1.28. Identification of the commodities	
Species Manufacturing plant Number of packages Net weight Batch num (Scientific name)	
	ber

С	OUNTRY		ts, colostrum and colostrum-base consumption for transit or storag
1		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 Par] / [colostrum-based products] (¹) (²) fo
	(a) come from a country or part thereof authorised for import based products as laid down in Annex I to Regulation (El		dairy products, colostrum or colostrum
	(b) comply with the relevant animal health conditions for II.1 of the model health certificates [Milk-RM] / [Milk-RN Regulation (EU) No 605/2010;		
	(c) was/were produced on and	(³) or between	(³)
٨	Notes		
F	Part I:		
_	 Box reference I.7: Provide name and ISO code of the cour No 605/2010 of 2 July 2010 laying down animal and public hea Union of raw milk and dairy products intended for human const 	alth and veterinary certification condition	
_	 Box reference I.11: Name, address and approval number of the same as the country of export. 	e establishment of dispatch. Name of t	he country of origin which must be the
	 Box reference I.15: Registration number (railway wagons or conta of transport in containers, the total number of containers and be indicated in box I.23. In case of unloading and reloading, European Union. 	I their registration number and where the	ere is a serial number of the seal it mus
_	 Box reference I.19: Use the appropriate Harmonised Syste 04.05; 04.06; 15.17; 17.02; 19.01; 21.05; 21.06.; 22.02; 2 		
_	- Box reference I.20: Indicate total gross weight and total net wei	ight.	
_	- Box reference I.23: For containers or boxes, the cont	ainer number and the seal numb	er (if applicable) should be included
-	 Box reference I.28: Manufacturing plant: introduce the standardization centre approved for exportation to the Eu 		tion holding(s), collection centre o
F	Part II:		
Ċ	 Raw milk, dairy products, colostrum and colostrum -based products for human consumption in transit or storage in accc December 1997 laying down the principles governing the orga countries (OJ L 24, 30.1.1998, p. 9). 	ordance 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 product obtained either prior to the date of authorisation for exporta I.7 and I.8, or during a period where restrictive measures h dairy products, colostrum and colostrum-based products fr	ation to the European Union of the third on the third on the been adopted by the European	country or part thereof mentioned unde
	 The colour of the signature shall be different to that of the printir 		

COUNTRY	Model Milk/Colostrum-T/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:					