EUROPEAN UNION

INTRA

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.8. Region of origin Code

I.9. Country of destination ISO 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
Document Type
Accompanying document reference
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.27. Total quantity	I.28. Total gross weight

I.30. Description of consignment

1. 05 PRODUCTS OF ANIMAL ORIGIN, NOT ELSEWHERE SPECIFIED OR INCLUDED

0511 Animal products not elsewhere specified or included; dead animals of Chapter 1 or 3, unfit for human consumption

051199 Other

05119985 Other

Part I: Description of consignment

#1.	Commodity	Identification Number	Quantity	Nature of commodity	Identification Mark
Species		Package count	Date of collection / production	Plant / Establishment / Centre	Type

Part II: Certification

II. Health information

I, the undersigned official veterinarian, hereby certify, that:

II.1. The ☐ [semen] (1) ☐ [in vivo derived embryos] (1) ☐ [oocytes] (1) ☐ [in vitro produced embryos] (1) ☐ [micromanipulated embryos] (1) of the consignment described in Part I:

II.1.1. has/have been collected or produced, processed and stored, and dispatched from the confined establishment (2) which is approved, assigned with a unique approval number and kept in a register by the competent authority;

II.1.2. has/have been collected or produced, processed and stored, and dispatched from the confined establishment which complies with the requirements as regards quarantine, isolation and other biosecurity measures, surveillance and control measures, facilities and equipment referred to in Article 16 of Commission Delegated Regulation (EU) 2019/2035;

(1) either ○ [II.1.3. is/are dispatched from the confined establishment (2) or zone not subject to movement restrictions affecting donor animals of the germinal products to be moved and established for reasons of listed diseases relevant for those species or diseases subject to emergency measures relevant for those species, or those restrictions do not apply to these germinal products because they were collected before the restrictions were established, and it/they has/have not been in contact with other germinal products of a lower health status for an adequate period.]

(1) or ○ [II.1.3. is/are dispatched from the confined establishment (2) or zone subject to movement restrictions affecting donor animals of the germinal products to be moved and established for (3), but derogations from movement restrictions have been granted, and:

(1) ☐ [it/they comply(ies) with the requirements set out in (4);]]

(1) ☐ [and, in particular, it/they is/are (5).]]

II.2. The ☐ [semen] (1) ☐ [oocytes] (1) ☐ [embryos] (1) described in Part I is/are intended for artificial reproduction and was/were obtained from 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;

II.2.2. have remained in a single confined establishment of origin for at least 30 days prior to the date of collection of the ☐ [semen] (1) ☐ [oocytes] (1) ☐ [embryos] (1);

(1) ☐ [II.2.3. are bovine animals and they are identified as provided for in Article 38 of Delegated Regulation (EU) 2019/2035.]

(1) ☐ [II.2.3. are porcine animals and they are identified as provided for in Article 52(1) or Article 54(2) of Delegated Regulation (EU) 2019/2035.]

(1) ☐ [II.2.3. are ovine or caprine animals and they are identified as provided for in Article 45(2) or (4), or Article 46(1), (2) or (3), of Delegated Regulation (EU) 2019/2035.]

(1) ☐ [II.2.3. are equine animals and they are identified as provided for in Article 58(1), Article 59(1) or Article 62(1) of Delegated Regulation (EU) 2019/2035.]

(1) ☐ [II.2.3. are terrestrial animals other than bovine, porcine, ovine, caprine and equine animals and they are identified and registered in accordance with the rules of the confined establishment.]

II.3. The ☐ [semen] (1) ☐ [oocytes] (1) ☐ [embryos] (1) described in Part I comes/come from the confined establishment indicated in box I.11 and is/are destined to another confined establishment.

II.4. According to official information, the ☐ [semen] (1) ☐ [oocytes] (1) ☐ [embryos] (1) described in Part I was/were obtained from donor animals which:

II.4.1. do not come from a confined establishment, nor have been in contact with

Part II: Certification

II. Health information

animals from a confined establishment, situated in a zone subject to movement restrictions established due to the occurrence of a category A disease, referred to in the Annex to Commission Implementing Regulation (EU) 2018/1882, or of an emerging disease relevant for species of those donor animals;

II.4.2. come from a confined establishment where no category D disease relevant for that species as referred to in the Annex to Implementing Regulation (EU) 2018/1882 has been reported for at least 30 days prior to the date of collection of the ☐ [semen] (1) ☐ [oocytes] (1) ☐ [embryos] (1).

II.5. To the best of my knowledge and as declared by the operator, the ☐ [semen] (1) ☐ [oocytes] (1) ☐ [embryos] (1) described in Part I was/were obtained from donor animals which:

II.5.1. have been clinically examined by the establishment veterinarian responsible for the activities carried out at the confined establishment and showed no disease symptoms on the day of collection of the ☐ [semen] (1) ☐ [oocytes] (1) ☐ [embryos] (1);

II.5.2. as much as possible, were not used for natural breeding during at least 30 days prior to the date of collection of the ☐ [semen] (1) ☐ [oocytes] (1) ☐ [embryos] (1) and during the collection period.

II.6. To the best of my knowledge and based on the documentary check of the data submitted by the establishment veterinarian responsible for the activities carried out at the confined establishment, the ☐ [semen] (1) ☐ [oocytes] (1) ☐ [embryos] (1) described in Part I is/are placed in straws or other packages on which the mark is applied in accordance with requirements provided for in:

(1) (6) either ○ [Article 10 of Commission Delegated Regulation (EU) 2020/686 and that mark is indicated in box I.30.]

(1) (7) or ○ [Article 11 of Delegated Regulation (EU) 2020/686 and that mark is indicated in box I.30.]

II.7. The ☐ [semen] (1) ☐ [oocytes] (1) ☐ [embryos] (1) described in Part I:

II.7.1. is/are transported in a container which:

II.7.1.1. was sealed and numbered prior to the date of dispatch by the establishment veterinarian responsible for the activities carried out at the confined establishment, or by an official veterinarian, and the seal bears the number as indicated in box I.19;

II.7.1.2. has been cleaned and either disinfected or sterilised before use, or is single-use container;

(1) (8) ☐ [II.7.1.3. has been filled in with a cryogenic agent which has not been previously used for other products;]

(1) (6) (9) ☐ [II.7.2. is/are placed in straws or other packages which are securely and hermetically sealed;

II.7.3. is/are transported in a container where the different types are separated from each other by physical compartments or by being placed in secondary protective bags.]

Notes

In accordance with the Agreement on the withdrawal of the United Kingdom of Great Britain and Northern Ireland from the European Union and the European Atomic Energy Community, and in particular Article 5(4) of the Protocol on Ireland/Northern Ireland in conjunction with Annex 2 to that Protocol, references to the Union in this animal health certificate include the United Kingdom in respect of Northern Ireland.

This animal health certificate shall be completed in accordance with 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 address and the unique approval number of the confined establishment of dispatch of the consignment of semen, oocytes or embryos.

Box “Place of destination”: Indicate the address and the unique approval number of the confined

Part II: Certification

II. Health information		

reference I.12: establishment of destination of the consignment of semen, oocytes or embryos.

Box reference I.30: “Type”: Specify if semen, 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 the semen, oocytes or embryos of the consignment is/are placed.

“Date of collection/production”: Indicate the date on which the semen, oocytes or embryos of the consignment was/were collected or produced.

“Approval or registration number of plant/establishment”: Indicate the unique approval number of the confined establishment of the collection or production of the semen, oocytes or embryos of the consignment.

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

Part II:

(1) Delete if not applicable.

(2) Confined establishment as defined in Article 4, point (48), of Regulation (EU) 2016/429 of the European Parliament and of the Council.

(3) Insert the name of the disease(s).

(4) Insert the specific reference to the article(s), title, and number of the relevant legal act(s) adopted by the Commission providing for those requirements.

(5) Insert the specific attestation(s) provided for in and required by the relevant legal act(s) adopted by the Commission as referred to in Article 159(2), points (a), (b) and (c), of Regulation (EU) 2016/429.

(6) Applicable for consignments of semen, oocytes or embryos of bovine, porcine, ovine, caprine or equine animals.

(7) Applicable for consignments of semen, oocytes or embryos of terrestrial animals other than bovine, porcine, ovine, caprine or equine animals.

(8) Applicable for frozen semen, oocytes or embryos.

(9) Applicable for consignments where oocytes, in vivo derived embryos, in vitro produced embryos and micromanipulated embryos of bovine, porcine, ovine, caprine or equine animals are placed and transported in one container.

Certifying Officer/Official veterinarian	
Name (in capital letters)	Qualification and title
Date of declaration	Signature
Stamp	