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.