II

(Non-legislative acts)

REGULATIONS

COMMISSION IMPLEMENTING REGULATION (EU) No 468/2012

of 1 June 2012

amending Regulation (EU) No 28/2012 laying down requirements for the certification for imports into and transit through the Union of certain composite products

(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 Article 8(5) thereof,

Whereas:

(1) Commission Regulation (EU) No 28/2012 [2] lays down rules on the certification of consignments of certain composite products introduced into the Union from third countries, including composite products containing processed egg products.

(2) Pursuant to Regulation (EU) No 28/2012, consignments of composite products introduced into or transited through the Union are to be accompanied by a health certificate in accordance with the models set out in Annexes I and II thereto and comply with the conditions established in that certificate.

(3) The model certificates set out in Annexes I and II to Regulation (EU) No 28/2012 do not currently include detailed conditions as regards processed egg products contained in composite products which are being introduced into or transited through the Union.

(4) Commission Regulation (EC) No 798/2008 of 8 August 2008 laying down a list of third countries, territories, zones or compartments from which poultry and poultry products may be imported into and transit through the Community and the veterinary certification requirements [3] lays down veterinary certification requirements for imports into and transit, through the Union of certain commodities, including egg products. It provides that commodities imported into or transited through the Union are to be accompanied by a veterinary certificate for the commodity concerned and comply with the conditions set out therein.

(5) Processed egg products present a potential risk for animal health, also when they are used to manufacture certain composite products. It is therefore appropriate that the same conditions which must be complied with by egg products pursuant to Regulation (EC) No 798/2008, when those products are introduced into or transited through the Union, apply also to processed egg products used to manufacture composite products.

(6) The model certificates set out in Annexes I and II to Regulation (EU) No 28/2012 include the condition that the country of origin of meat or milk products used to manufacture composite products imported into or transited through the Union is authorised by relevant Union legislation to export meat or milk products into the Union. In addition, those model certificates include the condition that the country of origin of the meat or milk products be the same as the country of export of the composite products.

(7) Those two conditions ensure that meat and milk products originating from third countries and used to manufacture composite products comply with Union

[1] OJ L 18, 23.1.2003, p. 11.
[2] OJ L 12, 14.1.2012, p. 1.
[3] OJ L 226, 23.8.2008, p. 1.

rules for human and animal health. However, the condition that the country of origin and the country of export be the same does not allow for the import into and transit through the Union of composite products exported from a third country but which contain meat and milk products originating in the Union.

(8) Meat and milk products originating in the Union are in compliance with the human and animal health conditions laid down in Union legislation. It is therefore appropriate to amend the conditions included in the model certificates set out in Annexes I and II to Regulation (EU) No 28/2012 to allow the use of meat and milk products originating in the Union to manufacture composite products in third countries authorised to export composite products to the Union.

(9) Commission Decision 2007/777/EC of 29 November 2007 laying down the animal and public health conditions and model certificates for imports of certain meat products and treated stomachs, bladders and intestines for human consumption from third countries [1] provides that Member States are to authorise imports into the Union of certain meat products that comply with the conditions concerning origin and treatment set out in Annex II thereto. That Annex sets out rules on a non-specific treatment (treatment A) to which the imported products are to be subjected when they originate in third countries where the animal health status does not present a risk for the animal health status in the Union. Since those products may be directly imported into the Union, it is appropriate to amend the conditions included in the model certificates set out in Annexes I and II to Regulation (EU) No 28/2012 to allow the use of such meat products to manufacture composite products in third countries authorised to export composite products to the Union, provided that the third country exporting the composite products ensures that those meat products comply with the health and origin requirements foreseen in Union legislation and that it is authorised to export itself the same meat products to the Union under the same conditions.

(10) 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 [2] provides that Member States are to authorise the importation of consignments of raw milk and dairy products from the third countries or parts thereof listed in column A of Annex I thereto. In addition, Regulation (EU) No 605/2010 provides that Member States are to authorise the importation of consignments of certain dairy products from the third countries or parts thereof not at risk from foot-and-mouth disease listed in column B of Annex I thereto, provided that such dairy products have undergone, or been produced from raw milk which has undergone a pasteurisation treatment involving a single heat treatment as laid down in that Regulation. Since those dairy products may be directly imported into the Union, it is appropriate to amend the conditions included in the model certificates set out in Annexes I and II to Regulation (EU) No 28/2012 to allow the use of such dairy products to manufacture composite products in third countries authorised to export composite products to the Union, provided that the third country exporting the composite products ensures that those milk products comply with the health and origin requirements foreseen in Union legislation and that it is authorised to export itself the same dairy products to the Union under the same conditions.

(11) Regulation (EU) No 28/2012 should therefore be amended accordingly.

(12) To avoid any disruption of trade, the use of certificates issued in accordance with Regulation (EU) No 28/2012 prior to the entry into force of this Regulation should be authorised for a transitional period.

(13) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on the Food Chain and Animal Health,

HAS ADOPTED THIS REGULATION:

Article 1

Annexes I and II to Regulation (EU) No 28/2012 are replaced by the text in the Annex to this Regulation.

Article 2

For a transitional period until 31 December 2012, consignments of composite products accompanied by certificates issued before 1 October 2012 in accordance with the models set out in Annexes I and II to Regulation (EU) No 28/2012 before the amendments introduced by 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*.

[1] OJ L 312, 30.11.2007, p. 49.
[2] OJ L 175, 10.7.2010, p. 1.

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

Done at Brussels, 1 June 2012.

For the Commission
The President
José Manuel BARROSO

ANNEX

'ANNEX I

Model Health Certificate for import into the European Union of composite products intended 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 Name Approval number Address 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	Nature of commodity	Net weight	Batch number

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Part II: Certification

II. **Health information**	II.a. Certificate reference No	II.b.

I, the undersigned official veterinarian/official inspector hereby certify that

II.1. I am aware of the relevant provisions of Regulations (EC) No 178/2002, (EC) No 852/2004 and (EC) No 853/2004, in particular Article 6.1(b) on the origin of the products of animal origin used in the production of the composite products described above and certify that the composite products described above were produced in accordance with those requirements, in particular that they come from (an) establishment(s) implementing a programme based on the HACCP principles in accordance with Regulation (EC) No 852/2004;

II.2. the composite products described above contain:

(1) either [II.2.A **Meat products, treated stomachs, bladders and intestines** (2) in any quantity which meet the animal health requirements in Commission Decision 2007/777/EC and contain the following meat constituents which meet the criteria indicated below:

Species (A)	Treatment (B)	Origin (C)	Approved Establishment(s) (D)

(A) Insert the code for the relevant species of meat product, treated stomachs, bladders and intestines where BOV = domestic bovine animals (*Bos taurus, Bison bison, Bubalus bubalis* and their crossbreds); OVI = domestic sheep (*Ovis aries*) and goats (*Capra hircus*); EQI = domestic equine animals (*Equus caballus, Equus asinus* and their crossbreds), POR = domestic porcine animals (*Sus scrofa*); RM = domestic rabbits, PFG = domestic poultry and farmed feathered game, RUF farmed non-domestic animals other than suidae and solipeds; RUW = wild non-domestic animals other than suidae and solipeds; SUW = wild non-domestic suidae: EQW = wild non-domestic solipeds, WL = wild lagomorphs, WGB = wild game birds.

(B) Insert A, B, C, D, E or F for the required treatment as specified and defined in Parts 2, 3 and 4 of Annex II to Decision 2007/777/EC.

(C) Insert the ISO code of the country of origin of the meat product, treated stomachs, bladders and intestines as listed in Annex II, Part 2 to Decision 2007/777/EC and, in the case of regionalization by Union legislation for the relevant meat constituents, the region as indicated in Part 1 of Annex II to Decision 2007/777/EC or a Member State of the European Union. The country of origin of the meat products must be one the following:

— the same as the country of export in box I.7,

— a Member State of the European Union,

— a third country or parts thereof authorised to export to the Union meat products treated with treatment A as set out in Annex II to Decision 2007/777/EC, where the third country where the composite product is produced is also authorised to export to the Union meat products treated with that treatment.

(D) Insert EU approval number of the establishments of origin of the meat products, treated stomachs, bladders and intestines contained in the composite product.

(E) If containing material from bovine, ovine or caprine animals, the fresh meat and/or intestines used in the preparation of the meat products and/or treated intestines shall be subject to the following conditions depending on the BSE risk category of the country of origin:

(1) (E.1) for imports from a country or a region with a negligible BSE risk as listed in Annex to Commission Decision 2007/453/EC as amended:

(1) the country or region is classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 of the European Parliament and of the Council as a country or region posing a negligible BSE risk;

(2) the animals from which the products of bovine, ovine and caprine animal origin were derived were born, continuously reared and slaughtered in the country with negligible BSE risk and passed ante-mortem and post-mortem inspections;

(1) (3) if in the country or region there have been BSE indigenous cases:

(1) (a) the animals were born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants had been enforced; or

(1) (b) the products of bovine, ovine and caprine animal origin do not contain and are not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001, or mechanically separated meat obtained from bones of bovine, ovine or caprine animals.

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[1] (E.2) for imports from a country or a region with a controlled BSE risk as listed in the Annex to Commission Decision 2007/453/EC as amended:

(1) the country or region is classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region posing a controlled BSE risk;

(2) the animals from which the products of bovine, ovine and caprine animal origin were derived passed ante-mortem and post-mortem inspections;

(3) animals from which the products of bovine, ovine and caprine animal origin destined for export were derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration after stunning of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity;

[1] [3] (4) the products of bovine, ovine and caprine animal origin do not contain and are not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001, or mechanically separated meat obtained from bones of bovine, ovine or caprine animals;

[1] [4] (5) in the case of intestines originally sourced from a country or a region with a negligible BSE risk, imports of treated intestines shall be subject to the following conditions:

(a) the country or region is classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region posing a controlled BSE risk;

(b) the animals from which the products of bovine, ovine and caprine animal origin were derived were born, continuously reared and slaughtered in the country or region with a negligible BSE risk and passed ante-mortem and post-mortem inspections;

[1] (c) if the intestines are sourced from a country or region where there have been BSE indigenous cases:

[1] (i) the animals were born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants had been enforced; or

[1] (ii) the products of bovine, ovine and caprine animal origin do not contain and are not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001.

[1] (E.3) for imports from a country or a region with an undetermined BSE risk as listed in the Annex to Commission Decision 2007/453/EC:

(1) the animals from which the products of bovine, ovine and caprine animal origin were derived have not been fed meat-and-bone meal or greaves derived from ruminants and passed ante-mortem and post-mortem inspections;

(2) the animals from which the products of bovine, ovine and caprine animal origin were derived have not been slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered by laceration after stunning of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity;

[1] [5] (3) the products of bovine, ovine and caprine animal origin are not derived from:

(i) specified risk material as defined in Annex V to Regulation (EC) No 999/2001;

(ii) nervous and lymphatic tissues exposed during the deboning process;

(iii) mechanically separated meat obtained from bones of bovine, ovine or caprine animals;

[1] [4] (4) in the case of intestines originally sourced from a country or a region with a negligible BSE risk, imports of treated intestines shall be subject to the following conditions:

(a) the country or region is classified in accordance with Article 5(2) of Regulation (EC) No 999/2001 as a country or region posing an undetermined BSE risk;

(b) the animals from which the products of bovine, ovine and caprine animal origin were derived were born, continuously reared and slaughtered in the country or region with a negligible BSE risk and passed ante-mortem and post-mortem inspections;

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(1) (c) if the intestines are sourced from a country or region where there have been BSE indigenous cases:

(1) (i) the animals were born after the date from which the ban on the feeding of ruminants with meat-and-bone meal and greaves derived from ruminants had been enforced; or

(1) (ii) the products of bovine, ovine and caprine animal origin do not contain and are not derived from specified risk material as defined in Annex V to Regulation (EC) No 999/2001.]

(1) *and/or* [II.2.B **Processed dairy products** (6) in an amount of half or more of the substance of the composite product or not shelf stable dairy products in any quantity that:

(a) have been produced in the country ..... in the establishment ..... (approval number of the establishments of origin of the dairy products contained in the composite product authorised at the time of production for export of dairy products to the EU). The country of origin of the dairy products must be one of the following:

— the same as the country of export in box I.7,

— a Member State of the European Union,

— a third country authorised to export to the Union milk and dairy products in Column A or B of Annex I to Regulation (EU) No 605/2010, where the third country where the composite product is produced is also authorised, under the same conditions, to export to the Union milk and dairy products.

The country of origin indicated in box I.7 must be listed in Annex I to Regulation (EU) No 605/2010 and the treatment applied must be conform to the treatment provided for in that list for the relevant country;

(b) have been produced from milk 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;

(c) are dairy products made from raw milk obtained from:

(1) *either* [cows, ewes, goats or buffaloes and prior to import into the territory of the European Union have undergone or been produced from raw milk which has undergone

(1) *either* [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;]

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

(1) *or* [an ultra high temperature (UHT) treatment at not less than 135 °C in combination with a suitable holding time;]

(1) *or* [a high temperature short time pasteurisation treatment (HTST) at 72 °C for 15 seconds, or a treatment with an equivalent pasteurisation effect, applied to milk with a pH lower than 7,0 achieving, where applicable, a negative reaction to an alkaline phosphatase test

(1) *or* [a high temperature short time pasteurisation treatment (HTST) at 72 °C for 15 seconds, or a treatment with an equivalent pasteurisation effect, 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, immediately followed by

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

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

(1) *or* [animals other than cows, ewes, goats or buffaloes and prior to import into the territory of the European Union have undergone or been produced from raw milk which has undergone

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(1) *either* [a sterilisation process, to achieve an F_0 value equal to or greater than three;]

(1) *or* [an ultra high temperature (UHT) treatment at not less than 135 °C in combination with a suitable holding time;]]

(d) were produced on ..... or between ..... and ..... (7).]

(1) *and/or* [II.2.C **Processed fishery products** that originate from the approved establishment No (8) ..... situated in the country (9) .....]

(1) *and/or* [II.2.D **Processed egg products** that originate from the approved country (9) .....]

were produced from eggs coming from an establishment which satisfies the requirements of Section X of Annex III to Regulation (EC) No 853/2004 which at the date of issue of the certificate is free from highly pathogenic avian influenza as defined in Regulation (EC) No 798/2008 and

either

(1) II.2.D.1 [within a 10 km radius of which [including, where appropriate, the territory of a neighbouring country,] there has been no outbreak of highly pathogenic avian influenza or Newcastle disease for at least the previous 30 days.]

or

(1) II.2.D.2 [the egg products were processed:

(1) *either* [liquid egg white was treated:

(1) *either* [with 55,6 °C for 870 seconds.]

(1) *or* [with 56,7 °C for 232 seconds.]

(1) *or* [10 % salted yolk was treated with 62,2 °C for 138 seconds.]

(1) *or* [dried egg white was treated:

(1) *either* [with 67 °C for 20 hours.]

(1) *or* [with 54,4 °C for 513 hours.]

(1) *or* [whole eggs were at least treated:

(1) *either* [with 60 °C for 188 seconds.]

(1) *or* [completely cooked.]

[whole egg blends were at least treated]:

(1) *either* [with 60 °C for 188 seconds.]

(1) *or* [with 61,1 °C for 94 seconds.]

Notes

Part I:

— Box reference I.7: Insert the ISO code of the country of origin of the composite product containing meat product, treated stomachs, bladders and intestines as listed in Annex II, Part 2 to Decision 2007/777/EC and/or for processed dairy products in Annex I to Commission Regulation (EU) No 605/2010 and/or for processed fishery products in Annex I and II to Commission Decision 2006/766/EC and/or for processed egg products in Annex I part 1 to Commission Regulation (EC) No 798/2008.

— Box reference I.11: Name, address and registration/approval number if available of the establishments of production of the composite product(s). Name of the country of origin which must be the same as the country of origin in box I.7.

— 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 of the World Customs Organisation such as: 16.01; 16.02; 16.03; 16.04; 16.05; 19.01; 19.02; 19.05; 20.04; 20.05; 21.03; 21.04; 21.05; 21.06.

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

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

— Box reference I.28: *Manufacturing plant*: insert the name and approval number if available of the establishments of production of the composite product(s). Nature of commodity: in case of composite products containing meat products, treated stomachs, bladders and intestines indicate "meat product", "treated stomachs", "bladders" or "intestines". In case of composite product containing dairy products indicate "dairy product". In case of composite product containing processed fishery products specify whether aquaculture or wild origin. In case of composite product containing egg products specify the egg content percentage.

Part II:

([1]) Keep as appropriate.

([2]) Meat products as laid down in point 7.1 of Annex I to Regulation (EC) No 853/2004 and treated stomachs, bladders and intestines as laid down in point 7.9 of Annex I to Regulation (EC) No 853/2004 that have undergone one of the treatments laid down in Annex II part 4 to Decision 2007/777/EC.

([3]) By way of derogation from point 4, carcasses, half carcasses or half carcasses cut into no more than three wholesale cuts, and quarters containing no specified risk material other than the vertebral column, including dorsal root ganglia, may be imported.

When removal of the vertebral column is not required, carcasses or wholesale cuts of carcasses of bovine animals containing vertebral column shall be identified by a blue stripe on the label referred to in Regulation (EC) No 1760/2000.

The number of bovine carcasses or wholesale cuts of carcasses, from which removal of the vertebral column is required as well as the number where removal of the vertebral column is not required shall be added to the document referred to in Article 2(1) of Regulation (EC) No 136/2004 in case of imports.

([4]) Only applicable to imports of treated intestines.

([5]) By way of derogation from point 3, carcasses, half carcasses or half carcasses cut into no more than three wholesale cuts, and quarters containing no specified risk material other than the vertebral column, including dorsal root ganglia, may be imported.

When removal of the vertebral column is not required, carcasses or wholesale cuts of carcasses of bovine animals containing vertebral column shall be identified by a clearly visible blue stripe on the label referred to in Regulation (EC) No 1760/2000.

Specific information on the number of bovine carcasses or wholesale cuts of carcasses, from which removal of the vertebral column is required and from which removal of the vertebral column is not required shall be added to the document referred to in Article 2(1) of Regulation (EC) No 136/2004 in case of imports.

([6]) Raw milk and dairy products means, raw milk and dairy products for human consumption as defined in point 7.2 of Annex I to Regulation (EC) No 853/2004.

([7]) Date or dates of production. Imports of raw milk and dairy 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 and dairy products from this third country or part thereof.

([8]) Number of the fishery product establishment authorised to export to the EU.

([9]) Country of origin authorised to export to the EU.

([10]) In case of composite products containing only egg or fishery products the signature of an official Inspector can be accepted.

— 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.

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II. **Health information**	II.a. Certificate reference No	II.b.
Official veterinarian/Official inspector ([10])		
Name (in capital letters):	Qualification and title:	
Date:	Signature:	
Stamp:		

ANNEX II

Model Health Certificate for transit through or storage in the European Union of composite products intended 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. Person responsible for the load in EU
Name
Address
Postcode
Tel.

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
Name Approval number
Address
Name Approval number
Address

I.12. Place of origin
Custom warehouse ☐ Ship supplier ☐
Name Approval number
Address
Postcode

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. For transit through EU to third country ☐
Third country ISO code

I.27.

I.28. Identification of the commodities
Manufacturing plant | Number of packages | Nature of commodity | Net weight | Batch number

COUNTRY

Composite products intended for human consumption
Transit/Storage

Part II: Certification

II. **Health information**	II.a. Certificate reference No	II.b.

I, the undersigned official veterinarian/official inspector hereby certify that the composite products described above contain:

(1) either II.1.A Meat products, treated stomachs, bladders and intestines (2) in any quantity and such meat products, treated stomachs, bladders and intestines have been produced according to Commission Decision 2007/777/EC and contain the following meat constituents and meet the criteria indicated below:

Species (A)	Treatment (B)	Origin (C)

(A) Insert the code for the relevant species of meat product, treated stomachs, bladders and intestines where BOV = domestic bovine animals (*Bos taurus, Bison bison, Bubalus bubalis* and their crossbreds); OVI = domestic sheep (*Ovis aries*) and goats (*Capra hircus*); EQI = domestic equine animals (*Equus caballus, Equus asinus* and their crossbreds), POR = domestic porcine animals (*Sus scrofa*); RM = domestic rabbits, PFG = domestic poultry and farmed feathered game, RUF farmed non-domestic animals other than suidae and solipeds; RUW = wild non-domestic animals other than suidae and solipeds; SUW = wild non-domestic suidae: EQW = wild non-domestic solipeds, WL = wild lagomorphs, WGB = wild game birds.

(B) Insert A, B, C, D, E or F for the required treatment as specified and defined in Parts 2, 3 and 4 of Annex II to Decision 2007/777/EC.

(C) Insert the ISO code of the country of origin of the meat product, treated stomachs, bladders and intestines as listed in Annex II, Part 2 to Decision 2007/777/EC and, in the case of regionalization by Union legislation for the relevant meat constituents, the region as indicated in Part 1 of Annex II to Decision 2007/777/EC or a Member State of the European Union. The country of origin of the meat products must be one the following:

— the same as the country of export in box I.7,

— a Member State of the European Union,

— a third country or parts thereof authorised to export to the Union meat products treated with treatment A as set out in Annex II to Decision 2007/777/EC, where the third country where the composite product is produced is also authorised to export to the Union meat products treated with that treatment.

(1) and/or [II.1.B Processed dairy products (3) in an amount of half or more of the substance of the composite product or not shelf stable dairy products in any quantity that

(a) have been produced in the country ..... The country of origin of the dairy products must be one of the following:

— the same as the country of export in box I.7,

— a Member State of the European Union,

— a third country or parts thereof authorised to export to the Union meat products treated with treatment A as set out in Annex II to Decision 2007/777/EC, where the third country where the composite product is produced is also authorised to export to the Union meat products treated with that treatment.

The country of origin indicated in box I.7 must be listed in Annex I to Regulation (EU) No 605/2010 and the treatment applied must be conform to the treatment provided for in that list for the relevant country;

(b) have been produced from milk 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;

(c) are dairy products made from raw milk obtained from

(1) either [cows, ewes, goats or buffaloes and prior to import into the territory of the European Union have undergone or been produced from raw milk which has undergone

(1) either [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;]

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

(1) *or* [an ultra high temperature (UHT) treatment at not less than 135 °C in combination with a suitable holding time;]

(1) *or* [a high temperature short time pasteurisation treatment (HTST) at 72 °C for 15 seconds, or a treatment with an equivalent pasteurisation effect, applied to milk with a pH lower than 7,0 achieving, where applicable, a negative reaction to an alkaline phosphatase test];

(1) *or* [a high temperature short time pasteurisation treatment (HTST) at 72 °C for 15 seconds, or a treatment with an equivalent pasteurisation effect, 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, immediately followed by

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

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

(1) *or* [animals other than cows, ewes, goats or buffaloes and prior to import into the territory of the European Union have undergone or been produced from raw milk which has undergone

(1) *either* [a sterilisation process, to achieve an F_0 value equal to or greater than three;]

(1) *or* [an ultra high temperature (UHT) treatment at not less than 135 °C in combination with a suitable holding time;]]

(d) were produced on ..... or between ..... and .....(4).]

and/or [II.1.C Processed egg products that originate from the approved country (5)

Were produced from eggs coming from an establishment which satisfies the requirements of Section X of Annex III to Regulation (EC) No 853/2004 which at the date of issue of the certificate is free from highly pathogenic avian influenza as defined in Regulation (EC) No 798/2008 and

either

(1) *[II.1.C.1 [within a 10 km radius of which [including, where appropriate, the territory of a neighbouring country,] there has been no outbreak of highly pathogenic avian influenza or Newcastle disease for at least the previous 30 days.]*

or

(1) *[II.1.C.2 [the egg products were processed:*

(1) *either* [liquid egg white was treated:

(1) *either* [with 55,6 °C for 870 seconds.]

(1) *or* [with 56,7 °C for 232 seconds.]

(1) *or* [10 % salted yolk was treated with 62,2 °C for 138 seconds.]

(1) *or* [dried egg white was treated:

(1) *either* [with 67 °C for 20 hours.]

(1) *or* [with 54,4 °C for 513 hours.]

(1) *or* [whole eggs were at least treated:

(1) *either* [with 60 °C for 188 seconds.]

(1) *or* [completely cooked.]

[whole egg blends were at least treated]:

(1) *either* [with 60 °C for 188 seconds.]

(1) *or* [with 61,1 °C for 94 seconds.]

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

Notes

Part I:

— Box reference I.7: Insert the ISO code of the country of origin of the meat product, treated stomachs, bladders and intestines as listed in Annex II, Part 2 to Decision 2007/777/EC and/or for processed dairy product in Annex I to Commission Regulation (EU) No 605/2010.

— Box reference I.11: Name, address of the establishments of production of the composite product(s). Name of the country of origin which must be the same as the country of origin in box I.7.

Approval number is not applicable.

— 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 of the World Customs Organisation such as: 16.01; 16.02; 16.03; 16.04; 16.05; 19.01; 19.02; 19.05; 20.04; 20.05; 21.03; 21.04; 21.05; 21.06.

— 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) must be included.

— Box reference I.28: *Manufacturing plant*: insert the name and approval number if available of the establishments of production of the composite product(s). Nature of commodity: in case of composite products containing meat products, treated stomachs, bladders and intestines indicate "meat product", "treated stomachs", "bladders" or "intestines". In case of composite product containing dairy products indicate "dairy product".

Part II:

(1) Keep as appropriate.

(2) Meat products as laid down in point 7.1 of Annex I to Regulation (EC) No 853/2004 and treated stomachs, bladders and intestines as laid down in point 7.9 of Annex I to Regulation (EC) No 853/2004 that have undergone one of the treatments laid down in Annex II part 4 to Decision 2007/777/EC.

(3) Raw milk and dairy products means, raw milk and dairy products for human consumption as defined in point 7.2 of Annex I to Regulation (EC) No 853/2004.

(4) Date or dates of production. Imports of raw milk and dairy 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 and dairy products from this third country or part thereof.

(5) Country of origin authorised to export to the EU.

— The colour of the signature shall be different to that of the printing. The same rule applies to stamps other than those embossed or watermark.

Official veterinarian/Official inspector

Name (in capital letters): Qualification and title:

Date: Signature:'

Stamp: