Part I: Description of consignment

I.1 Consignor/Exporter
Name
Address
Country ISO country code

I.2 Certificate reference
.....

I.2a

I.3 Central Competent Authority
DEPARTMENT FOR ENVIRONMENT, FOOD & RURAL AFFAIRS

I.4 Local Competent Authority
ANIMAL AND PLANT HEALTH AGENCY

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 Registration/Approval No
Name
Address
Country ISO country code

I.12 Place of destination Registration/Approval No
Name
Address
Country ISO country code

I.13 Place of loading

I.14 Date and time of departure

I.15 Means of transport
☐ Aircraft ☐ Vessel
☐ Railway ☐ Road vehicle
Identification

I.16 Entry Border Control Post

I.17 Accompanying documents
Type Code
Country ISO country code
Commercial document reference

I.18 Transport conditions ☐ Ambient ☐ Chilled ☐ Frozen

I.19 Container number/Seal number
Container No Seal No

I.20 Certified as or for
☒ Products for human consumption

I.21 ☐ **For transit**
Third country ISO country code

I.22 ☐ **For internal market**

I.23

I.24 Total number of packages	**I.25**	**I.26 Total net weight/gross weight (kg)**

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I.27 Description of consignment

1

CN Code	Species	Nature of commodity	Treatment type	Slaughterhouse	Manufacturing plant	Cold store
Date of collection /production	Batch No	Type of packaging	Number of packages	Net weight	Final consumer ☐	

2

CN Code	Species	Nature of commodity	Treatment type	Slaughterhouse	Manufacturing plant	Cold store
Date of collection /production	Batch No	Type of packaging	Number of packages	Net weight	Final consumer ☐	

3

CN Code	Species	Nature of commodity	Treatment type	Slaughterhouse	Manufacturing plant	Cold store
Date of collection /production	Batch No	Type of packaging	Number of packages	Net weight	Final consumer ☐	

4

CN Code	Species	Nature of commodity	Treatment type	Slaughterhouse	Manufacturing plant	Cold store
Date of collection /production	Batch No	Type of packaging	Number of packages	Net weight	Final consumer ☐	

5

CN Code	Species	Nature of commodity	Treatment type	Slaughterhouse	Manufacturing plant	Cold store
Date of collection /production	Batch No	Type of packaging	Number of packages	Net weight	Final consumer ☐	

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Part II: Certification

II. Health information

II.1. Public health attestation [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 is listed as a Union 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;

II.1.4. they have been marked with an identification mark in accordance with Section I of Annex II to Regulation (EC) No 853/2004;

II.1.5. 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, game handling establishment and cutting plants) approved for the entry into the Union;

II.1.6. they satisfy the relevant criteria laid down in Commission Regulation (EC) No 2073/2005;

II.1.7. the guarantees covering live animals and products thereof provided by the control plan submitted in accordance with Article 6(2) of Commission Delegated Regulation (EU) 2022/2292 are fulfilled and the concerned animals and products are listed in Annex -I to Commission Implementing Regulation (EU) 2021/405 for the concerned third country or territory;

II.1.8. the means of transport and the loading conditions of the meat products of this consignment meet the hygiene requirements laid down as regards the entry into the Union;

[1] [II.1.9.1. 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 recognised by the competent authorities as free from *Trichinella* in accordance with Annex IV to Implementing Regulation (EU) 2015/1375;]]

[1] [II.1.9.2. if obtained from meat of solipeds or wild boar, this meat fulfils the requirements of 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.1.9.3. 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.1.9.4. the rendered animal fats and greaves have been produced in accordance with Section XII of Annex III to Regulation (EC) No 853/2004.]

[1] [II.1.10. if containing material from bovine, ovine or caprine animals, with regard to bovine spongiform encephalopathy (BSE):

[1] *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

[1] *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;]]]

[1] *and/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;]]]

[1] *and/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:

(i) the meat products 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;

(ii) the meat products do not contain and are not derived from mechanically separated meat obtained from bones of bovine, ovine and caprine animals;

(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;]]]

[1] *and/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:

(i) the meat products 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;

(ii) the meat products do not contain and are not derived from mechanically separated meat obtained from bones of bovine, ovine and caprine animals;

(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;

(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;

(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;]]]

(1) *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:

(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;

(b) the meat products do not contain and are not derived from:

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

(ii) mechanically separated meat obtained from bones of bovine, ovine and caprine animals.

(1) *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;]]]

(1) *and/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:

(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;

(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;]]]

(1) *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:

(a) the animals from which the meat products are derived have not been:

(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;

(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;

(b) the meat products do not contain and are not derived from:

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

(ii) mechanically separated meat obtained from bones of bovine, ovine and caprine animals;

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

(1) [II.1.11. if containing material from domestic solipeds, the fresh meat used in the preparation of the meat products was obtained from domestic solipeds which immediately prior to the date of their slaughter had been kept:

(1) *either* [for at least six months in the third country of slaughter, if born in that third country or have entered that third country from another third country which is listed for the concerned animals and products in Annex -I to Implementing Regulation (EU) 2021/405, and]

(1) *or* [in the third country of slaughter, since birth, if slaughtered at an age of less than six months, and]

(1) *or* [in the third country of slaughter for six months or less if they entered that third country from a Member State as domestic solipeds for food production, and]

in a third country or territory of slaughter in which:

(a) the administration to domestic solipeds of:

(i) substances listed in Table 2 of the Annex to Commission Regulation (EU) No 37/2010 is prohibited;

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

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

– (1) *either* [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,]

– (1) *or* [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,]

(b) the domestic solipeds fulfilled, at least during six months prior to the date of their slaughter, guarantees provided by the control plan submitted in accordance with Article 6(2) of Commission Delegated Regulation (EU) 2022/2292 and the

concerned animals and products are listed in Annex -I to Implementing Regulation (EU) 2021/405 for the concerned third country or territory.]

(1) [II.1.12. (1) (10) *either* [if containing material from farmed cervidae, 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.]]

(1) (11) *or* [if containing material from wild cervidae, 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 prior to the date of issue of this animal health/official certificate or is officially suspected.]]

(1) (12) [**II.1.a. Attestation as regards Commission Delegated Regulation (EU) 2023/905** [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 (EU) 2019/6 of the European Parliament and of the Council and Commission Delegated Regulation (EU) 2023/905 and hereby certify that the meat products, 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, and in particular, that the animals from which the meat is derived have not been administered antimicrobial medicinal products for growth promotion or yield increase or antimicrobial medicinal products containing an antimicrobial that is included in the list of antimicrobials reserved for the treatment of certain infections in humans laid down in Commission Implementing Regulation (EU) 2022/1255 as set out in Article 3 of Delegated Regulation (EU) 2023/905 and originate from a third country or region thereof listed in accordance with Article 5(2) of Delegated Regulation (EU) 2023/905.]

II.2. Animal health attestation [Delete when the meat product is entirely derived from meat of domestic solipeds (*Equus caballus*, *Equus asinus* and their cross-breeds); wild game solipeds belonging to the subgenus *Hippotigris* (Zebra); wild leporidae; or wild land mammals other than ungulates and leporidae]

The **meat product**, including rendered animal fats and greaves, meat extracts and treated stomachs, bladders and intestines others than casings, described in Part I:

II.2.1. has been processed in and dispatched from the **zone** with code: ……(3), which, at the date of issue of this animal health/official certificate, is:

(a) authorised for the 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:

(1) *either* [Part 1 of Annex XIII to Commission Implementing Regulation (EU) 2021/404, in case of fresh meat of ungulates;]

(1) *or* [Part 1 of Annex XIV to Implementing Regulation (EU) 2021/404, in case of fresh meat of poultry and game birds;]

(b) 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";

II.2.2. has been processed from fresh meat from **the species of animals** with code/s ____, ____, ____ (4);

II.2.3. has been processed from fresh meat that has undergone a non-specific treatment (5);

II.2.4. 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

(1) *either* [the zone referred to in point II.2.1;]

(1) *or* [the zone/s with code/s _____, ____, ____ (6) which, at the date of issue of this animal health/official 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

(1) *either* [Part 1 of Annex XIII to Implementing Regulation (EU) 2021/404;]] (7)

(1) *or* [Part 1 of Annex XIV to Implementing Regulation (EU) 2021/404;]]

(1) *or* [a Member State;]]

II.2.5. has been processed from fresh meat obtained from:

(1) *either* [animals kept in an establishment that was not subject to national restriction measures for animal health reasons, including the listed diseases referred to in Annex I to Delegated Regulation (EU) 2020/692 relevant for the species of origin of the meat products and emerging diseases at the date 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 last 30 days prior to the date of slaughter of the animals;]

(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 Delegated Regulation (EU) 2020/692, has been reported during the last 30 days prior to the date of killing of the animals;]

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 or zone thereof 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

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have not been vaccinated against infection with Newcastle disease virus with a live vaccine during the last 30 days prior to the date of slaughter of the animals.]

II.3. Animal welfare attestation [Delete when the Union is not the final destination]

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.

Notes

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 the Union in this animal health/official certificate include the United Kingdom in respect of Northern Ireland.

This animal health/official certificate is intended for the 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.

This animal health/official certificate shall be completed in accordance with the notes for the completion of certificates provided for in Chapter 4 of Annex I to Commission Implementing Regulation (EU) 2020/2235.

Part I:

Box reference I.27.: Description of consignment:
"Slaughterhouse": Slaughterhouse or game handling establishment.

Part II:

(1) Delete if not applicable.
(2) Meat product as defined in point 7.1 of Annex I to Regulation (EC) No 853/2004.
(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= domestic bovine animals; OVI= domestic ovine animals and caprine animals; POR= domestic 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 may 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 the consignments intended for a Member State or zone thereof 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, may 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 Chapter F, point 1, of Annex IX to Regulation (EC) No 999/2001.
(11) Applicable when the meat has been obtained from a country mentioned in Chapter F, point 2, of Annex IX to Regulation (EC) No 999/2001.
(12) Applicable to consignments entering the Union as from 3 September 2026.

Official veterinarian

Name (in capital letters)

Date

Qualification and title

Stamp

Signature