UNITED KINGDOM **Animal health/Official certificate to the EU**

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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II.a Certificate reference

I.27 Description of consignment

1

CN code	Species	Cold store	Identification mark	Type of packaging	Net weight	Treatment type
Nature of commodity	Number of packages	Batch No	Date of collection/production	Manufacturing plant	Approval or registration number of plant/establishment/centre	Final consumer ☐

2

CN code	Species	Cold store	Identification mark	Type of packaging	Net weight	Treatment type
Nature of commodity	Number of packages	Batch No	Date of collection/production	Manufacturing plant	Approval or registration number of plant/establishment/centre	Final consumer ☐

3

CN code	Species	Cold store	Identification mark	Type of packaging	Net weight	Treatment type
Nature of commodity	Number of packages	Batch No	Date of collection/production	Manufacturing plant	Approval or registration number of plant/establishment/centre	Final consumer ☐

4

CN code	Species	Cold store	Identification mark	Type of packaging	Net weight	Treatment type
Nature of commodity	Number of packages	Batch No	Date of collection/production	Manufacturing plant	Approval or registration number of plant/establishment/centre	Final consumer ☐

5

CN code	Species	Cold store	Identification mark	Type of packaging	Net weight	Treatment type
Nature of commodity	Number of packages	Batch No	Date of collection/production	Manufacturing plant	Approval or registration number of plant/establishment/centre	Final consumer ☐

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II.a Certificate reference

Part II: Certification

II. Health information

II.1. Public health attestation [delete when the Union is not the final destination of the raw milk]

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[A], Regulation (EC) No 852/2004 of the European Parliament and of the Council[B], Regulation (EC) No 853/2004 of the European Parliament and of the Council[C] and Regulation (EU) 2017/625 of the European Parliament and of the Council[D] and Commission Implementing Regulation (EU) 2019/627[E] and hereby certify that the raw milk described in Part I was produced in accordance with these requirements, and in particular that:

(a) it comes from holdings registered in accordance with Regulation (EC) No 852/2004 and checked in accordance with Articles 49 and 50 of Implementing Regulation (EU) 2019/627;

(b) it was produced, collected, cooled, stored and transported in accordance with the hygiene conditions laid down in Annex III, Section IX, Chapter I, to Regulation (EC) No 853/2004;

(c) it meets the plate and somatic cell count criteria laid down in Annex III Section IX, Chapter I, to Regulation (EC) No 853/2004;

(d) it comes from animals belonging to herds free or officially free of brucellosis and tuberculosis;

(e) the guarantees on the residues status of raw milk provided by the monitoring plans for the detection of residues or substances submitted in accordance with Article 29 of Council Directive 96/23/EC[F], are fulfilled and milk is listed in Commission Decision 2011/163/EU[G] for the concerned country of origin;

(f) pursuant to testing for residues of antibacterial drugs carried out by the food business operator in accordance with the requirements of Annex III, Section IX, Chapter I, Part III, point 4,to Regulation (EC) No 853/2004, complies with the maximum residue limits for residues of antibacterial veterinary medicinal products laid down in the Annex to Commission Regulation (EU) No 37/2010[H];

(g) it 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 Council[I], and the maximum levels for contaminants laid down in Commission Regulation (EC) No 1881/2006[J].

II.2. Animal health attestation [Delete when the raw milk is derived from solipeds, leporidae or other wild land mammals others than ungulates]

The **raw milk** described in Part I:

II.2.1. Originates from the **zone** with code/: ……[(2)] which, at the date of issue of this animal health/official certificate isauthorised for the entry into the Union of raw milk and listed in Part 1 of Annex XVII to Commission Implementing Regulation (EU) 2021/404[K], and in which foot and mouth disease and infection with rinderpest virus have not been reported for the period of 12 months prior to the date of milking, and vaccination against these diseases has not been carried out during the same period.

II.2.2. has been obtained from **animals** of the species [Bos Taurus,] [(1)] [Ovis aries,] [(1)] [Capra hircus,] [(1)] [Bubalus bubalis,] [(1)] [Camelus dromedarius] [(1)] that:

[(1) either] [have remained in the zone referred to under point II.2.1. since birth, or for the period of at least 3 months prior to the date of milking;]

[(1) and/or] [were introduced in the zone referred to under point II.2.1. from:

A 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).

B 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).

C Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific hygiene rules for food of animal origin *(OJ L 139, 30.4.2004, p. 55).*

D Regulation (EU) 2017/625 of the European Parliament and of the Council of 15 March 2017 on official controls and other official activities performed to ensure the application of food and feed law, rules on animal health and welfare, plant health and plant protection products, amending Regulations (EC) No 999/2001, (EC) No 396/2005, (EC) No 1069/2009, (EC) No 1107/2009, (EU) No 1151/2012, (EU) No 652/2014, (EU) 2016/429 and (EU) 2016/2031 of the European Parliament and of the Council, Council Regulations (EC) No 1/2005 and (EC) No 1099/2009 and Council Directives 98/58/EC, 1999/74/EC, 2007/43/EC, 2008/119/EC and 2008/120/EC, and repealing Regulations (EC) No 854/2004 and (EC) No 882/2004 of the European Parliament and of the Council, Council Directives 89/608/EEC, 89/662/EEC, 90/425/EEC, 91/496/EEC, 96/23/EC, 96/93/EC and 97/78/EC and Council Decision 92/438/EEC (*OJ L 95, 7.4.2017, p. 1).*

E 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).

F 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).

G 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).

H Commission Regulation (EU) No 37/2010 of 22 December 2009 on pharmacologically active substances and their classification regarding maximum residue limits in foodstuffs of animal origin (OJ L 15, 20.1.2010, p. 1).

I 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).

J 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).

K Commission Implementing Regulation (EU) 2021/404 of 24 March 2021 laying down the lists of third countries, territories or zones thereof from which the entry into the Union of animals, germinal products and products of animal origin is permitted in accordance with Regulation (EU) 2016/429 of the European Parliament and of the Council (OJ L 114, 31.3.2021, p. 1).

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II.a Certificate reference

(1) either [another third country or territory, or zone thereof which is listed for the entry into the Union of raw milk, colostrum or colostrum-based products and the animals remained there for the period of at least 3 months prior to the date of milking;]]

(1) and/or [a Member State;]]

II.2.3. has been obtained from animals coming from **establishments**:

(a) registered by and under the control of the competent authority of the third country or territory and have a system in place to maintain and to keep records in accordance with Article 8 of Commission Delegated Regulation (EU) 2020/692[L];

(b) which receive regular animal health visits from a veterinarian for the purpose of the detection of, and information on, signs indicative of the occurrence of diseases, including the relevant listed diseases referred to in Annex I to Commission Delegated Regulation (EU) 2020/692 and emerging diseases;

(c) which were not subject to national restriction measures for animal health reasons, including the relevant listed diseases referred to in Annex I to Delegated Regulation (EU) 2020/692 and emerging diseases, at the date of milking.

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 is intended for the entry into the Union of milk, including when the Union is not the final destination of such raw milk.

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

Part I:

Box reference I.8:	Provide the code of the zone as appearing in column 2 of the table in Part 1 of Annex XVII to Implementing Regulation (EU) 2021/404.
Box reference I.11:	Name, address and approval number of the establishment of dispatch.
Box reference I.15:	Registration number (railway wagons or container and road vehicle), flight number (aircraft) or name (vessel) must be provided. In the case of unloading and reloading, the consignor must inform the border control post of the entry into the Union.
Box reference I.19:	For the containers or boxes, the container number and the seal number (if applicable) shall be included.
Box reference I.27:	Use the appropriate Harmonised System (HS) code under the following headings: 04.01; 04.02 or 04.03. Description of consignment: "*Manufacturing plant*": Introduce the approval number of the production holding(s), collection centre or standardization centre approved for exportation to the European Union.

Part II:

(1) Keep as appropriate.

(2) Code of the zone in accordance with column 2 of the table in Part 1 of Annex XVII to Implementing Regulation (EU) 2021/404.

(3) to be signed by :
- an official veterinarian when part II.2 Animal health attestation is not deleted
- a certifying officer or an official veterinarian when part II.2 Animal health attestation is deleted

[Official veterinarian][(1)(3)]/[Certifying officer][(1)(3)]

Name (in capital letters)

Date

Qualification and title

Stamp

Signature