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 from non-EU countries

GBHC097X v3.1 October 2022

Part I. Details of dispatched consignment		
I.1 Consignor Name: Address: 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 Name: Address: Tel:	**I.6 Person responsible for the load in Great Britain** Name: Address: 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 Name: Approval number: Address: Name: Approval number: Address: Name: Approval number: Address:	**I.12 Place of destination** ☐ Custom warehouse Name: Approval number: Address:
I.13 Place of loading	**I.14 Date of departure**
I.15 Means of transport ☐ Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other Identification: Documentation references:	**I.16 Entry BCP** **I.17** Not in use

MODEL CERTIFICATE ONLY

Treated blood products from non-EU countries
GBHC097X

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

I.18 Description of commodity		
I.19 Commodity code (HS code)	**I.21 Temperature of products** ☐ Ambient ☐ Chilled ☐ 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 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

I, the undersigned official veterinarian, declare that I 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, and certify that:

II.1 the blood products described above consist of blood products that satisfy the requirements below;

II.2 they consist exclusively of blood products not intended for human or animal consumption;

II.3 they have been prepared and stored in a plant supervised by the competent authority, exclusively with the following animal by-products:

[(1)]*either* [- 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;]

[(1)]*and/or* [- 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 carcasses 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;]

MODEL CERTIFICATE ONLY

(1)*and/or* [- 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;]

(1)*and/or* [- blood and blood products originating from live animals that did not show clinical signs of any disease communicable through these products to humans or animals;]

(1)*and/or* [- blood and blood products derived from the production of products intended for human consumption;]

(1)*and/or* [- animal by-products which have been 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;]

(1)*and/or* [- 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 levels laid down by retained EU law or, in the absence thereof, in national legislation;]

II.4 the blood that these products were manufactured from has been collected in slaughterhouses approved in accordance with retained EU law, 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.

(1)[**II.5** 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:

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

(1)*and/or* [irradiation at 25 kGy by gamma rays, followed by an effectiveness check;]

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

(1)*and/or* [heat treatment of at least 80 °C throughout their substance, followed by an effectiveness check]]

(1)[**II.6** 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:

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

(1)*and/or* [irradiation at 25 kGy by gamma rays, followed by an effectiveness check;]

(1)*and/or* [heat treatment of at least 80°C for *Suidae*/*Tayassuidae* (1) and at least 70°C for poultry and other avian species (1) throughout the substance of the product, followed by an effectiveness check]].

(1)[**II.7** In the case of blood products derived from species other than those listed in point II.5 or II.6, the products have undergone the following treatment (*please specify*): ……]

MODEL CERTIFICATE ONLY

II.8 The products were:

(1)*either* [packed in new or sterilised bags or bottles,]

(1)*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';

II.9 the products were stored in enclosed storage;

II.10 all precautions were taken to avoid the contamination of the products with pathogenic agents after treatment;

(1)(2)[**II.11** The treated blood products described above

(1)*either* [is derived from other ruminants than bovine, ovine or caprine animals.]]

(1)*or* [is derived from bovine, ovine or caprine animals and does not contain and is not derived from:

(1)(2)*either* [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.]]

(1)(2)*or* [**(a)** 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;

(b) 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, which there has been no indigenous BSE case,

(c) 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.]]]

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).

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

Part I:

MODEL CERTIFICATE ONLY

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, 35.02 or 35.04.

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: In case of Species: select from the following: *Aves, Ruminantia, Suidae, Mammalia* other than *Ruminantia* or *Suidae, Pesca, Reptilian.*

Part II:

(1) Delete as appropriate.

(2) 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, may be found here:

Animal health status of countries approved to export animals and animal products to Great Britain - data.gov.uk

MODEL CERTIFICATE ONLY

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

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

Note for the person responsible for the consignment in Great Britain: this certificate is only for veterinary purposes and must accompany the consignment until it reaches the border control post.

Official Veterinarian / Official Inspector

Name (in capital letters):	Qualification and title:
Date:	Signature:
Stamp:	

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