Part I : Details of dispatched consignment

I.1. Consignor
Name
Address
Country
Phone

I.2. Certificate reference number

I.2.a. TRACES reference number :

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Consignee
Name
Address
Country
Phone

I.6 Person responsible for the consignment in the EU

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 origin
Name Approval number
Address

I.12. Place of destination

I.13 Place of loading
Address

I.14 Date of departure

I.15. Means of transport
Aeroplane ☐ Ship ☐ Railway wagon ☐
Road vehicle ☐ Other ☐
Identification::
Document:

I.16. Entry BIP in EU
Name BIP unit no.:

I.17. No.(s) of CITES

I.21 Temperature of products
Ambient ☐ Chilled ☐ Frozen ☐

I.20.Quantity

I.22. Total Number of Packages

I.23. Seal / Container No.

I.25. Commodity certified for:

I.26. For transit to 3rd Country by EU

I.27. For import or admission into EU ☐
Definitive import ☐
Horses Re-entry ☐
Temporary admission horses ☐

I.28. Identification of the commodity

Part II: Certification

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

II.1. Animal health attestation

I, the undersigned official veterinarian certify that:

II.1.1. The meat product, treated stomachs, bladders and intestines(1) described in this certificate contain the following meat constituents and meet the criteria indicated below:

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

(A) Insert the code for the relevant species of meat product, treated stomachs, bladders and intestines where BOV = domestic bovine animals (Bos Taurus, Bison bison, Bubalus bubalis and their crossbreds); OVI = domestic sheep (Ovis aries) and goats (Capra hircus); EQI = domestic equine animals (Equus caballus, Equus asinus and their crossbreds), POR = domestic porcine animals (Sus scrofa); RAB = 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; WLP = 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 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.

(2) II.1.2. The meat product, treated stomachs, bladders and intestines described in point II.1.1. has been prepared from fresh meat from domestic bovine animals (Bos Taurus, Bison bison, Bubalus bubalis and their crossbreds); domestic sheep (Ovis aries) and goats (Capra hircus); domestic equine animals (Equus caballus, Equus asinus and their crossbreds), domestic porcine animals (Sus scrofa); farmed non-domestic animals other than suidae and solipeds; wild non-domestic animals other than suidae and solipeds; wild non- domestic suidae: wild non-domestic solipeds and the fresh meat used in the production of the meat products:

(2) either [II.1.2.1. has undergone a non-specific treatment as specified and defined under point A in Part 4 of Annex II to Decision 2007/777/EC and:

(2) either [II.1.2.1.1. satisfies the relevant animal and public health requirements laid down in the appropriate health certificate(s) in Part 2 of Annex II to Regulation (EU) No 206/2010 and originates in a third country, or part thereof in the case of regionalisation under Union legislation, as described in the relevant column of part 2 of Annex II to Decision 2007/777/EC].

(2) or [II.1.2.1.1. originates in a Member State of the European Union].

(2) or [II.1.2.1. meets any requirements agreed under Directive 2002/99/EC, is derived from animals coming from a holding not subject to restrictions for the specific diseases mentioned in the appropriate health certificate(s) in in Part 2 of Annex II to Regulation (EU)No 206/2010 and within a 10 km radius of which no outbreaks of such diseases have occurred in the last 30 days and has undergone the specific treatment laid down for the third country of origin or part thereof for the meat of the species concerned in Parts 2 or 3, as appropriate, of Annex II to Decision 2007/777/EC].

(2) II.1.3. The meat product, treated stomachs, bladders and intestines described under point II.1.1 has been prepared from fresh meat of domestic poultry, including farmed or wild game birds, that:

(2) either [II.1.3.1. has undergone a non-specific treatment as specified and defined under point A in Part 4 of Annex II to Decision 2007/777/EC] and:

(2) either [II.1.3.1.1. satisfies the animal health requirements laid down in Regulation (EC) No 798/2008,]

(2) or [II.1.3.1.1. originates in a Member State of the European Union satisfying the requirements of Article 3 of Directive 2002/99/EC,]

(2) or [II.1.3.1. originates in a third country referred to in Annex I part 1 to Regulation (EC) No 798/2008, comes from holdings or in the case of wild game-birds killed in territories where within a 10 km radius, 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 and has undergone the specific treatment laid down for the third country of origin or part thereof for the meat of the species concerned in Parts 2 or 3, as appropriate, of Annex II to Decision 2007/777/EC,]

(2) or [II.1.3.1. originates in a third country referred to in Annex I part 1 to Regulation (EC) No 798/2008, comes from holdings or in the case of wild game-birds killed in territories, where within a 10 km radius, 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 and has undergone the specific treatment referred to in points B, C or D in Part 4 of Annex II to Decision 2007/777/EC, provided that such treatment is more severe than that indicated in Parts 2 and 3 of Annex II to that Decision.]

(2) [II.1.4. in the case of meat product, treated stomachs, bladders and intestines derived from fresh meat from lagomorphs and other land mammals:

satisfies the relevant animal health and public health requirements laid down in Regulation (EC) No 119/2009 and has come from a holding not subject to restrictions for animal diseases affecting the animals concerned within a 10 km radius of which no outbreaks of such diseases have occurred in the last 30 days.]

II.1.5. the meat product, treated stomachs, bladders and intestines:

(2) either II.1.5.1. [consists of meat and/or meat products derived from a single species, and has undergone the treatment satisfying the relevant conditions laid down in Annex II to Decision 2007/777/EC,]

(2) or II.1.5.1. [consists of meat of more than one species and, after such meat has been mixed, the entire product has subsequently undergone a treatment at least as severe as that required for the meat components of the meat product as laid down in Annex II to Decision 2007/777/EC,]

(2) or II.1.5.1. [has been prepared from meat of more than one species and each meat component has previously undergone a treatment prior to mixing which meets the relevant treatment requirements for meat of that species as laid down in Annex II to Decision 2007/777/EC];

II.1.6. after treatment all precautions to avoid contamination have been taken

(2) [II.1.7. Additional guarantees:

in the case of poultry meat products which have not undergone a specific treatment and are destined for Member States or regions thereof, the status of which have been established as Newcastle disease non-vaccinating in accordance with Article 15 of Directive 2009/158/EC, the poultry meat was derived from poultry which had not been vaccinated with a live vaccine against Newcastle disease within 30 days prior to slaughter;]

(2) II.2. Public health attestation

I, the undersigned, declare that I am aware of the relevant provisions of Regulations (EC) No 999/2001, (EC) No178/2002, (EC) No 852/2004 and (EC) No 853/2004 and certify that the meat products, treated stomachs, bladders and intestines described above were produced in accordance with those requirements, in particular that:

II.2.1. they come from (an) establishment(s) implementing a programme based on the HACCP principles in accordance with Regulation (EC) No 852/2004;

II.2.2. 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;

Part II: Certification

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

(2) either II.2.3.1 the meat products have been obtained from domestic pig meat which either has been subject to an examination for trichinosis with negative results or has been subjected to a cold treatment in accordance with Regulation (EC) No 2075/2005;

(2)(5)or II.2.3.1 the meat products have been obtained from domestic porcine animals meat which is derived from domestic porcine animals either coming from a holding officially recognised as applying controlled housing conditions in accordance with Article 8 of Regulation (EC) No 2075/2005 or not weaned and less than 5 weeks of age;

(2) II.2.3.2. the meat products have been obtained from horse meat or wild boar meat which has been subject to an examination for trichinosis with negative results in accordance with Regulation (EC) No 2075/2005;

(2) II.2.3.3. the treated stomachs, bladders and intestines have been produced in accordance with Section XIII of Annex III, to Regulation (EC) No 853/2004.

II.2.4. they have been marked with an identification mark in accordance with Section I of Annex II to Regulation (EC) No 853/2004;

II.2.5. the label(s) affixed on the packaging of meat products described above, bear(s) a mark to the effect that the meat products come wholly from fresh meat from animals slaughtered in slaughterhouses approved for exporting to the European Union or, from animals slaughtered in a slaughterhouse specially for the delivery of meat for the required treatment as laid down in Part 2 and 3 of Annex II of Decision 2007/777/EC;

II.2.6. they satisfy the relevant criteria set out in Regulation (EC) No 2073/2005 on microbiological criteria for foodstuffs;

II.2.7. the guarantees covering live animals and products thereof provided by the residue plans submitted in accordance with Directive 96/23/EC, and in particular Article 29 thereof, are fulfilled;

II.2.8. the means of transport and the loading conditions of meat products of this consignment meet the hygiene requirements laid down in respect of export to the European Union;

(2) II.2.9. if containing material from bovine, ovine or caprine animals, the meat products and treated intestines are subject to the following conditions depending on the BSE risk category of the country of origin:

(2) either [(1) the country or region of dispatch is classified in accordance with Decision 2007/453/EC as a country or region posing a negligible BSE risk;
(2) the animals, from which the fresh meat and intestines used in the preparation of the meat products and treated intestines of bovine, ovine and caprine origin were derived, have passed ante mortem and post mortem inspections;

(2) either [(3) the animals, from which the fresh meat and intestines used in the preparation of the meat products and treated intestines of bovine, ovine and caprine origin were derived:

(a) 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;

(2) [(b) have 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;]]

(2) or [(3) the animals, from which the fresh meat and intestines used in the preparation of the meat products and treated intestines of bovine, ovine and caprine 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;]

(4) the meat products of bovine, ovine and caprine 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;

(2) either [(5) the meat products of bovine, ovine and caprine origin do not contain and are not derived from mechanically separated meat, obtained from bones of bovine, ovine and caprine animals;]

(2) or [(5) the meat products of bovine, ovine and caprine origin are derived from mechanically separated meat, obtained from bones of bovine, ovine and caprine animals which 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 and in which there has been no BSE indigenous cases;]

(2) [(6) (a) the animals, from which the fresh meat and intestines used in the preparation of the meat products and treated intestines of bovine, ovine and caprine origin were 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;

(b) the animals, from which the fresh meat and intestines used in the preparation of the meat products and treated intestines of bovine, ovine and caprine origin were derived, have not been fed with meat-and-bone meal or greaves, as defined in the World Organisation for Animal Health (OIE) Terrestrial Animal Health Code, and

(c) the meat products 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.]]

(2) or [(1) the country or region of dispatch is classified in accordance with Decision 2007/453/EC as a country or region posing a controlled BSE risk;
(2) the animals, from which the fresh meat and intestines used in the preparation of the meat products and treated intestines of bovine, ovine and caprine origin were derived, have passed ante mortem and post mortem inspections;

(3) the animals from which the fresh meat and intestines used in the preparation of the meat products and treated intestines of bovine, ovine and caprine origin were derived, have not been 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;

(4) the meat products of bovine, ovine and caprine 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 bones of bovine, ovine and caprine animals;

(2)(3) [(5) in the case of intestines originally sourced from a country or a region with a negligible BSE risk, the treated intestines are subject to the following conditions:

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

(b) for intestines sourced from a country or region where there have been BSE indigenous cases:

Part II: Certification

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

(2) either [(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 has been enforced;]

(2) or [(i) the meat products 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.]]]

(2) or [(1) the country or region of dispatch has not been classified in accordance with Decision 2007/453/EC or is classified as a country or region with an undetermined BSE risk;

(2) the animals, from which the fresh meat and intestines used in the preparation of the meat products and treated intestines of bovine, ovine and caprine origin were derived, have passed ante mortem and post mortem inspections;

(3) the animals from which the fresh meat and intestines used in the preparation of the meat products and treated intestines of bovine, ovine and caprine origin were derived have not been fed meat-and-bone meal or greaves derived from ruminants, as defined in the OIE Terrestrial Animal Health Code;

(4) the animals from which the fresh meat and intestines used in the preparation of the meat products and treated intestines of bovine, ovine and caprine origin were derived, have not been 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;

(5) the meat products of bovine, ovine and caprine origin do not contain and are not derived from:

(a) specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001;

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

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

(2)(3) [(6) in the case of intestines originally sourced from a country or a region with a negligible BSE risk, the treated intestines are subject to the following conditions:

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

(b) for intestines sourced from a country or region where there have been BSE indigenous cases:

(2) either [(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 has been enforced;]

(2) or [(i) the meat products 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.]]]

(2) II.2.10. if containing material from domestic equine animals, the fresh meat, stomachs, bladders or intestines used in the preparation of the meat products and/or treated stomachs, bladders and intestines

(2) either [was/were obtained from domestic equine animals 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 equidae from a Member State of the European Union, if imported less than six months prior to slaughter, in a third country:

(a) in which the administration to domestic equine animals:

(i) of thyrostatic substances, stilbenes, stilbene derivatives, their salts and esters, oestradiol 17β and its ester-like derivatives is prohibited;

(ii) of other substances having oestrogenic, androgenic or gestagenic action and of beta-agonists is only allowed for:

- therapeutic treatment as defined in Article 1(2)(b) of Directive 96/22/EC, where applied in conformity with Article 4(2) of that Directive, or
- zootechnical treatment as defined in Article 1(2)(c) of Directive 96/22/EC, where applied in conformity with Article 5 of that Directive; and

(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 equidae born in and imported into the third country and was approved in accordance with the fourth subparagraph of Article 29(1) of Directive 96/23/EC.]

(2) and/or [was/were imported from a Member State of the European Union.]

Notes

Part I:

Box reference I.8.: region (if appropriate) as appearing in Annex II to Decision 2007/777/EC (as last amended).

Box reference I.11: Place of origin: name and address of the dispatch establishment.

Box reference I.15: Registration number (railway wagons or container and road vehicle), flight number (aircraft) or name (ship). Separate information is to be provided in the event of unloading and reloading.

Box reference I.19: Use the appropriate Harmonised System (HS) code under the following headings: 02.10, 16.01, 16.02 and 05. 04.

Box reference I.23: Identification of container/Seal number: only where applicable.

Box reference I.28:

Species: select among species described in Part II 1.1. (A);

Nature of commodity: choose among the following: meat product, treated stomachs, bladders and intestines;

Abattoir: approval number of any abattoir or game-handling establishment;

Cold store: any storage facility;

Manufacturing plant: approval number.

Part II:

Part II: Certification

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

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

(2) Keep as appropriate.

(3) Only applicable to imports of treated intestines.

(4) 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 point 11.3(a) of Annex V to Regulation (EC) No 999/2001.

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.

(5) Only for third countries with the entry “K” in column “SG” in Part 1 of Annex II to Regulation (EU) No 206/2010.
The colour of the signature shall be different to that of the printing. The same rule applies to the stamp other than those embossed or watermarked.

Official veterinarian or official inspector

Name (in Capital): Qualification and title:

Local Veterinary Unit: LVU N°:

Date: Signature:

Stamp