

**COUNTRY**

**Official certificate to the EU**

<b>Part I : Details of consignment</b>	I.1. Consignor Name Address Country ISO Code		I.2. Certificate reference I.3. Central competent authority I.4. Local competent authority		I.2.a. IMSOC reference <b>Specimen not to be used for imports into the EU</b>		
	I.5. Consignee Name Address Country ISO Code			I.6. Responsible for the consignment in EU Name Address Country ISO Code			
	I.7. Country of origin ISO Code		I.8. Region of origin Code		I.9. Country of destination ISO Code		I.10. Region of destination Code
	I.11. Place of dispatch Name Address Approval Number Country ISO Code			I.12. Place of destination Name Address Approval Number Country ISO Code			
	I.13. Place of loading Name Address Approval Number Country ISO Code			I.14. Date and time of departure			
	I.15. Means of Transport Mode International transport document Identification			I.16. Entry BCP Authority Country			
	I.18. Transport conditions Ambient <input type="checkbox"/> Frozen <input type="checkbox"/> Chilled <input type="checkbox"/>			I.17. Accompanying documents Type Number			
	I.19. Container No / Seal No						
	I.20. Certified as Circu s/exh iberitio n <input type="checkbox"/> Appr oved Bodi es <input type="checkbox"/> Quar antin e <input type="checkbox"/> Man ufact ure of petfo od <input type="checkbox"/> Anim al Feedi ngstu ff <input type="checkbox"/> Phar mace utical use <input type="checkbox"/> Tech nical Use <input type="checkbox"/> Relay ing <input type="checkbox"/> Cann ing indus try <input type="checkbox"/> Slaug hter <input type="checkbox"/> Othe r <input type="checkbox"/> Trad e samp les <input type="checkbox"/> Petfo od <input type="checkbox"/> Pets <input type="checkbox"/> Regis tered equi dae <input type="checkbox"/> Fatte ning <input type="checkbox"/> Bree ding/ prod uctio n <input type="checkbox"/> Gam e resto cking <input type="checkbox"/> Furth er proc ess <input type="checkbox"/> Artifi cial repro ductio n <input type="checkbox"/> Hum an Cons umpt ion <input type="checkbox"/> Prod uctio n of petfo od <input type="checkbox"/>						
	I.21. For transit Non-EU ISO Code			I.22. For internal market			I.23. For re-entry
I.24. Total number of packages		I.25. Total net weight		I.26. Total gross weight			
I.27. Description of consignment <b>1. 19 PREPARATIONS OF CEREALS, FLOUR, STARCH OR MILK; PASTRYCOOKS' PRODUCTS</b> <b>1904</b> Prepared foods obtained by the swelling or roasting of cereals or cereal products (for example, corn flakes); cereals (other than maize (corn)) in grain form or in the form of flakes or other worked grains (except flour, groats and meal), pre-cooked or otherwise prepared, not elsewhere specified or included <b>190490</b> Other <b>19049010</b> Obtained from rice							
Commodity		Manufacturing plant	Package count	Net weight	Batch number		

II. Health information		
I, the undersigned official veterinarian/official inspector hereby certify that		
II.1	I am aware of the relevant provisions of Regulations (EC) 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 <input type="checkbox"/> 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)
_____	_____	_____
—	—	—
_____	_____	_____
—	—	—
_____	_____	_____
—	—	—
_____	_____	_____
—	—	—

Part II : Certification

SPECIMEN

II. Health information			
Part II : Certification	(A)	Insert the code for the relevant species of meat product, treated stomachs, bladders and intestines where BOV = domestic bovine animals ( <i>Bos taurus</i> , <i>Bison bison</i> , <i>Bubalus bubalis</i> and their crossbreds); OVI = domestic sheep ( <i>Ovis aries</i> ) and goats ( <i>Capra hircus</i> ); EQI = domestic equine animals ( <i>Equus caballus</i> , <i>Equus asinus</i> and their crossbreds), POR = domestic porcine animals ( <i>Sus scrofa</i> ); 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. The country of origin of the meat products must be the same as the country of export in box 1.7.	
	(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)	<input type="checkbox"/> (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)	(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 negligible BSE risk;	
(1)	(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)	<input type="checkbox"/> (3)	if in the country or region there have been BSE indigenous cases:	
(1)	(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)	(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.	
(1)	<input type="checkbox"/> (E.2)	for imports from a country or a region with a controlled BSE risk as listed in Annex to Commission Decision 2007/453/EC as amended:	
(1)	(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;	
(1)	(2)	the animals from which the products of bovine, ovine and caprine animal origin were derived passed ante-mortem and post-mortem inspections;	
(1)	(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)	<input type="checkbox"/> (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.	

II. Health information			
Part II : Certification	(1)(4)	<input type="checkbox"/> (5)	<p>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:</p> <p>(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;</p> <p>(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;</p>
	(1)	<input type="checkbox"/> (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)	<input type="checkbox"/> (E.3)	for imports from a country or a region with an undetermined BSE risk as listed in 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)	<input type="checkbox"/> (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)	<input type="checkbox"/> (4)	<p>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:</p> <p>(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;</p> <p>(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;</p>
	(1)	<input type="checkbox"/> (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	

II. Health information			
Part II : Certification	(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	<input type="checkbox"/> 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 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 the same as the country of export in box 1.7)
			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 is 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 at 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 F0 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 a 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 a 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 F0 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	<input type="checkbox"/> II.2.C	Processed fishery products that originate from the approved establishment N°(8) _____ situated in the country(9) _____	
(1)and/or	<input type="checkbox"/> II.2.D	Processed egg products that originate from the approved country(9) _____	
Notes			

Part II : Certification	II. Health information		
	<p>Part I :</p> <ul style="list-style-type: none"> <li>• 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.</li> <li>• 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 1.7.</li> <li>• 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.</li> <li>• Box reference I.19: Use the appropriate Harmonised System (HS) code of the World Customs Organisation: codes of the following headings: 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.</li> <li>• Box reference 20: Indicate total gross weight and total net weight.</li> <li>• Box reference I.23: for containers or boxes, the container number and the seal number (if applicable) must be included.</li> <li>• 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.</li> </ul> <p>Part II:</p> <ol style="list-style-type: none"> <li>(1) Keep as appropriate.</li> <li>(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.</li> <li>(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.</li> <li>(4) Only applicable to imports of treated intestines.</li> <li>(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.</li> </ol>		

Part II : Certification	II. Health information			
	(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.		
	Official veterinarian or Official inspector			
	Name (in capital letters)		Qualification and title	
	Date of signature		Signature	
Stamp				