ANIMAL HEALTH CERTIFICATE FOR THE MOVEMENT BETWEEN MEMBER STATES OF CONSIGNMENTS OF STOCKS OF OOCYTES AND EMBRYOS OF EQUINE ANIMALS COLLECTED OR PRODUCED, PROCESSED AND STORED IN ACCORDANCE WITH DIRECTIVE 92/65/EEC AFTER 30 SEPTEMBER 2014 AND BEFORE 21 APRIL 2021, DISPATCHED AFTER 20 APRIL 2021 BY THE EMBRYO COLLECTION OR PRODUCTION TEAM BY WHICH THE OOCYTES OR EMBRYOS WERE COLLECTED OR PRODUCED

(MODEL 'EQUI-OOCYTES-EMB-B-INTRA')

EUROPEAN UNION						**INTRA**
Part I: Description of consignment	**I.1**	**Consignor** Name Address Country	ISO country code	**I.2** **I.2a** **I.3** **I.4**	**IMSOC reference** **Local reference** **Central Competent Authority** **Local Competent Authority**	**QR CODE**
	I.5	**Consignee** Name Address Country	ISO country code	**I.6**	**Operator conducting assembly operations independently of an establishment** Name Address Country	Registration No ISO country code
	I.7	**Country of origin**	ISO country code	**I.9**	**Country of destination**	ISO country code
	I.8	**Region of origin**	Code	**I.10**	**Region of destination**	Code
	I.11	**Place of dispatch** Name Address Country	Registration/Approval No ISO country code	**I.12**	**Place of destination** Name Address Country	Registration/Approval No ISO country code
	I.13	**Place of loading**		**I.14**	**Date and time of departure**	
	I.15	**Means of transport** □ Vessel □ Railway Identification Document	□ Aircraft □ Road vehicle □ Other	**I.16**	**Transporter** Name Address Country	Registration/Authorisation No ISO country code
				I.17	**Accompanying documents** Type Country Commercial document reference	Code ISO country code
	I.18	**Transport conditions**	□ Ambient	□ Chilled	□ Frozen	
	I.19	**Container number/Seal number** Container No	Seal No			

I.20	**Certified as or for**		
□ Further keeping	□ Slaughter	□ Confined establishment	□ Germinal products
□ Registered equine animal	□ Travelling circus/animal act	□ Exhibition	□ Event or activity near borders
□ Release into the wild	□ Dispatch centre	□ Relaying area/purification centre	□ Ornamental aquaculture establishment
□ Further processing	□ Organic fertilizers and soil improvers	□ Technical use	□ Quarantine or similar establishment
□ Products for human consumption	□ Pollination	□ Live aquatic animals for human consumption	□ Other

I.21	**□ For transit through a third country**	
	Third country	ISO country code
	Exit point	BCP code
	Entry point	BCP code

I.22	**□ For transit through Member State(s)**		**I.23**	**□ For export**	
	Member State	ISO country code		Third country	ISO country code
	Member State	ISO country code		Exit point	BCP code
	Member State	ISO country code			

I.24	**Estimated journey time**	**I.25**	**Journey log**	□ yes	□ no
I.26	**Total number of packages**	**I.27**	**Total quantity**		
I.28	**Total net weight/gross weight (kg)**	**I.29**	**Total space foreseen for the consignment**		

I.30 Description of consignment

CN code	Species	Subspecies/Category	Sex	Identification system	Identification number	Age	Quantity Type
Region of origin		Cold store		Identification mark	Type of packaging		Net weight
Slaughterhouse		Treatment type		Nature of commodity	Number of packages		Batch No
		Date of collection/production		Manufacturing plant	Approval or registration number of plant/establishment/centre	Test	

Part II: Certification

II. Health information

II.a Certificate reference	II.b IMSOC reference

I, the undersigned official veterinarian, hereby certify that:

[(1)*either* [II.1. the *in vivo* derived embryos/*in vivo* derived ova[(1)] described in Part I were collected, processed and stored by an embryo collection team[(2)] approved and supervised in accordance with Chapter I(III)(1) of Annex D to Directive 92/65/EEC[(3)];]

[(1)*or* [II.1. the *in vitro* produced embryos/micromanipulated embryos[(1)] described in Part I were produced, processed and stored by an embryo production team[(2)], approved and supervised in accordance with Chapter I(III) (1) and (2) of Annex D to Directive 92/65/EEC;]

[(1)*either* [II.2. the *in vivo* derived embryos described in Part I meet the requirements of Chapter III(II)(1) of Annex D to Directive 92/65/EEC;]

[(1)*or* [II.2. the *in vivo* derived ova described in Part I meet the requirements of Chapter III(II)(2) of Annex D to Directive 92/65/EEC;]

[(1)*or* [II.2. the *in vitro* produced embryos described in Part I meet the requirements of Chapter III(II)(3) of Annex D to Directive 92/65/EEC;]

[(1)*or* [II.2. the micromanipulated embryos described in Part I meet the requirements of Chapter III(II)(4) of Annex D to Directive 92/65/EEC;]

II.3. the ova or embryos described in Part I come from donor mares which:

II.3.1. come from holdings fulfilling the conditions laid down in Article 4(5) of Directive 2009/156/EC[(4)] onto which only equidae satisfying the conditions laid down in Articles 4 and 5 or Articles 12 to 16 of Directive 2009/156/EC were admitted;

II.3.2. meet the requirements of Chapter IV(4) of Annex D to Directive 92/65/EEC;

II.3.3. were not used for natural breeding during a period of at least 30 days prior to the date of collection of the ova or embryos and between the date of the first sample referred to in points II.3.4.1 and II.3.4.2. and the date of the collection of the ova or embryos;

II.3.4. underwent the tests, which meet at least the requirements of the relevant Chapter 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 in accordance with Article 12 of Regulation (EC) No 882/2004[(5)], as follows:

II.3.4.1. 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 samples taken on ……[(6)], being not less than 14 days following the date of commencement of the period referred to in point II.3.3, and the test was last carried out on a sample of blood taken on ……[(6)]; being not more than 90 days prior to the date of the collection of the ova or embryos intended for trade;

II.3.4.2. 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.3.3 from at least the mucosal surfaces of the clitoral fossa and the clitoral sinuses of the donor mare;

[1]*either* [II.3.4.2.1. on two occasions with an interval of not less than 7 days on……[6] and on……..[6], in the case of isolation of *Taylorella equigenitalis* after cultivation under microaerophilic conditions for a period of at least 7 days, set up within the 24 hour period after taking the specimens from the donor animal, or the 48 hour period where the specimens are kept cool during transport;]

[1]*and/or* [II.3.4.2.2. on one occasion on……..[6], in the case of the detection of genome of *Taylorella equigenitalis* by a polymerase chain reaction (PCR) or real-time PCR test, carried out within the 48 hour period after taking the specimens from the donor animal.]

The samples referred to in points II.3.4.2.1. and II.3.4.2.2. were in no case taken earlier than 7 days (systemic treatment) or 21 days (local treatment) after antimicrobial treatment of the donor mare and were placed in transport medium with activated charcoal, such as Amies medium, before dispatch to the laboratory;

[1]*either* [II.4. the embryos described in Part I were conceived as a result of artificial insemination of the donor mares with semen which was collected, processed, stored and transported under conditions which comply with the requirements of Chapters I(I), II(I) and III(I) of Annex D to Directive 92/65/EEC;]

[1]*or* [II.4. the embryos described in Part I were conceived as a result of *in vitro* fertilisation of ova complying with the conditions set out in point 2 of Chapter III(II) of Annex D to Directive 92/65/EEC with semen which was collected, processed, stored and transported under conditions which comply with the requirements of Chapters I(I), II(I) and III(I) of Annex D to Directive 92/65/EEC;]

[1]*or* [II.4. the ova have not been in contact with semen of the equine species;]

II.5. the ova or embryos described in Part I were sent to the place of loading in a sealed container in accordance with point 6 of Chapter III(II) of Annex D to Directive 92/65/EEC and bearing the number detailed in Box I.19.

Notes

This animal health certificate shall be completed according to the notes for the completion of certificates provided for in Chapter 2 of Annex I to Commission Implementing Regulation (EU) 2020/2235.

Part I:

Box I.11: The place of dispatch shall correspond to the embryo collection team or embryo production team of ova/embryos collection/production.

Box I.12: The place of destination shall correspond to the embryo collection team, embryo production team or to the holding of ova/embryos destination.

Box I.19: The identification of container and Seal number shall be indicated.

Box I.30: "Type": specify if: *in vivo* derived embryos, *in vivo* derived oocytes, *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 embryo collection or production teams approved by the competent authority and listed in accordance with Article 11(4) of Directive 92/65/EEC.

(3) OJ L 268, 14.9.1992, p. 54.

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

(5) OJ L 165, 30.4.2004, p. 1.

(6) Insert date.

Official veterinarian	
Name (in capital letters)	Qualification and title
Local Control Unit name	Local Control Unit code
Date	
Stamp	Signature