**Model health certificate for blood products not intended for human consumption that could be used as feed material (BP) GBHC500 v1.0 May-23**

## Part I. Details of the dispatched consignment

| | | | |
|---|---|---|---|
| **I.1 Consignor**<br>Name:<br>Address:<br>Tel: | **I.2 Certificate reference no.** | | |
| | **I.2.a**<br>Not in use | **I.3 Central competent authority**<br>**APHIS-VS** | **I.4 Local competent authority** |
| **I.5 Consignee**<br>Name:<br>Address:<br>Tel: | | **I.6 Person responsible for the load in Great Britain**<br>Name:<br>Address:<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 |
|---|---|---|---|---|---|---|---|
| USA---------- | US--------- | ---------- | ---------- | UNITED KINGDOM--------- | GB----- | ---------- | --------- |

| | |
|---|---|
| **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 ☐ Ship<br>☐ Railway wagon ☐ Road vehicle<br>☐ Other<br><br>Identification:<br><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 ☐ 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** ☐ Animal feedingstuff ☐ Technical use ☐ Manufacture of petfood | | |
| **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) | Nature of commodity | Approval number of establishments / Manufacturing plant | Batch number |
|---|---|---|---|
| | | | |
| | | | |
| | | | |

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

## 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/E102 Establishment requirements (plant)**

have been prepared and stored in a plant approved and supervised by the competent authority in accordance with GB requirements;

**AH/P010 Product requirements (segregation)**

the product has undergone all precautions to avoid contamination with pathogenic agents after treatment;

(*)***AND*** [has been stored in dry warehouse conditions under room temperature for a period of at least 6 weeks;]

**AH/P104 Product requirements (composition)**

have been prepared exclusively with the following animal by-products as set out in the notes for completion (*)[A] and/or (*)[B];

**AH/P151A Product requirements**

consist exclusively of blood products not intended for human consumption;

**AH/P506 Packaging and labelling**

the end product was:

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

(*)***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 which bear labels indicating 'NOT FOR HUMAN CONSUMPTION';

**AH/P550B Storage**

the end product was stored in enclosed storage;

**AH/P700 Product treatment**

in order to inactivate pathogenic agents, have been submitted:

(*)***EITHER*** **(a)** [to processing in accordance with processing method …… in accordance with GB requirements;]

(*)***OR*** **(b)** [to a method and parameters which ensure that the product complies with the GB microbiological requirements;]

(*)***OR*** **(c)** [to treatment for blood products as set out in the notes for completion and meets the relevant requirements;]

**AH/P800B Testing**

the competent authority examined a random sample of the products immediately prior to dispatch and found it to comply with microbiological GB requirements for *Salmonella* and *Enterobacteriaceae;*

**AH/P901 Product requirements (statement)**

the product(s) described above contain or are derived from animal-by products of nonruminant origin, and are, according to the statement of the Consignor referred to in box reference I.1,

(*)***EITHER*** [not intended for the production of feed for farmed animals, other than fur animals;]

(*)***OR*** [intended for the production of feed for non-ruminant farmed animals, other than fur animals, and the Consignor has undertaken to ensure that the border control post of entry will be provided with the results of the analyses carried out in accordance with the methods set out in GB legislation;]

**AH/D200A TSE (scrapie)**

the animal by-products described above:

(*)***EITHER*** [does not contain ovine or caprine milk or milk products or is not intended for feed for farmed animals, other than fur animals;]

(*)***OR*** [contains ovine or caprine milk or milk products intended as feed for farmed animals, other than fur animals, and the milk or milk products:

**(a)** are derived from animals from countries which meet GB requirements in regards to scrapie controls;

**(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 the period of the preceding seven years or, following the confirmation of a case of classical scrapie:

(*)***EITHER*** [all ovine and caprine animals on the holding have been killed and destroyed or slaughtered, except for animals which meet GB requirements;]

(*)***OR*** [all animals in which classical scrapie was confirmed have been killed and destroyed, and the holding has been subjected to TSE monitoring which meets GB requirements;]]

## 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;]

(*)***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:

The signature and the stamp must be in a different colour to that of the printing.