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

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 Article 8(d) and Article 10 thereof, and Commission Regulation (EU) No 142/2011, and in particular Chapter II of Annex XIV thereto.

AH/E109 Establishment requirements

One or more options can be selected.

~~**A**: 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.~~

B: 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 carcases 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.

~~**C**: 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.~~

D: Blood and blood products derived from the production of products intended for human consumption.

~~**E**: Blood and blood products originating from live animals that did not show clinical signs of any disease communicable through that product to humans or animals.~~

F: 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.

G: 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 level laid down in retained EU law or, in the absence thereof, in national legislation.

AH/E302 Establishment requirements (slaughterhouse)

GB legislation refers to the approval in accordance with retained EU law.

AH/P007 Product requirements (segregation)

No further notes for completion.

AH/P151B Product requirements

No further notes for completion.

AH/P509 Packaging and labelling

No further notes for completion.

AH/P550B Storage

No further notes for completion.

AH/P701 Product requirements

No further notes for completion.

AH/P702 Product requirements

No further notes for completion.

AH/P703 Product requirements

No further notes for completion.

Public Health

PH/D011A Bovine spongiform encephalopathy (BSE)

The products described in Part I of the certificate:

EITHER are derived from other ruminants than bovine, ovine or caprine animals.

~~***OR*** are derived from bovine, ovine or caprine animals, and do not contain and are not derived from:~~

~~***EITHER*** **(a)** 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[‡];~~

~~***OR*** **(b)** the following:~~

~~**(i)** 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 and 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(‡), in which there have been no indigenous BSE cases.~~

~~**(ii)** 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.(‡)~~

(‡) 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, can be found at:

Animal health status of countries approved to export animals and animal products to Great Britain - data.gov.uk
(Available at: https://www.data.gov.uk/dataset/b7712d2e-debb-4996-8e79-d27ca7492a00/animal-health-status-of-countries-approved-to-export-animals-and-animal-products-to-great-britain)