Export Health Certificate

| | I.1. Consignor | | | | I.2. IMSOC Refe | erence | | | |
|--------------------------------|---|--|----------------------------------|---|---|------------|--|-------------------------------------|--|
| | Name | | I.2.a. Local Reference | | | | | | |
| | Address | | | | | | | | |
| | Country | | | | | | | | |
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| | I.S. Consignee | | | | I.3. Central competent authority | | | | |
| nt | Name | | | | I.4. Local competent authority | | | | |
| ne | Address | | ICO Cod | d. | | | | | |
| Щ | Country | | ISO Cod | ie – | | | | | |
| sig | I.7. Country of orig | in | | ISO Code | I.9. Country of destination ISO Code | | | ISO Code | |
| on | | | | | | | | | |
| f c | I.8. Region of origin | n | | Code | I.10. Region of | destinatio | nn. | | |
| S O | I.11. Place of Dispa | | | couc | I.12. Place of d | | | | |
| ail | Name | | | | Name | 000111011 | | | |
|)et | Address | | | | Address | | | | |
| $: \Gamma$ | Approval Number | | | | Approval Number | | | | |
| τI | Country | | ISO | Code | Country | ibei | | ISO Code | |
| Part I: Details of consignment | | | | | | | | | |
| Η | I.13. Place of Loadi | ng | | | I.14. Date and | time of de | parture | | |
| | Name | | | | | | | | |
| | Address | | | | | | | | |
| | Approval Number | • | | | | | | | |
| | Country | | ISO | Code | | | | | |
| | I.15. Means of Trai | nenort | | | I.16 Entry Poin | t | | | |
| | Mode | International | Identificati | on | 1.10 Entry 1 on | | | | |
| | Wode | transport | luelillicati | OII | | | | | |
| | | document | | | _ | | | | |
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| | I.18. Transport con | ditions | | | I 17 Accompar | wing door | ımonte | | |
| | | Controlled | Chilled \square | Ambient \square | I.17. Accompanying documents Commercial | | | | |
| | 110Zell 🗀 | temperature \square | симси 🗀 | Antibient 🗀 | document Date of issue | | | | |
| | tomporature <u>—</u> | | | | reference | | | | |
| | | | | | | | 70.1 | c | |
| | | | | | Country | | Place o issue | of | |
| | I.19. Container No | / Seal No | | | Country | | | of | |
| | I.19. Container No | / Seal No | | | Country | | | of | |
| | I.20. Certified as | | | | Country | | | of | |
| | | | | | Country | | | of | |
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| | I.20. Certified as Human consumpti I.21. For transit thi | on \square | | | | t through | | of | |
| | I.20. Certified as Human consumpti I.21. For transit the Country | on \square | ISO Code | | | t through | issue | | |
| | I.20. Certified as Human consumpti I.21. For transit the Country EU Exit Authority | on \square | | | | t through | issue | | |
| | I.20. Certified as Human consumpti I.21. For transit the Country EU Exit Authority EU Entry | on \square | ISO Code | | I.22. For transi | t through | issue Member State(s) | | |
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| | II. Health information | | | | | | | | | |
| | II.1. | Animal H | ealth Attesta | ıtion | ı | | | | | |
| | | | e undersigned official veterinarian, declare that I am aware of the relevant provisions of Directive 2/99/EC and of Regulation (EC) No 853/2004 and hereby certify that the dairy product described | | | | | | | |
| _ | | (a) | has been o | obtained fron | n animals: | | | | | |
| | | | (i) | under the control of the official veterinary service; | | | | | | |
| ımıca | | | (ii) | | o holdings v rinderpest: a | which were not under restrict and, | ions due to foot-and-mouth | | | |
| raitii. Cei micauoii | | | (iii) | health cond | ditions laid d | inary inspections to ensure t lown Chapter I of Section IX o ective 2002/99/EC; | | | | |
| | (1)(2)eithe r | ○ [(b) | where aut | the dairy product was made from raw milk sourced from cows, ewes, goats, buffaloes or, where authorised, countries with footnote (b) as set out in a document relating to 'milk and milk products' published on gov.uk, in accordance with Regulation (EC) No 605/2010, from camels of the species Camelus dromedarieus, and has undergone, prior to import into Great | | | | | | |
| | (1) | either | ○ [(i) | a sterilizati | on process, | to achieve an F0 value equal | to or greater than three;] | | | |
| | (1) | or | ○ [(ii) | | | ure (UHT) treatment at not leatable holding time;] | ss than 135 °C in | | | |
| | (1) | or | o [(iii) | seconds app | plied twice t icable, a neg | ort time pasteurisation treatm o milk with a pH equal to or g gative reaction to an alkaline leat treatment;] | greater than 7,0 achieving, | | | |
| | (1) | or | ○ [(iv) | where appl | icable, a neg | uivalent pasteurisation effect gative reaction to an alkaline leat treatment;] | | | | |
| | (1) | or | ○ [(v) | a HTST trea | atment of mi | lk with a pH below 7,0;] | | | | |
| | (1) | or | ○ [(vi) | a HTST trea | atment comb | oined with another physical t | reatment by | | | |
| | (1) | | either | o [(1) | lowering th | e pH below 6 for one hour;] | | | | |
| | (1) | | or | o [(2) | additional h | neating equal to or greater the | an 72 °C, combined with | | | |
| | (1)or | (b) the dairy product was made from goats, buffaloes or camels of the s import into Great Britain: | | | els of the sp | | | | | |
| | (1) | either | ○ [(i) | a sterilizati | on process, | to achieve an F0 value equal | to or greater than three;] | | | |
| | (1) | or | ○ [(ii) | | - | ure (UHT) treatment at not le table holding time;]] | ss than 135 °C in | | | |
| | II.2. | Public Hea | Health attestation | | | | | | | |
| | | (EC) No 17 | undersigned official inspector, declare that I am aware of the relevant provisions of Regulations to 178/2002, (EC) No 852/2004, (EC) No 853/2004 and (EU) 2019/627 and hereby certify that the product described above was produced in accordance with those provisions, and in particular | | | | | | | |
| | | (a) | it was mai | nufactured fr | om raw mil | k: | | | | |
| | | | (i) | | | lings registered in accordance n accordance with Article 49- | _ | | | |
| | | | (ii) | with the hy | | ollected, cooled, stored and tr tions laid down in Chapter I o '2004; | | | | |
| | | | (iii) | | _ | and somatic cell count criteria to Regulation (EC) No 853/200 | - | | | |

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EUROPEAN UNION

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|--------|---------------------|------------------------|------------|---|--|--|--|--|
| | | II. Health information | | | | | | |
| | | | (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; | | | | |
| 100 | ncation | | (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 I, Part III, point 4 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; | | | | |
| it II. | Part II: Ceruncauon | | (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; | | | | |
| 6 | Ĕ | (b) | | rom an establishment implementing a programme based on the HACCP principles ance with Regulation (EC) No 852/2004; | | | | |
| | | (c) | relevant h | n processed, stored, wrapped, packaged and transported in accordance with the hygiene conditions laid down in Annex II to Regulation (EC) No 852/2004 and of Section IX of Annex III to Regulation (EC) No 853/2004; | | | | |
| | | (d) | (EC) No 85 | ne relevant criteria laid down in Chapter II of Section IX of Annex III to Regulation 53/2004 and the relevant microbiological criteria laid down in Regulation (EC) No 5 on microbiological criteria for foodstuffs; | | | | |
| | | (e) | _ | ntees covering live animals and products thereof provided by the residue plans I in accordance with Directive 96/23/EC, and in particular Article 29 thereof, are | | | | |
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| 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 ce be viewed on the UK legislation website (legislation.gov.uk). References to Great Britain in this certificate include Channel Islands and Isle of Man. This certificate is intended for dairy products for human consumption from third countries or parts thereof authorised, where applicable, for milk from certain animal species only, listed in column C in a document relating to 'milk and milk products' published on gov.uk, in accordance with Regulation (EU) No 605/2010 intended for importation into Great Britain.(2) Part I: Box provide name and ISO code of the country or part thereof as set out in a document relating reference 1.11: Name, address and approval number of the establishment of dispatch. Box registration number (railway wagons or container and road vehicles), flight number reference 1.15: number of containers and their registration number and where there is a serial number the seal it must be indicated in box 1.19. In the case of unloading and reloading, the consignor must inform the border control post of introduction into Great Britain. Box for containers or boxes, the container number and the seal number (if applicable) should reference 1.23: Box use the appropriate Harmonised System (HS) code under the following headings: 04.01; reference 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 of 1.28: 35.04. Box use the appropriate Harmonised System (HS) code under the following headings: 04.01; reference 1.28: 35.04. Box use the appropriate Harmonised System (HS) code under the following headings: 04.01; reference 1.28: 35.04. Part II: (1) Keep as appropriate. | J | JROPEAN UNION | (GB) Dairy products from third countries authorised in column C (MIIR HTC) from EU countries GBHC067E (v3.0 | | | | | | | |
|---|--|--|--|--|--|--|--|--|--|--|
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