Norway

HEALTH CERTIFICATE

Covering export of raw milk from third countries or parts thereof authorised in column A of Annex I intended for further processing before being used for human consumption from Norway* to GB GBHC064E

Original ☐ Replacement ☐

Part I: Details of dispatched consignment

I.1. Consignor Name, Address and Tel.		I.2. Certificate reference number	I.2.a. Original certificate number
		I.3. Central Competent Authority NORWEGAIN FOOD SAFETY AUTHORITY, N-2381 BRUMUNDDAL, NORWAY. E-mail: postmottak@mattilsynet.no, Phone: +47 22400000	
		I.4. Local Competent Authority NORWEGIAN FOOD SAFETY AUTHORITY, REGIONAL OFFICE	
I.5. Consignee Name, Address, Postcode and Tel.		I.6.	
I.7. Country of origin	ISO code	I.8. Region of origin	Code
		I.9. Country of destination	ISO code
		I.10.	
I.11. Place of origin Name, Address and Approval number		I.12.	
I.13. Place of loading		I.14. Date of departure	
I.15. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification: Documentary references:		I.16. Entry BCP in Great Britain, Channel Islands or Isle of Man	
		I.17.	

I.18. Description of commodity		I.19. Commodity code (HS code)
I.21. Temperature of products Ambient ☐ Chilled ☐ Frozen ☐	I.20 Quantity	I.22 Number of packages
I.23. Seal/Container No.		I.24. Type of packaging
I.25. Commodities certified for: Human Consumption ☐		
I.26.		I.27. For import or admission into Great Britain, Channel Islands or Isle of Man ☐

Original ☐ Replacement ☐

	I.2. Certificate reference number	I.2.a. Original certificate no.

I.28. Identification of the commodities

Manufacturing plant (approval number)	Number of packages	Species (scientific name)	Net weight	Batch Number

Original ☐ Replacement ☐

Part II: Certification

II. Health information	II.a. Certificate reference number	II.b. Original certificate number

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) 2019/627and 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.

Original ☐ Replacement ☐

II. Health information	II.a. Certificate reference number	II.b. Original certificate number

Notes

(*) Those countries subject to the transitional import arrangements include: an EU member State; Iceland; Liechtenstein; Norway and Switzerland.

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

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).

This certificate is intended for raw milk from third countries or parts thereof authorised in column A of Annex 1 to Regulation (EU) No 605/2010 intended for further processing in Great Britain before being used for human consumption.

Part I

- Box reference I.7: Provide name and ISO code of the country or part thereof as appearing in Annex 1 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 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.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 or 04.03.
- 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 production holding(s), collection centre or standardization centre approved for exportation 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.

Official veterinarian

Qualification and title:

Name (in capital letters):

Signature:

Date:

Stamp: