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 (Text with EEA relevance)

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

(Text with EEA relevance)

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Council Directive 2002/99/EC of 16 December 2002 laying down the animal health rules governing the production, processing, distribution and introduction of products of animal origin for human consumption⁽¹⁾, and in particular the introductory phrase of Article 8, the first subparagraph of point (1) and point (4) of Article 8 and Article 9(4) thereof,

Having regard to Regulation (EC) No 852/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs⁽²⁾, and in particular Article 12 thereof,

Having regard to Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific hygiene rules for food of animal origin⁽³⁾, and in particular Article 9 thereof,

Having regard to Regulation (EC) No 854/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific rules for the organisation of official controls on products of animal origin intended for human consumption⁽⁴⁾, and in particular Articles 11(1) and 14 (4) and Article 16 thereof,

Having regard to Regulation (EC) No 882/2004 of the European Parliament and of the Council of 29 April 2004 on official controls performed to ensure the verification of compliance with feed and food law, animal health and animal welfare rules⁽⁵⁾, and in particular Article 48 (1) thereof,

Whereas:

(1) Council Directive 92/46/EEC of 16 June 1992 laying down the health rules for the production and placing on the market of raw milk, heat-treated milk and milk-based products⁽⁶⁾ provided for a list to be drawn up of third countries or parts thereof from which Member States were to authorise the introduction of milk or milk-based products and for such commodities to be accompanied by a health certificate and comply with certain requirements, including heat treatment requirements, and guarantees.

- (2) Accordingly, Commission Decision 2004/438/EC of 29 April 2004 laying down animal and public health and veterinary certifications conditions for introduction in the Community of heat-treated milk, milk-based products and raw milk intended for human consumption⁽⁷⁾ was adopted.
- (3) Since the date of adoption of that Decision, a number of new animal health and public health requirements have been laid down, constituting a new regulatory framework in this area, which should be taken into account in this Regulation. In addition, Directive 92/46/EEC was repealed by Directive 2004/41/EC of the European Parliament and of the Council of 21 April 2004 repealing certain Directive concerning food hygiene and health conditions for the production and placing on the market of certain products of animal origin intended for human consumption⁽⁸⁾.
- (4) Regulation (EC) No 178/2002 of the European Parliament and of the Council of 28 January 2002 laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety⁽⁹⁾ lays down the general principles governing food and feed in general, and food and feed safety in particular, at European Union and national level.
- (5) Directive 2002/99/EC lays down rules governing the introduction from third countries of products of animal origin intended for human consumption. It provides that such products are only to be introduced into the European Union if they comply with the requirements applicable to all stages of the production, processing and distribution of those products in the European Union or if they offer equivalent animal health guarantees.
- (6) Regulation (EC) No 852/2004 lays down the general rules for food business operators on the hygiene of foodstuffs at all stages of the food chain, including at primary production level.
- (7) Regulation (EC) No 853/2004 lays down specific rules for food business operators on the hygiene of food of animal origin. That Regulation provides that food business operators producing raw milk and dairy products intended for human consumption are to comply with the relevant provisions of Annex III thereto.
- (8) Regulation (EC) No 854/2004 lays down specific rules for the organisation of official controls on products of animal origin.
- (9) Commission Regulation (EC) No 2073/2005 of 15 November 2005 on microbiological criteria for foodstuffs⁽¹⁰⁾ lays down the microbiological criteria for certain microorganisms and the implementing rules to be complied with by food business operators when implementing the general and specific hygiene measures referred to in Article 4 of Regulation (EC) No 852/2004. Regulation (EC) No 2073/2005 provides that food business operators are to ensure that foodstuffs comply with the relevant microbiological criteria set out in that Regulation.
- (10) Under the scope of Council Directive 92/46/EEC, raw milk and products thereof could only be obtained from cows, ewes, goats or buffaloes. However, the definitions of raw milk and dairy products set out in Annex I to Regulation (EC) No 853/2004 broadens

the scope of milk hygiene rules to all mammalian species and defines raw milk as milk produced by the secretion of the mammary gland of farmed animals that has not been heated to more than 40 $^{\circ}$ C or undergone any treatment that has an equivalent effect. In addition, it defines dairy products as processed products resulting from the processing of raw milk or from further processing of such processed products.

- (11) In view of the entry into application of Regulations (EC) Nos 852/2004, 853/2004 and 854/2004 and the acts implementing those Regulations, it is necessary to amend and update European Union public and animal health conditions and certification requirements for the introduction into the European Union of raw milk and dairy products intended for human consumption.
- (12) In the interests of consistency of Union law, this Regulation should also take into account the rules laid down in Regulation (EC) No 470/2009 of the European Parliament and of the Council of 6 May 2009 laying down Community procedures for the establishment of residue limits of pharmacologically active substances in foodstuffs of animal origin, repealing Council Regulation (EEC) No 2377/90 and amending Directive 2001/82/EC of the European Parliament and of the Council and Regulation (EC) No 726/2004 of the European Parliament and of the Council⁽¹¹⁾ and its implementing rules laid down in Commission Regulation (EU) No 37/2010 of 22 December 2009 on pharmacologically active substances and their classification regarding maximum residue limits in foodstuffs of animal origin⁽¹²⁾ and Council Directive 96/23/EC of 29 April 1996 on measures to monitor certain substances and residues thereof in live animals and animal products and repealing Directives 85/358/EEC and 86/469/EEC and Decisions 89/187/EEC and 91/664/EEC⁽¹³⁾.
- (13) Council Directive 96/93/EC of 17 December 1996 on the certification of animals and animal products⁽¹⁴⁾ lays down the rules to be observed in issuing certificates required by veterinary legislation to prevent misleading or fraudulent certification. It is appropriate to ensure that certification requirements at least equivalent to those laid down in that Directive are applied by the competent authorities of exporting third countries.
- (14) In addition, Council Directive 90/425/EEC of 26 June 1990 concerning veterinary and zootechnical checks applicable in intra-Community trade in certain live animals and products with a view to the completion of the intenal market⁽¹⁵⁾, provides for a computerized system linking veterinary authorities which has been developed in the Europena Union. The format of all model health certificates need to be amended to take into account their compatibility with possible electronic certification under the Trade Control and Expert System (TRACES) provided for in Directive 90/425/EEC. According, the rules laid down in this Regulation should take account of TRACES.
- (15) Council Directive 97/78/EC of 18 December 1997 laying down the principles governing the organisation of veterinary checks on products entering the Community from third countries⁽¹⁶⁾ lays down rules concerning veterinary checks on products of animal origin introduced into the European Union from third countries for their importation or transit, including certain certification requirements. Those rules are applicable to the commodities covered by this Regulation.

- (16) Specific conditions for transit via the European Union of consignments to and from Russia should be provided for, owing to the geographical situation of Kaliningrad, which only concerns Latvia, Lithuania and Poland.
- (17) In the interests of clarity of European law, Commission Decision 2004/438/EC should be repealed and replaced by this Regulation.
- (18) To avoid any disruption in trade, the use of health certificates issued in accordance with Decision 2004/438/EC should be authorised during a transitional period.
- (19) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on the Food Chain and Animal Health,

HAS ADOPTED THIS REGULATION:

Article 1

Subject matter and scope

This Regulation lays down:

- (a) the public and animal health conditions and certification requirements for the introduction into the European Union of consignments of raw milk and dairy products;
- (b) the list of third countries from which the introduction into the European Union of such consignments is authorised.

Article 2

Imports of raw milk and dairy 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 and dairy products from the third countries or parts thereof listed in column A of Annex I.

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 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 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:

- 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 F₀ 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.

Article 6

Transit and storage conditions

The introduction into the European Union of consignments of raw milk and dairy products not intended for importation into the European Union but destined for a third country either by immediate transit or after storage in the European Union, in accordance with Articles 11, 12 or 13 of Council 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 authorised for the introduction into the European Union of consignments of raw milk or dairy products and comply with the appropriate heat 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 or dairy product concerned, as laid down in the animal health attestation in Part II.1 of the relevant model health certificate set out 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 Commission Regulation (EC) No 136/2004⁽¹⁷⁾, signed by the official veterinarian of the border inspection post of introduction into the European 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⁽¹⁸⁾, 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.

Article 8

Specific treatment

Consignments of dairy 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 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.

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 milkbased 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.

Done at Brussels, 2 July 2010.

For the Commission The President José Manuel BARROSO

ANNEX I

LIST OF THIRD COUNTRIES OR PARTS THEREOF AUTHORISED FOR THE INTRODUCTION INTO THE EUROPEAN UNION OF CONSIGNMENTS OF RAW MILK AND DAIRY PRODUCTS AND INDICATING THE TYPE OF HEAT TREATMENT REQUIRED FOR SUCH COMMODITIES

ISO code of third country	Third country or part thereof	Column A	Column B	Column C
AD	Andorra	+	+	+
AL	Albania	0	0	+
AN	Netherlands Antilles	0	0	+
AR	Argentina	0	0	+
AU	Australia	+	+	+
BR	Brazil	0	0	+
BW	Botswana	0	0	+
BY	Belarus	0	0	+
BZ	Belize	0	0	+
BA	Bosnia and Herzegovina	0	0	+
CA	Canada	+	+	+
СН	Switzerland ^a	+	+	+
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	+

in agricultural products (OJ L 114, 30.4.2002, p. 132).b 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.

c Not including Kosovo which is at present under international administration pursuant to United Nations Security Council Resolution 1244 of 10 June 1999.

·+' '0'	: third country	is authorised		
		is not authorised	0].
HK	Hong Kong	0	0	+
HN	Honduras	0	0	+
HR	Croatia	0	+	+
IL	Israel	0	0	+
IN	India	0	0	+
IS	Iceland	+	+	+
KE	Kenya	0	0	+
MA	Morocco	0	0	+
MG	Madagascar	0	0	+
МК ^ь	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	+
РҮ	Paraguay	0	0	+
RS ^c	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	+
	accordance with the Agreemer products (OJ L 114, 30.4.2002		Community and the Swiss	Confederation on trade
b The former Yu				

c Not including Kosovo which is at present under international administration pursuant to United Nations Security Council Resolution 1244 of 10 June 1999.

`+` `0`	third country is authorisedthird country is not authorised			
US	United States	+	+	+
UY	Uruguay	0	0	+
ZA	South Africa	0	0	+
ZW Zimbabwe 0 0		0	+	
	cordance with the Agreem oducts (OJ L 114, 30.4.20		ropean Community and	the Swiss Confederation on trade
The former Yugoslav Republic of Macedonia: the definitive nomenclature for this country will be agreed following the				

b 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.

c Not including Kosovo which is at present under international administration pursuant to United Nations Security Council Resolution 1244 of 10 June 1999.

ANNEX II

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.
'Milk-T/S'	Animal health certificate for raw milk or dairy products for human consumption, for transit/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 or dairy 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 both 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 column 2 of 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 or dairy products meet the health conditions laid down in Section IX, 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 Directive 96/93/EC⁽¹⁹⁾ 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

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

cou	NTR	Y	Veterinary certificate to EU
	l.1.	Consignor Name Address	I.2. Certificate reference number I.2.a. I.3. Central Competent Authority
lent		Tel. N°	I.4. Local Competent Authority
consignm	I.5. Consignee I Name		1.6.
of dispatched consignment		Address Postal code Tel. Nº	
tails of di	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of destination ISO code I.10.
De	1.11.	Place of origin	1.12.
Part I: Details		Name Approval number Address	
	l.13.	Place of loading	I.14. Date of departure
	l.15.	. Means of transport Aeroplane Ship Railway wagon Road vehicle Other	I.16. Entry BIP in EU
		tification: umentary references:	l.17.
	I.18.	Description of commodity	I.19. Commodity code (HS code)
			I.20. Quantity
		Temperature of product Ambient Chilled	I.22. Number of packages
	1.23.	Identification of container/Seal number	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	
		Species Manufacturing plant Numi (Scientific name)	ber of packages Net weight Batch number

οι	INTRY	1	Model Milk-RM Raw milk
	Health information	II.a. Certificate reference number	II.b.
1	Animal Health Attestation		
	I, the undersigned official veterinarian, declare that I am awar (EC) No 853/2004 and hereby certify that the raw milk desc		
	(a) under the control of the official veterinary service,		
	(b) which were in a country or part thereof that has been free of foot-and-mouth disease and of rinderpest for a period of at least months prior to the date of this certificate, and where vaccination against foot-and-mouth disease has not been carried out duri that period,		
	(c) belonging to holdings which were not under restrictions of	due to foot-and-mouth disease or rine	derpest, and
	(d) subject to regular veterinary inspections to ensure that the IX of Annex III to Regulation (EC) No 853/2004 and in D		s laid down in Chapter I of Section
.2	Public Health attestation		
	I, the undersigned official inspector, declare that I am awa No 852/2004, (EC) No 853/2004 and (EC) No 854/2004 ar accordance with those provisions, in particular that:		
 (a) it comes from holdings registered in accordance with Regulation (EC) No 852/2004 and checked in accordance with Anne Regulation (EC) No 854/2004, 			ed in accordance with Annex IV to
	(b) it was produced, collected, cooled, stored and transported in accordance with the hygiene conditions laid down in Chapter I o Section IX of Annex III to Regulation (EC) No 853/2004,		
	(c) it meets the plate and somatic cell count criteria laid down	n in Chapter I of Section IX of Annex I	II to Regulation (EC) No 853/2004,
	(d) it does not contain antibiotic residues exceeding the limit	ts authorised under the Annex to Re	gulation (EU) No 37/2010,
	(e) the guarantees covering live animals and products thereo 96/23/EC, and in particular Article 29 thereof, are fulfilled		nitted in accordance with Directive
	(f) it does not contain pesticide residues exceeding the limit	ts authorized by Regulation (EC) No	396/2005, and
	(g) it does not contain contaminants exceeding the maximum	n tolerances laid down by Regulatior	n (EC) No 1881/2006.
10	tes		
	s certificate is intended for raw milk from third countries or 605/2010 intended for further processing in the European Uni		
Pa	t I:		
	Box reference I.7: Provide name and ISO code of the co No 605/2010.	ountry or part thereof as appearing	in Annex I to Regulation (EU)
_	Box reference I.11: Name, address and approval number of the	he establishment of dispatch.	
	Box reference I.15: Registration number (railway wagons or cor of unloading and reloading, the consignor must inform the bor		
_	Box reference I.19: Use the appropriate Harmonised System	(HS) code of the World Customs Or	ganisation: 04.01; 04.02 or 04.03.

Part II: Certification

Model	Milk-RM

COUNTRY		Raw milk
II. Health information	II.a. Certificate reference number	II.b.
- Box reference I.20: Indicate total gross weight and total net w	veight.	
- Box reference I.23: For containers or boxes, the container nu	mber and the seal number (if applica	able) should be included.
 Box reference I.28: Manufacturing plant: introduce the a standardization centre approved for exportation to the Europe 		holding(s), collection centre or
Part II:		
 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		
Name (in capital letters):	C	Qualification and title:
Date:	s	Signature:
Stamp:		

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

cou	NTR	(Veterinary certificate to EU
	l.1.	Consignor	I.2. Certificate reference number I.2.a.
		Name	I.3. Central Competent Authority
		Address	
ent		Tel. Nº	I.4. Local Competent Authority
gnm	1.5.	Consignee	1.6.
onsi		Name	
o p		Address	
- Š		Postal code	
spat		Tel. Nº	
Part I: Details of dispatched consignment	1.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country of destination ISO code I.10.
Detai	1.11.	Place of origin	1.12.
art		Name Approval number	
۵.		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 D Other	l.17.
		tification:	
		umentary references: Description of commodity	I.19. Commodity code (HS code)
		Description of commonly	
			I.20. Quantity
	121	Temperature of product	I.22. Number of packages
		Ambient Chilled	Frozen
	1.23.	Identification of container/Seal number	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 Nu (Scientific name)	mber of packages Net weight Batch number

	cou	NTRY	Dairy products derived from raw	<i>Model Milk-RMP</i> milk for human consumption			
	II.	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 the relevant provisions of Directive 2003 (EC) No 853/2004 and hereby certify that the dairy products described above has been manufactured fro animals:							
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 at least 12 months prior to the date of this certificate, and where vaccination against foot-and-mouth disease has not been carried out during that period,					
Part		(c) belonging to holdings which were not under restrictions du	e to foot-and-mouth disease or rinde	erpest, and,			
		(d) subject to regular veterinary inspections to ensure that they IX of Annex III to Regulation (EC) No 853/2004 and in Direction (EC) No 853/		aid down in Chapter I of Section			
	11.2	Public Health attestation					
	I, the undersigned official inspector, declare that I am aware of the relevant provisions of Regulations (EC) No 178/200 No 852/2004, (EC) No 853/2004 and (EC) No 854/2004 and hereby certify that the dairy product made with raw milk describe was produced in accordance with those provisions, in particular that:						
(a) it was manufactured from raw milk:							
		 (i) which comes from holdings registered in accordance Annex IV to Regulation (EC) No 854/2004, 	with Regulation (EC) No 852/2004 a	and checked in accordance with			
		(ii) which was produced, collected, cooled, stored and tr Chapter I of Section IX of Annex III to Regulation (EC)		nygiene conditions laid down in			
		(iii) which meets the plate and somatic cell count criteria No 853/2004,	laid down in Chapter I of Section IX	of Annex III to Regulation (EC)			
		(iv) which does not contain antibiotic residues exceeding the	ne limits authorised under the Annex	to Regulation (EU) No 37/2010,			
		(v) which does not contain pesticide residues exceeding the	he limits authorized by Regulation (E	C) No 396/2005, and			
		(vi) which does not contain contaminants exceeding the	maximum tolerances laid down by	Regulation (EC) No 1881/2006.			
		 (b) it comes from an establishment implementing a programme No 852/2004, 	based on the HACCP principles in a	accordance with Regulation (EC)			
		(c) it has been obtained from raw milk that has not undergone manufacturing process,	any heat treatment or any physical	or chemical treatment during the			
		 (d) it has been wrapped, packaged and labelled in accordance No 853/2004, 	with Chapters III and IV of Section I	X of Annex III to Regulation (EC)			
		(e) it meets the relevant microbiological criteria laid down in Re and	egulation (EC) No 2073/2005 on micr	obiological criteria for foodstuffs,			
		(f) the guarantees covering live animals and products thereof 96/23/EC, and in particular Article 29 thereof, are fulfilled.	provided by the residue plans submit	tted in accordance with Directive			

COUNTRY	Dairy products derived from raw	<i>Model Milk-RMP</i> milk for human consumption		
II. Health information	II.a. Certificate reference number	II.b.		
Notes				
This certificate is intended for dairy products derived from raw milk for in column A of Annex I to Regulation (EU) No 605/2010 intended f				
Part I:				
- Box reference I.7: Provide name and ISO code of the country of	r part thereof as appearing Annex I	to Regulation (EU) No 605/2010.		
- Box reference I.11: Name, address and approval number of the	establishment of dispatch.			
case of transport in containers, the total number of containers an	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 (04.04; 04.05; 04.06 or 21.05. 	HS) code of the World Customs O	rganisation: 04.01; 04.02; 04.03;		
- Box reference I.20: Indicate total gross weight and total net wei	ght.			
- Box reference I.23: For containers or boxes, the container number	per and the seal number (if applicat	ble) should be included.		
 Box reference I.28: Manufacturing plant: introduce the app standardization centre approved for exportation to the European 		holding(s), collection centre or		
Part II:				
 The colour of the signature shall be different to that of the printi watermark. 	ing. The same rule applies to stamp	os other than those embossed or		
Official veterinarian	Official veterinarian			
Name (in capital letters): Qualification and title:				
Date:	Si	gnature:		
Stamp:				

ModelHealth Certificate for dairy products derived from milk of cows, ewes, goatsMilk-and buffaloes for human consumption from third countries or parts thereofHTBauthorised in column B of Annex I to Regulation (EU) No 605/2010 intended forimportation into the European Union

cou	NTRY	1	Veterinary certificate to E	
	l.1.	Consignor Name	I.2. Certificate reference number I.2.a.	
		Address	I.3. Central Competent Authority	
ŧ		Tel. Nº	I.4. Local Competent Authority	
a a	-			
of dispatched consignment	1.5.	Consignee Name	1.6.	
8				
hed		Address		
atc		Postal code		
isp		Tel. Nº		
ils of d	1.7.	Country of origin ISO code	I.9. Country of destination ISO code I.10.	
Part I: Details	1.11.	Place of origin	1.12.	
Ţ		Name Approval number		
å		Address		
		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		
	Iden	tification:	l.17.	
	Doc	umentary references:		
	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.	Identification of container/Seal number	I.24. Type of packaging	
	1.25.	Commodities certified for:	1	
	Human consumption			
	1.26.		I.27. For import or admission into EU	
	1.28.	Identification of the commodities	1	
		Species Manufacturing plant Nu (Scientific name)	mber of packages Net weight Batch number	

				Mode	l Milk-HTB
Dairy products	derived	from milk	of cows,	ewes,	goats and
buffaloes for	human	consumpt	ion from	third	countries
authorised in co	olumn B				

	COUNTRY		authorised in column B			
	II.	Health information	II.a. Certificate reference number	II.b.		
	II.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 Regulati (EC) No 853/2004 and hereby certify that the dairy product described above:				
E		(a) has been obtained from animals:				
(i) under the control of the official veterinary service,						
 (i) under the control of the official veterinary service, (ii) which were in a country or part thereof that has been free of foot-and-mouth disease and of rinderpest for a period 12 months prior to the date of this certificate, and where vaccination against foot-and-mouth disease has not been during that period, 						
		(iii) belonging to holdings which were not under restrictions	due to foot-and-mouth disease or	rinderpest, and,		
		(iv) subject to regular veterinary inspections to ensure that Section IX of Annex III to Regulation (EC) No 853/2004		nditions laid down in Chapter I of		
(b) has undergone or been produced from raw milk which has been submitted to a pasteurisation treatment involving a si treatment with a heating effect at least equivalent to that achieved by a pasteurisation process of at least 72 °C for at 15 and where applicable, sufficient to ensure a negative reaction to an alkaline phosphatase test applied immediately after treatment.			of at least 72 °C for at 15 seconds			
II.2 Public Health attestation						
I, the undersigned official veterinarian, declare that I am aware of the No 852/2004, (EC) No 853/2004 and (EC) No 854/2004 and hereby accordance with those provisions, in particular that:						
(a) it was manufactured from raw milk:						
 (i) which comes from holdings registered in accordance with Regulation (EC) N Annex IV to Regulation (EC) No 854/2004, 			vith Regulation (EC) No 852/2004	and checked in accordance with		
	 (ii) which was produced, collected, cooled, stored and t Chapter I of Section IX of Annex III to Regulation (EC) 			hygiene conditions laid down in		
		(iii) which meets the plate and somatic cell count criteria la No 853/2004,	aid down in Chapter I of Section I	X of Annex III to Regulation (EC)		
		(iv) which does not contain antibiotic residues exceeding th	e limits authorised under the Anne	x to Regulation (EU) No 37/2010;		
	(v) which does not contain pesticide residues exceeding the		e limits authorized by Regulation ((EC) No 396/2005, and		
	(vi) which does not contain contaminants exceeding the		naximum tolerances laid down by	Regulation (EC) No 1881/2006.		
	 (b) it comes from an establishment implementing a programm No 852/2004, 		based on the HACCP principles in	accordance with Regulation (EC)		
		(c) it has been processed, stored, wrapped, packaged and trans in Annex II to Regulation (EC) No 852/2004 and Chapter II				
		(d) it meets the relevant criteria laid down in Chapter II of Sect microbiological criteria laid down in Regulation (EC) No 207	ion IX of Annex III to Regulation (E 73/2005 on microbiological criteria	EC) No 853/2004 and the relevant for foodstuffs,		
		(e) the guarantees covering live animals and products thereof p 96/23/EC, and in particular Article 29 thereof, are fulfilled.	provided by the residue plans subm	nitted in accordance with Directive		

Model Milk-HTB Dairy products derived from milk of cows, ewes, goats and buffaloes for human consumption from third countries authorised in column B

COUNTRY	authorised in column B				
II. Health information	II.a. Certificate reference number	II.b.			
lotes					
This certificate is intended for dairy products for human consumption from third countries or parts thereof authorised in column B of Annex of Regulation (EU) No 605/2010 intended for importation into the European Union.					
Part I:					
 Box reference I.7: Provide name and ISO code of the cour No 605/2010. 	ntry or part thereof as appearing	in Annex I to Regulation (EU)			
- Box reference I.11: Name, address and approval number of est	ablishment of dispatch.				
In the case of transport in containers, the total number of co	Box reference I.15: Registration number (railway wagons or container and road vehicle), flight number (aircraft) or name (ship). In the case of transport in containers, the total number of containers and their registration number and where there is a serial number of the seal it must be indicated in box I.23. In case of unloading and reloading, the consignor must inform the border inspection post of introduction into the European Union.				
 Box reference I.19: Use the appropriate Harmonised System (04.04; 04.05; 04.06 or 21.05. 	 Box reference I.19: Use the appropriate Harmonised System (HS) code of the World Customs Organisation: 04.01; 04.02; 04.03; 04.04; 04.05; 04.06 or 21.05. 				
- Box reference I.20: Indicate total gross weight and total net weight	ght				
- Box reference I.23: For containers or boxes, the container number	per and the seal number (if applica	ble) should be included.			
 Box reference I.28: Manufacturing plant: introduce the approval r for export to the European Union. 	 Box reference I.28: Manufacturing plant: introduce the approval number of the treatment and/or processing establishment(s) approved for export to the European Union. 				
Part II:					
 The colour of the signature shall be different to that of the printi watermark. 	- 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):	a	ualification and title:			
Date:	S	ignature:			
Stamp:					

ModelHealth Certificate for dairy products for human consumption from thirdMilk-countries or parts thereof authorised in column C of Annex I to Regulation (EU)HTCNo 605/2010 intended for importation into the European Union

I.1. Consignor I.2. Certificate reference number Name I.3. Central Competent Authority I.3. Consignee I.4. Local Competent Authority I.5. Consignee I.6. Name Address Postal code I.6. I.7. Country of origin ISO I.7. Country of origin I.8. Region of origin Code I.1. Place of origin I.1.2. Name Approval number I.12.	Veterinary certificate to EU		
Address	l.2.a.		
14 Local Compotent Authority	I.3. Central Competent Authority		
Tel. N° I.5. Consignee I.6. Name I.6. Address Postal code Tel. N° I.6. I.7. Country of origin ISO I.7. Country of origin ISO I.8. Region of origin Code I.9. Country of destination ISO I.9. Country of destination ISO I.1. I.1.			
I.o. Consigned I.o. Name Name Address Postal code Tel. N° I.7. Country of origin ISO I.8. Region of origin Code I.9. Country of destination ISO ISO Tel.			
So Address Postal code Tel. N° 1.7. Country of origin ISO code I.8. Region of origin Code I.9. Country of destination ISO code ISO			
Postal code Tel. N° I.7. Country of origin ISO code I.8. Region of origin Code I.9. Country of destination ISO code I.9. Country of destination ISO			
Tel. N° I.7. Country of origin ISO code I.7. Country of origin I.8. Region of origin Code			
iiiiiiiiiiiiiiiiiiiiiiiiiiiiiiiiiiii			
I.11. Place of origin I.12.			
T Name Approval number			
Address			
I.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 I			
Identification: Documentary references:			
I.18. Description of commodity I.19. Commodity cod	le (HS code)		
	I.20. Quantity		
	I.22. Number of packages		
Ambient Chilled Frozen			
	I.24. Type of packaging		
I.25. Commodities certified for:			
Human consumption			
I.26. I.27. For import or admission into B			
I.28. Identification of the commodities			
Species Manufacturing plant Number of packages Net weig (Scientific name)	ht Batch number		

Model M	liik-HTC
---------	----------

	co	UNTF	RY			Dairy products from third cou	untries authorised in column C
	II.		Hea	lth ir	nformation	II.a. Certificate reference number	II.b.
	II.1	An	imal	Hea	alth Attestation		
					signed official veterinarian, declare that I am aware 3/2004 and hereby certify that the dairy product dea		e 2002/99/EC and of Regulation
		(a)	has	bee	en obtained from animals:		
ation			(i)	uno	der the control of the official veterinary service,		
ertific	(ii) belonging to holdings which were not under restrictions due to foot-and-mouth disease or rinderpest, and,						rinderpest, and,
Part II: Certification			(iii)		pject to regular veterinary inspections to ensure that ction IX of Annex III to Regulation (EC) No 853/200		ditions laid down in Chapter I of
8	(¹)	eithe	<i>r</i> [(b		the case of dairy products made from raw milk sou port into the territory of the European Union:	rced from cows, ewes, goats or bu	ffaloes have undergone, prior to
		(¹) ei	ther	[(i)	a sterilisation process, to achieve an F_0 value equa	al to or greater than three;]	
		(¹) 01		[(ii)	an ultra high temperature (UHT) treatment at not le	ess than 135 °C in combination with	a suitable holding time;]
(1) or [(iii) a high temperature short time pasteurisation treatment (HTST) at 72 °C for 15 equal to or greater than 7.0 achieving, where applicable, a negative reaction immediately after the heat treatment;]				[(iii)	equal to or greater than 7.0 achieving, where app		
(1) or [(iv) a treatment with an equivalent pasteurisation effe alkaline phosphatase test, applied immediately after			[(iv)	a treatment with an equivalent pasteurisation effec alkaline phosphatase test, applied immediately after		licable, a negative reaction to a	
	 (¹) or [(v) a HTST treatment with a pH below 7.0;] (¹) or [(vi) a HTST treatment combined with another physical (¹) either [(vi) (1) lowering the pH below 6 for one hour;] 			[(v)	a HTST treatment with a pH below 7.0;]		
				[(vi)	a HTST treatment combined with another physical	treatment by	
				[(vi)	(1) lowering the pH below 6 for one hour;]		
		(¹) 01		[(vi)	(2) additional heating equal to or greater than 72 $^{\circ \rm c}$	C or more, combined with desiccati	ion;]]
 (¹) or [(b) in the case of dairy products made from raw milk sourced from animals other than cows, ewes, goats or build undergone, prior to import into the territory of the European Union: (¹) either [(i) a sterilisation process, to achieve an F₀ value equal to or greater than three;] 				s, ewes, goats or buffaloes have			
				[(i)	a sterilisation process, to achieve an F_0 value equa	al to or greater than three;]	
	(¹) or [(ii) an ultra high temperature (UHT) treatment at not less than 135 °C in combination with a suitable holding time;]				a suitable holding time;]]		
	II.2 Public Health attestation						
 I, the undersigned official inspector, declare that I am aware of the relevant provisions of Regulations (EC) No 178/2002, No 852/2004, (EC) No 853/2004 and (EC) No 854/2004 and hereby certify that the dairy product described above was product accordance with provisions, in particular that: (a) it was manufactured from raw milk: 							
			(i)		ich comes from holdings registered in accordance v nex IV to Regulation (EC) No 854/2004;	with Regulation (EC) No 852/2004 a	and checked in accordance with
			(ii)		ich was produced, collected, cooled, stored and tr apter I of Section IX of Annex III to Regulation (EC)		hygiene conditions laid down in

Model Milk-H OUNTRY Dairy products from third countries authorised in column					
II. Health information	II.a. Certificate reference number	II.b.			
(iii) which meets the plate and somatic cell count crite No 853/2004;	ria laid down in Chapter I of Section I	IX of Annex III to Regulation (EC)			
(iv) which does not contain antibiotic residues exc No 37/2010;	eeding the limits authorised under	the Annex to Regulation (EU)			
(v) which does not contain pesticide residues exceeding the limits authorized by Regulation (EC) No 396/2005, and					
(vi) which does not contain contaminants exceeding t	ne maximum tolerances laid down by	Regulation (EC) No 1881/2006.			
 (b) it comes from an establishment implementing a programme No 852/2004, 	based on the HACCP principles in	accordance with Regulation (EC)			
(c) it has been processed, stored, wrapped, packaged and train in Annex II to Regulation (EC) No 852/2004 and Chapter					
(d) it meets the relevant criteria laid down in Chapter II of Sec microbiological criteria laid down in Regulation (EC) No 20					
(e) the guarantees covering live animals and products thereof 96/23/EC, and in particular Article 29 thereof, are fulfilled.	provided by the residue plans submi	tted in accordance with Directive			
Notes					
This certificate is intended for dairy products for human consumption I to Regulation (EU) No 605/2010 intended for importation into the		authorised in column C of Annex			
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 establishments of dispatch.					
— Box reference I.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 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 (04.04; 04.05; 04.06; 19.01; 21.05; 21.06.90; 35.01 or 35.02. 	HS) code of the World Customs Or	rganisation: 04.01; 04.02; 04.03;			
- Box reference I.20: Indicate total gross weight and total net weight.					
- Box reference I.23: For containers or boxes, the container number and the seal number (if applicable) should be included.					
 Box reference I.28: Manufacturing plant: introduce the approval for export to the European Union. 	Box reference I.28: Manufacturing plant: introduce the approval number of the treatment and/or processing establishment(s) approved for export to the European Union.				
Part II:					
(1) Keep as appropriate.					

— 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.

COUNTRY	Model Milk-HTC Dairy products from third countries authorised in column C
II. Health information	II.a. Certificate reference number II.b.
Official veterinarian	
Name (in capital letters):	Qualification and title:
Date:	Signature:
Stamp:	
Stamp:	

PART 3

Model Animal Health Certificate for raw milk or dairy products for human *Milk-T/* consumption, for [transit]/[storage] $\binom{1}{2}$ in the European Union S

l.1.	Consignor Name		I.2. Certificate reference number	
	Nomo		1.2. Certificate reference number	I.2.a.
	Name			
	Address		I.3. Central Competent Authority	
			I.4. Local Competent Authority	
	Tel. Nº			
1.5 .	Consignee		I.6. Person responsible for the loa	d in EU
lign	Name		Name	
Suc	Address		Address	
2	Postal code		Postal code	
tche	Tel. Nº		Tel. Nº	
of dispatched consignment	Country of origin ISO	I.8. Region of origin Co	le I.9. Country of destination ISO	1.10.
di di	code		code	
s				
in 1.11.	. Place of origin		I.12. Place of origin	
Part I: Details	Name	Approval number	Customs warehouse	Ship supplier 🗌
art	Address		Name	Approval number
–			Address	Approval number
			, lau eee	
			Postal code	
I.13.	. Place of loading		I.14. Date of departure	
1.15.	. Means of transport		I.16. Entry BIP in EU	
	Aeroplane 🗌 Ship	_ , .		
	Road vehicle	Other	1.17.	
	ntification:			
	cumentary references:			
1.18.	. Description of commodity		I.19. Commodity code	e (HS code)
			L I.	20. Quantity
1.21.	. Temperature of product	_		22. Number of packages
	Ambient	Chilled	Frozen 🗌	
1.23.	. Identification of container/Sea	al number	1.	24. Type of packaging
1.25	. Commodities certified for:			
	н	uman consumption 🗌		
1.00	. For transit through EU to 3rd		□ 1.27.	
1.20.	-	ISO code		
	3rd country			
1.28.	. Identification of the commodi	ties		
	Species (Scientific name)	Manufacturing plant	Number of packages Net weigh	nt Batch number

	col	JNTRY	Raw milk or dairy products in for transit or storage	<i>Model Milk-T/S</i> tended for human consumption			
	11.	Health information	II.a. Certificate reference number	II.b.			
	11.1	Animal Health Attestation					
		I, the undersigned official veterinarian, hereby certify that the European Union described above:	[raw milk] / [dairy products] (1) (2)	for [transit] / [storage] $(^{2})$ in the			
ation		 (a) come from a country or part thereof authorised for imports to the European Union of raw milk or dairy products as laid down in Annex I to Regulation (EU) No 605/2010, 					
Part II: Certification		(b) comply with the relevant animal health conditions for the pro- II.1 of the model certificates [Milk-RM] / [Milk-RMP] / [M No 605/2010;					
Part		(c) was produced on	or between				
	No	tes					
	Pa	rt 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 esta the same as the country of export.	ablishments of dispatch. Name of th	ne country of origin which must be			
	— Box reference I.15: Registration number (railway wagons or container and road vehicles), flight number (aircraft) or name (ship). In case of transport in containers, the total number of containers and their registration number and where there is a serial number of seal it must be indicated in box I.23. In case of unloading and reloading, the consignor must inform the border inspection posintroduction into the European Union.			ere there is a serial number of the			
	-	Box reference I.19: Use the appropriate Harmonised System (I 04.04; 04.05; 04.06; 19.01; 21.05; 21.06.90; 35.01 or 35.02.	HS) code of the World Customs (Drganisation: 04.01; 04.02; 04.03;			
	-	Box reference I.20: Indicate total gross weight and total net wei	ight.				
	-	Box reference I.23: For containers or boxes, the container num	ber and the seal number (if applica	able) should be included.			
 Box reference I.28: Manufacturing plant: introduce the approval number of the production holding(s), col standardization centre approved for exportation to the European Union. 				holding(s), collection centre or			
	Part II:						
	(1)	Raw milk and dairy products means, raw milk and dairy produ Article 12(4) or Article 13 of Council Directive 97/78/EC.	cts for human consumption in tran	sit or storage in accordance with			
	(²)	Keep as appropriate.					
	(3) Date or dates of production. Imports of raw milk and dairy products shall not be allowed when obtained either prior to the date authorisation for exportation to the European Union of the third country or part thereof mentioned under I.7 and I.8, or during a perior where restrictive measures have been adopted by the European Union against imports of raw milk and dairy products from this thi country or part thereof.						
	-	The colour of the signature shall be different to that of the printi watermark.	ing. The same rule applies to stam	ps other than those embossed or			

Model Milk-T/S Raw milk or dairy products intended for human consumption for transit or storage

COUNTRY	transit or storage	
II. Health information	II.a. Certificate reference number	II.b.
Official veterinarian		
Name (in capital letters):	C	Qualification and title:
Date:	٤	Signature:
Stamp:		

(**1**) OJ L 18, 23.1.2003, p. 11.

- (2) OJ L 139, 30.4.2004, p. 1.
- (**3**) OJ L 139, 30.4.2004, p. 55.
- (4) OJ L 139, 30.4.2004, p. 206.
- (5) OJ L 165, 30.4.2004, p. 206.
- (6) OJ L 268, 14.9.1992, p. 1.
- (7) OJ L 154, 30.4.2004, p. 72.
- (8) OJ L 157, 30.4.2004, p. 33.
- **(9)** OJ L 31, 1.2.2002, p. 1.
- (**10**) OJ L 338, 22.12.2005, p. 1.
- (11) OJ L 152, 16.6.2009, p. 11.
- (12) OJ L 15, 20.1.2010, p. 1.
- (13) OJ L 125, 23.5.1996, p. 10.
- (14) OJ L 13, 16.1.1997, p. 28.
- (15) OJ L 224, 18.8.1990, p. 29.
- (16) OJ L 24, 30.1.1998, p. 9.
- (17) OJ L 21, 28.1.2004, p. 11.
- (18) OJ L 296, 12.11.2009, p. 1
- (19) OJ L 13, 16.1.1997, p. 28.