Model health certificate for dairy products derived from milk of cows, ewes, goats and buffaloes for human consumption from third countries or parts thereof authorised in column B from non-EU countries

GBHC066X v3.1 August 2022

Part I. Details o	t dispatcr	iea cons					
I.1 Consignor			I.2 Cer	rtificate re	ference no.	I.3 Cent	ral competent authority
Name:							
Address:			L2.a N	I.2.a Not in use		L4 Loca	Il competent authority
			1121011	01 111 000		2000	a compositing dutility
Tel:							
I.5 Consignee					I.6 Not in use		
Name:							
Address:							
Tel:							
I.7 Country of	ISO	I.8 Regi		Code	I.9 Country of	ISO	I.10 Not in use
origin	code	origi	n		destination	code	
I.11 Place of or	igin				I.12 Not in use		
Name:							
Approval number	er:						
Address:							
I.13 Place of loa	ading				I.14 Date of depart	arture	
I.15 Means of to	ransport				I.16 Entry BCP		
☐ Aeroplane	•				, ,		
☐ Ship							
☐ Railway wagon							
☐ Road vehicle					I.17 Not in use		
Other					I.II NOUTH USE		
Identification:							
Documentation references:							

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II.a. Certificate reference no.	II.b.

I.18 Description of commodity							
I.19 Commodity code (HS code)	I.21 Temperature of products			I.23 Seal / Container No.			
	☐ Ambient	Ambient					
	Chilled						
	Frozen					4	
I.20 Quantity	I.22 Number	I.22 Number of packag		I.24 Type of packaging		ng	
I.25 Commodity certified for	<u> </u>			I			
☐ Human consumption							
I.26 Not in use			I.27 For imp	ort or adm	ission into C	Freat Britain	
I.28 Identification of the commodities							
Manufacturing Plant	Number of Packages		Species (Scientific na	me)	Net Weight	Batch Number	
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Part II. Certification

II.1 Animal health attestation

I, the undersigned official veterinarian, declare that I am aware of the relevant provisions of Directive 2002/99/EC and of Regulation (EC) No 853/2004 and hereby certify that the dairy product described above:

- (a) has been obtained from animals:
 - (i) under the control of the official veterinary service,
 - which were in a country or part thereof that has been free of foot-and-mouth disease and of rinderpest for a period of at least 12 months prior to the date of this certificate, and where vaccination against foot-and-mouth disease has not been carried out during that period,
 - (iii) belonging to holdings which were not under restrictions due to foot-and-mouth disease or rinderpest, and,
 - (iv) subject to regular veterinary inspections to ensure that they satisfy the animal health conditions laid down in Chapter I of Section IX of Annex 3 to Regulation (EC) No 853/2004 and in Directive 2002/99/EC,

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II.a. Certificate reference no.	II.b.

(b) has undergone or been produced from raw milk which has been submitted to a pasteurisation treatment involving a single heat treatment with a heating effect at least equivalent to that achieved by a pasteurisation process of at least 72°C for 15 seconds and, where applicable, sufficient to ensure a negative reaction to an alkaline phosphatase test applied immediately after the heat treatment.

II.2 Public health attestation

I, the undersigned official veterinarian, declare that I am aware of the relevant provisions of Regulations (EC) No 178/2002, (EC) No 852/2004, (EC) No 853/2004 and (EU) 2019/627 and hereby certify that the dairy product described above was produced in accordance with those provisions, in particular that:

- (a) it was manufactured from raw milk:
 - (i) which comes from holdings registered in accordance with Regulation (EC) No 852/2004 and checked in accordance with Article 49-50 of Regulation (EU) 2019/627,
 - (ii) which was produced, collected, cooled, stored and transported in accordance with the hygiene conditions laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004.
 - which meets the plate and somatic cell count criteria laid down in Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004,
 - (iv) which complies with 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 Council Directive 96/23/EC, and in particular, Article 29 thereof,
 - (v) which, pursuant to testing for residues of antibacterial drugs carried out by the food business operator in accordance with the requirements of Chapter I of Section IX of Annex III to Regulation (EC) No 853/2004, it complies with the maximum residue limits for residues of antibacterial veterinary medicinal products laid down in the Annex to Regulation (EU) No 37/2010,
 - (vi) which has been produced under conditions guaranteeing compliance with the maximum residue levels for pesticides laid down in Regulation (EC) No 396/2005, and maximum levels for contaminants laid down in Regulation (EC) No 1881/2006.
- (b) it comes from an establishment implementing a programme based on the HACCP principles in accordance with Regulation (EC) No 852/2004,
- (c) it has been processed, stored, wrapped, packaged and transported in accordance with the relevant hygiene conditions laid down in Annex II to Regulation (EC) No 852/2004 and Chapters II and III of Section IX of Annex III to Regulation (EC) No 853/2004;
- (d) meets the relevant criteria laid down in Chapter II of Section IX of Annex III to Regulation (EC) No 853/2004 and the relevant microbiological criteria laid down in Regulation (EC) No 2073/2005 on microbiological criteria for foodstuffs,
- the guarantees covering live animals and products thereof provided by the residue plans submitted in accordance with Directive 96/23/EC, and in particular Article 29 thereof, are fulfilled.

Notes

References to European Union legislation within this certificate are references to direct EU legislation which has been retained in Great Britain (retained EU law as defined in the European Union (Withdrawal) Act 2018) and can be viewed on the UK legislation website (legislation.gov.uk).

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II.a. Certificate reference no.	II.b.

References to Great Britain in this certificate include Channel Islands and Isle of Man.

This certificate is intended for dairy products for human consumption from third countries or parts thereof authorised in Column B as set out in a document relating to 'milk and milk products' published on gov.uk, in accordance with EU Retained Regulation 605/2010 intended for importation into Great Britain.⁽¹⁾

Part I:

Box reference I.7: Provide name and ISO code of the country or part thereof as set out in a

document relating to 'milk and milk products' published on gov.uk, in accordance

with Regulation (EU) No 605/2010.(1)

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 vehicles), flight

number (aircraft) or name (ship). In the case of transport in containers, the total number of containers and their registration number and where there is a serial number of the seal it must be indicated in box I.23. In the case of unloading and reloading, the consignor must inform the border control post of introduction into

Great Britain.

Box reference I.19 Use the appropriate Harmonised System (HS) code under the following

headings: 04.01; 04.02; 04.03; 04.04; 04.05; 04.06; 15.17; 17.02; 19.01; 21.05;

21.06; 22.02; 28.35; 35.01; 35.02 or 35.04.

Box reference I.20 Indicate total gross weight and total net weight.

Box reference I.23 For containers or boxes, the container number and the seal number (if

applicable) should be included.

Box reference I.28 Manufacturing plant: introduce the approval number of the treatment and/or

processing establishment(s) approved for export to Great Britain.

Part II:

A document relating to 'milk and milk products' for non-EU countries published by the Secretary of State, with the consent of the Scottish and Welsh Ministers, may be found here:

Non-EU countries approved to export animals and animal products to Great Britain - data.gov.uk

The colour of the signature shall be different to that of the printing. The same rule applies to stamps other than those embossed or watermark.

Official Veterinarian	
Name (in capital letters):	Qualification and title:
Date:	Signature:
Stamp:	