Part I: Description of consignment

I.1. Consignor
Name
Address
Country ISO Code

I.2. IMSOC reference

I.2.a. Local reference

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Consignee
Name
Address
Country ISO Code

I.6. Operator conducting assembly operations independently of an establishment
Name
Address
Approval Number
Country ISO Code

I.7. Country of origin ISO Code

I.9. Country of destination ISO Code

I.8. Region of origin Code

I.10. Region of destination Code

I.11. Place of dispatch
Name
Address
Approval Number
Country ISO Code

I.12. Place of destination
Name
Address
Approval Number
Country ISO Code

I.13. Place of loading
Name
Address
Approval Number
Country ISO Code

I.14. Date and time of departure

I.15. Means of Transport

Mode	International transport document	Identification

I.16. Transporter
Name
Address
Approval Number
Country ISO Code

I.17. Accompanying documents
[en] accompanying.document.number
Date of issue
Country
Place of issue

I.18. Transport conditions
Chilled ☐ Ambient ☐ Frozen ☐

I.19. Container No / Seal No

I.20. Certified as
Germinal products ☐

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

I.22. For transit through Member State(s) ☐
Member State ISO Code

I.23. For export ☐
Third country ISO Code
Exit point BCP code

I.24. Estimated journey time

I.25. Journey Log

I.26. Total number of packages
I.28. Total gross weight

I.27. Total quantity

I.30. Description of consignment

Commodity	Species	Identification Number	Quantity	Nature of commodity

Identification Mark	Package count	Date of collection / production	Plant / Establishment / Centre

Part II: Certification

II. Health information

I, the undersigned official veterinarian, hereby certify that:

(1) ○ [II.1. The in vivo derived embryos of porcine animals described in Part I have been collected, processed and stored, and dispatched by the embryo collection team(2) which

II.1.1. is approved and kept in a register by the competent authority;

II.1.2. complies with requirements as regards responsibilities, operational procedures, facilities and equipment set out in Part 2 of Annex I to Delegated Regulation (EU) 2020/686.]

(1) ○ [II.1. The oocytes(1)/ in vitro produced embryos(1)/ micromanipulated embryos(1) of porcine animals described in Part I have been collected or produced, processed and stored, and dispatched by the embryo production team(2) which

II.1.1. is approved and kept in a register by the competent authority;

II.1.2. complies with requirements as regards responsibilities, operational procedures, facilities and equipment set out in Parts 2 and 3 of Annex I to Delegated Regulation (EU) 2020/686.]

II.2. The oocytes(1)/ embryos(1) described in Part I are intended for artificial reproduction and were obtained from the donor animals which

II.2.1. have been born and remained since birth in the Union, or have entered the Union in accordance with the requirements for entry into the Union;

(1)(3) ☐ [II.2.2. come from a Member State or zone thereof which is free from infection with Aujeszky's disease virus or where an approved eradication programme for infection with Aujeszky's disease virus is carried out;]

II.2.3. come from establishments in a Member State or zone thereof, or from establishments under official control by the competent authority in a third country or territory, or a zone thereof

II.2.3.1. in which infection with Brucella abortus, B. melitensis and B. suis in porcine animals has not been reported during the last 42 days prior to collection of the oocytes(1)/ embryos(1), and in which during at least the 12 month period prior to collection of the oocytes(1)/ embryos(1)

(1) ☐ either [II.2.3.2.1. biosecurity and risk mitigating measures set out in Article 19(1)(f)(i) of Delegated Regulation 2020/688 have been introduced;

(1) ☐ and/or [II.2.3.2.2. surveillance for infection with Brucella abortus, B. melitensis and B. suis has been carried out on the porcine animals kept on the establishments in accordance with Article 19(1)(f)(ii) of Delegated Regulation 2020/688;]

II.2.3.2. where no clinical, serological, virological or pathological evidence of infection with Aujeszky's disease virus had been detected during the period of at least 12 months prior to collection of the oocytes(1)/ embryos(1).

II.2.4. were examined by the team veterinarian or its team member and did not show symptoms or clinical signs of transmissible animal diseases on the day of collection of the oocytes(1)/ embryos(1);

II.2.5. are identified as provided for in Article 52 or 54(2) of Delegated Regulation (EU) 2019/2035;

II.2.6. for a period of at least 30 days prior to the date of first collection of the oocytes(1)/ embryos(1) and during the collection period;

II.2.6.1. were kept on establishments not situated in a restricted zone established due to the occurrence of foot-and-mouth disease, infection with rinderpest virus, classical swine fever and African swine fever or of an emerging disease relevant for the porcine animals;

II.2.6.2. were kept on a single establishment where infection with Brucella abortus, B. melitensis and B. suis, infection with rabies virus, anthrax, infection with Aujeszky's disease virus and infection with porcine reproductive and respiratory syndrome virus have not been reported;

II.2.6.3. were not in contact with animals from establishments situated in a restricted zone due to the occurrence of diseases referred to in point II.2.6.1. or from establishments which do not meet the conditions referred to in point II.2.6.2.;

II.2.6.4. were not used for natural breeding;

Part II: Certification

II. Health information

II.2.7. comply with the following conditions as regards foot-and-mouth disease

II.2.7.1. they come from establishments

- situated in an area where foot-and-mouth disease has not been reported within a 10-km radius centred on the establishment for a period of at least 30 days immediately prior to the date of collection of the oocytes(1)/ embryos(1);

- in which foot-and-mouth disease has not been reported during a period of at least 3 months immediately prior to the date of collection of the oocytes(1)/ embryos(1);

(1) ○ either [II.2.7.2. they were not vaccinated against foot-and-mouth disease;]

(1)(4) ○ or [II.2.7.2. they were vaccinated against foot-and-mouth disease during the period of 12 months prior to the date of collection or production of the embryos and

II.2.7.2.1. have not been vaccinated against foot-and-mouth disease within the period of at least 30 days immediately prior to the date of collection of the embryos;

II.2.7.2.2. the semen used for fertilisation was collected from a male donor that complies with the conditions set out in point 1(b) or the semen complies with the conditions set out in point 2 of Chapter I of Part 5 of Annex II to Delegated Regulation (EU) 2020/686;

II.2.7.2.3. prior to freezing, the embryos have been subjected to trypsin washing carried out in accordance with the recommendations of the IETS Manual(5);

II.2.7.2.4. the embryos were stored deep frozen for a period of at least 30 days from the date of collection, and during this period the donor animal has not shown clinical signs of foot-and-mouth disease;]

(1)(6) ☐ [II.2.8. were subjected to a serological test for infection with porcine reproductive and respiratory syndrome virus, with negative results, on two occasions, at an interval of not less than 21 days, the second test being performed within a period of 15 days prior to embryo collection.]

II.3. The oocytes(1)/ embryos(1) described in Part I

II.3.1. has been collected, processed and stored in accordance with animal health requirements set out in Part 2(1)/Part 3(1)/Part 4(1)/Part 5(1) and Part 6 of Annex III to Delegated Regulation (EU) 2020/686;

II.3.2. are placed in straws or other packages on which the mark is applied in accordance with requirements provided for in Article 10 of Delegated Regulation (EU) 2020/686 and that mark is indicated in Box I.30;

II.3.3. are transported in a container which:

II.3.3.1. was sealed and numbered prior to the dispatch by the embryo collection or production team under responsibility of the team veterinarian, or by an official veterinarian, and the seal bears the number as indicated in Box I.19;

II.3.3.2. has been cleaned and either disinfected or sterilised before use, or is single-use container;

(1)(7)II.3.3.3. has been filled in with the cryogenic agent which not have been previously used for other products;

(1)(8) ☐ [II.3.4. are placed in straws or other packages which are securely and hermetically sealed;

II.3.5. are transported in a container where they are separated from each other by physical compartments or by being placed in secondary protective bags.]

(1)(9) ☐ [II.4. The in vivo derived embryos(1)/ in vitro produced embryos(1)/ micromanipulated embryos(1) described in Part I were conceived by artificial insemination using semen coming from a semen collection centre, germinal product processing establishment or germinal product storage centre approved for the collection, processing and/or storage of semen by the competent authority of a Member State or by the competent authority of a third country, territory or zone thereof listed in Annex XI to Commission Implementing Regulation (EU) 2021/404].]

Part II: Certification

II. Health information		
(1)(10) ☐ [II.5.	The following antibiotic or mixture of antibiotics(10) has been added to the collection, processing, washing or storage media: _____]	
II.6.	This certificate is valid for 10 days from the date of issuing.	

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 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.
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.26:	Total number of packages shall correspond to the number of containers.
Box reference I.30:	"Type": specify if in vivo derived embryos, in vivo derived oocytes, in vitro produced embryos or micromanipulated embryos.

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

"Identification mark": indicate 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 was 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 were collected or produced.

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

Part II:

(1)	Delete if not applicable.
(2)	Only embryo collection or production teams approved by the competent authority and included in the register referred to in Article 101(1)(b) of Regulation (EU) 2016/429 and Article 7 of Delegated Regulation (EU) 2020/686.
(3)	Not applicable for in vivo derived embryos subject to trypsin treatment.
(4)	Option available only for the consignment of in vivo derived embryos.
(5)	Manual of the International Embryo Transfer Society — A procedural guide and general information for the use of embryo transfer technology emphasising sanitary procedures, published by the International Embryo Transfer Society 1 111 North Dunlap Avenue, Savoy, Illinois 61 874 USA (http://www.iets.org/).
(6)	Applicable for in vivo derived embryos.
(7)	Applicable for frozen oocytes or embryos.
(8)	Applicable for the consignment where in one container oocytes, in vivo derived embryos, in vitro produced embryos and micromanipulated embryos of porcine animals are placed and transported.
(9)	Does not apply to oocytes.
(10)	Mandatory attestation in case antibiotics were added.
(11)	Insert the name(s) of the antibiotic(s) added and its(their) concentration.

Certifying Officer/Official veterinarian

Part II: Certification

II. Health information		
Name (in capital letters) Date of signature Stamp	Qualification and title Signature	