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| Part I : Details of consignment | I.1. Consignor | | I.2. IMSOC Reference | | |
| | Name | | Specimen not to be used for exports from EU | | |
| | 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 | | | | | |
| 1. 04 DAIRY PRODUCE; BIRDS' EGGS; NATURAL HONEY; EDIBLE PRODUCTS OF ANIMAL ORIGIN, NOT ELSEWHERE SPECIFIED OR INCLUDED | | | | | |
| 0405 Butter and other fats and oils derived from milk; dairy spreads | | | | | |
| Commodity | Species | Manufacturing plant | Package count | Net weight | |
| | | | | | |
| Batch number | | | | | |
| | | | | | |

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| II. Health information | | |
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Part II: Certification

I, the undersigned Official veterinarian hereby certify that:

1. The dairy product described above, which is exported to the Republic of Moldova, 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 requirements of the EU.
2. It has undergone pasteurisation or been produced from raw milk which has been submitted to a pasteurisation treatment fulfilling the provisions of Regulation (EC) No 853/2004 laying down specific rules or requirements for heat treatment.
3. It was manufactured from raw milk:
 - a) which comes from holdings registered in accordance with Regulation (EC) No 852/2004 and checked in accordance with Article 49 and Article 50 of Regulation (EU) 2019/627,
 - b) which was produced, collected, cooled, stored and transported in accordance with the hygiene conditions laid down in Annex III to Regulation (EC) No 853/2004,
 - c) which meets the plate and somatic cell count criteria laid down in Annex III to Regulation (EC) No 853/2004,
 - d) which, pursuant to testing for residues of antibacterial drugs carried out by the food business operator in accordance with the requirements of the Regulation (EC) No 853/2004, it complies with the maximum residue limits for residues of antibacterial veterinary medicinal products laid down in Commission Regulation (EU) No 37/2010,
 - e) which has been produced under conditions guaranteeing compliance with the maximum residue levels for pesticides in accordance with the requirements of the EU.
4. It comes from an establishment implementing a programme based on the HACCP principles in accordance with Regulation (EC) No 852/2004.
5. It has been processed, stored, wrapped, packaged and transported in accordance with the relevant hygiene requirements of the EU.
6. It meets the relevant microbiological criteria of Commission Regulation (EC) No 2073/2005.
7. The guarantees covering live animals and products thereof provided by the residue plans submitted in accordance with the requirements of Regulation (EU) 2017/625 are fulfilled.

Notes:

Part I:

- Box I.19: Indicate total gross weight and total net weight
- Box I.21: Either seal- or container number or both is to be indicated in this box.
- Box I.25: Custom code and title: Use the appropriate Harmonized System (HS) code under the following headings: 04.01; 04.02; 04.03; 04.04; 04.05; 04.06; 21.05

Signature and stamp must be different color that in the printed certificate.

Certifying Officer

Name (in capital letters)
 Date of signature
 Stamp

Qualification and title
 Signature