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(Non-legislative acts)

REGULATIONS

COMMISSION IMPLEMENTING REGULATION (EU) No 914/2011

of 13 September 2011

amending Regulation (EU) No 605/2010 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 and point (b) of Article 9(4) thereof,

Whereas:

- (1) Regulation (EU) No 605/2010 of 2 July 2010 (²) 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 provides that consignments of raw milk and dairy products intended for human consumption, authorised for importation into the Union, are to be accompanied by a health certificate drawn up in accordance with the appropriate model set out in Part 2 of Annex II thereto for the commodity concerned ('the model health certificates').
- (2) It should be clarified that the requirement regarding the use of the model health certificates provided for in that Regulation is without prejudice to specific certification requirements laid down in other Union acts or in agreements concluded by the Union with third countries.
- (1) OJ L 18, 23.1.2003, p. 11.

- (3) The model health certificates specify the commodity code for the commodities covered by Regulation (EU) No 605/2010 on the basis of the Harmonised Commodity Description and Coding System ('HS codes') of tariff nomenclature maintained by the World Customs Organization (WCO).
- (4) Certain dairy products covered by Regulation (EU) No 605/2010 do not fall within the commodity codes in the model health certificates. In order to allow a more precise identification of those commodities in the model health certificates, it is necessary to amend those models and add the missing HS codes, in particular as regards HS codes 35.01 and 35.02 (casein, caseinates and albumines).
- (5) In addition, it should be clarified in the model health certificates that the requirements regarding antibiotic residues, contaminants and pesticide residues may be based on the findings of official monitoring programmes which are at least equivalent to those provided for in Union legislation.
- (6) For reasons of clarity and transparency of Union legislation, the model health certificates should be replaced by the model health certificates set out in the Annex to this Regulation.
- (7) Regulation (EU) No 605/2010 should therefore be amended accordingly.
- (8) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on the Food Chain and Animal Health,

⁽²⁾ OJ L 175, 10.7.2010, p. 1.

HAS ADOPTED THIS REGULATION:

Article 1

Regulation (EU) No 605/2010 is amended as follows:

(1) In Article 1, the following second paragraph is added:

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.'

(2) Annex II is amended in accordance with the Annex to this Regulation.

Article 2

For a transitional period until 30 November 2011, consignments of raw milk and dairy products for which the relevant health certificates have been issued in accordance with Regulation (EU) No 605/2010 before the entry into force of this Regulation may continue to be introduced into the Union.

Article 3

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

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

Done at Brussels, 13 September 2011.

For the Commission The President José Manuel BARROSO COUNTRY:

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ANNEX

In Annex II, parts 2 and 3 to Regulation (EU) No 605/2010 are replaced by the following:

'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

Veterinary certificate to EU

	l.1.	Consignor	I.2. Certificat	te reference No	l.2.a.
		Name Address	I.3. Central of	competent authority	
lent		Tel.	I.4. Local co	mpetent authority	
Part I: Details of dispatched consignment	1.5.	Consignee Name Address	1.6.		
spatcheo		Postcode Tel.			
ils of dis	I.7.	Country of origin ISO code I.8. Region of origin Code	I.9. Country destinati	of ISO code ion	1.10.
Deta	1.11.	Place of origin	I.12.	I	
Part I:		Name Approval number Address			
	I.13.	Place of loading	I.14. Date of	departure	
	l.15.	Means of transport	I.16. Entry BI	P in EU	
		Aeroplane 🗌 Ship 🛄 Railway wagon 🗌			
		Road vehicle Other Other I Identification Documentation references	l.17.		
	l.18.	Description of commodity		I.19. Commodity cod	e (HS code)
					I.20. Quantity
	1.21.	Temperature of product			I.22. Number of packages
		Ambient Chilled		Frozen	
		Seal/Container No			I.24. Type of packaging
	1.25.	Commodities certified for:			
	1.26.		I.27. For impo	ort or admission into E	U 🗆
	1.28.	Identification of the commodities			
		Manufacturing plant Number of packages	Species (Scientific name	Net we	eight Batch number

Model Milk-RM

	cou	NTRY		Raw milk
	11.	Health information	II.a. Certificate reference number	II.b.
	11.1.	Animal Health Attestation		
_		I, the undersigned official veterinarian, declare that 853/2004 and hereby certify that the raw milk desc		
atior		(a) under the control of the official veterinary service	Ce,	
Part II: Certification		(b) which were in a country or part thereof that has prior to the date of this certificate, and where v		
art II		(c) belonging to holdings which were not under res	strictions due to foot-and-mouth disease or	rinderpest, and
<u>а</u>		(d) subject to regular veterinary inspections to ensu Annex III to Regulation (EC) No 853/2004 and		tions laid down in Chapter I of Section IX of
	II.2.	Public Health attestation		
		I, the undersigned official inspector, declare that I a (EC) No 853/2004 and (EC) No 854/2004 and her provisions, in particular that:	m aware of the relevant provisions of Regula reby certify that the raw milk described abo	itions (EC) No 178/2002, (EC) No 852/2004, ve was produced in accordance with those
		 (a) it comes from holdings registered in accordar Regulation (EC) No 854/2004, 	nce with Regulation (EC) No 852/2004 and	I checked in accordance with Annex IV to
		(b) it was produced, collected, cooled, stored and tr of Annex III to Regulation (EC) No 853/2004,	ransported in accordance with the hygiene co	onditions laid down in Chapter I of Section IX
		(c) it meets the plate and somatic cell count crite	ria 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 mi in accordance with Council Directive 96/23/EC,		
		(e) pursuant to testing for residues of antibacterial Annex III, Section IX, Chapter I, Part III, point 4 of of antibacterial veterinary medicinal products la	of Regulation (EC) No 853/2004, it complies	with the maximum residue limits for residues
		(f) it has been produced under conditions guarante (EC) No 396/2005, and maximum levels for co		
	Note	35		
		certificate is intended for raw milk from third countr ided for further processing in the European Union b		of Annex I to Regulation (EU) No 605/2010
	Part	1:		
	— Е	Box reference I.7: Provide name and ISO code of	the country or part thereof as appearing in	Annex I to Regulation (EU) No 605/2010.
	— Е	Box reference I.11: Name, address and approval nur	mber of the establishment of dispatch.	
		Box reference I.15: Registration number (railway wa nloading and reloading, the consignor must inform t		
	— E	Box reference I.19: Use the appropriate Harmonised	System (HS) code under the following hear	dings: 04.01; 04.02 or 04.03.
	— Е	Box reference I.20: Indicate total gross weight and to	otal net weight.	
	— Е	Box reference I.23: For containers or boxes, the con	tainer number and the seal number (if appli	cable) should be included.
		Box reference I.28: Manufacturing plant: introduce the proved for exportation to the European Union.	approval number of the production holding(s	s), collection centre or standardization centre

Model Milk-RM

COUN	ITRY		Raw mile
11.	Health information	II.a. Certificate reference number	II.b.
Part	II:		
— т	he colour of the signature shall be different to that o	f the printing. The same rule applies to stamp	os other than those embossed or watermark.
Offici	ial veterinarian		
	Name (in capital letters):	Qualificati	on and title:
	Date:	Signature	:
	Stamp:		

Model Milk-RMP

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

COUN	ITRY	:			Veterinary certificate to EU
	l.1.	Consignor		I.2. Certificate reference No	l.2.a.
		Name		I.3. Central competent authority	
		Address			
lent		Tel.		I.4. Local competent authority	
Part I: Details of dispatched consignment	l.5.	Consignee		1.6.	
suos		Name Address			
ed o					
atch		Postcode Tel.			
disp	17	Country of origin ISO code	I.8. Region of origin Code	I.9. Country of ISO code	9 1.10.
s of	1.7.	Country of origin 150 code		destination	1.10.
etail					
D :	1.11.	Place of origin		l.12.	
Part		Name Address	Approval number		
	l.13.	Place of loading		I.14. Date of departure	
		-			
	l.15.	Means of transport		I.16. Entry BIP in EU	
		Aeroplane Ship	Railway wagon		
		Road vehicle D Other			
		Identification		1.17.	
		Documentation references			
	l.18.	Description of commodity		I.19. Commodity co	ode (HS code)
					I.20. Quantity
	I.21.	Temperature of product			I.22. Number of packages
		Ambient	Chilled	Frozen	
	1.23.	Seal/Container No			I.24. Type of packaging
	I.25.	Commodities certified for:			
		Human consumption			
	I.26.			I.27. For import or admission into	EU 🗌
	1.28.	Identification of the commodities	3		
		Manufacturing plant	Number of packages	Species Net v	weight Batch number
				(Scientific name)	

Part II: Certification

c	ou	NTRY	Dairy products deriv	<i>Model Milk-RMF</i> red from raw milk for human consumptior
ſ	11.	Health information	II.a. Certificate reference number	II.b.
	II.1.	Animal Health Attestation		
		I, the undersigned official veterinarian, declare that 853/2004 and hereby certify that the dairy produc		
		(a) under the control of the official veterinary servi	ice,	
		(b) which were in a country or part thereof that has prior to the date of this certificate, and where		
		(c) belonging to holdings which were not under re	estrictions due to foot-and-mouth disease or	rinderpest, and
		(d) subject to regular veterinary inspections to ens Annex III to Regulation (EC) No 853/2004 and		tions laid down in Chapter I of Section IX of
	11.2.	Public Health attestation		
		I, the undersigned official inspector, declare that I a (EC) No 853/2004 and (EC) No 854/2004 and her accordance with those provisions, in particular tha	reby certify that the dairy product made with	
		(a) it was manufactured from raw milk:		
		(i) which comes from holdings registered in ac Regulation (EC) No 854/2004,	cordance with Regulation (EC) No 852/2004	and checked in accordance with Annex IV to
		(ii) which was produced, collected, cooled, sto Section IX of Annex III to Regulation (EC)		hygiene conditions laid down in Chapter I of
		(iii) which meets the plate and somatic cell c 853/2004,	count criteria laid down in Chapter I of Sec	ation IX of Annex III to Regulation (EC) No
		(iv) which complies with the guarantees on the or substances submitted in accordance wit	residues status of raw milk provided by the r th Council Directive 96/23/EC, and in particu	01
		(v) which, pursuant to testing for residues of requirements of Annex III, Section IX, Cha residue limits for residues of antibacterial	apter I, Part III, point 4 of Regulation (EC) N	business operator in accordance with the No 853/2004, it complies with the maximum the Annex to Regulation (EU) No 37/2010;
		(vi) which has been produced under condition: Regulation (EC) No 396/2005, and maximu	s guaranteeing compliance with the maximu um levels for contaminants laid down in Reg	
		(b) it comes from an establishment implementing No 852/2004,	g a programme based on the HACCP prir	nciples in accordance with Regulation (EC)
		(c) it has been obtained from raw milk that has manufacturing process,	not undergone any heat treatment or any	physical or chemical treatment during the
		(d) it has been wrapped, packaged and labeled No 853/2004,	in accordance with Chapters III and IV of	Section IX of Annex III to Regulation (EC)
		(e) it meets the relevant microbiological criteria la	id down in Regulation (EC) No 2073/2005 o	on microbiological criteria for foodstuffs, and
		(f) the guarantees covering live animals and pro 96/23/EC, and in particular Article 29 thereof, a		ans submitted in accordance with Directive

Model Milk-RMP

COUNTR	TRY Dairy products derived from raw milk for		ed from raw milk for human consumption
11.	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:

Stamp:

Signature:

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

COUN	ITRY	:			Veterinary certificate to EU
	l.1.	Consignor		I.2. Certificate reference No	l.2.a.
		Name		I.3. Central competent authority	
		Address			
Part I: Details of dispatched consignment		Tel.		I.4. Local competent authority	
ignn	l.5.	Consignee		1.6.	
suo		Name Address			
o pa		Address			
atche		Postcode			
lispa		Tel.			
of d	1.7.	Country of ISO code origin	I.8. Region of origin Code	I.9. Country of ISO code destination	l.10.
ails					
Det	l.11.	Place of origin	· ·	I.12.	
ut I:		Name	Approval number		
Ра		Address			
	l.13.	Place of loading		I.14. Date of departure	
	l.15.	Means of transport		I.16. Entry BIP in EU	
		Aeroplane 🗌 Ship	Railway wagon		
			er 🔲	1.17.	
		Identification		1.17.	
		Documentation references			
	l.18.	Description of commodity		I.19. Commodity co	de (HS code)
					I.20. Quantity
	I.21.	Temperature of product			I.22. Number of packages
		Ambient 🔲	Chilled	Frozen 🗖	
	1.23.	Seal/Container No			I.24. Type of packaging
	1.25.	Commodities certified for:			
		Human consumption 🔲			
	1.26.			I.27. For import or admission into I	
	1.28.	Identification of the commoditie	96		
		Manufacturing plant	Number of packages	Species Net w	veight Batch number
		manaraotan'ny piant		(Scientific name)	Daton humbol

Model Milk-HTB

	coui		Dairy products derived from milk of cov consumption from third countries autho	vs, ewes, goats and buffaloes for human rised in column B
	11.	Health information	II.a. Certificate reference number	II.b.
	11.1.	Animal Health Attestation		
		I, the undersigned official veterinarian, declare that I 853/2004 and hereby certify that the dairy product		ctive 2002/99/EC and of Regulation (EC) No
		(a) has been obtained from animals:		
tion		(i) under the control of the official veterinary se	ervice,	
Part II: Certification		 (ii) which were in a country or part thereof that months prior to the date of this certificate, ar period, 		and of rinderpest for a period of at least 12 disease has not been carried out during that
Part II		(iii) belonging to holdings which were not under	r restrictions due to foot-and-mouth disease	or rinderpest, and,
		 (iv) subject to regular veterinary inspections to e Annex III to Regulation (EC) No 853/2004 a 		ditions laid down in Chapter I of Section IX of
		(b) has undergone or been produced from raw milk with a heating effect at least equivalent to the applicable, sufficient to ensure a negative react	at achieved by a pasteurisation process o	f at least 72°C for 15 seconds and where
	11.2.	Public Health attestation		
		I, the undersigned official veterinarian, declare the No 852/2004, (EC) No 853/2004 and (EC)	54/2004 and hereby certify that the dairy	
		(a) it was manufactured from raw milk:		
		 (i) which comes from holdings registered in acc Regulation (EC) No 854/2004, 	ordance with Regulation (EC) No 852/2004	and checked in accordance with Annex IV to
		(ii) which was produced, collected, cooled, stor Section IX of Annex III to Regulation (EC) №		hygiene conditions laid down in Chapter I of
		(iii) which meets the plate and somatic cell co 853/2004,	ount criteria laid down in Chapter I of Sec	tion IX of Annex III to Regulation (EC) No
		 (iv) which complies with the guarantees on the r or substances submitted in accordance with 		
			oter I, Part III, point 4 of Regulation (EC) N	I business operator in accordance with the lo 853/2004, it complies with the maximum the Annex to Regulation (EU) No 37/2010,
		(vi) which has been produced under conditions Regulation (EC) No 396/2005, and maximu	0 1	
		(b) it comes from an establishment implementing No 852/2004,	a programme based on the HACCP prin	ciples in accordance with Regulation (EC)
		(c) it has been processed, stored, wrapped, packa Annex II to Regulation (EC) No 852/2004 and (
		(d) it meets the relevant criteria laid down in Cha microbiological criteria laid down in Regulation		
		(e) the guarantees covering live animals and pro 96/23/EC, and in particular Article 29 thereof, a		ns submitted in accordance with Directive

Model Milk-HTB

COUNTRY	Dairy products derived from milk o consumption from third countries a	f cows, ewes, goats and buffaloes for human authorised in column B
II. Health information	II.a. Certificate reference number	II.b.
Notes		
This certificate is intended for dairy products for humar Regulation (EU) No 605/2010 intended for importation i		arts thereof authorised in column B of Annex I of
Part I:		
- Box reference I.7: Provide name and ISO code of	the country or part thereof as appear	ing in Annex I to Regulation (EU) No 605/2010.
- Box reference I.11: Name, address and approval nu	umber of the establishment of dispatch.	
 Box reference I.15: Registration number (railway wag transport in containers, the total number of container indicated in box I.23. In the case of unloading and European Union. 	s and their registration number and whe	ere there is a serial number of the seal it must be
 Box reference I.19: Use the appropriate Harmonised 17.02; 21.05; 22.02; 35.01; 35.02 or 35.04. 	System (HS) code under the following h	eadings: 04.01; 04.02; 04.03; 04.04; 04.05; 04.06;
- Box reference I.20: Indicate total gross weight and t	otal net weight.	
- Box reference I.23: For containers or boxes, the con	ntainer number and the seal number (if	applicable) should be included.
 Box reference I.28: Manufacturing plant: introduce export to the European Union. 	the approval number of the treatment	and/or processing establishment(s) approved for
Part II:		
 The colour of the signature shall be different to that of 	of the printing. The same rule applies to	stamps other than those embossed or watermark.
Official veterinarian		
Name (in capital letters):	Qua	lification and title:
Date:	Sigr	nature:
Stamp:		

Model Milk-HTC

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

COUN	ITRY	:								Vet	erinary certificate	e to EU
	l.1.	Consignor				1.2. (Certificate	e referenc	e No	l.2.a.		
		Name					<u> </u>					
		Address				1.3. (Central c	ompetent	authority			
Part I: Details of dispatched consignment		Tel.			I.4. I	Local cor	mpetent a	uthority				
ignn	l.5.	Consignee				I.6.						
suo		Name Address										
ed c		Address										
Itche		Postcode			_							
lispa		Tel.		I								
of d	I.7.	Country of origin	ISO code	I.8. Region of origin	Code	1.9.	Country o destinatio	of	ISO code	I.10.		
ails							uestinatio	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	1			
Det	l.11.	Place of origin				l.12.			•			
μ		Name		Approval numb	er							
Ра		Address										
	l.13.	Place of loading				I.14.	Date of c	departure				
	l.15.	Means of transport				I.16.	Entry BIF	in EU				
		Aeroplane 🔲	Ship	Railway w	/agon 🔲							
		Road vehicle 🗌	Other			1.17.						
		Identification				1.17.						
		Documentation refe	rences									
	l.18.	Description of comr	nodity					I.19. Cor	mmodity coo	de (HS coo	de)	
							L			1.20. Quar	ntitv	
	I.21.	Temperature of proc	duct							I.22. Num	ber of packages	
		Ambient 🔲		Chilled 🗌				Frozen	ן כ			
	1.23.	Seal/Container No								1.24. Туре	e of packaging	
	I.25.	Commodities certifie	ed for:						1			
		Human consumption	n 🗖									
						1						
	1.26.					1.27. I	For impo	rt or adm	ission into E	U		
	I.28.	Identification of the	commodities	3								
		Manufacturing plant		Number of packages		Sp	ecies		Net w	eight	Batch numb	er
						(Scient	ific name)				

	COUNTRY		Dairy products fro	<i>Model Milk-HTC</i> om third countries authorised in column C
	II.	Health information	II.a. Certificate reference number	II.b.
	II.1.	Animal Health Attestation		
		I, the undersigned official veterinarian, dec (EC) No 853/2004 and hereby certify that	lare that I am aware of the relevant provision the dairy product described above:	s of Directive 2002/99/EC and of Regulation
_		(a) has been obtained from animals:		
icatio		(i) under the control of the official v	eterinary service,	
Certif		(ii) belonging to holdings which were	e not under restrictions due to foot-and-mouth	n disease or rinderpest, and,
Part II: Certification			ections to ensure that they satisfy the anima tion (EC) No 853/2004 and in Directive 2002	
	(¹) either	[(b) in the case of dairy products made from into the territory of the European Unio	om raw milk sourced from cows, ewes, goats n:	or buffaloes have undergone, prior to import
	(¹) eith	er [(i) a sterilisation process, to achieve	e 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 combinat	ion with a suitable holding time;]
	(¹) or		steurisation treatment (HTST) at 72°C for 15 s where applicable, a negative reaction to a all	
	(¹) or	(iv) a treatment with an equivalent pa phosphatase test, applied immed	steurisation effect to point (iii) achieving, where liately after the heat treatment;]	e applicable, a negative reaction to a alkaline
	(¹) or	[(v) a HTST treatment with a pH belo	vw 7.0;]	
	(¹) or	[[(vi) a HTST treatment combined with	another physical treatment by	
	(¹) eith	er [(vi) (1) lowering the pH below 6 for a	one hour;]	
	(¹) or	[(vi) (2) additional heating equal to or	greater than 72 $^\circ\mathrm{C}$ or more, combined with	desiccation;]]
	(¹) or	[(b) in the case of dairy products made undergone, prior to import into the ter	from raw milk sourced from animals other ritory of the European Union:	than cows, ewes, goats or buffaloes have
	(¹) eith	er [(i) a sterilisation process, to achieve	an F_0 value equal to or greater than three;]	
	(¹) or	[(ii) an ultra high temperature (UHT) tr	eatment at not less than 135°C in combination	on with a suitable holding time;]]
	11.2.	Public Health attestation		
			clare that I am aware of the relevant provisi No 854/2004 and hereby certify that the dair at:	
		(a) it was manufactured from raw milk:		
		 (i) which comes from holdings register Annex IV to Regulation (EC) No 85 	ered in accordance with Regulation (EC) No 54/2004;	852/2004 and checked in accordance with
		(ii) which was produced, collected, coo I of Section IX of Annex III to Regu	led, stored and transported in accordance with Ilation (EC) No 853/2004;	n the hygiene conditions laid down in Chapter
		(iii) which meets the plate and somatic No 853/2004;	e cell count criteria laid down in Chapter I o	f Section IX of Annex III to Regulation (EC)
			s on the residues status of raw milk provided n accordance with Council Directive 96/23/E0	
		requirements of Annex III, Section IX	ues of antibacterial drugs carried out by the fc K, Chapter I, Part III, point 4 of Regulation (EC sterial veterinary medicinal products laid down) No 853/2004, it complies with the maximum

:001	NTRY		Dairy products	s from third countries authorised in column		
II.	Health information	II.a. Certificate refe	rence number	II.b.		
				aximum residue levels for pesticides 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 c microbiological criteria laid down in			Regulation (EC) No 853/2004 and the releva criteria for foodstuffs,		
	(e) the guarantees covering live anima 96/23/EC, and in particular Article 2		led by the residue	e plans submitted in accordance with Directi		
Note	95					
	certificate is intended for dairy products ulation (EU) No 605/2010 intended for ir			arts thereof authorised in column C of Annex I		
Part	l:					
— в	ox reference I.7: Provide name and IS	O code of the country or part	thereof as appearir	ng in Annex I to Regulation (EU) No 605/201		
— в	ox reference I.11: Name, address and a	approval number of the establish	ment of dispatch.			
р о	rovided. In the case of transport in conta	iners, the total number of contain	ers and their regist	flight number (aircraft) or name (ship) is to ration number and where there is a serial numb signor must inform the border inspection post		
	Box reference I.19: Use the appropriate H 7.02; 19.01; 21.05; 21.06.90.98; 22.02;		der the following he	eadings: 04.01; 04.02; 04.03; 04.04; 04.05; 04.0		
— В	lox reference I.20: Indicate total gross w	reight and total net weight.				
— В	ox reference I.23: For containers or box	kes, the container number and th	ne seal number (if	applicable) should be included.		
	ox reference I.28: Manufacturing plant: xport to the European Union.	introduce the approval number	of the treatment a	and/or processing establishment(s) approved t		
Part	II:					
(1) K	keep as appropriate.					
— т	he colour of the signature shall be differe	ent to that of the printing. The sa	me rule applies to s	stamps other than those embossed or waterma		
Offic	ial veterinarian					
	Name (in capital letters):		Quali	ification and title:		
	Date:		Signa	ature:		
	Stamp:					

PART 3

Model Milk-T/S

Animal Health Certificate for raw milk or dairy products for human consumption, for [transit] / [storage] (¹) (²) in the European Union

COUNTRY	1
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Veterinary certificate to EU

									101011	mary continuate to	
	I.1. Consignor					I.2.	Certificate reference No		l.2.a.		
		Name Address					I.3. Central competent authority				
Part I: Details of dispatched consignment		Tel.				I.4. Local competent authority					
	1.5.	Name Address					B. Person responsible for the load in EU Name Address				
		Postcode Tel.					Postal code Tel. N°				
	1.7.	Country of I origin	ISO code	I.8. Region of origin	Code	1.9.	Country of ISC destination) code	I.10.		/
	l.11.	Place of origin	I	<u></u>	1	I.12.	Place of destination				
		Name Approval number Address				Customs warehouse	3		Ship supplier 🔲 roval number		
а.							Address		1,661		
					Postal code						
	1.13.	Place of loading				1.14.	Date of departure				
	l.15.	Means of transport				I.16.	Entry BIP in EU				
		Aeroplane	Ship		wagon 🗌						
		Road vehicle	Other			l.17.					
	Documentation references I.18. Description of commodity										
				I.19. Commodity code (HS code)							
									I.20. Quantit	у	
	l.21.	Temperature of produ	Jot	Chilled 🗌			Frozen 🗌		I.22. Numbe	r of packages	
	1.23.	Seal/Container No							I.24. Type of	f packaging	
	I.25. Commodities certified for:							I			
	Human consumption										
	1.26.	For transit through El	U to 3rd Co	ountry		1.27.					\sim
		3rd country ISO Code								-	
	1.28.	Identification of the co	ommodities	3							
		Manufacturing plant		Number of packages			pecies tific name)	Net w	eight	Batch number	

	Health information Animal Health Attestation I, the undersigned official veterinarian, he	II.a. Certificate reference number	II.b.									
	I the undersigned official veterinarian he											
		reby certify for [transit] / [storage](²) in the European Union d	escribed above:									
	Regulation (EU) No 605/2010,	thorised for imports to the European Union of raw										
	the model certificates [Milk-RM] / [Mill	conditions for the products concerned as laid do <-RMP] / [Milk-HTB] / [Milk-HTC](²) in Part 2 of A	nnex II to Regulation (EU) No 605/2010;									
	(c) was produced on	or between	and									
Notes	s											
Part	l:											
— Во	ox reference I.7: Provide name and ISO	code of the country or part thereof as appearing	g in Annex I to Regulation (EU) No 605/2010.									
	ox reference I.11: Name, address and app ame as the country of export.	proval number of the establishment of dispatch. N	lame of the country of origin which must be the									
 Box reference I.15: Registration number (railway wagons or container and road vehicles), flight number (aircraft) or name (ship). In the case 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 Europea 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; 04.0 17.02; 19.01; 21.05; 21.06.90.98; 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 ce approved for exportation to the European Union. 					
(¹) Raw milk and dairy products means, raw milk and dairy products for human consumption in transit or storage in accordance with Article 1: or Article 13 of Council Directive 97/78/EC.												
(²) K	eep as appropriate.											
fo	or exportation to the European Union of the	milk and dairy products shall not be allowed when third country or part thereof mentioned under bean Union against imports of raw milk and dairy	I.7 and I.8, or during a period where restrictive									
- 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.												
Offici	al veterinarian											
	Name (in capital letters):	Qualif	ication and title:									
	Date:	Signa	ture:'									
	Stamp:											