UNITED KINGDOM **Animal health certificate to the EU**

Part I: Description of consignment

I.1 Consignor/Exporter
Name
Address

Country ISO country code

I.2 Certificate reference
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I.2a

I.3 Central Competent Authority
DEPARTMENT FOR ENVIRONMENT, FOOD & RURAL AFFAIRS

I.4 Local Competent Authority
ANIMAL AND PLANT HEALTH AGENCY

I.5 Consignee/Importer
Name
Address

Country ISO country code

I.6 Operator responsible for the consignment
Name
Address

Country ISO country code

I.7 Country of origin ISO country code

I.8 Region of origin Code

I.9 Country of destination ISO country code

I.10 Region of destination Code

I.11 Place of dispatch Registration/Approval No
Name
Address

Country ISO country code

I.12 Place of destination Registration/Approval No
Name
Address

Country ISO country code

I.13 Place of loading

I.14 Date and time of departure

I.15 Means of transport

☐ Aircraft ☐ Vessel

☐ Railway ☐ Road vehicle

Identification

I.16 Entry Border Control Post

I.17 Accompanying documents
Type Code
Country ISO country code
Commercial document reference

I.18 Transport conditions ☐ Ambient ☐ Chilled ☐ Frozen

I.19 Container number/Seal number
Container No Seal No

I.20 Certified as or for

☒ Germinal products

I.21 ☐ **For transit**
Third country ISO country code

I.22 ☐ **For internal market**

I.23

I.24 Total number of packages

I.25 Total quantity

I.26

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I.27 Description of consignment

1

CN code	Species	Subspecies/Category	Identification number	Quantity
Type	Approval or registration number of plant/establishment/centre		Identification mark	Date of collection/production

2

CN code	Species	Subspecies/Category	Identification number	Quantity
Type	Approval or registration number of plant/establishment/centre		Identification mark	Date of collection/production

3

CN code	Species	Subspecies/Category	Identification number	Quantity
Type	Approval or registration number of plant/establishment/centre		Identification mark	Date of collection/production

4

CN code	Species	Subspecies/Category	Identification number	Quantity
Type	Approval or registration number of plant/establishment/centre		Identification mark	Date of collection/production

5

CN code	Species	Subspecies/Category	Identification number	Quantity
Type	Approval or registration number of plant/establishment/centre		Identification mark	Date of collection/production

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Part II: Certification

II. Health information

I, the undersigned, official veterinarian, of the exporting country (1) _____ hereby certify that: (*name of exporting country*)

II.1. The [ova] (2) [embryos] (2) described in Part I:

II.1.2. were [collected] (2) [produced] (2) by the team (3) described in box I.11, which had been approved and supervised in accordance with Chapter I(III) of Annex D to Directive 92/65/EEC and was subject to inspection by an official veterinarian at least once every calendar year;

II.1.3. were [collected] (2) [produced] (2), 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.1.6., 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 3 months (or since entry if they were directly imported from a Member State during the 3 months period) in the exporting country or, in the case of regionalisation according to Article 13 of Directive 2009/156/EC (4), 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 at least 2 years,

– free from glanders and dourine for at least 6 months;

(2) *either* [II.1.6.2. originated from a country of export which was on the day of collection free of vesicular stomatitis for at least 6 months;]

(2) *or* [II.1.6.2. were tested by a virus neutralisation test for vesicular stomatitis on a blood sample taken on _____ (5) within 30 days prior to collection, with negative result at a serum dilution of 1 in 12;]

(2) *either* [II.1.6.3. during the past 30 day period prior to collection have been located in holdings under veterinary supervision which fulfilled from the day of collection of [ova] (2) [embryos] (2) 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:]

(2) *or* [II.1.6.3. during the past 30 day period prior to collection have been located in holdings under veterinary supervision which fulfilled from the day of collection of [ova] (2) [embryos] (2) until, in the case of frozen [ova] (2) [embryos] (2), the period of 30 days mandatory storage at approved premises elapsed, the conditions for a holding laid down in Article 4(5) of Directive 2009/156/EC and in particular:]

(2) *either* [II.1.6.3.1. following a case of a disease mentioned below not all the animals of species susceptible to the disease located on the holding were slaughtered or killed and the holding has been free:

– from any type of equine encephalomyelitis for 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 (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 at least 6 months from the last recorded case,

– from rabies for at least one month from the last recorded case,

– from anthrax for at least 15 days from the last recorded case,]

(2) *or* [II.1.6.3.1. following a case of a disease mentioned below all the animals of species susceptible to the disease located in the holding have been slaughtered or killed and the premises disinfected, the holding has been free for at least 30 days from any type of equine encephalomyelitis, equine infectious anaemia, vesicular stomatitis and rabies or 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;]

II.1.6.4. during the past 30 days prior to collection have been kept in holdings each of them having been free from clinical signs of contagious equine metritis for at least 60 days;

II.1.6.5. have not been used for natural breeding during at least 30 days prior to the date of collection of ova or embryos and between the date of the first samples referred to in points II.1.6.6 and II.1.6.7 and the date of the collection of ova and embryos;

II.1.6.6. have been subjected with negative result to an agar-gel immuno-diffusion test (Coggins test) or an ELISA for equine infectious anaemia carried out on a blood sample taken on _____ (5) being during the past 30 days prior to the date of the first collection of ova or embryos and not more than 90 days before the ova or embryos were collected (6);

II.1.6.7. have been subjected to an agent identification test for contagious equine metritis by isolation of *Taylorella equigenitalis* after a cultivation of 7 to 14 days carried out with negative results in each case on samples taken during the past 30 days prior to the date of the first collection of ova or embryos from mucosal surfaces of the clitoral fossa and clitoral sinuses

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on two consecutives oestrus periods on _____ (5) and on _____ (5), and on an additional culture specimen taken during one of the oestrus periods from the endometrial cervix on _____ (5);

II.1.6.8. to the best of my knowledge and as far as I could ascertain, have not been in contact with equidae suffering from an infectious or contagious disease during the 15 days immediately preceding the collection;

II.1.6.9. have on the day of collection of [ova] (2) [embryos] (2) not shown clinical signs of an infectious or contagious disease;

II.1.7. were [collected] (2) [produced] (2) after the date on which the embryo [collection] (2) [production] (2) team described in box I.11 was approved by the competent authority of the exporting country;

II.1.8. were processed and stored under approved conditions for at least 30 days immediately after their [collection] (2) [production] (2), and were transported under conditions which satisfy the terms laid down in Chapter III(II) of Annex D to Directive 92/65/EEC;

II.2. The embryos described in Part I were conceived [by artificial insemination] (2) [as a result of *in vitro* fertilisation] (2) 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 and located respectively in a Member State of the European Union or in a third country or parts of the territory of third country listed in columns 2 and 4 of the table in Annex I 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 I thereto. (7)(8);

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 animal health certificate (2).

Notes

This animal health certificate is intended for the entry into the Union of oocytes and embryos of equine animals, including when the Union is not the final destination of the oocytes and embryos.

In accordance with the Agreement on the withdrawal of the United Kingdom of Great Britain and Northern Ireland from the European Union and the European Atomic Energy Community, and in particular Article 5(4) of the Windsor Framework (see Joint Declaration No 1/2023 of the Union and the United Kingdom in the Joint Committee established by the Agreement on the withdrawal of the United Kingdom of Great Britain and Northern Ireland from the European Union and the European Atomic Energy Community of 24 March 2023, OJ L 102, 17.4.2023, p. 87) in conjunction with Annex 2 to that Framework, references to the Union in this animal health certificate include the United Kingdom in respect of Northern Ireland.

This animal health certificate shall be completed in accordance with the notes for the completion of certificates provided for in Chapter 4 of Annex I to Commission Implementing Regulation (EU) 2020/2235.

Part I:

Box reference I.11: "Place of dispatch": Indicate the unique approval number and the name and address of the embryo collection or production team of dispatch of the consignment of oocytes or embryos. Only embryo collection or production teams approved in accordance with Article 17(3), point (b), of Council Directive 92/65/EEC and listed on the Commission website:

http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm.

Box reference I.12: "Place of destination": Indicate the address and unique registration or approval number of the establishment of destination of the consignment of oocytes or embryos.

Box reference I.19: Seal number shall be indicated.

Box reference I.24: Total number of packages shall correspond to the number of containers.

Box reference I.27: "Type": specify if *in vivo* derived embryos, *in vivo* derived oocytes, *in vitro* produced embryos or micromanipulated embryos.

"Identification number": Indicate the identification number of each donor animal.

"Identification mark": Indicate the mark on the straw or other packages where oocytes or embryos of the consignment are placed.

"Date of collection/production": Indicate the date on which oocytes or embryos of the consignment were collected or produced.

"Approval or registration number of plant/establishment/centre": Indicate the unique approval number of the embryo collection or production team by which the oocytes or embryos of the consignment were collected or produced.

"Quantity": Indicate the number of straws or other packages with the same mark.

Part II:

(1) Only third countries or territories, or zones thereof listed in column 1 of the table in Part 1 of Annex XII to Commission Implementing Regulation (EU) 2021/404 from which the entry into the Union of equine animals, other than for slaughter, is also authorised and as indicated in column 3 of the table in Part 1 of that Annex.

(2) Delete if not applicable.

(3) Only embryo collection or production teams listed in accordance with Article 17(3), point (b), of Directive 92/65/EEC on the Commission website:

https://ec.europa.eu/food/animals/semen/equine_en.

(4) OJ L 192, 23.7.2010, p. 1.

(5) Insert date.

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(6) The agar gel immunodiffusion test (Coggins test) or the ELISA for equine infectious anaemia are not required for donor equine animals 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, oocytes and embryos have been introduced into Iceland from outside prior to and during the period the semen was collected.

(7) Only semen collection centres approved by the competent authority of a third country or territory, or zone thereof listed in Part 1 of Annex XII to Implementing Regulation (EU) 2021/404 for semen of equine animals or by the competent authority of a Member State.

(8) Does not apply to ova.

Official veterinarian

Name (in capital letters)

Date

Qualification and title

Stamp

Signature