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 31 AUGUST 2010 AND BEFORE 1 OCTOBER 2014, 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-C-INTRA')

EUROPEAN UNION					INTRA
Part I: Description of consignment	I.1	**Consignor** Name Address Country ISO country code	I.2	**IMSOC reference**	
			I.2a	**Local reference**	
			I.3	**Central Competent Authority**	**QR CODE**
			I.4	**Local Competent Authority**	
	I.5	**Consignee** Name Address Country ISO country code	I.6	**Operator conducting assembly operations independently of an establishment** Name Registration No Address Country 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 Registration/Approval No Address Country ISO country code	I.12	**Place of destination** Name Registration/Approval No Address Country ISO country code	
	I.13	**Place of loading**	I.14	**Date and time of departure**	
	I.15	**Means of transport** □ Vessel □ Aircraft □ Railway □ Road vehicle Identification □ Other Document	I.16	**Transporter** Name Registration/Authorisation No Address Country ISO country code	
			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**		
□ 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	

EUROPEAN UNION **Certificate model EQUI-OOCYTES-EMB-C-INTRA**

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;]

(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. coming 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 have been admitted;

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

II.3.3. 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 sample referred to in points II.3.4. and II.3.5. and the date of the collection of ova and embryos;

II.3.4. 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 samples taken on ……(3), being during the past 30 days prior to the date of the first collection of ova or embryos and the last test was carried out on a sample of blood taken on ……(3); being not more than 90 days before the ova and embryos were collected;

II.3.5. 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 on two consecutives oestrus periods on……(3) and on……..(3), and on an additional culture specimen taken during one of the oestrus periods from the endometrial cervix on……(3);

(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 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: Place of dispatch shall correspond to the embryo collection team or embryo production team of ova/embryos collection/production.

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

Box I.19: 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.

Donor identity shall correspond to the official identification of the animal.

Date of collection shall be indicated in the following format: dd/mm/yyyy.

Approval number of the team shall correspond to the embryo collection team or embryo production team of ova/embryos collection/production.

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 Council Directive 92/65/EEC.

(3) Insert date.

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

Official veterinarian

Name (in capital letters) Qualification and title

Local Control Unit name Local Control Unit code

Date

Stamp Signature