

Model health certificate for 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 (Regulation 2018/659) GBHC050E

COUNTRY: Countries subject to transitional import arrangements (*)

Health certificate to Great Britain, Channel Islands and Isle of Man

Part I: Details of dispatched consignment	I.1. Consignor Name Address		I.2. Certificate reference number		I.2.a. UNN			
	Tel.		I.3. Central Competent Authority					
			I.4. Local Competent Authority					
	I.5. Consignee Name Address Postal Code Tel.		I.6. Person responsible for the load in Great Britain, Channel Islands and Isle of Man Name Address Postal Code 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 Embryo team Names Address Postal Code		I.12. Place of destination Holding Name Address Postal Code					
	I.13. Place of loading		I.14. Date of departure					
	I.15. Means of transport Aeroplane Ship Road vehicle Railway wagon Other		I.16. Entry BCP in Great Britain, Channel Islands and Isle of Man					
	Identification: Documentary references:		I.17.					
	I.18. Description of commodity		I.19. Commodity code (HS code) 05 11 99 85					
I.21.		I.20. Quantity		I.22. Number of packages				
I.23. Seal/Container No.		I.24.						
I.25. Commodity certified for: Artificial reproduction								
I.26. For transit through Great Britain, Channel Islands and Isle of Man to third country Third country		I.27. For import or admission into Great Britain, Channel Islands and Isle of Man ISO code						
I.28. Identification of the commodities								
Species (Scientific Name)	Category	Donor identity	Date of collection	Quantity				

**COUNTRY: Countries subject to
transitional import arrangements (*)**

Equine ova/embryos - Section A

Part II: Certification

II. Health information	II.a. Certificate reference number	II.b. UNN
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I, the undersigned, official veterinarian, of the exporting country ⁽²⁾
(name of exporting country)

hereby certify that:

II.1. The ova ⁽¹⁾/embryos ⁽¹⁾ described above:

II.1.2. were collected ⁽¹⁾/produced ⁽¹⁾ by the team ⁽³⁾ described in Box I.11,
which has been approved and supervised in accordance with Chapter I(III) of
Annex D to Directive 92/65/EEC ⁽⁴⁾ and is subject to inspection by an official
veterinarian at least once every calendar year;

II.1.3. were collected ⁽¹⁾/produced ⁽¹⁾, processed and stored in accordance with
the requirements of Chapter III(II) of Annex D to Directive 92/65/EEC;

II.1.4. 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;

II.1.5. 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
Box II.I.6, 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;

II.1.6. come from donor mares which:

II.1.6.1. 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 Article 13 of Directive 2009/156/EC⁽⁵⁾, in that part of
the territory of the exporting country which was during that period:

- not considered to be infected with African horse sickness in
accordance with Article 5(2)(a) and (b) of Directive 2009/156/EC,
- free from Venezuelan equine encephalomyelitis for a period of at
least 2 years,
- free from glanders and dourine for a period of at least 6 months;

⁽¹⁾ either [II.1.6.2. 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:]

⁽¹⁾ or [II.1.6.2. 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
relevant Chapter of the Manual of Diagnostic Tests and Vaccines for
Terrestrial Animals of the OIE on a blood sample taken on
.....⁽⁶⁾ within 30 days prior to the collection of the ova ⁽¹⁾
/ embryos ⁽¹⁾ :]

⁽¹⁾ either [II.1.6.3. 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 ⁽¹⁾ / embryos ⁽¹⁾ until the
date of their dispatch the conditions for a holding laid down in Article
4(5) of Directive 2009/156/EC, and in particular:]

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Equine ova/embryos - Section A

II. Health information	II.a. Certificate reference number	II.b. UNN
<p>(1) or [II.1.6.3.</p> <p>(1) either [II.1.6.3.1.</p> <p>(1) or [II.1.6.3.1.</p> <p>II.1.6.4.</p> <p>II.1.6.5.</p> <p>II.1.6.6.</p>	<p>in the case of frozen ova⁽¹⁾ / embryos⁽¹⁾, 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 ⁽¹⁾ / embryos ⁽¹⁾ until the end of the period of 30 days mandatory storage at approved premises, the conditions for a holding laid down in Article 4(5) of Directive 2009/156/EC, and in particular:]</p> <p>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:</p> <ul style="list-style-type: none"> - 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, - 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, - from vesicular stomatitis for a period of at least 6 months from the last recorded case, - from rabies for a period of at least one month from the last recorded case, - from anthrax for a period of at least 15 days from the last recorded case,] <p>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 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;]</p> <p>during a period of the past 30 days prior to the collection of the ova ⁽¹⁾ / embryos⁽¹⁾ 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;</p> <p>were not used for natural breeding during a period of at least 30 days prior to the date of the collection of the ova ⁽¹⁾ / embryos ⁽¹⁾ and between the date of the first samples referred to in points II.1.6.6.1 and II.1.6.6.2 and the date of the collection of the ova ⁽¹⁾ / embryos ⁽¹⁾;</p> <p>have undergone the tests, which meet at least the requirements of the relevant Chapters of the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals of the OIE, 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 that provided for in Article 37 of Regulation (EU) 2017/625⁽⁷⁾, as follows:</p>	

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Equine ova/embryos - Section A

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<p>(⁸) [II.1.6.6.1. 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 (⁶), being not less than 14 days following the date of commencement of the period referred to in point II.1.6.5, and the test was last carried out on a blood sample taken on (⁶); being not more than 90 days prior to the date of the collection of the ova(¹)/ embryos(¹)intended for imports into Great Britain);]</p> <p>II.1.6.6.2 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 II.1.6.5 from at least the mucosal surfaces of the clitoral fossa and the clitoral sinuses of the donor mare;</p> <p>(¹)either [II.1.6.6.2.1. on two occasions with an interval of not less than 7 days on (⁶) and on (⁶), in the case of isolation of <i>Taylorella equigenitalis</i> 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,]</p> <p>(¹)and/or [II.1.6.6.2.2. on one occasion on (⁶), in the case of detection of genome of <i>Taylorella equigenitalis</i> by a polymerase chain reaction (PCR) or real-time PCR, carried out within 48 hours after taking the specimens from the donor animal,]</p> <p>The samples referred to in points II.1.6.6.2.1 and II.1.6.6.2.2 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.</p> <p>II.1.6.7. 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;</p> <p>II.1.6.8. on the day of the collection of the ova (¹)/ embryos (¹) did not show clinical signs of an infectious or contagious disease;</p> <p>II.1.7. were collected (¹)/produced (¹) after the date on which the embryo collection (¹) /production (¹) team described in Box I.11 was approved by the competent authority of the exporting country;</p> <p>II.1.8. were processed and stored under approved conditions for a period of at least 30 days immediately after their collection (¹)/production (¹), and were transported under conditions which satisfy the terms laid down in Chapter III (II) of Annex D to Directive 92/65/EEC;</p> <p>II.2. The embryos described above were conceived by artificial insemination (¹)/as a result of <i>in vitro</i> fertilisation (¹) using semen meeting the requirements of Directive 92/65/EEC and coming from semen collection centres approved in accordance with Article 11(2) or 17(3)(b) of Directive 92/65/EEC (⁹) and located respectively in Great Britain or in a third country or parts of the territory of a third country listed in columns 2 and 4 of Annex 1 to Commission Implementing Regulation (EU) 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 Commission Implementing Regulation (EU) 2018/659 and indicated in columns 11,12 and 13 of Annex 1 thereto. (¹⁰) (¹¹);</p>		

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Equine ova/embryos - Section A

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(¹²) [II.3. The ova used for *in vitro* production of the embryos described above comply with the requirements of Annex D to Directive 92/65/EEC and in particular the requirements set up in points II.1.1 to II.1.8 of this certificate.]

Notes

(*) Those countries subject to the transitional import arrangements include: an EU member State; Liechtenstein; Norway and Switzerland.

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).

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

Part I:

Box 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 I.16: Do not use this box until the end of the transitional staging period.

Box I.22: The number of packages shall correspond to the number of containers.

Box I.23: The identification of container and seal number shall be indicated.

Box 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:

(1) Delete as appropriate.

(2) Only third countries or parts of the territory of third countries listed in columns 2 and 4 of Annex 1 to Commission Implementing Regulation (EU) 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 Annex 1 thereto.

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

(4) Council Directive 92/65/EEC of 13 July 1992 laying down animal health requirements governing trade in and imports into the Community of animals, semen, ova and embryos not subject to animal health requirements laid down in specific Community rules referred to in Annex A(I) to Directive 90/425/EEC.

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

(6) Insert date.

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Equine ova/embryos - Section A

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- (7) Regulation (EU) 2017/625 of the European Parliament and of the Council of 15 March 2017 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).
- (8) 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.
- (9) Only approved semen collection centres listed in accordance with Article 11(4) or Article 17(3) (b) of Directive 92/65/EEC.
- (10) Imports of equine semen are authorised from third countries listed in column 2 of Annex 1 to Commission Implementing Regulation (EU) 2018/659 provided that the semen was collected in the part of the territory of the third country detailed in column 4 from a donor stallion of the category of equidae positively indicated in column 11,12 or 13 of Annex I thereto.
- (11) Does not apply to ova.
- (12) Delete if none of the embryos in the consignment was produced by *in vitro* fertilisation of ova.
- The signature and the stamp must be in a different colour to that of the printing.

Official Veterinarian

Name (in capital letters):

Qualification and title:

Date:

Signature:

Stamp: