

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			I.9. Country of destination		
				ISO Code		
	I.8. Region of origin			I.10. Region of destination		
	Code					
	I.11. Place of Dispatch			I.12. Place of destination		
Name			Name			
Address			Address			
Approval Number			Approval Number			
Country			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			
Ambient <input type="checkbox"/> Chilled <input type="checkbox"/> Frozen <input type="checkbox"/> Controlled temperature <input type="checkbox"/>			Commercial document reference			
			Date of issue			
			Country			
			Place of issue			
I.19. Container No / Seal No						
I.20. Certified as						
Further process <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			Country			
ISO Code			ISO Code			
EU Exit Authority			BCP 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. 04 DAIRY PRODUCE; BIRDS' EGGS; NATURAL HONEY; EDIBLE PRODUCTS OF ANIMAL ORIGIN, NOT ELSEWHERE SPECIFIED OR INCLUDED						
0401 Milk and cream, not concentrated nor containing added sugar or other sweetening matter						
Commodity	Batch number	Package count	Manufacturing plant	Species		
Net weight						

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 raw milk described above has been obtained from animals:</p> <ul style="list-style-type: none">(a) under the control of the official veterinary service,(b) 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,(c) belonging to holdings which were not under restrictions due to foot-and-mouth disease or rinderpest, and(d) 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>II.2. Public Health attestation</p> <p>I, the undersigned official inspector, 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 raw milk described above was produced in accordance with those provisions, in particular that:</p> <ul style="list-style-type: none">(a) it 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(b) it 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,(c) it meets the plate and somatic cell count criteria laid down in Chapter I of Section IX of Annex 3 to Regulation (EC) No 853/2004,(d) 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, are fulfilled;(e) 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 of 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;(f) it 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.		

Part II: Certification	II. Health information			
	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.			
	(e) pursuant to testing for residues of antibacterial drugs carried out by the food business operator in accordance with the requirements of Annex III, +B21+B21+B21 Section IX, Chapter I, Part III, point 4 of 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;			
	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' as 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 vehicle), flight number (aircraft) or name (ship). In 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 or 04.03.			
Box reference I.28:	Manufacturing plant: introduce the approval number of the production holding(s), collection centre or standardization centre approved for exportation 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: 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				