



ANIMAL HEALTH/OFFICIAL CERTIFICATE FOR THE ENTRY INTO THE UNION OF MEAT PRODUCTS INTENDED FOR HUMAN CONSUMPTION, INCLUDING RENDERED ANIMAL FATS AND GREAVES, MEAT EXTRACTS AND TREATED STOMACHS, BLADDERS AND INTESTINES OTHERS THAN CASINGS, THAT ARE NOT REQUIRED TO UNDERGO A SPECIFIC RISK- MITIGATING TREATMENT (MPNT)

COUNTRY: United States		Animal health/Official certificate to the EU	
Part I: Description of consignment	I.1 Consignor/Exporter Name Address Country ISO country code	I.2 Certificate reference	I.2a IMSOC reference
		I.3 Central Competent Authority Food Safety Inspection Service	
		I.4 Local Competent Authority	
	I.5 Consignee/Importer Name Address Country ISO country code	I.6 Operator responsible for the consignment Name Address Country ISO country code	
	I.7 Country of origin ISO country code	I.9 Country of destination ISO country code	
	I.8 Region of origin Code	I.10 Region of destination Code	
	I.11 Place of dispatch Name Registration/Approval No Address Country ISO country code		
	I.13 Place of loading		
	I.15 Means of transport <input type="checkbox"/> Aircraft <input type="checkbox"/> Vessel <input type="checkbox"/> Railway <input type="checkbox"/> Road vehicle Identification	I.14 Date and time of departure	
		I.16 Entry Border Control Post	
		I.17	
	I.18 Transport conditions <input type="checkbox"/> Ambient <input type="checkbox"/> Chilled <input type="checkbox"/> Frozen		
	I.19 Container number/Seal number Container No Seal No		
	I.20 Certified as or for <input type="checkbox"/> Products for human consumption		
	I.21 <input type="checkbox"/> For transit Third country ISO country code	I.22 X For internal market	
	I.23		

Signature of Official Veterinarian or Official Inspector

Certificate reference:

I.24	Total number of packages	I.25	Total quantity	I.26	Total net weight/gross weight (kg)
I.27 Description of consignment					
CN code		Species			
Cold store		Identification mark	Type of packaging	Net weight	
Slaughterhouse		Treatment type	Nature of commodity	Number of packages	Batch No
<input type="checkbox"/> Final consumer		Date of collection/production	Manufacturing plant	Approval or registration number of plant/establishment/centre	

Signature of Official Veterinarian or Official Inspector

II. Health information		Certificate reference
Part II : Certification	II.1.	Public health attestation [to delete when the Union is not the final destination of the meat products]
	I, the undersigned, declare that I am aware of the relevant requirements of Regulation (EC) No 999/2001 of the European Parliament and of the Council , Regulation (EC) No 178/2002 of the European Parliament and of the Council , Regulation (EC) No 852/2004 of the European Parliament and of the Council , Regulation (EC) No 853/2004 of the European Parliament and of the Council and Regulation (EU) 2017/625 of the European Parliament and of the Council and hereby certify that the meat products(2), including rendered animal fats and greaves, meat extracts and treated stomachs, bladders and intestines others than casings, described in Part I were produced in accordance with these requirements, in particular that:	
	II.1.1.	they come from (an) establishment(s) applying general hygiene requirements and implementing a programme based on the hazard analysis and critical control points (HACCP) principles in accordance with Article 5 of Regulation (EC) No 852/2004, regularly audited by the competent authorities, and being listed as an EU approved establishment;
	II.1.2.	(1) either [the animals from which the meat products were derived have passed ante-mortem and post-mortem inspections;] (1) or [the wild game from which the meat products were derived have passed post-mortem inspection;]
	II.1.3.	they have been produced from raw material which met the requirements of Sections I to VI of Annex III to Regulation (EC) No 853/2004;
	(1) [II.1.4.1	if obtained from meat of domestic porcine animals, this meat fulfills the requirements of Commission Implementing Regulation (EU) 2015/1375 , and in particular: (1) either [has been subjected to an examination by a digestion method for Trichinella with negative results;] (1) or [has been subjected to a freezing treatment in accordance with Annex II to Implementing Regulation (EU) 2015/1375;] (1)(9) or [in the case of meat from domestic porcine animals kept solely for fattening and slaughter, comes from a holding or category of holdings that has been officially recognized by the competent authorities as free from Trichinella in accordance with Annex IV to Implementing Regulation (EU) 2015/1375;]]
	(1) [II.1.4.2.	if obtained from meat of solipeds or wild boar, this meat fulfils the requirements of Commission Implementing Regulation (EU) 2015/1375, and in particular, has been subject to an examination by a digestion method for Trichinella with negative results;]
	(1) [II.1.4.3.	the treated stomachs, bladders and intestines and meat extracts have been produced in accordance with Section XIII of Annex III to Regulation (EC) No 853/2004.]
	(1) [II.1.4.4.	the rendered animal fats and greaves have been produced in accordance with Section XII of Annex III to Regulation (EC) No 853/2004.]
	II.1.5.	they have been marked with an identification mark in accordance with Section I of Annex II to Regulation (EC) No 853/2004;
	II.1.6.	the label(s) affixed on the packaging of meat products described in Part I, bear(s) an identification mark to the effect that the meat products come wholly from fresh meat from establishments (slaughterhouses and cutting plants) approved for exporting to the European Union;
	II.1.7.	they satisfy the relevant criteria laid down in Commission Regulation (EC) No 2073/2005 ;
	II.1.8.	the guarantees covering live animals and products thereof provided by the residue plans submitted in accordance with Article 29 of Council Directive 96/23/EC , are fulfilled and the concerned animals and products are listed in Commission Decision 2011/163/EU for the concerned country of origin;
	II.1.9.	they have been produced under conditions guaranteeing compliance with the maximum residue levels for pesticides laid down in Regulation (EC) No 396/2005 of the European Parliament and of the Council , and the maximum levels for contaminants laid down in Commission Regulation (EC) No 1881/2006 .
	II.1.10.	the means of transport and the loading conditions of the meat products of this consignment meet the hygiene requirements laid down in respect of export to the European Union;

Part II : Certification	II. Health information	Certificate reference	
	<p>(1) [II.1.11. if containing material from bovine, ovine or caprine animals, with regard to bovine spongiform encephalopathy (BSE):</p> <p>(2) either [the country or region of origin is classified in accordance with Commission Decision 2007/453/EC as a country or region posing a negligible BSE risk, and</p> <p>(2) either [the animals from which the meat products are derived 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 have been no BSE indigenous cases;]</p> <p>(2) or [the animals from which the meat products are derived originate from 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 at least one BSE indigenous case, and the meat products do not contain and are not derived from mechanically separated meat obtained from bones of bovine, ovine and caprine animals;]</p> <p>(2) or [the animals from which the meat products are derived originate from a country or region classified in accordance with Decision 2007/453/EC as a country or region posing a controlled BSE risk and:</p> <p>(i) the meat products do not contain and are not derived from specified risk material as defined in Annex V, point 1, to Regulation (EC) No 999/2001;</p> <p>(ii) the meat products do not contain and are not derived from mechanically separated meat obtained from bones of bovine, ovine and caprine animals;</p> <p>(iii) the animals from which the meat products are 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;]</p> <p>(2) or [the animals from which the meat products 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:</p> <p>(i) the meat products do not contain and are not derived from specified risk material as defined in Annex V, point 1, to Regulation (EC) No 999/2001;</p> <p>(ii) the meat products do not contain and are not derived from mechanically separated meat obtained from bones of bovine, ovine and caprine animals;</p> <p>(iii) the animals from which the meat products are 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;]</p> <p>(iv) the animals from which the meat products are derived have not been fed with meat and bone meal or greaves, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health;</p> <p>(v) the meat products were produced and handled in a manner which ensures that they do not contain and were not contaminated with nervous and lymphatic tissues exposed during the deboning process;]]</p>		

II. Health information	Certificate reference
<p>(2) or {the country or region of origin is classified in accordance with Decision 2007/453/EC as a country or region posing a controlled BSE risk, and</p> <p>(a) the animals from which the meat products are 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;</p> <p>(b) the meat products do not contain and are not derived from:</p> <p>(i) specified risk material as defined in Annex V, point 1, to Regulation (EC) No 999/2001;</p> <p>(ii) mechanically separated meat obtained from bones of bovine, ovine and caprine animals.</p> <p>(2) either — {(c) the animals from which the meat products are derived originate from a country or region classified in accordance with Decision 2007/453/EC as a country or region posing a negligible or a controlled BSE risk;}</p> <p>(2) or — {(c) the animals from which the meat products 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</p> <p>(i) the animals from which the meat products are derived have not been fed with meat and bone meal or greaves, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health;</p> <p>(ii) the meat products were produced and handled in a manner which ensures that they do not contain and were not contaminated with nervous and lymphatic tissues exposed during the deboning process;}}</p> <p>(2) or {the country or region of origin is classified in accordance with Decision 2007/453/EC as a country or region with an undetermined BSE risk, and</p> <p>(a) the animals from which the meat products are derived have not been:</p> <p>(i) 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;</p> <p>(ii) 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;</p> <p>(b) the meat products do not contain and are not derived from:</p> <p>(i) specified risk material as defined in Annex V, point 1, to Regulation (EC) No 999/2001;</p> <p>(ii) mechanically separated meat obtained from bones of bovine, ovine and caprine animals;</p> <p>(iii) nervous and lymphatic tissues exposed during the deboning process.}}</p> <p>(1){H.1.12. if containing material from domestic solipeds, the fresh meat used in the preparation of the meat products:</p>	

Part II : Certification	II. Health information	Certificate reference	
	<p>either (1) {was obtained from domestic solipeds which immediately prior to slaughter had been kept for at least six months or since birth, if slaughtered at an age of less than six months, or since importation as food producing domestic solipeds from a Member State of the European Union, if imported less than six months prior to slaughter, in a third country:</p> <p>(a) in which the administration to domestic solipeds:</p> <p style="padding-left: 40px;">(i) of thyrostatic substances, stilbenes, stilbene derivatives, their salts and esters, oestradiol 17β and its ester like derivatives is prohibited;</p> <p style="padding-left: 40px;">(ii) of other substances having oestrogenic, androgenic or gestagenic action and of beta agonists is only allowed for:</p> <p>therapeutic treatment as defined in Article 1(2), point (b), of Council Directive 96/22/EC, where applied in conformity with Article 4(2) of that Directive, or</p> <p>zootechnical treatment as defined in Article 1(2), point (c), of Directive 96/22/EC, where applied in conformity with Article 5 of that Directive; and</p> <p>(b) which has had, at least during the six months prior to slaughter of the animals, a plan for the monitoring of the groups of residues and substances referred to in Annex I to Directive 96/23/EC which covers domestic solipeds born in and imported into the third country and was approved in accordance with Article 29(1), fourth subparagraph, of Directive 96/23/EC.</p> <p>and/or (1) {was imported from a Member State of the European Union.}}</p> <p>(1)(10) if containing material from farmed cervidae: {H.1.13.</p> <p>the product contains or is derived exclusively from meat, excluding offal and spinal cord, of farmed cervid animals which have been examined for chronic wasting disease by histopathology, immunohistochemistry or other diagnostic method recognised by the competent authorities with negative results and is not derived from animals coming from a herd where chronic wasting disease has been confirmed or is officially suspected.}</p> <p>(1)(11) if containing material from wild cervidae: {H.1.14.</p> <p>the product contains or is derived exclusively from meat, excluding offal and spinal cord, of wild cervid animals which have been examined for chronic wasting disease by histopathology, immunohistochemistry or other diagnostic method recognised by the competent authorities with negative results and is not derived from animals coming from a region where chronic wasting disease has been confirmed in the last three years or is officially suspected.}</p> <p>II.2. Animal health attestation [to delete when the meat product is entirely derived from meat of solipeds, leporidae or other wild land mammals others than ungulates]</p> <p>The meat product, including rendered animal fats and greaves, meat extracts and treated stomachs, bladders and intestines others than casings, described in Part I:</p> <p>II.2.1. has been processed in and dispatched from the zone with code: _____ (3), which, at the date of issue of this certificate, is authorised:</p> <p>(a) for entry into the Union of fresh meat of the species of animals from which the meat product described in Part I has been processed and listed in</p> <p style="padding-left: 40px;">(1) either [Part 1 of Annex XIII to Commission Implementing Regulation (EU) 2021/404 , in case of fresh meat of ungulates];</p> <p style="padding-left: 40px;">(1) or [Part 1 of Annex XIV to Commission Implementing Regulation (EU) 2021/404 , in case of fresh meat of poultry and game birds];</p> <p style="padding-left: 40px;">and</p> <p style="padding-left: 40px;">(b) and listed in Part 1 of Annex XV to Implementing Regulation (EU) 2021/404 for entry into the Union of the meat products described in Part I under the non-specific treatment "A";</p> <p>II.2.2. has been processed from fresh meat from the species of animals with code/s _____, _____ (4);</p> <p>II.2.3. has been processed from fresh meat that has undergone a non-specific treatment (5);</p>		

II. Health information		Certificate reference
Part II : Certification	II.2.4.	<p>has been processed from fresh meat that complied with all the relevant requirements for entry into the Union of fresh meat laid down in Commission Delegated Regulation (EU) 2020/692 and, therefore, was eligible for entry into the Union as such and was obtained from animals that complied with the residency period in an establishment located in:</p> <p>(1) either [the zone referred to in point II.2.1;]</p> <p>(1) or [the zone/s with code/s _____, _____, _____]</p> <p>(6) which, at the date of issue of this certificate is/are authorised for the entry into the Union of fresh meat of the species from which the meat product has been processed and listed in</p> <p>(1) either [Part 1 of Annex XIII to Implementing Regulation (EU) 2021/404;] (7)</p> <p>(1) or [Part 1 of Annex XIV to Implementing Regulation (EU) 2021/404;]</p> <p>(1) or [a Member State;]</p>
	II.2.5.	<p>has been processed from fresh meat obtained from:</p> <p>(1) either [animals kept in an establishment that was not subject to national restriction measures for animal health reasons, including the relevant listed diseases referred to in Annex I to Commission Delegated Regulation (EU) 2020/692 and emerging diseases at the time of dispatch of the animals to the slaughterhouse and in and around the establishment, in an area of 10 km radius, including where appropriate the territory of a neighbouring country, such diseases have not been reported during the period of 30 days prior to the date of dispatch of the animals to the Union;]</p> <p>(1) or [wild animals which originate from a place in and round which none of the listed diseases relevant for the species of origin of the meat products in accordance with Annex I to Commission Delegated Regulation (EU) 2020/692, has been reported during the period of 30 days prior to the date of dispatch of the meat product to the Union;]</p>
	II.2.6.	after processing has been handled until packaging in a way to prevent cross contamination that could introduce an animal health risk;
	(8) [II.2.7.	is intended for a Member State which has been granted the status free from infection with Newcastle disease virus without vaccination in accordance with Commission Delegated Regulation (EU) 2020/689 , and has been obtained from poultry which have not been vaccinated against infection with Newcastle disease virus with a live vaccine during the period of 30 days prior to the date of slaughter]
	II.3.	Animal welfare attestation [to delete when the Union is not the final destination]
	<p>I, the undersigned official veterinarian, hereby certify, that the meat products described in Part I derive from animals which have been treated in the slaughterhouse in accordance with the requirements of the Union legislation on the protection of animals at the time of killing or at least equivalent requirements.</p> <p>Notes</p> <p>In accordance with the Agreement on the withdrawal of the United Kingdom of Great Britain and Northern Ireland from the European Union and the European Atomic Energy Community, and in particular Article 5(4) of the Protocol on Ireland / Northern Ireland in conjunction with Annex 2 to that Protocol, references to European Union in this certificate include the United Kingdom in respect of Northern Ireland.</p> <p>This certificate is intended for entry into the Union of meat products coming from zones authorised to enter fresh meat of the relevant species and therefore are not required to undergo a specific risk-mitigating treatment, including when the Union is not the final destination of such meat product.</p> <p>This animal health/official certificate shall be completed according to the notes for the completion of certificates provided for in Chapter 4 of Annex I to Implementing Regulation (EU) 2020/2235.</p>	
	Part II:	
	(1)	Keep as appropriate.
	(2)	Meat product as defined in Point 7.1 of Annex I to Regulation (EC) No 853/2004.

Country: United States

Part II : Certification	II. Health information	Certificate reference							
	(3)	Code of the zone in accordance with column 2 of the table in Part 1 of Annex XV to Implementing Regulation (EU) 2021/404.							
	(4)	BOV= bovine animals; OVI= ovine animals and caprine animals; POR= porcine animals; RUF= animals of the family Bovidae (other than domestic bovine, ovine and caprine animals), camelid animals and cervid animals kept as farmed game; RUW: wild animals of the family Bovidae (other than domestic bovine, ovine and caprine animals), wild camelid animals and wild cervid animals; SUF: animals kept as farmed game of wild breeds of porcine animals and animals of the family Tayassuidae; SUW: wild animals of wild breeds of porcine animals and animals of the family Tayassuidae; POU= poultry other than ratites; RAT= Ratites; GB= game birds.							
	(5)	This can be certified only when treatment “A” is assigned in Part 1 of Annex XV to Implementing Regulation (EU) 2021/404 to the species of origin of the fresh meat and to the zone referred to in point II.2.1.							
	(6)	Code of the zone in accordance with column 2 of the table in Part 1 of Annex XIII or column 2 of the table in Part 1 of Annex XIV to Implementing Regulation (EU) 2021/404.							
	(7)	Not for zones with entry related to specific conditions ‘Maturation, pH and de-boning’ in column 5 of the table in Part 1 of Annex XIII to Implementing Regulation (EU) 2021/404.							
	(8)	This guarantee is required only for consignments intended for a Member State which has been granted the status free from infection with Newcastle disease virus without vaccination in accordance with Delegated Regulation (EU) 2020/689.							
	(9)	The derogation for domestic porcine animals coming from a holding officially recognised as applying controlled housing conditions, can only be applied in countries listed in Annex VII to Implementing Regulation (EU) 2015/1375.							
	(10)	Applicable when the meat has been obtained from a country mentioned in Annex IX, Chapter F, point 1, to Regulation (EC) No 999/2001.							
	(11)	Applicable when the meat has been obtained from a country mentioned in Annex IX, Chapter F, point 2, to Regulation (EC) No 999/2001.							
	Official veterinarian or Official inspector <table border="0" style="width: 100%;"> <tr> <td style="width: 50%;">Name (in capital letters)</td> <td style="width: 50%;">Qualification and title</td> </tr> <tr> <td>Date of signature</td> <td>Signature</td> </tr> <tr> <td>Stamp</td> <td></td> </tr> </table>				Name (in capital letters)	Qualification and title	Date of signature	Signature	Stamp
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