ANIMAL HEALTH CERTIFICATE FOR THE MOVEMENT BETWEEN MEMBER STATES OF CONSIGNMENTS OF STOCKS OF OOCYTES AND EMBRYOS OF OVINE AND CAPRINE ANIMALS COLLECTED OR PRODUCED, PROCESSED AND STORED IN ACCORDANCE WITH DIRECTIVE 92/65/EEC AFTER 31 AUGUST 2010 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 'OV/CAP-OOCYTES-EMB-B-INTRA')

EUROPEAN UNION					**INTRA**
Part I: Description of consignment	**I.1**	**Consignor** Name Address Country ISO country code	**I.2**	**IMSOC reference**	**QR CODE**
			I.2a	**Local reference**	
			I.3	**Central Competent Authority**	
			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	

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(1)/*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(1)/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 consignment consists of embryos of the ovine or caprine species which comply with the following conditions as regards classical scrapie:

(1)*either* [they were collected from animals which have been kept continuously since birth on a holding or holdings recognised as having a negligible or a controlled risk of classical scrapie in accordance with point 1 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001;]

(1)*or* [they were collected from animals which have been kept continuously for the last three years before the collection on a holding or holdings which have complied for the last three years before collection with the requirements laid down in points (a) to (f) of point 1.3. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001;]

(1)*or* [they were collected from animals which have been kept continuously since birth in a Member State or zone of a Member State with a negligible risk status for classical scrapie approved in accordance with the first subparagraph of point 2.2. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001;]

(1)*or* [they were collected from ovine animals and

(1)*either* [are of the ARR/ARR prion protein genotype;]

(1)*or* [carry at least one ARR allele and were collected after the date of 1 January 2015;]]

II.4. the ova or embryos described in Part I come from female donors of the ovine(1)/caprine species(1) which meet the requirements of Chapter IV(3) of Annex D to Directive 92/65/EEC;

[(1)*either*	[II.5.	the embryos described in Part I were conceived as a result of artificial insemination of the donor females 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.5.	the embryos described in Part I were conceived as a result of *in vitro* fertilisation of ova complying with the conditions in Chapter III(II)(2) 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.5.	the ova have not been in contact with semen of the ovine and caprine species;]
	II.6.	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 embryos collection/production.

Box I.12: Place of destination shall correspond to the embryo collection team, embryo production team, germinal product processing establishment, germinal product storage centre or to the establishment 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.

Identification number 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 Directive 92/65/EEC.

Official veterinarian

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