

## Part I : Details of consignment

I.1. Consignor Name _____ Address _____ Country _____ ISO Code _____			I.2. IMSOC Reference I.2.a. Local Reference _____																		
I.5. Consignee Name _____ Address _____ Country _____ ISO Code _____			I.3. Central competent authority I.4. Local competent authority _____																		
I.7. Country of origin _____		ISO Code _____	I.9. Country of destination _____		ISO Code _____																
I.8. Region of origin _____		Code _____	<del>I.10. Region of destination _____</del>																		
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 <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 20%;">Mode</td> <td style="width: 20%;">International transport document</td> <td style="width: 60%;">Identification</td> </tr> <tr><td> </td><td> </td><td> </td></tr> <tr><td> </td><td> </td><td> </td></tr> <tr><td> </td><td> </td><td> </td></tr> <tr><td> </td><td> </td><td> </td></tr> </table>			Mode	International transport document	Identification													I.16 Entry Point _____			
Mode	International transport document	Identification																			
I.18. Transport conditions Frozen <input type="checkbox"/> Chilled <input type="checkbox"/> Controlled temperature <input type="checkbox"/> Ambient <input type="checkbox"/>			I.17. Accompanying documents Document Type _____ Accompanying document reference _____ Date of Issue _____ Country _____ Place of issue _____																		
I.19. Container No / Seal No _____																					
I.20. Certified as <table style="width: 100%;"> <tr> <td>Other <input type="checkbox"/></td> <td>Pharmaceutical use <input type="checkbox"/></td> <td>Technical use <input type="checkbox"/></td> <td>Animal Feedingstuff <input type="checkbox"/></td> </tr> <tr> <td>Breeding and production <input type="checkbox"/></td> <td>Production <input type="checkbox"/></td> <td>Fattening <input type="checkbox"/></td> <td>Production of petfood <input type="checkbox"/></td> </tr> <tr> <td>Human consumption <input type="checkbox"/></td> <td>Relaying <input type="checkbox"/></td> <td>Artificial reproduction <input type="checkbox"/></td> <td>Breeding <input type="checkbox"/></td> </tr> <tr> <td>Slaughter <input type="checkbox"/></td> <td colspan="3"></td> </tr> </table>						Other <input type="checkbox"/>	Pharmaceutical use <input type="checkbox"/>	Technical use <input type="checkbox"/>	Animal Feedingstuff <input type="checkbox"/>	Breeding and production <input type="checkbox"/>	Production <input type="checkbox"/>	Fattening <input type="checkbox"/>	Production of petfood <input type="checkbox"/>	Human consumption <input type="checkbox"/>	Relaying <input type="checkbox"/>	Artificial reproduction <input type="checkbox"/>	Breeding <input type="checkbox"/>	Slaughter <input type="checkbox"/>			
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Slaughter <input type="checkbox"/>																					
I.21. For transit through a third country <input type="checkbox"/> <table style="width: 100%;"> <tr> <td>Country _____</td> <td>ISO Code _____</td> </tr> <tr> <td>EU Exit Authority _____</td> <td>BCP code _____</td> </tr> <tr> <td>EU Entry Authority _____</td> <td>BCP code _____</td> </tr> </table>			Country _____	ISO Code _____	EU Exit Authority _____	BCP code _____	EU Entry Authority _____	BCP code _____	I.22. For transit through Member State(s) <input type="checkbox"/> <table style="width: 100%;"> <tr> <td>Country _____</td> <td>ISO Code _____</td> </tr> </table>			Country _____	ISO Code _____								
Country _____	ISO Code _____																				
EU Exit Authority _____	BCP code _____																				
EU Entry Authority _____	BCP code _____																				
Country _____	ISO Code _____																				
I.23. Total number of packages _____		I.24. Total quantity _____		I.25. Total net weight _____																	
I.28. Description of consignment <b>1. 03 FISH AND CRUSTACEANS, MOLLUSCS AND OTHER AQUATIC INVERTEBRATES</b> <b>0302 Fish, fresh or chilled, excluding fish fillets and other fish meat of heading  0304</b>																					
#1.	Commodity	Quantity	Net weight	Package count																	
	Species	Identification number	Identification system																		

## II. Health information

## II.1. Public health attestation

I, the undersigned, declare that I am aware of the relevant provisions of Regulation (EC) No 178/2002 of the European Parliament and of the Council of 28 January 2002 laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety Regulation (EC) No 852/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific hygiene rules for food of animal origin and Regulation (EU) 2017/625 of the European Parliament and of the Council of 15 March 2017 on official controls and other official activities performed to ensure the application of food and feed law, rules on animal health and welfare, plant health and plant protection products, amending Regulations (EC) No 999/2001, (EC) No 396/2005, (EC) No 1069/2009, (EC) No 1107/2009, (EU) No 1151/2012, (EU) No 652/2014, (EU) 2016/429 and (EU) 2016/2031 of the European Parliament and of the Council, Council Regulations (EC) 1/2005 and (EC) No 1099/2009 and Council Directives 98/58/EC, 1999/74/EC, 2007/43/EC, 2008/119/EC and 2008/120/EC and repealing Regulations (EC) No 854/2004 and (EC) No 882/2004 of the European Parliament and of the Council, Council Directives 89/608/EEC, 89/662/EEC, 90/425/EEC, 91/496/EEC, 96/23/EC, 96/93/EC and 97/78/EC and Council Decision 92/438/EEC (Official Controls Regulation) and

I certify that the raw materials described above comply with these requirements, in particular that:

(1) ☐ [bones, hides and skins of domestic and farmed ruminant animals, pigs and poultry, as well as tendons and sinews described above are derived from animals which were slaughtered in a slaughterhouse and the carcasses of which were found to be fit for human consumption following ante- and post-mortem inspection;]

and/or

(1) ☐ [wild game hides, skins and bones described above are derived from killed animals whose carcasses have been found to be fit for human consumption following post-mortem inspection;]

and/or

(1) ☐ [fish skins and bones described above are derived from plants that manufacture fishery products for human consumption which are authorised for export;]

(1) and, if of bovine, ovine and caprine animal origin,

they have been derived from animals which passed ante-mortem and post-mortem inspections,

(1) and, except for hides and skins of ruminants,

(1) either

- [they come from a country or a region classified in accordance with Commission Decision 2007/453/EC of 29 June 2007 establishing the BSE status of Member States or third countries or regions thereof according to their BSE risk as a country or region posing a negligible BSE risk;

they do not contain and are not derived from specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001 of the European Parliament and of the Council of 22 May 2001 laying down rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies

they do not contain and are not derived from mechanically separated meat obtained from bones of bovine, ovine or caprine animals, except for raw materials derived from animals that were born, continuously reared and slaughtered in a country or region classified in accordance with Decision 2007/453/EC as a country or region posing a negligible BSE risk in which there has been no indigenous BSE cases;

the animals, from which the raw materials are derived, were not slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered, after stunning, by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, except if the animals were born, continuously reared and slaughtered in a country or region classified in accordance with Decision 2007/453/EC as a country or region posing a negligible BSE risk;

(1) ☐ [the animals, from which the raw materials are derived, originate from a country or

## II. Health information

region classified in accordance with Decision 2007/453/EC as a country or region posing an undetermined BSE risk, and the animals were not fed with meat-and-bone meal or greaves, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health];

(1) ☐ [the animals, from which the raw materials are derived, originate from a country or region classified in accordance with Decision 2007/453/EC as a country or region posing an undetermined BSE risk, and the raw materials were produced and handled in a manner which ensures that they did not contain and were not contaminated with nervous and lymphatic tissues exposed during the deboning process.]

(1) or

- [they come from a country or a region classified in accordance with Commission Decision 2007/453/EC of 29 June 2007 establishing the BSE status of Member States or third countries or regions thereof according to their BSE risk as a country or region posing a controlled BSE risk;

the animals, from which the raw materials of bovine, ovine and caprine animal origin intended for export are derived, were not killed, after stunning, by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, or by means of gas injected into the cranial cavity;

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

(1) or

- [they come from a country or a region classified in accordance with Commission Decision 2007/453/EC of 29 June 2007 establishing the BSE status of Member States or third countries or regions thereof according to their BSE risk as a country or region with an undetermined BSE risk;

the animals, from which the raw materials are derived, were not fed meat-and-bone meal or greaves derived from ruminants, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health;

the animals from which the raw materials of bovine, ovine and caprine animal origin are derived, were not killed, after stunning, by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, or by means of gas injected into the cranial cavity; the raw materials are not derived from:

- (i) specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001;
- (ii) nervous and lymphatic tissues exposed during the deboning process;
- (ii) mechanically separated meat obtained from the bones of bovine, ovine or caprine animals.]

## II.2. Animal Health Attestation (1)

I, the undersigned official veterinarian, certify that the raw materials described above:

II.2.1. consist of animal products that satisfy the animal health requirements below;

II.2.2. have been obtained in the country(ies) or region(s) thereof of (1)

either ○ [ ] (1) or ○ [ ] (2)(3)(4) from:

(1) either ○ [II.2.2.1 animals that come from holdings and have remained in that territory since birth or for at least the last three months before slaughter; and

(1) either ○ [(i) are derived from the species referred to in Commission Regulation (EU No 206/2010 of 12 March 2010 laying down lists of third countries, territories or parts thereof authorised for the introduction into the European Union of certain animals and fresh meat and the veterinary certification requirements satisfying all the relevant animal health import requirements laid down in that Regulation,

## Part II: Certification

## II. Health information

and that were slaughtered for human consumption on a date for which import into the Great Britain of fresh meat from animals of those species was authorised from the country or territory thereof in accordance with Column 8 of Part 1 of Annex 2 to that Regulation:]

- (1) or      ○ [(ii) are derived from the species referred to in Commission) Regulation (EC) No 119/2009 laying down a list of third countries or parts thereof, for imports into, or transit through, the Community of meat of wild leporidae, of certain wild land mammals and of farmed rabbits and the veterinary certification requirements, satisfying all the relevant animal health import requirements laid down in that Regulation.]]

- (1) or      ○ [II.2.2.1 poultry that have remained in that territory since hatching or have been imported as day-old chicks or slaughter poultry from (a) third country(ies) listed for that commodity in Part 1 of Annex 1 to 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, under conditions at least equivalent to those in that Regulation satisfying all the relevant animal health import requirements laid down in that Regulation and were slaughtered for human consumption on a date for which import into Great Britain of meat from animals of those species was authorised from the country or territory thereof in accordance with Column 6 B of Part 1 to Annex 1 to that Regulation.

- (1) or      ○ [II.2.2.1 animals that have been killed in the wild in that territory ( 5 ) and captured and killed in an area:
- (i) in which within 25 km there has been no case/outbreak of any of the following diseases for which the animals are susceptible: foot and mouth disease, rinderpest, Newcastle disease or highly pathogenic avian influenza during the prior 30 days, nor of classical or African swine fever during the prior 40 days, and
  - (ii) that is situated at a distance that exceeds 20 km from the borders separating another territory of a country or part thereof, which is not authorised on these dates to export these raw materials into Great Britain, and
  - (iii) in which after killing were transported within 12 hours for chilling either to a collection centre and immediately afterwards to a game-handling establishment, or directly to a game-handling establishment;]

- II.2.3. have been obtained in an establishment around which, within a radius of 10 km, there has been no case/outbreak of the following diseases that the animals are susceptible to: foot and mouth disease, rinderpest, Newcastle disease or highly pathogenic avian influenza, and classical or African swine fever during the prior 30 days or, in the event of a case of one of those diseases, the preparation of raw materials for export to Great Britain has been authorised only after the removal of all meat and the total cleaning and disinfection of the establishment under the control of an official veterinarian;

- II.2.4. have been obtained and prepared without contact with other materials that do not comply with the conditions required above, and have been handled so as to avoid contamination with pathogenic agents; and

- II.2.5. have been transported in clean and sealed containers or lorries.

## Notes

(\*) Those countries subject to the transitional import arrangements include: an EU member State; Liechtenstein; Norway; Iceland and Switzerland.

References to European Union legislation within this certificate are references to direct EU legislation which has

Part II: Certification	II. Health information			
	<p>been retained in Great Britain (retained EU law as defined in the European Union (Withdrawal) Act 2018). References to Great Britain in this certificate include Channel Islands and Isle of Man.</p>			
	Part I:			
	Box reference I.8:	provide the code of territory as appearing in Part 1 of Annex 1 to Regulation (EC) No 798/2008 and/or in Part 1 of Annex 1 to Regulation (EC) No 119/2009 and/or Part 1 of Annex 2 to Regulation (EU) No 206/2010.		
	Box reference I.16:	Do not use this box until the end of the transitional staging period.		
	Box reference I.25:	Insert the appropriate Harmonised System (HS) code(s) such as 0206, 0207, 0208, 0302, 0303, 0305, 0505, 0506, 0511 91, 0511 99, 4101, 4102 or 4103.		
	Box reference I.25:	Nature of commodity: hides, skins, bones, tendons and sinews;		
		Manufacturing plant: includes slaughterhouse, factory vessel, cutting plant, game-handling establishment and processing plant		
	Part II:			
	(1)	Delete as appropriate. In the case of products derived from fishery products, the whole section 11.2 should be deleted.		
(2)	<p>The name and ISO code number of the exporting country or territory or zone as laid down in:</p> <ul style="list-style-type: none"> <li>— the Annex 2 of Commission Delegated Regulation (EU) 2019/625 of 4 March 2019 supplementing Regulation (EU) 2017/625 of the European Parliament and of the Council with regard to requirements for the entry into the Union of consignments of certain animals and goods intended for human consumption);</li> <li>— Annex 1 to Regulation (EC) No 798/2008;</li> <li>— Part 1 of Annex 1 to Regulation (EC) No 119/2009;</li> <li>— Part 1 of Annex 2 to Regulation (EC) No 206/2010.</li> </ul>			
(3)	If parts of the materials were derived from animals originating from (an)other third country(ies) listed in Annex 2 to Regulation (EU) No 206/2010 for import of that commodity into Great Britain, then the code(s) of country(ies) or territory(ies) and of the third country slaughtering the animals shall be stated (the material cannot come from a country or territory that has supplementary guarantees A or F as indicated in column 5 of that Annex).			
(4)	If the meat comes from slaughter poultry originating from an(other) third country(ies) listed in Part 1 of Annex 1 to Regulation (EC) No 798/2008 for imports of that commodity into Great Britain, then the code(s) of country(ies) or territory(ies) and of the third country slaughtering the poultry shall be stated.			
(5)	Only for countries from where game meat intended for human consumption of the same animal species is authorised for importation into the Great Britain.			
(6)	The removal of specified risk material is not required if the raw materials derive from animals that are born, continuously reared and slaughtered in a third country or region of a third country classified in accordance with Decision 2007/453/EC as posing a negligible BSE risk			
—	The signature and the stamp must be in a different colour to that of the printing.			
NB	Note for the person responsible for the consignment in Great Britain: this certificate is only for veterinary purposes and has to accompany the consignment until it reaches the border control post. The consignment must be transported directly to the manufacturing plant of destination.			
Certifying Officer				
Name (in capital letters)		Qualification and title		
Date of signature		Signature		
Stamp				

Μέρος I

I.1. Αποστολέας Όνομα Διεύθυνση Χώρα Κωδικός ISO			I.2. Κωδικός αναφοράς IMSOC I.2.a. Local Reference	
I.5. Παραλήπτης Όνομα Διεύθυνση Χώρα Κωδικός ISO			I.3. Κεντρική αρμόδια αρχή (ΚΑΑ) I.4. Local competent authority	
I.7. Χώρα προέλευσης		Κωδικός ISO	I.9. Country of destination Κωδικός ISO	
I.8. Region of origin Κωδικός			<del>I.10. Περιφέρεια προορισμού</del>	
I.11. Place of Dispatch Όνομα Διεύθυνση Αριθμός έγκρισης Χώρα Κωδικός ISO			I.12. Τόπος προορισμού Όνομα Διεύθυνση Αριθμός έγκρισης Χώρα Κωδικός ISO	
I.13. Τόπος φόρτωσης Όνομα Διεύθυνση Αριθμός έγκρισης Χώρα Κωδικός ISO			I.14. Date and time of departure	
I.15. Μέσο μεταφοράς Τύπος Έγγραφο Ταυτοποίηση			I.16 Entry Point	
I.18. Transport conditions Κατεψυγμένα <input type="checkbox"/> Σε ψύξη <input type="checkbox"/> Controlled temperature <input type="checkbox"/> σε θερμοκρασία περιβάλλοντος <input type="checkbox"/>			I.17. Συνοδευτικά έγγραφα Document Type Κωδικός αναφοράς του εμπορικού εγγράφου Ημερομηνία έκδοσης Χώρα Τόπος έκδοσης	
I.19. Εμπορευματοκιβώτιο αριθ./ Σφραγίδα αριθ.				
I.20. Certified as Άλλο <input type="checkbox"/> Φαρμακευτική χρήση <input type="checkbox"/> Τεχνική χρήση <input type="checkbox"/> Κτηνοτροφές <input type="checkbox"/> Breeding and production <input type="checkbox"/> Production <input type="checkbox"/> Πάχυνση <input type="checkbox"/> Production of petfood <input type="checkbox"/> Κατανάλωση από τον άνθρωπο <input type="checkbox"/> Αναμετάδοση <input type="checkbox"/> Τεχνητή αναπαραγωγή <input type="checkbox"/> Breeding <input type="checkbox"/> Σφαγή <input type="checkbox"/>				
I.21. For transit through a third country <input type="checkbox"/> Country EU Exit Authority EU Entry Authority Κωδικός ISO BCP code BCP code			I.22. For transit through Member State(s) <input type="checkbox"/> Country Κωδικός ISO	
I.23. Συνολικός αριθμός δεμάτων		I.24. Συνολική ποσότητα	I.25. Συνολικό καθαρό βάρος	I.25. Συνολικό μεικτό βάρος
I.28. Description of consignment 1. 03 ΨΑΡΙΑ ΚΑΙ ΜΑΛΑΚΟΣΤΡΑΚΑ, ΜΑΛΑΚΙΑ ΚΑΙ ΑΛΛΑ ΑΣΠΟΝΔΥΛΑ ΥΔΡΟΒΙΑ 0302 Ψάρια νωπά ή διατηρημένα με απλή ψύξη, με εξαίρεση τα φιλέτα και άλλη σάρκα ψαριών της κλάσης 0304				
#1.	Εμπόρευμα	Ποσότητα	Καθαρό βάρος	Πλήθος πακέτων
Είδος		Αναγνωριστικός αριθμός	Σύστημα ταυτοποίησης	

## II. Υγειονομικές πληροφορίες

## II.1. Public health attestation

I, the undersigned, declare that I am aware of the relevant provisions of Regulation (EC) No 178/2002 of the European Parliament and of the Council of 28 January 2002 laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety Regulation (EC) No 852/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific hygiene rules for food of animal origin and Regulation (EU) 2017/625 of the European Parliament and of the Council of 15 March 2017 on official controls and other official activities performed to ensure the application of food and feed law, rules on animal health and welfare, plant health and plant protection products, amending Regulations (EC) No 999/2001, (EC) No 396/2005, (EC) No 1069/2009, (EC) No 1107/2009, (EU) No 1151/2012, (EU) No 652/2014, (EU) 2016/429 and (EU) 2016/2031 of the European Parliament and of the Council, Council Regulations (EC) 1/2005 and (EC) No 1099/2009 and Council Directives 98/58/EC, 1999/74/EC, 2007/43/EC, 2008/119/EC and 2008/120/EC and repealing Regulations (EC) No 854/2004 and (EC) No 882/2004 of the European Parliament and of the Council, Council Directives 89/608/EEC, 89/662/EEC, 90/425/EEC, 91/496/EEC, 96/23/EC, 96/93/EC and 97/78/EC and Council Decision 92/438/EEC (Official Controls Regulation) and

I certify that the raw materials described above comply with these requirements, in particular that:

(1) ☐ [bones, hides and skins of domestic and farmed ruminant animals, pigs and poultry, as well as tendons and sinews described above are derived from animals which were slaughtered in a slaughterhouse and the carcasses of which were found to be fit for human consumption following ante- and post-mortem inspection;]

and/or

(1) ☐ [wild game hides, skins and bones described above are derived from killed animals whose carcasses have been found to be fit for human consumption following post-mortem inspection;]

and/or

(1) ☐ [fish skins and bones described above are derived from plants that manufacture fishery products for human consumption which are authorised for export;]

(1) and, if of bovine, ovine and caprine animal origin,

they have been derived from animals which passed ante-mortem and post-mortem inspections,

(1) and, except for hides and skins of ruminants,

(1) either

- [they come from a country or a region classified in accordance with Commission Decision 2007/453/EC of 29 June 2007 establishing the BSE status of Member States or third countries or regions thereof according to their BSE risk as a country or region posing a negligible BSE risk;

they do not contain and are not derived from specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001 of the European Parliament and of the Council of 22 May 2001 laying down rules for the prevention, control and eradication of certain transmissible spongiform encephalopathies

they do not contain and are not derived from mechanically separated meat obtained from bones of bovine, ovine or caprine animals, except for raw materials derived from animals that were born, continuously reared and slaughtered in a country or region classified in accordance with Decision 2007/453/EC as a country or region posing a negligible BSE risk in which there has been no indigenous BSE cases;

the animals, from which the raw materials are derived, were not slaughtered after stunning by means of gas injected into the cranial cavity or killed by the same method or slaughtered, after stunning, by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, except if the animals were born, continuously reared and slaughtered in a country or region classified in accordance with Decision 2007/453/EC as a country or region posing a negligible BSE risk;

(1) ☐ [the animals, from which the raw materials are derived, originate from a country or

## II. Υγειονομικές πληροφορίες

region classified in accordance with Decision 2007/453/EC as a country or region posing an undetermined BSE risk, and the animals were not fed with meat-and-bone meal or greaves, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health];

(1) ☐ [the animals, from which the raw materials are derived, originate from a country or region classified in accordance with Decision 2007/453/EC as a country or region posing an undetermined BSE risk, and the raw materials were produced and handled in a manner which ensures that they did not contain and were not contaminated with nervous and lymphatic tissues exposed during the deboning process.]

(1) or

- [they come from a country or a region classified in accordance with Commission Decision 2007/453EC of 29 June 2007 establishing the BSE status of Member States or third countries or regions thereof according to their BSE risk as a country or region posing a controlled BSE risk;

the animals, from which the raw materials of bovine, ovine and caprine animal origin intended for export are derived, were not killed, after stunning, by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, or by means of gas injected into the cranial cavity;

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

(1) or

- [they come from a country or a region classified in accordance with Commission Decision 2007/453EC of 29 June 2007 establishing the BSE status of Member States or third countries or regions thereof according to their BSE risk as a country or region with an undetermined BSE risk;

the animals, from which the raw materials are derived, were not fed meat-and-bone meal or greaves derived from ruminants, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health;

the animals from which the raw materials of bovine, ovine and caprine animal origin are derived, were not killed, after stunning, by laceration of central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, or by means of gas injected into the cranial cavity; the raw materials are not derived from:

- (i) specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001;
- (ii) nervous and lymphatic tissues exposed during the deboning process;
- (ii) mechanically separated meat obtained from the bones of bovine, ovine or caprine animals.]

## II.2. Animal Health Attestation (1)

I, the undersigned official veterinarian, certify that the raw materials described above:

II.2.1. consist of animal products that satisfy the animal health requirements below;

II.2.2. have been obtained in the country(ies) or region(s) thereof of (1)

either ○ [ ] (1) or ○ [ ] (2)(3)(4) from:

(1) either ○ [II.2.2.1 animals that come from holdings and have remained in that territory since birth or for at least the last three months before slaughter; and

(1) either ○ [(i) are derived from the species referred to in Commission Regulation (EU No 206/2010 of 12 March 2010 laying down lists of third countries, territories or parts thereof authorised for the introduction into the European Union of certain animals and fresh meat and the veterinary certification requirements satisfying all the relevant animal health import requirements laid down in that Regulation,



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and that were slaughtered for human consumption on a date for which import into the Great Britain of fresh meat from animals of those species was authorised from the country or territory thereof in accordance with Column 8 of Part 1 of Annex 2 to that Regulation:]

(1) or ○ [(ii) are derived from the species referred to in Commission) Regulation (EC) No 119/2009 laying down a list of third countries or parts thereof, for imports into, or transit through, the Community of meat of wild leporidae, of certain wild land mammals and of farmed rabbits and the veterinary certification requirements, satisfying all the relevant animal health import requirements laid down in that Regulation.]]

(1) or ○ [II.2.2.1 poultry that have remained in that territory since hatching or have been imported as day-old chicks or slaughter poultry from (a) third country(ies) listed for that commodity in Part 1 of Annex 1 to 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, under conditions at least equivalent to those in that Regulation satisfying all the relevant animal health import requirements laid down in that Regulation and were slaughtered for human consumption on a date for which import into Great Britain of meat from animals of those species was authorised from the country or territory thereof in accordance with Column 6 B of Part 1 to Annex 1 to that Regulation.

(1) or ○ [II.2.2.1 animals that have been killed in the wild in that territory ( 5 ) and captured and killed in an area:

- (i) in which within 25 km there has been no case/outbreak of any of the following diseases for which the animals are susceptible: foot and mouth disease, rinderpest, Newcastle disease or highly pathogenic avian influenza during the prior 30 days, nor of classical or African swine fever during the prior 40 days, and
- (ii) that is situated at a distance that exceeds 20 km from the borders separating another territory of a country or part thereof, which is not authorised on these dates to export these raw materials into Great Britain, and
- (iii) in which after killing were transported within 12 hours for chilling either to a collection centre and immediately afterwards to a game-handling establishment, or directly to a game-handling establishment;]

II.2.3. have been obtained in an establishment around which, within a radius of 10 km, there has been no case/outbreak of the following diseases that the animals are susceptible to: foot and mouth disease, rinderpest, Newcastle disease or highly pathogenic avian influenza, and classical or African swine fever during the prior 30 days or, in the event of a case of one of those diseases, the preparation of raw materials for export to Great Britain has been authorised only after the removal of all meat and the total cleaning and disinfection of the establishment under the control of an official veterinarian;

II.2.4. have been obtained and prepared without contact with other materials that do not comply with the conditions required above, and have been handled so as to avoid contamination with pathogenic agents; and

II.2.5. have been transported in clean and sealed containers or lorries.

## Notes

(\*) Those countries subject to the transitional import arrangements include: an EU member State; Liechtenstein; Norway; Iceland and Switzerland.

References to European Union legislation within this certificate are references to direct EU legislation which has

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