ANIMAL HEALTH CERTIFICATE FOR THE MOVEMENT BETWEEN MEMBER STATES OF CONSIGNMENTS OF STOCKS OF OOCYTES AND EMBRYOS OF PORCINE 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 'POR-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**	
				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	

Part II: Certification

II. Health information

II.a Certificate reference	II.b IMSOC reference

I, the undersigned official veterinarian, hereby certify that the ova/embryos(1) described in Part I:

II.1. were produced/collected(1), processed and stored by an embryo collection/production(1) team(2) approved and supervised in accordance with Chapter I(III) of Annex D to Directive 92/65/EEC;

II.2. meet the requirements of Chapter III(II) of Annex D to Directive 92/65/EEC;

II.3. come from donor females of the porcine species which meet the requirements of Chapter IV(2) of Annex D to Directive 92/65/EEC;

(1)*either* [II.4. are *in vivo* derived embryos which:

II.4.1. were conceived as a result of artificial insemination with semen meeting the requirements of Directive 90/429/EEC,

II.4.2. originate from a Member State or region thereof:

(1)*either* [listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC;]

(1)*or* [listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;]

(1)*or* [listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC and have been washed with trypsin;]

(1)*or* [listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex II to Decision 2008/185/EC;]

(1)*or* [not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I or II to Decision 2008/185/EC and have been washed with trypsin;]

(1)*or* [not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;];]

(1)*or* [II.4. are *in vitro* produced/micromanipulated(1) embryos which:

II.4.1. were conceived as a result of *in vitro* fertilisation with semen meeting the requirements of Directive 90/429/EEC,

II.4.2. originate from a Member State or region thereof:

(1)*either* [listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC;]

			(1)*or*	[listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;]
			(1)*or*	[listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC and the donor females of the ova used for their production comply with the conditions of Article 1 of Decision 2008/185/EC;]
			(1)*or*	[listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex II to Decision 2008/185/EC;]
			(1)*or*	[not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I or II to Decision 2008/185/EC and the donor females of the ova used for their production comply with the conditions of Article 1 of Decision 2008/185/EC;]
			(1)*or*	[not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;]]
	(1)*or*	[II.4.	are *in vivo* derived ova which originate from a Member State or region thereof:	
			(1)*either*	[listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC;]
			(1)*or*	[listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;]
			(1)*or*	[listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC and which come from donor females complying with the conditions of Article 1 of Decision 2008/185/EC;]
			(1)*or*	[listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex II to Decision 2008/185/EC;]
			(1)*or*	[not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I or II to Decision 2008/185/EC and which come from donor females complying with the conditions of Article 1 of Decision 2008/185/EC;]
			(1)*or*	[not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;]]
		II.5.	were sent to the place of loading in a sealed container under conditions complying with point 6 of Chapter III(II) of Annex D to Directive 92/65/EEC and bearing the number detailed in Box I.23.	

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, 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 indicated in Box I.11.

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.

Official veterinarian

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