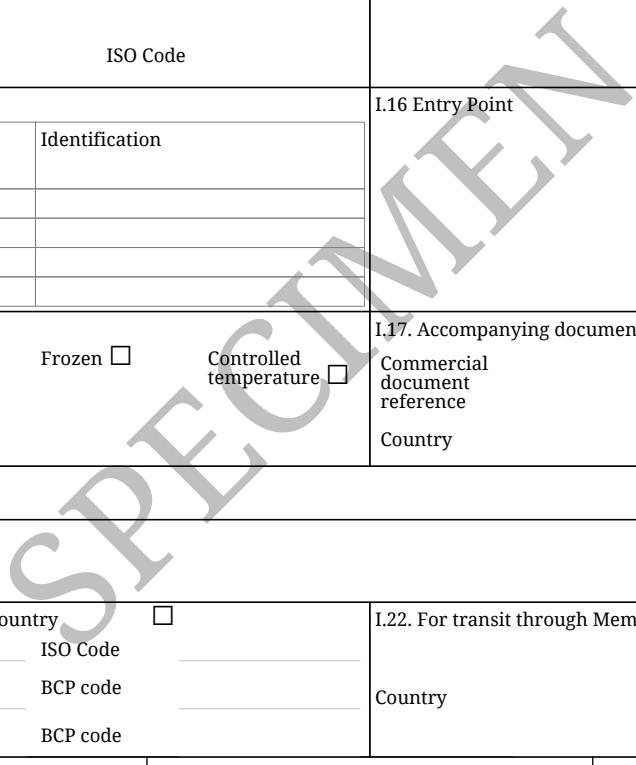


Part I : Details of consignment	I.1. Consignor		I.2. IMSOC Reference		
	Name		<b>Specimen not to be used for exports from EU</b>		
	Address		I.2.a. Local Reference		
	Country	ISO Code			
	I.5. Consignee		I.3. Central competent authority		
	Name		I.4. Local competent authority		
	Address				
	Country	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	
				Code	
	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			
Chilled <input type="checkbox"/> Ambient <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 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	Country		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					
<b>1. 04 DAIRY PRODUCE; BIRDS' EGGS; NATURAL HONEY; EDIBLE PRODUCTS OF ANIMAL ORIGIN, NOT ELSEWHERE SPECIFIED OR INCLUDED</b>					
<b>0404</b> Whey, whether or not concentrated or containing added sugar or other sweetening matter; products consisting of natural milk constituents, whether or not containing added sugar or other sweetening matter, not elsewhere specified or included					
Commodity	Manufacturing plant	Package count	Species	Net weight	
Batch number					



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 III 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 (EC) No 854/2004 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 Annex III, Section IX, Chapter 1, 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,</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 Chapter II of Section IX of Annex III to Regulation (EC) No 853/2004,</p>		

Part II: Certification	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		
(*)Those countries subject to the transitional import arrangements include: an EU member State; Liechtenstein; Norway; Iceland and Switzerland.		
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).		
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 appearing in Annex I to 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.23. In the case of unloading and reloading, the consignor must inform the border control post of introduction into Great Britain.		
Box reference I.16: Do not use this box until the end of the transitional staging period.		
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; 17.02; 21.05; 22.02; 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:		
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		