**Model health certificate for untreated blood products, excluding those of equidae, for the manufacture of derived products for purposes outside the feed chain for farmed animals (BP-U)**

**GBHC502 v1.1 Aug-23**

**Part I. Details of dispatched consignment**

| **I.1 Consignor**<br>Name:<br>Address:<br><br>Tel: | **I.2 Certificate reference no.** | **I.3 Central competent authority** |
|---|---|---|
| | **I.2.a** Not in use | **I.4 Local competent authority** |

| **I.5 Consignee**<br>Name:<br>Address:<br><br>Tel: | **I.6 Person responsible for the load in Great Britain**<br>Name:<br>Address:<br><br>Tel: |
|---|---|

| **I.7 Country of origin** | **ISO code** | **I.8 Region of origin** | **Code** | **I.9 Country of destination** | **ISO code** | **I.10 Region of destination** | **Code** |
|---|---|---|---|---|---|---|---|
| | | | | | | | |

| **I.11 Place of origin**<br>Name:<br>Approval number:<br>Address:<br><br>Name:<br>Approval number:<br>Address:<br><br>Name:<br>Approval number:<br>Address: | **I.12 Place of destination**<br>☐ Custom warehouse<br>Name:<br>Approval number:<br>Address: |
|---|---|
| **I.13 Place of loading** | **I.14 Date of departure** |
| **I.15 Means of transport**<br>☐ Aeroplane<br>☐ Ship<br>☐ Railway wagon<br>☐ Road vehicle<br>☐ Other<br>Identification:<br>Documentation references: | **I.16 Entry BCP**<br><br>**I.17** Not in use |

| **II.a. Certificate reference no.** | **II.b.** |
|---|---|
| | |

| **I.18 Description of commodity** | | |
|---|---|---|
| **I.19 Commodity code (HS code)** | **I.21 Temperature of products**<br>☐ Ambient<br>☐ Chilled<br>☐ Frozen | **I.23 Seal / Container No.** |
| **I.20 Quantity** | **I.22 Number of packages** | **I.24 Type of packaging** |

**I.25 Commodity certified for**

☐ Technical use

| **I.26** ☐ **For transit through Great Britain to third country**<br>**Third country** **ISO Code** | **I.27** ☐ **For import or admission into Great Britain** |
|---|---|

**I.28 Identification of the commodities**

| Species (Scientific name) | Approval number of establishments / Manufacturing plant | Batch number |
|---|---|---|
| | | |
| | | |
| | | |
| | | |

## Part II. Certification

### Animal Health

I, the undersigned official veterinarian, declare that I have read and understood the requirements of the relevant GB regulations and certify that the blood products described in Part I of this certificate consist of blood products that satisfy the health requirements below:

**AH/T111 Territory requirements**

(*)[**(a)** in the case of blood products obtained from animals belonging to the taxa *Artiodactyla, Perissodactyla* and *Proboscidea*, including crossbreds between species of those taxa, the blood was collected in a country or region where no case of rinderpest, peste des petits ruminants and Rift Valley fever has been recorded and in which vaccination has not been carried out in accordance with GB requirements set out in the notes for completion ; and

(*)***EITHER*** [in third countries, territories or parts thereof …… where no case of foot-and-mouth disease has been recorded for a period of at least the preceding 12 months and in which vaccination has not been carried out against this disease for a period of at least the preceding 12 months; and]

(*)***OR*** [in third countries, territories or parts thereof …… where no case of foot-and-mouth disease has been recorded for a period of at least the preceding 12 months and in which vaccination programmes against foot-and-mouth disease are being officially carried out and controlled in domestic ruminant animals for a period of at least the preceding 12 months; and]]

(*)[**(b)** in the case of animals other than *Suidae* and *Tayassuidae*, in third countries or regions in which:

(*)***EITHER*** [no case of (*)[vesicular stomatitis] and (*)[bluetongue] (including the presence of seropositive animals) has been recorded in accordance with GB requirements as set out in the notes for completion;]

(*)***OR*** [(*)[vesicular stomatitis] and (*)[bluetongue] seropositive animals are present;]]

(*)[**(c)** in the case of *Suidae* and *Tayassuidae*, in third countries or regions in which no case of swine vesicular disease, classical swine fever and African swine fever has been recorded for a period of at least the preceding 12 months and vaccination has not been carried out against those diseases for a period of at least the preceding 12 months in the susceptible species and:

(*)***EITHER*** [no case of vesicular stomatitis (including the presence of seropositive animals) has been recorded in accordance with GB requirements as set out in the notes for completion;]

(*)***OR*** [vesicular stomatitis seropositive animals are present;]]

**AH/T112 Territory requirements**

(*)[in the case of blood products derived from poultry or other avian species the animals and the products come from the territory of the country or region with code …… which has been free from Newcastle disease and highly pathogenic avian influenza in accordance with GB requirements as set out in the notes for completion;]

**AH/E103 Establishment requirements**

have been prepared and stored in a plant and supervised by the competent authority or in the establishment of collection, exclusively with the following animal by-products as set out in the notes for completion: (*)[A] (*)[B] (*)[C] (*)[D] (*)[E] (*)[F] (*)[G];

**AH/E302 Establishment requirements (slaughterhouse)**

the blood, that these products were manufactured from, was collected in slaughterhouses approved in accordance with GB regulations, in slaughterhouses approved and supervised by the competent authority of the country of collection or from live animals in facilities approved and supervised by the competent authority of the country of collection;

**AH/P011 Product requirements (segregation)**

all precautions were taken to avoid contamination of the products with pathogenic agents during transport;

**AH/P151B Product requirements**

consist exclusively of blood products not intended for human or animal consumption;

**AH/P509 Packaging and labelling**

the products were:

(*)***EITHER*** [packed in new or sterilised bags or bottles;]

(*)***OR*** [transported in bulk in containers or other means of transport that were thoroughly cleaned and disinfected with a disinfectant approved by the competent authority before use;]

and the outer packaging or containers bear labels indicating 'NOT FOR HUMAN OR ANIMAL CONSUMPTION';

**AH/P550A Storage**

the product was stored in enclosed storage;

## Public Health

**PH/D011A Bovine spongiform encephalopathy (BSE)**

the products described in Part I

(*)***EITHER*** [come from other ruminants than bovine, ovine or caprine animals;]

| **II.a. Certificate reference no.** | **II.b.** |
|---|---|
| | |

(*)***OR*** [come from bovine, ovine or caprine material:

(*)***EITHER*** [**(a)** derived from animals that were born, continuously reared and slaughtered in a country or region with a negligible BSE risk as set out in a document published on GOV.UK in accordance with GB regulations;]

(*)***OR*** [**(b)** that does not contain and is not derived from:

**(i)** specified risk material and mechanically separated meat, in compliance with GB regulations;

**(ii)** animal by-product or derived product obtained from animals which have not been killed in compliance with GB regulations in regards laceration of certain tissues after stunning;]]

(*) Keep as appropriate.

**Official Veterinarian / Official Inspector**

**By signing this certificate, I certify that the requirements laid out above and in the accompanying notes for completion have been met.**

Name (in capital letters):

Qualification and title:

Date:

Signature:

Stamp:

## Part III. Notes for completion

These notes for completion must be read and understood by the certifying officer before signing the certificate. Notes are set out in sections that correspond to the sections in the certificate. By signing this certificate, certifiers are verifying that the consignment meets the requirements set out in the certificate and any relevant corresponding notes for completion.

These notes do not need to be printed as part of a paper certificate that accompanies the consignment or in any electronic copy of the certificate.

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.

References to GB requirements refer to the requirement(s) of Great Britain as set out in the accompanying notes for completion.

### Part I

Box reference I.6: *Person responsible for the consignment in Great Britain*: this box is required to be filled in only if it is a certificate for a commodity to be transited through Great Britain; it may be filled in if the certificate is for a commodity that is to be imported into Great Britain.

Box reference I.11 and I.12: *Approval number*: the registration number of the establishment or plant, which has been issued by the competent authority.

Box reference I.12: *Place of destination*: this box is to be filled in only if it is a certificate for transit commodity. Products in transit may only be stored in free zones, free warehouses and custom warehouses.

Box reference I.15: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) is to be provided. In the case of unloading and reloading, the consignor must inform the border control post of the point of entry into Great Britain.

Box reference I.19: Use the appropriate Harmonized System (HS) code under the following headings: 05.11; 30.02 or 35.02.

Box reference I.23: For bulk containers, the container number and the seal number (if applicable) must be included.

Box reference I.25: *Technical use*: any use other than feeding of farmed animals, other than fur animals, and the production or manufacturing of pet food.

Box reference I.26 and I.27: Fill in according to whether it is a transit or an import certificate.

Box reference I.28: *Species*: select from the following: *Aves, Ruminantia, Suidae, Mammalia,* other than *Ruminantia* or *Suidae, Pesca, Reptilian.*

### Part II

### Animal Health

By signing this certificate, you, the official veterinarian, are certifying that you have read and understood Regulation (EC) No 1069/2009, and in particular Article 8(c) and Article 8(d) and Article 10 thereof, and Commission Regulation (EU) No 142/2011, and in particular Chapter II of Annex XIV thereto.

**AH/T111 Territory requirements**

Delete the whole attestation if not applicable.

If applicable, certify attestation **(a)** and either **(b)** or **(c)** as applicable.

**When certifying any of the 'or' attestations:** Following the veterinary checks at the border control post, the consignment must be monitored and transported directly to the premises of destination in accordance with the relevant requirements of Regulation (EU) 2019/1666.

**(a)** Insert ISO country code in the case of a country, or codes in the case of territories or parts thereof as set out in a document relating to 'fresh meat of ungulates' published on GOV.UK, in accordance with Regulation (EU) No 206/2010.(†)

**Rinderpest, peste des petits ruminants and Rift Valley fever requirements**

The blood was collected in a country or region where no case of rinderpest, peste des petits ruminants and Rift Valley fever has been recorded for a period of at least the preceding 12 months and in which vaccination has not been carried out against those diseases for a period of at least the preceding 12 months.

**(b) Vesicular stomatitis and bluetongue requirements**

No case of vesicular stomatitis and bluetongue (including the presence of seropositive animals) has been recorded for a period of at least the preceding 2 months and in which vaccination has not been carried out against those diseases for a period of at least the preceding 12 months.

**(c) Vesicular stomatitis requirements**

No case of vesicular stomatitis (including the presence of seropositive animals) has been recorded for a period of at least the preceding 12 months and in which vaccination has not been carried out against this disease for a period of at least the preceding 12 months.

**AH/T112 Territory requirements**

Insert code of the territory as set out in a document relating to 'poultry and poultry products' published on GOV.UK, in accordance with Commission Regulation (EC) No 798/2008.(†)

**GB requirements**

The animals and the products come from the territory of the country or region which has been free from Newcastle disease and highly pathogenic avian influenza as defined in the Terrestrial Animal Health Code of the WOAH, which for a period of at least the preceding 12 months has not carried out vaccination against avian influenza, where the animals from which the products are derived, have not been vaccinated against Newcastle disease with vaccines prepared from a Newcastle disease master strain showing a higher pathogenicity than lentogenic virus strains.

**AH/E103 Establishment requirements**

One or more options can be selected.

**A**: Blood of slaughtered animals, which is fit for human consumption in accordance with retained EU law, but is not intended for human consumption for commercial reasons.

**B**: Blood of slaughtered animals, which is rejected as unfit for human consumption in accordance with retained EU law, but which did not show any signs of diseases communicable to humans or animals, derived from carcases that have been slaughtered in a slaughterhouse and were considered fit for human consumption following an ante-mortem inspection in accordance with retained EU law.

**C**: Blood of slaughtered animals, which did not show any signs of diseases communicable to humans or animals, obtained from animals that have been slaughtered in a slaughterhouse after having been considered fit for human consumption following an ante-mortem inspection in accordance with retained EU law.

**D**: Blood and blood products derived from the production of products intended for human consumption.

**E**: Blood and blood products originating from live animals that did not show signs of any disease communicable through that product to humans or animals.

**F**: Animal by-products derived from animals which have been submitted to illegal treatment as defined in Article 1(2)(d) of Council Directive 96/22/EC or Article 2(b) of Council Directive 96/23/EC.

**G**: Animal by-products containing residues of other substances and environmental contaminants listed in Group B(3) of Annex I to Directive 96/23/EC, if such residues exceed the permitted level laid down in retained EU law or, in the absence thereof, in national legislation.

**AH/E302 Establishment requirements (slaughterhouse)**

GB legislation refers to the approval in accordance with retained EU law.

**AH/P011 Product requirements (segregation)**

No further notes for completion.

**AH/P151B Product requirements**

No further notes for completion.

**AH/P509 Packaging and labelling**

No further notes for completion.

**AH/P550A Storage**

No further notes for completion.

## Public Health

**PH/D011A Bovine spongiform encephalopathy (BSE)**

The products described in Part I of the certificate:

***EITHER*** are derived from other ruminants than bovine, ovine or caprine animals.

***OR*** are derived from bovine, ovine or caprine animals, and do not contain and are not derived from:

***EITHER*** **(a)** bovine, ovine and caprine materials other than those derived from animals born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk as set out in a document relating to 'BSE risk status' published on GOV.UK, in accordance with Regulation (EC) No 999/2001(‡);

***OR*** **(b)** the following:

**(i)** specified risk material as defined in point 1 of Annex V to Regulation (EC) No 999/2001 of the European Parliament and of the Council and mechanically separated meat obtained from bones of bovine, ovine or caprine animals, except from those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk as set out in a document relating to 'BSE risk status' published on GOV.UK, in accordance with Regulation (EC) No 999/2001(‡), in which there have been no indigenous BSE cases.

**(ii)** animal by-product or derived product obtained from bovine, ovine or caprine animals which have been killed, after stunning, by laceration of the central nervous tissue by means of an elongated rod-shaped instrument introduced into the cranial cavity, or by means of gas injected into the cranial cavity, except for those animals that were born, continuously reared and slaughtered in a country or region classified as posing a negligible BSE risk as set out in a document relating to 'BSE risk status' published on GOV.UK, in accordance with Regulation (EC) No 999/2001.(‡)

(†) The document(s) referred to above can be found at:

EU and EFTA countries approved to export animals and animal products to Great Britain
(Available at: https://www.data.gov.uk/dataset/4698a65d-1a3b-42d1-981e-df869e04185b/eu-and-efta-countries-approved-to-export-animals-and-animal-products-to-great-britain)

Non-EU countries approved to export animals and animal products to Great Britain
(Available at: https://www.data.gov.uk/dataset/b92627b0-dd7b-4e1d-ba36-e25424f55eeb/non-eu-countries-approved-to-export-animals-and-animal-products-to-great-britain)

(‡) A document relating to the 'Bovine Spongiform Encephalopathy (BSE) risk status' of approved trading partners published by the Secretary of State, with the consent of the Scottish and Welsh Ministers, can be found at:

Animal health status of countries approved to export animals and animal products to Great Britain - data.gov.uk
(Available at: https://www.data.gov.uk/dataset/b7712d2e-debb-4996-8e79-d27ca7492a00/animal-health-status-of-countries-approved-to-export-animals-and-animal-products-to-great-britain)