

▼ **M1**

## 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 Address Approval number Name Address Approval number Name Address Approval number			I.12. Place of origin  Custom warehouse <input type="checkbox"/> Ship supplier <input type="checkbox"/> Name Address Approval number Postcode			
	I.13. Place of loading			I.14. Date of departure			
	I.15. Means of transport Aeroplane <input type="checkbox"/> Ship <input type="checkbox"/> Railway wagon <input type="checkbox"/> Road vehicle <input type="checkbox"/> Other <input type="checkbox"/> 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 <input type="checkbox"/> Chilled <input type="checkbox"/> Frozen <input type="checkbox"/>						I.22. Number of packages	
I.23. Seal/Container No						I.24. Type of packaging	
I.25. Commodities certified for: Human consumption <input type="checkbox"/>							
I.26. For transit through EU to third country <input type="checkbox"/>			I.27.				
Third country			ISO code				
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.						
	<p>I, the undersigned official veterinarian/official inspector hereby certify that the composite products described above contain:</p> <p>(<sup>1</sup>) either II.1.A Meat products, treated stomachs, bladders and intestines (<sup>2</sup>) 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:</p> <table border="1"> <thead> <tr> <th>Species (A)</th> <th>Treatment (B)</th> <th>Origin (C)</th> </tr> </thead> <tbody> <tr> <td colspan="3"> <p>(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 crossbreeds); 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 crossbreeds), 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.</p> <p>(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.</p> <p>(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:</p> <ul style="list-style-type: none"> <li>— the same as the country of export in box I.7,</li> <li>— a Member State of the European Union,</li> <li>— 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.</li> </ul> </td> </tr> </tbody> </table>	Species (A)	Treatment (B)	Origin (C)	<p>(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 crossbreeds); 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 crossbreeds), 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.</p> <p>(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.</p> <p>(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:</p> <ul style="list-style-type: none"> <li>— the same as the country of export in box I.7,</li> <li>— a Member State of the European Union,</li> <li>— 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.</li> </ul>				
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	<p>(<sup>1</sup>) and/or II.1.B Processed dairy products (<sup>3</sup>) in an amount of half or more of the substance of the composite product or not shelf stable dairy products in any quantity that</p> <p>(a) have been produced in the country ..... The country of origin of the dairy products must be one of the following:</p> <ul style="list-style-type: none"> <li>— the same as the country of export in box I.7,</li> <li>— a Member State of the European Union,</li> <li>— 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.</li> </ul> <p>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;</p> <p>(b) have been produced from milk obtained from animals:</p> <ul style="list-style-type: none"> <li>(i) under the control of the official veterinary service;</li> <li>(ii) belonging to holdings which were not under restrictions due to foot-and-mouth disease or rinderpest; and</li> <li>(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;</li> </ul> <p>(c) are dairy products made from raw milk obtained from</p> <p>(<sup>1</sup>) 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</p> <p>(<sup>1</sup>) 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;]</p>								



## COUNTRY

Composite products intended for human consumption  
Transit/Storage

II. Health information	II.a. Certificate reference number	II.b.
<p>(<sup>1</sup>) or [a sterilisation process, to achieve an F<sub>0</sub> value equal to or greater than three;]</p> <p>(<sup>1</sup>) or [an ultra high temperature (UHT) treatment at not less than 135 °C in combination with a suitable holding time;]</p> <p>(<sup>1</sup>) 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;]</p> <p>(<sup>1</sup>) 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</p> <p>(<sup>1</sup>) either [lowering the pH below 6 for one hour;]</p> <p>(<sup>1</sup>) or [additional heating equal to or greater than 72 °C, combined with desiccation;]</p> <p>(<sup>1</sup>) 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</p> <p>(<sup>1</sup>) either [a sterilisation process, to achieve an F<sub>0</sub> value equal to or greater than three;]</p> <p>(<sup>1</sup>) or [an ultra high temperature (UHT) treatment at not less than 135 °C in combination with a suitable holding time;]</p> <p>(d) were produced on ..... or between ..... and .....(<sup>4</sup>.)</p>		
<p>and/or III.1.C Processed egg products that originate from the approved country (<sup>5</sup>)</p> <p>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</p> <p>either</p> <p>(<sup>1</sup>) [III.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.]</p> <p>or</p> <p>(<sup>1</sup>) [III.1.C.2 [the egg products were processed:</p> <p>(<sup>1</sup>) either [liquid egg white was treated:</p> <p>(<sup>1</sup>) either [with 55,6 °C for 870 seconds.]</p> <p>(<sup>1</sup>) or [with 56,7 °C for 232 seconds.]</p> <p>(<sup>1</sup>) or [10 % salted yolk was treated with 62,2 °C for 138 seconds.]</p> <p>(<sup>1</sup>) or [dried egg white was treated:</p> <p>(<sup>1</sup>) either [with 67 °C for 20 hours.]</p> <p>(<sup>1</sup>) or [with 54,4 °C for 513 hours.]</p> <p>(<sup>1</sup>) or [whole eggs were at least treated:</p> <p>(<sup>1</sup>) either [with 60 °C for 188 seconds.]</p> <p>(<sup>1</sup>) or [completely cooked.]</p> <p>[whole egg blends were at least treated]:</p> <p>(<sup>1</sup>) either [with 60 °C for 188 seconds.]</p> <p>(<sup>1</sup>) or [with 61,1 °C for 94 seconds.]</p>		



## COUNTRY

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II. Health information	II.a. Certificate reference number	II.b.						
<p><b>Notes</b></p> <p><b>Part I:</b></p> <ul style="list-style-type: none"> <li>— 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.</li> <li>— 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.</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 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.</li> <li>— Box reference I.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: <i>Manufacturing plant</i>: 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".</li> </ul> <p><b>Part II:</b></p> <ul style="list-style-type: none"> <li>(<sup>1</sup>) Keep as appropriate.</li> <li>(<sup>2</sup>) 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>(<sup>3</sup>) 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.</li> <li>(<sup>4</sup>) 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.</li> <li>(<sup>5</sup>) Country of origin authorised to export to the EU.</li> </ul> <p>— 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.</p>								
<p>Official veterinarian/Official inspector</p> <table style="width: 100%; border: none;"> <tr> <td style="width: 60%; border: none;">Name (in capital letters):</td> <td style="width: 40%; border: none;">Qualification and title:</td> </tr> <tr> <td style="border: none;">Date:</td> <td style="border: none;">Signature:</td> </tr> <tr> <td style="border: none;">Stamp:</td> <td style="border: none;"></td> </tr> </table>			Name (in capital letters):	Qualification and title:	Date:	Signature:	Stamp:	
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