

Part I : Details of consignment	I.1. Consignor		I.2. IMSOC Reference	
	Name		I.2.a. Local Reference	
	Address			
	Country	ISO Code		
	I.5. Consignee		I.3. Central competent authority	
	Name		I.4. Local competent authority	
	Address			
	Country	ISO Code		
	I.7. Country of origin	ISO Code	I.9. Country of destination	ISO Code
	I.8. Region of origin	Code	I.10. Region of destination	
	I.11. Place of Dispatch		I.12. Place of destination	
	Name		Name	
	Address		Address	
	Approval Number		Approval Number	
Country	ISO Code	Country	ISO Code	
I.13. Place of Loading		I.14. Date and time of departure		
Name				
Address				
Approval Number				
Country	ISO Code			
I.15. Means of Transport		I.16 Entry Point		
Mode	International transport document	Identification		
I.18. Transport conditions		I.17. Accompanying documents		
Frozen <input type="checkbox"/>	Chilled <input type="checkbox"/>	Ambient <input type="checkbox"/>	Controlled temperature <input type="checkbox"/>	
		Accompanying document reference	Date of issue	
		Country	Place of issue	
I.19. Container No / Seal No				
I.20. Certified as				
Human consumption <input type="checkbox"/>				
I.21. For transit through a third country <input type="checkbox"/>		I.22. For transit through Member State(s) <input type="checkbox"/>		
Country	ISO Code			
EU Exit Authority	BCP code	Country	ISO Code	
EU Entry Authority	BCP code			
I.23. Total number of packages	I.25. Total net weight	I.25. Total gross weight		
I.28. Description of consignment				
1. 21 MISCELLANEOUS EDIBLE PREPARATIONS				
2106 Food preparations not elsewhere specified or included				
Commodity	Batch number	Manufacturing plant	Package count	
Species				

Part II: Certification	II. Health information		
	<p>II.1. Animal Health Attestation</p> <p>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:</p> <p>(a) has been obtained from animals:</p> <p>(i) under the control of the official veterinary service,</p> <p>(ii) 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,</p> <p>(iii) belonging to holdings which were not under restrictions due to foot-and-mouth disease or rinderpest, and,</p> <p>(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,</p> <p>(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.</p> <p>II.2. Public Health attestation</p> <p>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:</p> <p>(a) it was manufactured from raw milk :</p> <p>(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,</p> <p>(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,</p> <p>(iii) 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,</p> <p>(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,</p> <p>(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,</p> <p>(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.</p> <p>(b) it comes from an establishment implementing a programme based on the HACCP principles in accordance with Regulation (EC) No 852/2004,</p> <p>(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;</p>		

II. Health information	
(d)	it 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,
(e)	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).

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

- 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.
 - 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.19. In the case of unloading and reloading, the consignor must inform the border control post of introduction into Great Britain.
 - Box reference I.19: For containers or boxes, the container number and the seal number (if applicable) should be included.
 - Box reference I.25: Indicate total gross weight and total net weight.
 - Box reference I.28: 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.28: Manufacturing plant: introduce the approval number of the treatment and/or processing establishment(s) approved for export to Great Britain.

Part II:

(1) A document relating to 'milk and milk products' for EU and EFTA states published by the Secretary of State, with the consent of the Scottish and Welsh Ministers, may be found here: data.gov.uk

EU and EFTA states 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.

Certifying Officer	
Name (in capital letters)	Qualification and title
Date of signature	Signature
Stamp	

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