	UNITED KINGDOM			Animal health/Official certificate to the EU				
	I.1 Consignor/Exporter			I.2 Certificate ref	ference	I.2a		
	Name							
	Address			I.3 Central Comp				
				DEPARTMENT FOOD & RURAL	OR ENVIRONMENT, AFFAIRS			
				I.4 Local Competent Authority		7 /		
	Country	ISO co	untry code	ANIMAL AND P	LANT HEALTH AGENCY			
	I.5 Consignee/Importer			I.6 Operator resp	onsible for the consignmen	nt		
	Name			Name				
nt	Address			Address				
Part I: Description of consignment								
n of	Country	ISO co	untry code	Country	·	ISO country code		
otio	I.7 Country of origin		untry code	I.9 Country of destination		ISO country code		
criț								
: Des	I.8 Region of origin Code			I.10 Region of de	stination	Code		
Part I	I.11 Place of dispatch Ro		gistration/Approval No	I.12 Place of destination		Registration/Approval No		
	Name			Name				
	Address			Address				
				*				
	Country	150 coi	untry code	Country		ISO country code		
	I.13 Place of loading			I.14 Date and tim	e of departure			
	I.15 Means of transport			I.16 Entry Borde	r Control Post			
	П A' С							
	☐ Aircraft	□ Vessel		I.17 Accompanyi	ng documents			
				Туре	C	ode		
	□ Railway □ R		icle	Country	IS	O country code		
	Identification			Commercial document reference				
	I.18 Transport conditions Ambient			☐ Chilled ☐ Frozen				
	I.19 Container number/Seal nu				2 110			
	Container No		Seal No					
	I.20 Certified as or for							
	☐ Products for huma	an consumption						
	I.21			I.22				
				1.23				
	I.24 Total number of packages		I.25		I.26 Total net weight/gro	ss weight (kg)		

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27 Description	n 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	Date of collection /production	Manufacturing plant	Approval or regiplant/establi	istration number of ishment/centre	Final consume
CN code	Species	Cold store	Identification mark	Type of packaging	Net weight	Slaughterhouse	Treatment type
Nature of commodity	Number of packages	Batch No	Date of collection /production	Manufacturing plant	Approval or reg plant/establ	istration number of ishment/centre	Final consume
CN code	Species	Cold store	Identification mark	Type of packaging	Net weight	Slaughterhouse	Treatment type
Nature of commodity	Number of packages	Batch No	Date of collection /production	Manufacturing plant		istration number of ishment/centre	Final consume
CN code	Species	Cold store	Identification mark	Type of packaging	Net weight	Slaughterhouse	Treatment typ
Nature of commodity	Number of packages	Batch No	Date of collection /production	Manufacturing plant		istration number of ishment/centre	Final consume
CN code	Species	Cold store	Identification mark	Type of packaging	Net weight	Slaughterhouse	Treatment typ
Nature of commodity	Number of packages	Batch No	Date of collection /production	Manufacturing plant		istration number of ishment/centre	Final consume

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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 Council¹, Regulation (EC) No 852/2004 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) 2019/627³ 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 being listed as an EU 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) 2015/1375⁴, and in particular, has been subject to an examination by a digestion method for *Trichinella* with negative results;
- 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;
- (1) 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 2073/2005⁵;
- II.1.7. the meat 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 equine animals 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 solipeds:
 - (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:
 - the rapeutic treatment, as defined in Article 1(2), point (b), of Council Directive $96/22/EC^6$, where applied in conformity with Article 4(2) of that Directive, 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; 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 Council Directive 96/23/EC⁷ which covers equine born in and imported into the third country and was approved in accordance with Article 29(1), fourth subparagraph, of Directive 96/23/EC and the concerned animals and products are listed in Commission Decision 2011/163/EU⁸ for the concerned country of origin.

Regulation (EC) No 178/2002 of the European Parliament and of the Council of 28 January 2002 laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety (OJ L 31, 1.2.2002, p. 1).

Regulation (EC) No 852/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs (OJ L 139, 30.4.2004, p. 1).

Commission Implementing Regulation (EU) 2019/627 of 15 March 2019 laying down uniform practical arrangements for the performance of official controls on products of animal origin intended for human consumption in accordance with Regulation (EU) 2017/625 of the European Parliament and of the Council and amending Commission Regulation (EC) No 2074/2005 as regards official controls (OJ L 131, 17.5.2019, p. 51).

Commission Implementing Regulation (EU) 2015/1375 of 10 August 2015 laying down specific rules on official controls for Trichinella in meat (OJ L 212, 11.8,2015, p. 7).

Commission Regulation (EC) No 2073/2005 of 15 November 2005 on microbiological criteria for foodstuffs (OJ L 338, 22.12.2005, p. 1).

⁶ Council Directive 96/22/EC of 29 April 1996 concerning the prohibition on the use in stockfarming of certain substances having a hormonal or thyrostatic action and of beta-agonists, and repealing Directives 81/602/EEC, 88/146/EEC and 88/299/EEC (OJ L 125, 23.5.1996, p. 3).

Council Directive 96/23/EC of 29 April 1996 on measures to monitor certain substances and residues thereof in live animals and animal products and repealing Directives 85/358/EEC and 86/469/EEC and Decisions 89/187/EEC and 91/664/EEC (OJ L 125, 23.5.1996, p. 10).

Commission Decision 2011/163/EU of 16 March 2011 on the approval of plans submitted by third countries in accordance with Article 29 of Council Directive 96/23/EC (OJ L 70, 17.3.2011, p. 40).

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II.1.8. the meat has 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 Council⁹, and the maximum levels for contaminants laid down in Commission Regulation (EC)
No 1881/2006¹⁰:

II.1.9. 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.

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 European Union in this certificate include the United Kingdom in respect of Northern Ireland.

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.

The exclusion of minced meat and mechanically separated meat is expressly mentioned in the title to avoid any confusion as these products cannot be imported using this fresh meat certificate. This certificate is meant for fresh meat, excluding 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: Use the appropriate Harmonised System (HS) code: 02.05, 02.06 or 05.04.

Box reference I.27: Description of consignment:

"Nature of commodity": Indicate "carcase-whole", "carcase-side", "carcase-quarters" 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) Keep as appropriate.

Official veterinarian

Name (in capital letters)

Date Qualification and title

Stamp Signature

Regulation (EC) No 396/2005 of the European Parliament and of the Council of 23 February 2005 on maximum residue levels of pesticides in or on food and feed of plant and animal origin and amending Council Directive 91/414/EEC (OJ L 70, 16.3.2005, p. 1).

Commission Regulation (EC) No 1881/2006 of 19 December 2006 setting maximum levels for certain contaminants in foodstuffs (OJ L 364, 20.12.2006, p. 5).