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

GBHC503 v1.0 May-23

Part I. Details of the dispatched consignment I.1 Consignor I.2 Certificate reference no. Name: Address: Tel: I.3 Central competent I.4 Local competent I.2.a authority authority Not in **APHIS-VS** use 1.6 Person responsible for the load in Great I.5 Consignee Britain Name: Address: Name: Tel: Address: Tel: I.7 Country of origin ISO code I.8 Region of Code I.9 Country of ISO Code I.10 Region of origin destination code destination I.11 Place of origin 1.12 Place of destination Name: Custom warehouse Approval number: Address: Approval number: Address: Name: Approval number: Address: Name: Approval number: Address: I.13 Place of loading I.14 Date of departure I.15 Means of transport I.16 Entry BCP ☐ Aeroplane ☐ Ship I.17 Not in use ☐ Railway wagon Road vehicle Other Identification: Documentation references:

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II.a. Certificate reference no.	N.b.

I.18 Description of commodity		
I.19 Commodity code (HS code)	I.21 Temperature of products	I.23 Seal / Container No.
, ,	Ambient Chilled	
	— —	
	Frozen	
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 Brita	ain to third country	I.27 For import or admission into
Third country	ISO Code	Great Britain
Tillia country	ISO Code	
I.28 Identification of the commodities	S	
Species (Scientific name)	Approval number of establishments / Manufacturing plant	Batch number

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II.a. Certificate reference no.



Part II. Certification

Animal Health

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

AH/E109 Establishment requirements

have been prepared and stored in a plant and supervised by the competent authority, exclusively with the following animal by-products as set out in the notes for completion: (-)[A] (*)[B] (-)[C] (*)[D] (-)[C] (*)[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/P007 Product requirements (segregation)

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

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 Itransported in bulk in containers or other me

[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/P550B Storage

the end product was stored in enclosed storage;

AH/P701 Product requirements

(*)[in the case of blood products derived from *Artiodactyla*, *Perissodactyla* and *Proboscidea* including their crossbreeds, other than *Suidae* and *Tayassuidae*, the products have undergone one of the following treatments, guaranteeing the absence of pathogens of foot-and-mouth disease, vesicular stomatitis, rinderpest, peste des petits ruminants, Rift Valley fever and bluetongue:

(*) **EITHER** [heat treatment at a temperature of 65°C for at least three hours, followed by an effectiveness check;]

(*)AND/OR [irradiation at 25 kGy by gamma rays, followed by an effectiveness check;]

(*)AND/OR [change in pH to pH 5 for two hours, followed by an effectiveness check;]

*AND/OR [heat treatment of at least 80 °C throughout their substance, followed by an effectiveness check;]]

AH/P702 Product requirements

(*)[in the case of blood products derived from Suidae, Tayassuidae, poultry and other avian species, the products have undergone one of the following treatments guaranteeing the absence of pathogens of the following diseases: foot-and-mouth disease, vesicular stomatitis, swine vesicular disease, classical swine fever, African swine fever, Newcastle disease and highly pathogenic avian influenza, as appropriate to the species:

(heat treatment at a temperature of 65°C for at least three hours, followed by an effectiveness check;)

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И.b.

AND/OR [irradiation at 25 kGy by gamma rays, followed by an effectiveness check;]

(*) AND/OR [heat treatment of (*)[at least 80°C for Suidae/Tayassuidae] and (*)[at least 70°C for poultry and other avian species] throughout the substance of the product, followed by an effectiveness

check;]]

AH/P703 Product requirements

(*)[in the case of blood products derived from species other than those listed in AH/P701 or AH/P702, the products have undergone the following treatment (please specify).....;]

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;]]]

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.

^(*) Keep as appropriate.