

Imports of consignments of ova and embryos of equidae collected or produced in accordance with Directive 92/65/EEC after 30 September 2014 and dispatched by an approved embryo collection or production team of origin of the ova or embryos

GBHC814 v1.0 Aug-23

Part I. Details of dispatched consignment GERMINAL							
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 <input type="checkbox"/> Embryo team Name: _____ Approval number: _____ Address: _____				I.12 Place of destination <input type="checkbox"/> Embryo team <input type="checkbox"/> Holding Name: _____ Approval number: _____ Address: _____			
I.13 Place of loading _____				I.14 Date of departure _____			
I.15 Means of transport <input type="checkbox"/> Aeroplane <input type="checkbox"/> Ship <input type="checkbox"/> Railway wagon <input type="checkbox"/> Road vehicle <input type="checkbox"/> Other Identification: _____ Documentation references: _____				I.16 Entry BCP			
				I.17 Not in use			

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I.18 Description of commodity				
I.19 Commodity code (HS code) 05 11 99 85		I.23 Seal / Container No.		
I.20 Quantity	I.22 Number of packages			
I.25 Commodity certified for: <input type="checkbox"/> Artificial reproduction				
I.26 <input type="checkbox"/> For transit through Great Britain to third country Third country _____ ISO Code _____		I.27 <input type="checkbox"/> For import or admission into Great Britain		
I.28 Identification of the commodities				
Species (Scientific name)	Category	Donor identity	Date of collection	Quantity

Part II. Certification

Animal Health

I, the undersigned, official veterinarian, of the exporting country..... (*name of exporting country*) hereby certify that:

AH/E353D Establishment requirement (Collection centre)

(*)[the embryos described above were conceived by (*)[artificial insemination] (*)[as a result of in vitro fertilisation] using semen meeting the GB requirements and coming from semen collection centres approved in accordance with GB requirements and located respectively in Great Britain or in a third country or parts thereof listed on gov.uk;]

the ova used for in vitro production of the embryos described above comply with the GB requirements and in particular the requirements set up in points (a) to (n) of this certificate;]

AH/E373A Establishment requirement

the (*)[ova] (*)[embryos] described above:

- (a) were (*)[collected] (*)[produced] by the team described in Box I.11, which has been approved and supervised in accordance with GB requirements and is subject to inspection by an official veterinarian at least once every calendar year;
- (b) were (*)[collected] (*)[produced], processed and stored in accordance with the GB requirements;

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- (c) were collected at a place separated from other parts of the premises or holding which is in good repair and was cleaned and disinfected prior to the collection;
- (d) were examined, processed and packed in laboratory facilities which are not situated in a zone subject to prohibition or quarantine measures as set out in points (e) to (l), in a section which is separated from the section for storing equipment and materials used in contact with donor animals and from the area where the donor animals are handled;
- (e) come from donor mares which were continuously resident for a period of 3 months (or since entry if they were directly imported from Great Britain during the 3 months period) in the exporting country or, in the case of regionalisation in accordance with GB requirements, in that part of the territory of the exporting country which was during that period:
 - (i) not considered to be infected with African horse sickness in accordance with GB requirements,
 - (ii) free from Venezuelan equine encephalomyelitis for a period of at least 2 years,
 - (iii) free from glanders and dourine for a period of at least 6 months;
- (f) **Vesicular stomatitis** - come from donor mares which:
 - (*)**EITHER** [(i) originated from a country of export which was on the day of collection free from vesicular stomatitis (VS) for a period of at least 6 months;]
 - (*)**OR** [(ii) were subjected to a virus neutralisation test for vesicular stomatitis (VS) carried out with a negative result at a serum dilution of 1 in 32 or a VS ELISA carried out with a negative result in accordance with the WOAHP Manual on a blood sample taken on within 30 days prior to the collection of the (*)[ova] (*)[embryos];]
- (g) **Holding requirements** - come from donor mares which:
 - (*)**EITHER** [during a period of the past 30 days prior to the date of the collection were located in holdings under veterinary supervision which fulfilled, from the day of the collection of the (*)[ova] (*)[embryos] until the date of their dispatch the GB requirements for a holding, and in particular:]
 - (*)**OR** [(in the case of frozen (*)[ova] (*)[embryos], during a period of the past 30 days prior to the date of the collection were kept in holdings under veterinary supervision which fulfilled, from the day of the collection of the (*)[ova] (*)[embryos] until the end of the period of 30 days mandatory storage at approved premises, the GB requirements for a holding, and in particular:]
 - (*)**EITHER** [(i) following a case of a disease mentioned below not all the animals of species susceptible to that disease located in the holding were slaughtered or killed and the holding has been free:
 - (1) from any type of equine encephalomyelitis for a period of at least 6 months, beginning on the day on which the equidae suffering from the disease are slaughtered,
 - (2) from equine infectious anaemia for at least the period required to obtain a negative result in an agar gel immunodiffusion test (AGID or Coggins tests) carried out on samples taken after the infected animals were slaughtered on two occasions 3 months apart from each of the remaining equidae,
 - (3) from vesicular stomatitis for a period of at least 6 months from the last recorded case,
 - (4) from rabies for a period of at least one month from the last recorded case,
 - (5) from anthrax for a period of at least 15 days from the last recorded case,]
 - (*)**OR** [(ii) following a case of a disease mentioned below all the animals of species susceptible to that disease located in the holding were slaughtered or killed and the premises disinfected, the holding was free for a period of at least 30 days

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from any type of equine encephalomyelitis, equine infectious anaemia, vesicular stomatitis and rabies or a period of at least 15 days in the case of anthrax, beginning on the day on which following the destruction of the animals the disinfection of the premises was satisfactorily completed;]

- (h) come from donor mares which during a period of the past 30 days prior to the collection of the (*)[ova] (*)[embryos] were kept in holdings in which none of the equidae has shown clinical signs of contagious equine metritis for a period of at least 60 days;
- (i) come from donor mares which were not used for natural breeding during a period of at least 30 days prior to the date of the collection of the (*)[ova] (*)[embryos] and between the date of the first samples referred to in (j)(i) and (j)(ii) and the date of the collection of the (*)[ova] (*)[embryos];
- (j) come from donor mares which have undergone the tests, which meet at least the requirements of the WOAHA Manual carried out in a laboratory which is recognised by the competent authority and has the tests referred to hereinafter included in its accreditation equivalent to GB requirements, as follows:
 - (*)[(i)for equine infectious anaemia (EIA), an agar-gel immuno-diffusion test (AGID or Coggins test) or an enzyme-linked immunosorbent assay (ELISA) with a negative result carried out on a blood sample taken on, being not less than 14 days following the date of commencement of the period referred to in point (i), and the test was last carried out on a blood sample taken on ; being not more than 90 days prior to the date of the collection of the (*)[ova] (*)[embryos] intended for imports into Great Britain;]
 - (ii) for contagious equine metritis (CEM) an agent identification test carried out with a negative result on at least two specimens (swabs) taken during the period referred to in point (h) from at least the mucosal surfaces of the clitoral fossa and the clitoral sinuses of the donor mare;
 - (*)**EITHER** [(iii) on two occasions with an interval of not less than 7 days on and on, in the case of isolation of *Taylorella equigenitalis* after cultivation under microaerophilic conditions for a period of at least 7 days, set up within 24 hours after taking the specimens from the donor animal, or 48 hours where the specimens are kept cool during transport,]
 - (*)**AND/OR** [(iv) on one occasion on, in the case of detection of genome of *Taylorella equigenitalis* by a polymerase chain reaction (PCR) or real-time PCR, carried out within 48 hours after taking the specimens from the donor animal,]

The samples referred to in points (j)(iii) and (j)(iv) were in no case taken earlier than 7 days (systemic treatment) or 21 days (local treatment) after antimicrobial treatment of the donor stallion and were placed in transport medium with activated charcoal, such as Amies medium, before dispatch to the laboratory.

- (k) come from donor mares which to the best of my knowledge and as far as I could ascertain, were not in contact with equidae suffering from an infectious or contagious disease during the period of 15 days immediately preceding the collection;
- (l) come from donor mares which on the day of the collection of the (*)[ova] (*)[embryos] did not show clinical signs of an infectious or contagious disease;
- (m) were (*)[collected] (*)[produced] after the date on which the embryo (*)[collection] (*)[production] team described in Part I of this certificate Box I.11 was approved by the competent authority of the exporting country;
- (n) were processed and stored under approved conditions for a period of at least 30 days immediately after their (*)[collection](*) [production], and were transported in accordance with GB requirements;

(*) Keep as appropriate.

II.a. Certificate reference no.	II.b.
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Official Veterinarian

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.11: The *place of origin* shall correspond to the embryo collection team or embryo production team by which the ova/embryos were collected/produced, processed, stored and approved in accordance with Article 17(3)(b) of Directive 92/65/EEC.
- Box reference I.22: The *number of packages* shall correspond to the number of containers.
- Box reference I.23: The identification of container and seal number shall be indicated.
- Box reference I.28: The *category*: specify if *in vivo* derived embryos, *in vivo* derived ova, *in vitro* produced embryos or micromanipulated embryos.
- The *donor identity* shall correspond to the official identification of the animal.
- The *date of collection* shall be indicated in the following format: dd/mm/yyyy

Part II

Animal Health

Only third countries or parts of the territory of third countries listed in columns 2 and 4 in a document relating to 'equidae' published on gov.uk, in accordance with Implementing Regulation 2018/659, respectively from which imports of registered equidae and equidae for breeding and production are also authorised and as indicated in column 14 of that document. (†)

AH/E353D Establishment requirement (Collection centre)

Delete if consignment is of ova only.

GB requirements refers to semen compliant to conditions laid down in Directive 92/65.

GB requirements refer to conditions for approval of semen collection centres laid down in Article 11(2) or 17(3)(b) of Directive 92/65. Semen must be collected from approved semen collection centres listed in accordance with Article 11(4) or Article 17(3)(b) of Directive 92/65.

Gov.uk listing refers to columns 2 and 4 of a document relating to 'equidae' published on gov.uk, in accordance with Implementing Regulation 2018/659 from which the import of equine semen collected from registered horses, registered equidae or equidae for breeding and production is authorised in accordance with Article 4 of Implementing Regulation 2018/659 and indicated in columns 11,12 and 13 of that document. (†)

Imports of equine semen are authorised from third countries listed in column 2 in a document relating to 'equidae' published on gov.uk, in accordance with Implementing Regulation 2018/659 provided that the semen was collected in the part of the territory of the third country detailed in column 4 of that document from a donor stallion of the category of equidae positively indicated in column 11,12 or 13 of that document. (†)

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AH/E373A Establishment requirement

Directive 92/65 lays down animal health requirements governing - imports into Great Britain of animals, semen, ova and embryos not subject to animal health requirements laid down in legislation referred to in Annex F to the Directive.

Directive 2009/156 of 30 November 2009 on animal health conditions governing the movement and importation from third countries of equidae.

Regulation 2017/625 on official controls and other official activities performed to ensure the application of food and feed law, rules on animal health and welfare, plant health and plant protection products (Official Controls Regulation).

Point (a)

Only approved embryo collection teams and embryo production teams listed in accordance with Article 17(3)(b) of Directive 92/65.

GB requirements refers to Chapter I(III) of Annex D to Directive 92/65.

(b) GB requirements refer to Chapter III(II) of Annex D to Directive 92/65.

(e) GB requirements refer to Article 13 of Directive 2009/156.

(e) (i) GB requirements refer to Article 5(2)(a) and (b) of Directive 2009/156.

Points (f)(ii) and (j) – WOAHA Manual refers to the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals of the WOAHA.

Points (f)(ii) and (j)(i), (j)(iii), (j)(iv) - *Insert date*.

(g) GB requirements refer to conditions for a holding laid down in Article 4(5) of Directive 2009/156.

(j) GB requirements refer to conditions provided for in Article 37 of Regulation 2017/625.

(j) (i) The agar gel immunodiffusion test (AGID or Coggins test) or the ELISA for equine infectious anaemia are not required for donor equidae which have continuously resided in Iceland since birth, provided that Iceland has remained officially free of equine infectious anaemia and no equidae and their semen, ova and embryos have been introduced into Iceland from outside prior to and during the period the ova or embryos were collected and the semen was used for fertilisation.

(n) GB requirements refer to conditions which satisfy the terms laid down in Chapter III(II) of Annex D to Directive 92/65.

(^t) The document(s) referred to above can be found at:

[EU and EFTA countries approved to export animals and animal products to Great Britain](https://www.data.gov.uk/dataset/4698a65d-1a3b-42d1-981e-df869e04185b/eu-and-efta-countries-approved-to-export-animals-and-animal-products-to-great-britain)

(Available at: <https://www.data.gov.uk/dataset/4698a65d-1a3b-42d1-981e-df869e04185b/eu-and-efta-countries-approved-to-export-animals-and-animal-products-to-great-britain>)

[Non-EU countries approved to export animals and animal products to Great Britain](https://www.data.gov.uk/dataset/b92627b0-dd7b-4e1d-ba36-e25424f55eeb/non-eu-countries-approved-to-export-animals-and-animal-products-to-great-britain)

(Available at: <https://www.data.gov.uk/dataset/b92627b0-dd7b-4e1d-ba36-e25424f55eeb/non-eu-countries-approved-to-export-animals-and-animal-products-to-great-britain>)