

Part I : Details of consignment

I.1. Consignor Name _____ Address _____ Country _____ ISO Code _____			I.2. IMSOC Reference I.2.a. Local Reference _____																	
I.5. Consignee Name _____ Address _____ Country _____ ISO Code _____			I.3. Central competent authority I.4. Local competent authority _____																	
I.7. Country of origin _____ ISO Code _____		I.9. Country of destination _____ ISO Code _____		I.10. Region of destination																
I.8. Region of origin _____ Code _____			I.10. Region of destination																	
I.11. Place of Dispatch Name _____ Address _____ Approval Number _____ Country _____ ISO Code _____			I.12. Place of destination Name _____ Address _____ Approval Number _____ Country _____ ISO Code _____																	
I.13. Place of Loading Name _____ Address _____ Approval Number _____ Country _____ ISO Code _____			I.14. Date and time of departure _____																	
I.15. Means of Transport <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 20%;">Mode</td> <td style="width: 20%;">International transport document</td> <td style="width: 60%;">Identification</td> </tr> <tr><td> </td><td> </td><td> </td></tr> <tr><td> </td><td> </td><td> </td></tr> <tr><td> </td><td> </td><td> </td></tr> <tr><td> </td><td> </td><td> </td></tr> </table>			Mode	International transport document	Identification													I.16 Entry Point _____		
Mode	International transport document	Identification																		
I.18. Transport conditions Frozen <input type="checkbox"/> Chilled <input type="checkbox"/> Controlled temperature <input type="checkbox"/> Ambient <input type="checkbox"/>			I.17. Accompanying documents Document Type _____ Accompanying 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"/> Country _____ ISO Code _____ EU Exit Authority _____ BCP code _____ EU Entry Authority _____ BCP code _____			I.22. For transit through Member State(s) <input type="checkbox"/> Country _____ ISO 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 0403 Buttermilk, curdled milk and cream, yogurt, kephir and other fermented or acidified milk and cream, whether or not concentrated or containing added sugar or other sweetening matter or flavoured or containing added fruit, nuts or cocoa																				
#1.	Commodity	Batch number	Package count	Manufacturing plant	Net weight															
	Species																			

II. Health information

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 raw milk described above has been obtained from animals:

- (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;

II.2. Public Health attestation

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:

- (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.

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.

Part II: Certification

II. Health information

- 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)

Date of signature

Stamp

Qualification and title

Signature

Μέρος I

I.1. Αποστολέας Όνομα Διεύθυνση Χώρα Κωδικός ISO			I.2. Κωδικός αναφοράς IMSOC I.2.a. Local Reference		
I.5. Παραλήπτης Όνομα Διεύθυνση Χώρα Κωδικός ISO			I.3. Κεντρική αρμόδια αρχή (ΚΑΑ) I.4. Local competent authority		
I.7. Χώρα προέλευσης		Κωδικός ISO	I.9. Country of destination		Κωδικός ISO
I.8. Region of origin			I.10. Περιφέρεια προορισμού		
I.11. Place of Dispatch Όνομα Διεύθυνση Αριθμός έγκρισης Χώρα Κωδικός ISO			I.12. Τόπος προορισμού Όνομα Διεύθυνση Αριθμός έγκρισης Χώρα Κωδικός ISO		
I.13. Τόπος φόρτωσης Όνομα Διεύθυνση Αριθμός έγκρισης Χώρα Κωδικός ISO			I.14. Date and time of departure		
I.15. Μέσο μεταφοράς Τύπος Έγγραφο Ταυτοποίηση			I.16 Entry Point		
I.18. Transport conditions Κατεψυγμένα <input type="checkbox"/> Σε ψύξη <input type="checkbox"/> Controlled temperature <input type="checkbox"/> σε θερμοκρασία περιβάλλοντος <input type="checkbox"/>			I.17. Συνοδευτικά έγγραφα Document Type Κωδικός αναφοράς του εμπορικού εγγράφου Ημερομηνία έκδοσης Χώρα Τόπος έκδοσης		
I.19. Εμπορευματοκιβώτιο αριθ./ Σφραγίδα αριθ.					
I.20. Certified as Περαιτέρω επεξεργασία <input type="checkbox"/>					
I.21. For transit through a third country <input type="checkbox"/> Country EU Exit Authority EU Entry Authority Κωδικός ISO BCP code BCP code			I.22. For transit through Member State(s) <input type="checkbox"/> Country Κωδικός ISO		
I.23. Συνολικός αριθμός δεμάτων		I.25. Συνολικό καθαρό βάρος		I.25. Συνολικό μεικτό βάρος	
I.28. Description of consignment 1. 04 ΓΑΛΑ ΚΑΙ ΠΡΟΪΟΝΤΑ ΓΑΛΑΚΤΟΚΟΜΙΑΣ. ΑΥΓΑ ΠΤΗΝΩΝ. ΜΕΛΙ ΦΥΣΙΚΟ. ΠΡΟΪΟΝΤΑ ΒΡΩΣΙΜΑ ΖΩΙΚΗΣ ΠΡΟΕΛΕΥΣΗΣ, ΠΟΥ ΔΕΝ ΚΑΤΟΝΟΜΑΖΟΝΤΑΙ ΟΥΤΕ ΠΕΡΙΛΑΜΒΑΝΟΝΤΑΙ ΑΛΛΟΥ 0403 Βουτυρόγαλα, πηγμένο γάλα και πηγμένη κρέμα, γιαούρτι, κεφίρ και άλλα γάλατα και κρέμες που έχουν υποστεί ζύμωση ή έχουν καταστεί όξινα, έστω και συμπυκνωμένα ή με προσθήκη ζάχαρης ή άλλων γλυκαντικών, ή αρωματισμένα ή με προσθήκη φρούτων ή κακαού					
#1.	Εμπόρευμα	Αριθμός παρτίδας	Πλήθος πακέτων	Μονάδα μεταποίησης	Καθαρό βάρος
Είδος					

Part II: Certification	II. Υγειονομικές πληροφορίες							
	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 raw milk described above has been obtained from animals: <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; 						
	II.2.	Public Health attestation 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: <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. 						
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	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		
Ημερομηνία υπογραφής		Υπογραφή		
Σφραγίδα				