►B ►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 ◄

(Text with EEA relevance)

(OJ L 175, 10.7.2010, p. 1)

Amended by:

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		No	page	date
►M1	Commission Implementing Regulation (EU) No 914/2011 of 13 September 2011	L 237	1	14.9.2011
►M2	Commission Implementing Regulation (EU) No 957/2012 of 17 October 2012	L 287	5	18.10.2012
►M3	Commission Implementing Regulation (EU) No 300/2013 of 27 March 2013	L 90	71	28.3.2013
►M4	Commission Regulation (EU) No 519/2013 of 21 February 2013	L 158	74	10.6.2013
►M5	Commission Implementing Regulation (EU) No 556/2013 of 14 June 2013	L 164	13	18.6.2013
►M6	Commission Implementing Regulation (EU) No 209/2014 of 5 March 2014	L 66	11	6.3.2014

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

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COMMISSION REGULATION (EU) No 605/2010

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(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.

([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.

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(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 ([1]) 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 ([2]).

(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 ([3]) 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.

([1]) OJ L 154, 30.4.2004, p. 72.
([2]) OJ L 157, 30.4.2004, p. 33.
([3]) OJ L 31, 1.2.2002, p. 1.

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(9) Commission Regulation (EC) No 2073/2005 of 15 November 2005 on microbiological criteria for foodstuffs ([1]) lays down the microbiological criteria for certain micro-organisms 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 °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 ([2]) 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 ([3]) 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 ([4]).

([1]) OJ L 338, 22.12.2005, p. 1.
([2]) OJ L 152, 16.6.2009, p. 11.
([3]) OJ L 15, 20.1.2010, p. 1.
([4]) OJ L 125, 23.5.1996, p. 10.

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(13) Council Directive 96/93/EC of 17 December 1996 on the certification of animals and animal products [1] 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 [2], 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 [3] 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:

[1] OJ L 13, 16.1.1997, p. 28.
[2] OJ L 224, 18.8.1990, p. 29.
[3] OJ L 24, 30.1.1998, p. 9.

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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;

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

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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.

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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.

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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.

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Article 4

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

1. ►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: ◄

(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.

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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.

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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.

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Article 7

Derogation concerning transit and storage conditions

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

(a) the consignment is sealed with a serially numbered seal at the border inspection post of introduction into the European Union by the veterinary services of the competent authority;

(b) the documents accompanying the consignment and referred to in Article 7 of Directive 97/78/EC are stamped 'ONLY FOR TRANSIT TO RUSSIA VIA THE EU' on each page by the official veterinarian of the competent authority responsible for the border inspection post of introduction into the European Union;

(c) the procedural requirements provided for in Article 11 of Directive 97/78/EC are complied with;

(d) the consignment is certified as acceptable for transit on the common veterinary entry document by the official veterinarian of the border inspection post of introduction into the European Union.

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

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

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Article 7a

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

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

(a) the consignment is sealed with a serially numbered seal by the official veterinarian at the border inspection post of entry;

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

([1]) OJ L 296, 12.11.2009, p. 1

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(c) the procedural requirements provided for in Article 11 of Directive 97/78/EC are complied with;

(d) the consignment is certified as acceptable for transit on the Common Veterinary Entry Document referred to in Article 2(1) of Regulation (EC) No 136/2004 by the official veterinarian at the border inspection post of entry.

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

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

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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.

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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
AE	The Emirate of Dubai of the United Arab Emirates (1)	0	0	+ (2)
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	+
BA	Bosnia and Herzegovina	0	0	+
CA	Canada	+	+	+
CH	Switzerland (**)	+	+	+
CL	Chile	0	+	+
CN	China	0	0	+
CO	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	+
HK	Hong Kong	0	0	+
HN	Honduras	0	0	+
IL	Israel	0	0	+
IN	India	0	0	+

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ISO code of third country	Third country or part thereof	Column A	Column B	Column C
IS	Iceland	+	+	+
KE	Kenya	0	0	+
MA	Morocco	0	0	+
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.

(**) 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.

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

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

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ANNEX II

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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.

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(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.

([1]) OJ L 13, 16.1.1997, p. 28.

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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

COUNTRY: **Veterinary certificate to EU**

Part I: Details of dispatched consignment

I.1. Consignor Name Address Tel.		I.2. Certificate reference No	I.2.a.
		I.3. Central competent authority	
		I.4. Local competent authority	
I.5. Consignee Name Address Postcode Tel.		I.6.	
I.7. Country of origin ISO code	I.8. Region of origin Code	I.9. Country of destination ISO code	I.10.
I.11. Place of origin Name Approval number Address		I.12.	
I.13. Place of loading		I.14. Date of departure	
I.15. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification Documentation references		I.16. Entry BIP in EU	
		I.17.	

I.18. Description of commodity	I.19. Commodity code (HS code)	
		I.20. Quantity
I.21. Temperature of product Ambient ☐ Chilled ☐ Frozen ☐		I.22. Number of packages
I.23. Seal/Container No		I.24. Type of packaging
I.25. Commodities certified for: Further process ☐		
I.26.	I.27. For import or admission into EU ☐	

I.28. Identification of the commodities

Manufacturing plant	Number of packages	Species (Scientific name)	Net weight	Batch number

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Model Milk-RM
Raw milk

COUNTRY

II. Health information	II.a. Certificate reference number	II.b.

Part II: Certification

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 Regulation (EC) No 853/2004 and hereby certify that the raw milk described above has been obtained from animals:

(a) under the control of the official veterinary service,

(b) which were in a country or part thereof that has been free of foot-and-mouth disease and of rinderpest for a period of at least 12 months prior to the date of this certificate, and where vaccination against foot-and-mouth disease has not been carried out during that period,

(c) belonging to holdings which were not under restrictions due to foot-and-mouth disease or rinderpest, and

(d) subject to regular veterinary inspections to ensure that they satisfy the animal health conditions laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004 and in Directive 2002/99/EC;

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, (EC) No 852/2004, (EC) No 853/2004 and (EC) No 854/2004 and hereby certify that the raw milk described above was produced in 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 Annex IV to Regulation (EC) No 854/2004,

(b) it was produced, collected, cooled, stored and transported in accordance with the hygiene conditions laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004,

(c) it meets the plate and somatic cell count criteria laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004,

(d) the guarantees on the residues status of raw milk provided by the monitoring plans for the detection of residues or substances submitted in accordance with Council Directive 96/23/EC, and in particular, Article 29 thereof, are fulfilled;

(e) pursuant to testing for residues of antibacterial drugs carried out by the food business operator in accordance with the requirements of Annex III, Section IX, Chapter I, Part III, point 4 of Regulation (EC) No 853/2004, it complies with the maximum residue limits for residues of antibacterial veterinary medicinal products laid down in the Annex to Regulation (EU) No 37/2010;

(f) it has been produced under conditions guaranteeing compliance with the maximum residue levels for pesticides laid down in Regulation (EC) No 396/2005, and maximum levels for contaminants laid down in Regulation (EC) No 1881/2006.

Notes

This certificate is intended 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.

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 establishment of dispatch.

— Box reference I.15: Registration number (railway wagons or container and road vehicle), flight number (aircraft) or name (ship). In case of unloading and reloading, the consignor must inform the border inspection post of introduction into the European Union.

— Box reference I.19: Use the appropriate Harmonised System (HS) code under the following headings: 04.01; 04.02 or 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 number of the production holding(s), collection centre or standardization centre approved for exportation to the European Union.

▼**M1**

Model Milk-RM
Raw milk

COUNTRY

II. Health information	II.a. Certificate reference number	II.b.
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): Qualification and title: Date: Signature: Stamp:		

▼M1

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**

Part I: Details of dispatched consignment

I.1. Consignor Name Address Tel.		I.2. Certificate reference No	I.2.a.
		I.3. Central competent authority	
		I.4. Local competent authority	
I.5. Consignee Name Address Postcode Tel.		I.6.	
I.7. Country of origin ISO code	I.8. Region of origin Code	I.9. Country of destination ISO code	I.10.
I.11. Place of origin Name Approval number Address		I.12.	
I.13. Place of loading		I.14. Date of departure	
I.15. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification Documentation references		I.16. Entry BIP in EU	
		I.17.	

I.18. Description of commodity	I.19. Commodity code (HS code)	
		I.20. Quantity
I.21. Temperature of product Ambient ☐ Chilled ☐ Frozen ☐		I.22. Number of packages
I.23. Seal/Container No		I.24. Type of packaging
I.25. Commodities certified for: Human consumption ☐		
I.26.	I.27. For import or admission into EU ☐	

I.28. Identification of the commodities

Manufacturing plant	Number of packages	Species (Scientific name)	Net weight	Batch number

▼M1

Model Milk-RMP

COUNTRY **Dairy products derived from raw milk for human consumption**

II. Health information	II.a. Certificate reference number	II.b.

Part II: Certification

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 Regulation (EC) No 853/2004 and hereby certify that the dairy products described above has been manufactured from raw milk obtained from animals:

(a) under the control of the official veterinary service,

(b) which were in a country or part thereof that has been free of foot-and-mouth disease and of rinderpest for a period of at least 12 months prior to the date of this certificate, and where vaccination against foot-and-mouth disease has not been carried out during that period,

(c) belonging to holdings which were not under restrictions due to foot-and-mouth disease or rinderpest, and

(d) subject to regular veterinary inspections to ensure that they satisfy the animal health conditions laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004 and in Directive 2002/99/EC;

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, (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 accordance with Regulation (EC) No 852/2004 and checked in accordance with Annex IV to Regulation (EC) No 854/2004,

(ii) which was produced, collected, cooled, stored and transported in accordance with the hygiene conditions laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004,

(iii) which meets the plate and somatic cell count criteria laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004,

(iv) which complies with the guarantees on the residues status of raw milk provided by the monitoring plans for the detection of residues or substances submitted in accordance with Council Directive 96/23/EC, and in particular, Article 29 thereof,

(v) which, pursuant to testing for residues of antibacterial drugs carried out by the food business operator in accordance with the requirements of Annex III, Section IX, Chapter I, Part III, point 4 of Regulation (EC) No 853/2004, it complies with the maximum residue limits for residues of antibacterial veterinary medicinal products laid down in the Annex to Regulation (EU) No 37/2010;

(vi) which has been produced under conditions guaranteeing compliance with the maximum residue levels for pesticides laid down in Regulation (EC) No 396/2005, and maximum levels for contaminants laid down in Regulation (EC) No 1881/2006.

(b) it comes from an establishment implementing a programme based on the HACCP principles in accordance with Regulation (EC) No 852/2004,

(c) it has been obtained from raw milk that has not undergone any heat treatment or any physical or chemical treatment during the manufacturing process,

(d) it has been wrapped, packaged and labeled in accordance with Chapters III and IV of Section IX of Annex III to Regulation (EC) No 853/2004,

(e) it meets the relevant microbiological criteria laid down in Regulation (EC) No 2073/2005 on microbiological criteria for foodstuffs, and

(f) the guarantees covering live animals and products thereof provided by the residue plans submitted in accordance with Directive 96/23/EC, and in particular Article 29 thereof, are fulfilled.

▼M1

Model Milk-RMP

COUNTRY **Dairy products derived from raw 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 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.

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 establishment of dispatch.

— Box reference I.15: Registration number (railway wagons or container and road vehicles), flight number (aircraft) or name (ship). In the case of transport in containers, the total number of containers and their registration number and where there is a serial number of the seal it must be indicated in box I.23. In the case of unloading and reloading, the consignor must inform the border inspection post of introduction into the European Union.

— Box reference I.19: Use the appropriate Harmonised System (HS) code under the following headings: 04.01; 04.02; 04.03; 04.04; 04.05; 04.06; 17.02; 21.05; 22.02; 35.01; 35.02 or 35.04.

— Box reference I.20: Indicate total gross weight and total net weight.

— Box reference I.23: For containers or boxes, the container number and the seal number (if applicable) should be included.

— Box reference I.28: Manufacturing plant: introduce the approval number of the production holding(s), collection centre or standardization centre approved for exportation to the European Union.

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): Qualification and title:

Date: Signature:

Stamp:

▼M1

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

COUNTRY: **Veterinary certificate to EU**

Part I: Details of dispatched consignment

I.1. Consignor Name Address Tel.		I.2. Certificate reference No	I.2.a.
		I.3. Central competent authority	
		I.4. Local competent authority	
I.5. Consignee Name Address Postcode Tel.		I.6.	
I.7. Country of origin ISO code	I.8. Region of origin Code	I.9. Country of destination ISO code	I.10.
I.11. Place of origin Name Approval number Address		I.12.	
I.13. Place of loading		I.14. Date of departure	
I.15. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification Documentation references		I.16. Entry BIP in EU	
		I.17.	

I.18. Description of commodity	I.19. Commodity code (HS code)	
		I.20. Quantity
I.21. Temperature of product Ambient ☐ Chilled ☐ Frozen ☐		I.22. Number of packages
I.23. Seal/Container No		I.24. Type of packaging
I.25. Commodities certified for: Human consumption ☐		
I.26.	I.27. For import or admission into EU ☐	

I.28. Identification of the commodities

Manufacturing plant	Number of packages	Species (Scientific name)	Net weight	Batch number

▼M1

Model Milk-HTB

COUNTRY

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

II. Health information	II.a. Certificate reference number	II.b.

Part II: Certification

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 Regulation (EC) No 853/2004 and hereby certify that the dairy product described above:

(a) has been obtained from animals:

(i) under the control of the official veterinary service,

(ii) which were in a country or part thereof that has been free of foot-and-mouth disease and of rinderpest for a period of at least 12 months prior to the date of this certificate, and where vaccination against foot-and-mouth disease has not been carried out during that period,

(iii) belonging to holdings which were not under restrictions due to foot-and-mouth disease or rinderpest, and,

(iv) subject to regular veterinary inspections to ensure that they satisfy the animal health conditions laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004 and in Directive 2002/99/EC,

(b) has undergone or been produced from raw milk which has been submitted to a pasteurisation treatment involving a single heat treatment with a heating effect at least equivalent to that achieved by a pasteurisation process of at least 72°C for 15 seconds and where applicable, sufficient to ensure a negative reaction to an alkaline phosphatase test applied immediately after the heat treatment.

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, (EC) No 852/2004, (EC) No 853/2004 and (EC) No 854/2004 and hereby certify that the dairy product described above was produced in accordance with those provisions, in particular that:

(a) it was manufactured from raw milk:

(i) which comes from holdings registered in accordance with Regulation (EC) No 852/2004 and checked in accordance with Annex IV to Regulation (EC) No 854/2004,

(ii) which was produced, collected, cooled, stored and transported in accordance with the hygiene conditions laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004,

(iii) which meets the plate and somatic cell count criteria laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004,

(iv) which complies with the guarantees on the residues status of raw milk provided by the monitoring plans for the detection of residues or substances submitted in accordance with Council Directive 96/23/EC, and in particular, Article 29 thereof,

(v) which, pursuant to testing for residues of antibacterial drugs carried out by the food business operator in accordance with the requirements of Annex III, Section IX, Chapter I, Part III, point 4 of Regulation (EC) No 853/2004, it complies with the maximum residue limits for residues of antibacterial veterinary medicinal products laid down in the Annex to Regulation (EU) No 37/2010,

(vi) which has been produced under conditions guaranteeing compliance with the maximum residue levels for pesticides laid down in Regulation (EC) No 396/2005, and maximum levels for contaminants laid down in Regulation (EC) No 1881/2006.

(b) it comes from an establishment implementing a programme based on the HACCP principles in accordance with Regulation (EC) No 852/2004,

(c) it has been processed, stored, wrapped, packaged and transported in accordance with the relevant hygiene conditions laid down in Annex II to Regulation (EC) No 852/2004 and Chapter II of Section IX of Annex III to Regulation (EC) No 853/2004,

(d) it meets the relevant criteria laid down in Chapter II of Section IX of Annex III to Regulation (EC) No 853/2004 and the relevant microbiological criteria laid down in Regulation (EC) No 2073/2005 on microbiological criteria for foodstuffs,

(e) the guarantees covering live animals and products thereof provided by the residue plans submitted in accordance with Directive 96/23/EC, and in particular Article 29 thereof, are fulfilled.

▼M1

Model Milk-HTB

COUNTRY

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

II. Health information	II.a. Certificate reference 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 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 establishment of dispatch.

— Box reference I.15: Registration number (railway wagons or container and road vehicles), flight number (aircraft) or name (ship). In the case of transport in containers, the total number of containers and their registration number and where there is a serial number of the seal it must be indicated in box I.23. In the case of unloading and reloading, the consignor must inform the border inspection post of introduction into the European Union.

▶(1)— Box reference I.19: Use the appropriate Harmonised System (HS) code under the following headings: 04.01; 04.02; 04.03; 04.04; 04.05; 04.06; 15.17; 17.02; 21.05; 22.02; 28.35; 35.01; 35.02 or 35.04.◀

— Box reference I.20: Indicate total gross weight and total net weight.

— Box reference I.23: For containers or boxes, the container number and the seal number (if applicable) should be included.

— Box reference I.28: Manufacturing plant: introduce the approval number of the treatment and/or processing establishment(s) approved for export to the European Union.

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): Qualification and title:

Date: Signature:

Stamp:

▶(1) M3

▼M3

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

COUNTRY **Veterinary certificate to EU**

Part I: Details of dispatched consignment

I.1. Consignor Name Address Tel.	I.2. Certificate reference No	I.2.a.
	I.3. Central competent authority	
	I.4. Local competent authority	
I.5. Consignee Name Address Postcode Tel.	I.6.	
I.7. Country of origin ISO code	I.8. Region of origin Code	I.9. Country of destination ISO code / I.10.
I.11. Place of origin Name Approval number Address	I.12.	
I.13. Place of loading	I.14. Date of departure	
I.15. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification Documentary references	I.16. Entry BIP in EU	
	I.17.	
I.18. Description of commodity	I.19. Commodity code (HS code)	
	I.20. Quantity	
I.21. Temperature of product Ambient ☐ Chilled ☐ Frozen ☐	I.22. Number of packages	
I.23. Seal/Container No	I.24. Type of packaging	
I.25. Commodities certified for: Human consumption ☐		
I.26.	I.27. For import or admission into EU ☐	

I.28. Identification of the commodities

Species (scientific name)	Manufacturing plant	Number of packages	Net weight	Batch number

▼M3

Model Milk-HTC

COUNTRY — **Dairy products from third countries authorised in column C**

II. Health information	II.a. Certificate reference number	II.b.

Part II: Certification

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 Regulation (EC) No 853/2004 and hereby certify that the dairy product described above:

(a) has been obtained from animals:

(i) under the control of the official veterinary service;

(ii) belonging to holdings which were not under restrictions due to foot-and-mouth disease or rinderpest; and

(iii) subject to regular veterinary inspections to ensure that they satisfy the animal health conditions laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004 and in Directive 2002/99/EC;

either [(b) the dairy product was made from raw milk sourced from cows, ewes, goats, buffaloes or, where authorised in accordance with footnote (2) of Annex I to Regulation (EC) No 605/2010, from -camels of the species *Camelus dromedarius*, and has undergone, prior to import into the territory of the European Union:

([1]) *either* [(i) a sterilisation process, to achieve an F_0 value equal to 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;]

([1]) *or* [(iii) 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 an alkaline phosphatase test, applied immediately after the heat treatment;]

([1]) *or* [(iv) a treatment with an equivalent pasteurisation effect to point (iii) achieving, where applicable, a negative reaction to an alkaline phosphatase test, applied immediately after the heat treatment;]

([1]) *or* [(v) a HTST treatment of milk with a pH below 7,0;]

([1]) *or* [(vi) a HTST treatment combined with another physical treatment by

([1]) *either* [(1) lowering the pH below 6 for one hour;]

([1]) *or* [(2) additional heating equal to or greater than 72 °C, combined with desiccation;]]

([1]) *or* [(b) the dairy product was made from raw milk sourced from animals other than cows, ewes, goats, buffaloes or camels of the species *Camelus dromedarius*, and has undergone, prior to import into the territory of the European Union:

([1]) *either* [(i) a sterilisation process, to achieve an F_0 value equal to 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;]]

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, (EC) No 852/2004, (EC) No 853/2004 and (EC) No 854/2004 and hereby certify that the dairy product described above was produced in accordance with those provisions, and in particular that:

(a) it was manufactured from raw milk:

(i) which comes from holdings registered in accordance with Regulation (EC) No 852/2004 and checked in accordance with Annex IV to Regulation (EC) No 854/2004;

(ii) which was produced, collected, cooled, stored and transported in accordance with the hygiene conditions laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004;

(iii) which meets the plate and somatic cell count criteria laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004;

(iv) which complies with the guarantees on the residues status of raw milk provided by the monitoring plans for the detection of residues or substances submitted in accordance with Council Directive 96/23/EC, and in particular, Article 29 thereof;

▼M3

Model Milk-HTC

COUNTRY **Dairy products from third countries authorised in column C**

II. Health information	II.a. Certificate reference number	II.b.

(v) which, pursuant to testing for residues of antibacterial drugs carried out by the food business operator in accordance with the requirements of Annex III, Section IX, Chapter I, Part III, point 4 to Regulation (EC) No 853/2004, it complies with the maximum residue limits for residues of antibacterial veterinary medicinal products laid down in the Annex to Regulation (EU) No 37/2010;

(vi) which has been produced under conditions guaranteeing compliance with the maximum residue levels for pesticides laid down in Regulation (EC) No 396/2005, and maximum levels for contaminants laid down in Regulation (EC) No 1881/2006;

(b) it comes from an establishment implementing a programme based on the HACCP principles in accordance with Regulation (EC) No 852/2004;

(c) it has been processed, stored, wrapped, packaged and transported in accordance with the relevant hygiene conditions laid down in Annex II to Regulation (EC) No 852/2004 and Chapter II of Section IX of Annex III to Regulation (EC) No 853/2004;

(d) it meets the relevant criteria laid down in Chapter II of Section IX of Annex III to Regulation (EC) No 853/2004 and the relevant microbiological criteria laid down in Regulation (EC) No 2073/2005 on microbiological criteria for foodstuffs;

(e) the guarantees covering live animals and products thereof provided by the residue plans submitted in accordance with Directive 96/23/EC, and in particular Article 29 thereof, are fulfilled.

Notes

This certificate is intended for dairy products for human consumption from third countries or parts thereof authorised, where applicable for milk from certain animal species only, in column C of Annex I to 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 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 establishment 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 (HS) code under the following headings: 04.01; 04.02; 04.03; 04.04; 04.05; 04.06; 15.17; 17.02; 19.01; 21.05; 21.06; 22.02; 28.35; 35.01; 35.02 or 35.04.

— Box reference I.20: indicate total gross weight and total net weight.

— Box reference I.23: for containers or boxes, the container number and the seal number (if applicable) should be included.

— Box reference I.28: manufacturing plant: introduce the approval number of the treatment and/or processing establishment(s) approved for export to the European Union.

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.

Official veterinarian

Name (in capital letters): Qualification and title:

Date: Signature:

Stamp:

▼M6

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

COUNTRY **Veterinary certificate to EU**

Part I: Details of dispatched consignment

I.1. Consignor Name Address Tel.	I.2. Certificate reference No	I.2.a.
	I.3. Central competent authority	
	I.4. Local competent authority	
I.5. Consignee Name Address Postcode Tel.	I.6.	
I.7. Country of origin ISO code	I.8.	I.9. Country of destination ISO code / I.10.
I.11. Place of origin Name Approval number Address	I.12.	
I.13. Place of loading	I.14. Date of departure	
I.15. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification Documentary references	I.16. Entry BIP in EU	
	I.17.	
I.18. Description of commodity	I.19. Commodity code (HS code)	
	I.20. Quantity	
I.21. Temperature of product Ambient ☐ Chilled ☐ Frozen ☐	I.22. Number of packages	
I.23. Seal/Container No	I.24. Type of packaging	
I.25. Commodities certified for: Human consumption ☐		
I.26.	I.27. For import or admission into EU ☐	

I.28. Identification of the commodities

Species (Scientific name)	Manufacturing plant	Number of packages	Net weight	Batch number

▼M6

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

II. Health information	II.a. Certificate reference number	II.b.

Part II: Certification

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 Regulation (EC) No 853/2004 and hereby certify that the colostrum /colostrum-based products [1] described in Part I:

have been obtained or manufactured from colostrum obtained from animals:

(i) under the control of the official veterinary service;

(ii) which were in a third country or part thereof that has been free of foot-and-mouth disease and of rinderpest for a period of at least 12 months prior to the date of this certificate, and where vaccination against foot-and-mouth disease has not been carried out during that period;

(iii) belonging to holdings which were not under restrictions due to foot-and-mouth disease or rinderpest, and

(iv) subject to regular veterinary inspections to ensure that they satisfy the animal health conditions laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004 and in Directive 2002/99/EC.

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, (EC) No 852/2004, (EC) No 853/2004 and (EC) No 854/2004 and hereby certify that the colostrum/colostrum-based products made with colostrum [1] described in Part I were produced in accordance with those provisions, and in particular that:

(a) they were manufactured from colostrum:

(i) which comes from holdings registered in accordance with Regulation (EC) No 852/2004 and checked in accordance with Annex IV to Regulation (EC) No 854/2004;

(ii) which was produced, collected, cooled, stored and transported in accordance with the hygiene conditions laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004;

(iii) which complies with the guarantees on the residues status of colostrum provided by the monitoring plans for the detection of residues or substances submitted in accordance with Directive 96/23/EC, and in particular, Article 29 thereof;

(iv) which, pursuant to testing for residues of antibacterial drugs carried out by the food business operator in accordance with the requirements of point 4 in Part III of Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004, complies with the maximum residue limits for residues of antibacterial veterinary medicinal products laid down in the Annex to Regulation (EU) No 37/2010;

(v) which has been produced under conditions guaranteeing compliance with the maximum residue levels for pesticides laid down in Regulation (EC) No 396/2005, and maximum levels for contaminants laid down in Regulation (EC) No 1881/2006;

(b) they come from an establishment implementing a programme based on the HACCP principles in accordance with Regulation (EC) No 852/2004;

(c) they have been processed, stored, wrapped, packaged and labeled in accordance with Chapters III and IV of Section IX of Annex III to Regulation (EC) No 853/2004;

(d) they meet the relevant requirements laid down in Chapter II, Section IX of Annex III to Regulation (EC) N0 853/2004 and the relevant microbiological criteria laid down in Regulation (EC) No 2073/2005 on microbiological criteria for foodstuffs, and

(e) the guarantees covering live animals and products thereof provided by the residue plans submitted in accordance with Directive 96/23/EC, and in particular Article 29 thereof are fulfilled.

▼M6

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

II. Health information	II.a. Certificate reference number	II.b.

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 establishment of dispatch.

— Box reference I.15: Registration number (railway wagons or container and road vehicle), flight number (aircraft) or name (ship). In case of unloading and reloading, the consignor must inform the border inspection post of introduction into the European Union.

— Box reference I.19: Use the appropriate Harmonised System (HS) code under the following headings: 04.01; 04.02; 04.03; 04.04; 04.05; 04.06; 04.10; 15.17; 17.02; 19.01; 21.05; 21.06; 22.02; 28.35; 30.01; 35.01; 35.02 or 35.04.

— Box reference I.20: Indicate total gross weight and total net weight.

— Box reference I.23: For containers or boxes, the container number and the seal number (if applicable) should be included.

— Box reference I.28: Manufacturing plant: introduce the approval number of the production holding(s), collection centre or standardization centre approved for exportation 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.

Official veterinarian

Name (in capital letters): Qualification and title:

Date: Signature:

Stamp:

▼M6

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 **Veterinary certificate to EU**

Part I: Details of dispatched consignment

I.1. Consignor Name Address Tel.		I.2. Certificate reference No	I.2.a.
		I.3. Central competent authority	
		I.4. Local competent authority	
I.5. Consignee Name Address Postcode Tel.		I.6. Person responsible for the load in EU Name Address Postal code Tel. No	
I.7. Country of origin ISO code	I.8. Region of origin Code	I.9. Country of destination ISO code	I.10.
I.11. Place of origin Name Approval number Address		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 Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification Documentary references		I.16. Entry BIP in EU I.17.	
I.18. Description of commodity		I.19. Commodity code (HS code)	I.20. Quantity
I.21. Temperature of product Ambient ☐ Chilled ☐ Frozen ☐			I.22. Number of packages
I.23. Seal/Container No			I.24. Type of packaging
I.25. Commodities certified for: Human consumption ☐			
I.26. For transit through EU to 3rd Country ☐ 3rd country ISO code		I.27.	

I.28. Identification of the commodities

Species (Scientific name)	Manufacturing plant	Number of packages	Net weight	Batch number

▼M6

Model Milk/Colostrum-T/S

Raw milk, dairy products, colostrum and colostrum-based products for human consumption for transit or storage

COUNTRY

II. Health information	II.a. Certificate reference number	II.b.

Part II: Certification

II.1 Animal Health Attestation

I, the undersigned official veterinarian, hereby certify that the [raw milk] / [dairy products] / [colostrum] / [colostrum-based products] (1) (2) for [transit] / [storage] (2) in the European Union described in Part I:

(a) come from a country or part thereof authorised for imports to the European Union of raw milk, dairy products, colostrum or colostrum-based products as laid down in Annex I to Regulation (EU) No 605/2010;

(b) comply with the relevant animal health conditions for the products concerned as laid down in the animal health attestation in Part II.1 of the model health certificates [Milk-RM] / [Milk-RMP] / [Milk-HTB] / [Milk-HTC]/[Colostrum-C/CBP](2) in Part 2 of Annex II to Regulation (EU) No 605/2010;

(c) was/were produced on ..... (3) or between ..... (3) and ..... (3).

Notes

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 establishment of dispatch. Name of the country of origin which must be the same as the country of export.

— 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 case of unloading and reloading, the consignor must inform the border inspection post of introduction into the European Union.

— Box reference I.19: Use the appropriate Harmonised System (HS) code under the following headings: 04.01; 04.02; 04.03; 04.04; 04.05; 04.06; 15.17; 17.02; 19.01; 21.05; 21.06.; 22.02; 28.35; 30.01; 35.01; 35.02; 35.04 or 04.10.

— Box reference I.20: Indicate total gross weight and total net weight.

— Box reference I.23: For containers or boxes, the container number and the seal number (if applicable) should be included.

— Box reference I.28: Manufacturing plant: introduce the approval number of the production holding(s), collection centre or standardization centre approved for exportation to the European Union.

Part II:

(1) Raw milk, dairy products, colostrum and colostrum –based products means raw milk, dairy products, colostrum and colostrum-based products for human consumption in transit or storage in accordance with Article 12(4) or Article 13 of 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 (OJ L 24, 30.1.1998, p. 9).

(2) Keep as appropriate.

(3) Date or dates of production. Imports of raw milk, dairy products, colostrum and colostrum-based products shall not be allowed when obtained either prior to the date of authorisation for exportation to the European Union of the third country or part thereof mentioned under I.7 and I.8, or during a period where restrictive measures have been adopted by the European Union against imports of raw milk, dairy products, colostrum and colostrum-based products from this third country or part thereof.

— 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.

▼**M6**

Model Milk/Colostrum-T/S

Raw milk, dairy products, colostrum and colostrum-based products for human consumption for transit or storage

COUNTRY

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