Model health certificate for blood and blood products from equidae for purposes outside the feed chain (BP-E)

GBHC501 v1.0 May-23

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

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

Part II. Certification

Animal Health

I, the undersigned official veterinarian, declare that I am aware of the relevant requirements of the GB legislation,and certify that the animal by-products described in Part I of this certificate consist of blood or blood products from equidae that satisfy the health requirements below:

AH/T115 Territory requirements

have been obtained from animals that originate from the EU Member States or from a third country, territory or part thereof listed as per GB requirements where the following diseases are compulsorily notifiable: African horse sickness, dourine, glanders (*Burkholderia mallei*), equine encephalomyelitis (all types including Venezuelan equine encephalomyelitis), equine infectious anaemia, vesicular stomatitis, rabies, anthrax;

AH/T116 Territory requirements

blood products have been produced from blood which fulfils the conditions referred in AH/E008, AH/E303, AH/A702, AH/A051 and AH/A610 and:

(*)***EITHER*** [has been collected from equidae which have been kept for a period of at least three months, or since birth if less than three months old, prior to the date of collection on holdings under veterinary supervision in the country of collection which during that period and the period of blood collection has been free of:

(a) African horse sickness for two years;

(b) Venezuelan equine encephalomyelitis for a period of at least two years;

(c) glanders

(*)***EITHER*** [for a period of three years;]

(*)***OR*** [for a period of six months where the animals have passed the post-mortem inspection for glanders in the slaughterhouse referred to in AH/E303, including a careful examination of mucous membranes from the trachea, larynx, nasal cavities and sinuses and their ramifications, after splitting the head in the median plane and excising the nasal septum;]

(d) in the case of blood products other than serum and plasma, vesicular stomatitis for six months;]

(*)***OR*** [has been subjected to at least one of the following treatments, followed by an effectiveness check, for the inactivation of possible causative pathogens for African horse sickness, equine encephalomyelitis of all types including Venezuelan equine encephalomyelitis, equine infectious anaemia, vesicular stomatitis and glanders (*Burkholderia mallei*):

(*)***EITHER*** [heat treatment at a temperature of 65°C for at least three hours;]

(*)***AND/OR*** [irradiation at 25 kGy by gamma rays;]

(*)***AND/OR*** [change in pH to pH 5 for two hours;]

(*)***AND/OR*** [heat treatment of at least 80°C throughout their substance;]]

AH/E008 Establishment requirements (holding)

have been derived from blood which was collected from equidae which have been kept for at least 30 days prior to the date of and during blood collection on holdings under veterinary supervision which were not subject to a prohibition as per GB legislation;

AH/E111 Establishment requirements

blood products come from an establishment or plant approved or registered by the competent authority of the third country meeting the specific conditions required by GB legislation;

AH/E303 Establishment requirement (slaughterhouse)

have been derived from blood from equidae which was collected under the supervision of a veterinarian in slaughterhouses approved in accordance with GB requirements, in slaughterhouses approved and supervised by the competent authority of the country of collection and in facilities approved and supervised by the competent authority of the country of collection for the purpose of collecting blood from equidae for the production of blood products for purposes other than feeding for farmed animals;

AH/A051 Animal requirements

which had no contact with equidae from a holding which was subject to a prohibition order for animal health reasons pursuant to GB legislation;

AH/A610 Animal requirements

for which the period for the prohibition order referred to in AH/E008 and AH/A051 has been determined as follows:

(*)***EITHER*** [not all the animals of species susceptible to the disease located on the holding have been slaughtered, in which case the period of prohibition must be at least as per GB requirements set out in the notes for completion;]

(*)***OR*** [all the animals of species susceptible to the disease located on the holding have been slaughtered and the premises were disinfected, in which case the period of prohibition must be 30 days beginning on the date on which the animals were slaughtered, and the premises disinfected, except in the case of anthrax, where the period of prohibition shall be 15 days;]

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

AH/A702 Animal requirements

have been derived from blood which was collected from equidae which on inspection on the date of blood collection did not show clinical signs of any of the compulsorily notifiable diseases listed as per GB requirements;

AH/P017 Product requirements

all precautions have been taken to avoid contamination of the blood and blood products with pathogenic agents during production, handling and packaging;

AH/P159 Product requirements

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

AH/P522 Product requirements

blood and blood products were packed in sealed impermeable containers clearly labelled "NOT FOR HUMAN OR ANIMAL CONSUMPTION" and bearing:

(a) in the case of blood, the approval number of the establishment of collection;

(b) in the case of blood products, the approval number of the establishment of production;

AH/P550A Storage

the product was stored in enclosed storage;

(*) 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 heading: 30.02.

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 for animal consumption.

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

Box reference I.28: (a) *Manufacturing plant*:

(i) in the case of blood, provide the approval number of the registered establishment of collection;

(ii) in the case of blood products, provide the approval number of the establishment of production;

(b) *Species*: select amongst the following: *Equus cabalus, Equus asinus, Equus cabalus*asinus*.

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 (d) and Article 10 thereof, and Commission Regulation (EU) No 142/2011, and in particular Chapter IV of Annex XIII thereto.

AH/T115 Territory requirements

EU Member States or a third country, territory or part thereof must be listed in the column "third countries lists" of row No 3 of Table 2 in Section 1 of Chapter II of Annex XIV to Regulation (EU) No 142/2011.

AH/T116 Territory requirements

No further notes for completion.

AH/E008 Establishment requirements (holding)

Equidae must be kept on holdings, under veterinary supervision, which were not subject to a prohibition order pursuant to Article 4(5) or restrictions for African horse sickness in accordance with Article 5 of Directive 2009/156/EC.

AH/E111 Establishment requirements

Establishment or plant approved or registered by the competent authority of the third country must meet the specific conditions set out in Article 23 or 24 of Regulation (EC) No 1069/2009.

AH/E303 Establishment requirement (slaughterhouse)

GB requirements:

Approval in accordance with Regulation (EC) No 853/2004.

AH/A051 Animal requirements

Relevant GB legislation:

Article 4(5) of Directive 2009/156/EC.

AH/A610 Animal requirements

GB requirements for the period of prohibition:

(a) Six months in the case of glanders (*Burkholderia mallei*), beginning on the date on which the equidae infected with the disease are slaughtered.

(b) Six months in the case of equine encephalomyelitis of any type, including Venezuelan equine encephalomyelitis, beginning on the date on which the equidae infected with the disease are slaughtered, in the case of equine infectious anaemia, until the date on which, the infected animals having been slaughtered, and the remaining animals have shown a negative reaction to two Coggins tests carried out three months apart.

(c) Six months from the date of the last recorded case of vesicular stomatitis.

(d) One month from the date of the last recorded case of rabies.

(e) 15 days from the date of the last recorded case of anthrax.

AH/A702 Animal requirements

GB requirements:

The applicable compulsorily notifiable diseases as listed in Annex I to Council Directive 2009/156/EC , and of equine influenza, equine piroplasmosis, equine rhinopneumonitis and equine viral arteritis listed in point 4 of Article 1.2.3 of the Terrestrial Animal Health Code of the World Organisation for Animal Health (WOAH (formerly OIE)), 2010 edition.

AH/P017 Product requirements

No further notes for completion.

AH/P159 Product requirements

No further notes for completion.

AH/P522 Product requirements

No further notes for completion.

AH/P550A Storage

No further notes for completion.