

**Model health certificate for milk, milk-based products and milk-derived products not for human consumption from non-EU countries GBHC091X v3.1 October 2022**

**Part I. Details of the dispatched consignment**

<b>I.1 Consignor</b> Name: Address: Tel:				<b>I.2 Certificate reference no.</b>			
				<b>I.2.a</b> Not in use	<b>I.3 Central competent authority</b>  APHIS-VS	<b>I.4 Local competent authority</b>	
<b>I.5 Consignee</b> Name: Address: Tel:				<b>I.6 Person responsible for the load in Great Britain</b> Name: Address: Tel:			
<b>I.7 Country of origin</b>	<b>ISO code</b>	<b>I.8 Region of origin</b>	<b>Code</b>	<b>I.9 Country of destination</b>	<b>ISO code</b>	<b>I.10 Region of destination</b>	<b>Code</b>
<b>I.11 Place of origin</b> Name: Approval number: Address:       Name: Approval number: Address:				<b>I.12 Place of destination</b> <input type="checkbox"/> Custom warehouse Name: Approval number: Address:			
<b>I.13 Place of loading</b>				<b>I.14 Date of departure</b>			
<b>I.15 Means of transport</b> <input type="checkbox"/> Aeroplane <input type="checkbox"/> Ship <input type="checkbox"/> Railway wagon <input type="checkbox"/> Road vehicle <input type="checkbox"/> Other Identification:    Documentation references:				<b>I.16 Entry BCP</b>			
				<b>I.17 No(s) of CITES</b>			



**Part II. Certification**

I, the undersigned official veterinarian, declare that I have read and understood Regulation (EC) No 1069/2009, and in particular Article 10 thereof, and Commission Regulation (EU) No 142/2011, and in particular Section 4 of Chapter II of Annex X, and Chapter I of Annex XIV thereto, and certify that the milk,<sup>(1)</sup> the milk-based products<sup>(1)</sup> and milk-derived products<sup>(1)</sup> referred to in box I.28 comply with the following conditions:

<sup>(4)</sup>**II.1** they were produced and derived in ..... (*insert name of exporting country*),<sup>(2)</sup> ..... (*insert name of region*)<sup>(2)</sup>, which is listed in a document relating to 'milk and milk products' published on gov.uk, in accordance with Commission Regulation (EU) No 605/2010, and which has been free from foot-and mouth disease (FMD) and rinderpest for a period of 12 months immediately prior to export and has not practised vaccination against rinderpest during that period;

**II.2** they were produced from raw milk derived from animals which at the time of milking did not show clinical signs of any disease transmissible through milk to humans or animals, and which had been kept for a period of at least 30 days prior to production on holdings that were not subject to official restrictions due to foot-and-mouth disease or rinderpest;

**II.3** they are milk or milk products that:

<sup>(1)</sup>*either* [have undergone one of the treatments or combinations thereof described in point II.4;]

~~<sup>(4)</sup>*or* [comprise whey to be fed to animals of species susceptible to foot and mouth disease, and that whey was collected from milk subjected to one of the treatments described in point II.4 and:~~

~~<sup>(4)</sup>*either* [the whey was collected at least 16 hours after clotting and has a pH below 6;]~~

~~<sup>(4)</sup><sup>(3)</sup>*or* [the whey has been produced at least 21 days before the shipping and during that period no cases of FMD have been detected in the exporting country;]~~

~~<sup>(4)</sup><sup>(3)</sup>*or* [the whey has been produced on ...../...../....., this date, in consideration of the foreseen voyage duration, being at least 21 days before the consignment is presented to a border control post of the point of entry into Great Britain;]~~

**II.4** they have been subject to one of the following treatments:

<sup>(1)</sup>*either* [high temperature short time pasteurisation at 72°C for at least 15 seconds, or an equivalent pasteurisation achieving a negative reaction to a phosphatase test in bovine milk, in combination with:

~~<sup>(4)</sup>*either* [a subsequent second high temperature short time pasteurisation at 72°C for at least 15 seconds or an equivalent pasteurisation which itself achieves a negative reaction to a phosphatase test in bovine milk;]~~

~~<sup>(4)</sup>*or* [a subsequent drying process that in the case of milk intended for feeding is combined with additional heating to 72°C or higher;]~~

~~<sup>(4)</sup>*or* [a subsequent process by which the pH is reduced and kept for at least one hour at a level below 6;]~~

- ~~(<sup>1</sup>)(<sup>3</sup>) or [the condition that the milk/milk product has been produced at least 21 days prior to the date of shipping and during that period no cases of FMD have been detected in the exporting country;]~~
- ~~(<sup>1</sup>)(<sup>3</sup>) or [the milk/milk product has been produced on ...../...../....., (*insert the date*), this date, in consideration of the foreseen voyage duration, being at least 21 days prior to the date that the consignment is presented to a border control post of the point of entry into Great Britain;]~~
- ~~(<sup>4</sup>) or [sterilisation at a level of at least F03;]~~
- ~~(<sup>4</sup>) or [ultra high temperature treatment at 132°C for at least one second in combination with:~~
- ~~(<sup>4</sup>) either [a subsequent drying process that in the case of milk intended for feeding is combined with additional heating to 72°C or higher;]~~
- ~~(<sup>4</sup>) or [a subsequent process by which the pH is reduced and kept for at least one hour at a level below 6;]~~
- ~~(<sup>1</sup>)(<sup>3</sup>) or [the condition that the milk/milk product has been produced at least 21 days prior to the date of shipping and during that period no cases of FMD has been detected in the exporting country;]~~
- ~~(<sup>4</sup>)(<sup>3</sup>) or [the milk/milk product has been produced on ...../...../....., (*insert the date*), this date, in consideration of the foreseen voyage the duration, being at least 21 days prior to the date that consignment is presented to a border control post of the point of entry into Great Britain;]~~

II.5 every precaution was taken to avoid contamination of the milk/milk-based product/milk-derived product after processing;

II.6 the milk/milk-based product/milk-derived product was packed:

~~(<sup>1</sup>) either [in new containers;]~~

~~(<sup>4</sup>) or [in vehicles or bulk containers disinfected prior to loading using a product approved by the competent authority;]~~

and the containers are marked so as to indicate the nature of the milk/milk-based product/milk-derived product and bear labels indicating that the product is Category 3 material and not intended for human consumption;

II.7 the milk, milk-based products and milk-derived products described above:

~~(<sup>1</sup>) either [does not contain milk or milk products of ovine or caprine animal origin or is not intended for feed for farmed animals, other than fur animals.]~~

~~(<sup>4</sup>) or [contains milk or milk products of ovine or caprine animal origin and is intended for feed for farmed animals, other than fur animals, and the milk or milk products:~~

~~(a) [are derived from ovine and caprine animals which have been kept continuously since birth in a country where the following conditions are fulfilled:~~

- ~~(i) — classical scrapie is compulsorily notifiable;~~
- ~~(ii) — an awareness, surveillance and monitoring system is in place for classical scrapie;~~
- ~~(iii) — official restrictions apply to holdings of ovine or caprine animals in the case of a suspicion of TSE or the confirmation of classical scrapie;~~
- ~~(iv) — ovine and caprine animals affected with classical scrapie are killed and destroyed;~~
- ~~(v) — the feeding to ovine and caprine animals of meat and bone meal or greaves, as defined in the Terrestrial Animal Health Code of the World Organisation for Animal Health (WOAH (formerly OIE)), of ruminant origin has been banned and effectively enforced in the whole country for a period of at least the preceding seven years;~~
- ~~(b) — originate from holdings where no official restrictions are imposed due to a suspicion of TSE;~~
- ~~(c) — originate from holdings where no case of classical scrapie has been diagnosed during a period of at least the preceding seven years or, following the confirmation of a case of classical scrapie:
  - ~~<sup>(+)</sup> either [all ovine and caprine animals on the holding have been killed and destroyed or slaughtered, except for breeding rams of the ARR/ARR genotype, breeding ewes carrying at least one ARR allele and no VRQ allele and other ovine animals carrying at least one ARR allele;]~~
  - ~~<sup>(+)</sup> or [all animals in which classical scrapie was confirmed have been killed and destroyed, and the holding has been subjected for a period of at least two years since the date of confirmation of the last classical scrapie case to intensified TSE monitoring, including testing with negative results for the presence of TSE in accordance with the laboratory methods set out in point 3.2 of Chapter C of Annex X to Regulation (EC) No 999/2001, of all of the following animals which are over the age of 18 months, except ovine animals of the ARR/ARR genotype:
    - ~~— animals which have been slaughtered for human consumption; and~~
    - ~~— animals which have died or been killed on the holding but which were not killed in the framework of a disease eradication campaign.]]~~~~~~

## 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](http://legislation.gov.uk)).

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

