

UNITED KINGDOM

Official certificate to the EU

Part I: Description of consignment	I.1 Consignor/Exporter		I.2 Certificate reference		I.2a	
	Name				
	Address		I.3 Central Competent Authority DEPARTMENT FOR ENVIRONMENT, FOOD & RURAL AFFAIRS			
	Country		ISO country code		I.4 Local Competent Authority ANIMAL AND PLANT HEALTH AGENCY	
	I.5 Consignee/Importer			I.6 Operator responsible for the consignment		
	Name			Name		
	Address			Address		
	Country			ISO country code		Country
	ISO country code			Country		ISO country code
	I.7 Country of origin			I.9 Country of destination		ISO country code
ISO country code			I.10 Region of destination		Code	
I.8 Region of origin			I.12 Place of destination		Registration/Approval No	
Code			Name		Address	
I.11 Place of dispatch			Name		Address	
Registration/Approval No			Address		Country	
Name			Country		ISO country code	
Address			ISO country code		Country	
Country			ISO country code		Country	
ISO country code			ISO country code		Country	
I.13 Place of loading			I.14 Date and time of departure			
I.15 Means of transport			I.16 Entry Border Control Post			
<input type="checkbox"/> Aircraft			<input type="checkbox"/> Vessel			
<input type="checkbox"/> Railway			<input type="checkbox"/> Road vehicle			
Identification			I.17 Accompanying documents			
			Type			
			Code			
			Country			
			ISO country code			
			Commercial document reference			
I.18 Transport conditions			<input type="checkbox"/> Ambient			
<input type="checkbox"/> Chilled			<input type="checkbox"/> Frozen			
I.19 Container number/Seal number			Seal No			
Container No						
I.20 Certified as or for						
<input type="checkbox"/> Products for human consumption						
I.21			I.22 <input type="checkbox"/> 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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1.27 Description of consignment

1						
CN code	Species	Cold store	Type of packaging	Net weight	Slaughterhouse	Treatment type
Nature of commodity	Number of packages	Batch No	Date of collection /production	Manufacturing plant	Final consumer <input type="checkbox"/>	
2						
CN code	Species	Cold store	Type of packaging	Net weight	Slaughterhouse	Treatment type
Nature of commodity	Number of packages	Batch No	Date of collection /production	Manufacturing plant	Final consumer <input type="checkbox"/>	
3						
CN code	Species	Cold store	Type of packaging	Net weight	Slaughterhouse	Treatment type
Nature of commodity	Number of packages	Batch No	Date of collection /production	Manufacturing plant	Final consumer <input type="checkbox"/>	
4						
CN code	Species	Cold store	Type of packaging	Net weight	Slaughterhouse	Treatment type
Nature of commodity	Number of packages	Batch No	Date of collection /production	Manufacturing plant	Final consumer <input type="checkbox"/>	
5						
CN code	Species	Cold store	Type of packaging	Net weight	Slaughterhouse	Treatment type
Nature of commodity	Number of packages	Batch No	Date of collection /production	Manufacturing plant	Final consumer <input type="checkbox"/>	

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II.a Certificate reference

II. Health information**II.1. Public health attestation**

I, the undersigned, declare that I am aware of the relevant requirements of 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, Regulation (EU) 2017/625 of the European Parliament and of the Council, Commission Delegated Regulation (EU) 2019/624 and Commission Implementing Regulation (EU) 2019/627 and hereby certify that the fresh meat of domestic solipeds (*Equus caballus*, *Equus asinus* and their cross-breeds) described in Part I was produced in accordance with these requirements, in particular that:

- ◆ II.1.1. the meat comes 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. the meat has been obtained in compliance with the conditions set out in Section I of Annex III to Regulation (EC) No 853/2004;
- II.1.3. the 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;
- II.1.4. the meat has been found fit for human consumption following *ante-mortem* and *post-mortem* inspections carried out in accordance with Articles 8 to 17, 22, 24, 31 to 35, 37, 38 of Implementing Regulation (EU) 2019/627 and Articles 3, 4, 5, 7 and 8 of Delegated Regulation (EU) 2019/624;
- II.1.5. ⁽¹⁾ either [the carcase or parts of the carcase have been marked in accordance with Article 48 of and Annex II to Implementing Regulation (EU) 2019/627;]
⁽¹⁾ or [the packages of meat 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 meat satisfies the relevant criteria laid down in Commission Regulation (EC) No 2073/2005;
- II.1.7. the meat 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 Commission 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 of slaughter:
(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 the 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.
- II.1.8. the meat has been stored and transported in accordance with the relevant requirements of Section I of Annex III to Regulation (EC) No 853/2004.

(1) (3)

[II.1.a. Attestation as regards Commission Delegated Regulation (EU) 2023/905 [Delete when the Union is not the final destination of the fresh meat]

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 fresh meat of domestic solipeds (*Equus caballus*, *Equus asinus* and their cross-breeds) described in Part I was 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.]

Part II: Certification

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II.a Certificate reference

II.2. Animal welfare attestation

I, the undersigned official veterinarian, hereby certify, that the meat described in Part I derives 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 official certificate include the United Kingdom in respect of Northern Ireland.

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

The exclusion of minced meat and mechanically separated meat is expressly mentioned in the title to avoid any confusion as these products shall not enter into the Union using this fresh meat certificate.

This official certificate is meant for fresh meat, excluding fresh blood, minced meat and mechanically separated meat, of domestic solipeds (*Equus caballus*, *Equus asinus* and their cross-breeds).

Fresh meat as defined in point 1.10. of Annex I to Regulation (EC) No 853/2004.

Part I:

Box reference I.27.: Description of consignment:
 "CN code": Indicate the appropriate Harmonised System (HS) code(s) of the World Customs Organisation under the following headings: 02.05, 02.06 or 05.04.
 "Nature of commodity": Indicate "carcase-whole", "carcase-side", "carcase-quarters" "offal" ⁽²⁾ or "cuts".
 "Treatment type": If appropriate, indicate "de-boned", "bone in" and/or "matured". If frozen, indicate the date of freezing (mm/yy) of the cuts/pieces.

Part II:

- ⁽¹⁾ Delete if not applicable.
⁽²⁾ Excluding fresh blood entry into the Union of which is not permitted in accordance with Article 130 of Commission Delegated Regulation (EU) 2020/692.
⁽³⁾ Applicable to consignments entering the Union as from 3 September 2026.

Official veterinarian

Name (in capital letters)

Date

Qualification and title

Stamp

Signature