

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	
	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		I.12 Place of destination	
	Name				Registration/Approval No	
	Address				Name	
					Address	
	Country		ISO country code		Country	
					ISO country code	
	I.13 Place of loading		I.14 Date and time of departure			
	I.15 Means of transport		I.16 Entry Border Control Post			
	☐ Aircraft		☐ Vessel		I.17 Accompanying documents	
	☐ Railway		☐ Road vehicle		Type	
					Code	
	Identification				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			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	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 ☐	

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 ☐	

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 ☐	

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 ☐	

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 ☐	

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II.a Certificate reference

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Part II: Certification

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 establishments 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 listed as Union approved establishments;
- 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 meat is a carcass or part thereof which has been marked in accordance with Article 48 of and Annex II to Implementing Regulation (EU) 2019/627;]
⁽¹⁾ *or* [the meat is in packages which 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 6 months in the third country or territory of slaughter, if born in that third country or territory, or have entered that third country or territory from another third country or territory which is listed for the concerned animals and products in Annex -I to Commission Implementing Regulation (EU) 2021/405, and where;]
⁽¹⁾ *or* [in the third country or territory of slaughter, since birth, if slaughtered at an age of less than 6 months, and where;]
⁽¹⁾ *or* [in the third country or territory of slaughter for 6 months or less if they entered that third country or territory from a Member State as domestic solipeds for food production, and where;]
 - (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 17 β 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⁾ *and/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 6 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 third country or region thereof of their origin is listed in Annex -I to Implementing Regulation (EU) 2021/405 and marked with an "X" for the category "equine";
- 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.1a. Attestation as regards Commission Delegated Regulation (EU) 2023/905**

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 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 the Annex to Commission Implementing Regulation (EU) 2024/2598.]

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.

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II.a Certificate reference

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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 Windsor Framework (see Joint Declaration No 1/2023 of the Union and the United Kingdom in the Joint Committee established by 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 of 24 March 2023, OJ L 102, 17.4.2023, p. 87) in conjunction with Annex 2 to that Framework, 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 Commission 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 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: “CN code”: Indicate the appropriate Harmonised System (HS) code(s) of the World Customs Organisation under the following headings: 0205, 0206 or 0504.

“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