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

II. Health information

I, the undersigned official veterinarian, hereby certify that:

II.1. The germinal product storage centre (1) described in box I.11 at which the ☐ [semen] (2) ☐ [oocytes] (2) ☐ [in vivo derived embryos] (2) ☐ [in vitro produced embryos] (2) ☐ [micromanipulated embryos] (2) was/were stored:

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 5 of Annex I to Commission Delegated Regulation (EU) 2020/686;

II.2. The ☐ [semen] (2) ☐ [oocytes] (2) ☐ [in vivo derived embryos] (2) ☐ [in vitro produced embryos] (2) ☐ [micromanipulated embryos] (2) described in Part I is/are dispatched from:

(2) ○ either [the germinal product storage centre described in box I.11 or a zone not subject to movement restrictions affecting ovine and caprine animals 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.]

(2) ○ or [the germinal product storage centre described in box I.11 or a zone subject to movement restrictions affecting ovine and caprine animals and established for (3), but derogations from movement restrictions have been granted, and:

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

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

II.3. The ☐ [semen] (2) ☐ [oocytes] (2) ☐ [in vivo derived embryos] (2) ☐ [in vitro produced embryos] (2) ☐ [micromanipulated embryos] (2) described in Part I is/are intended for artificial reproduction, and:

(2) ☐ either [II.3.1. has/have been ☐ [collected] (2) ☐ [produced] (2) ☐ [processed] (2) ☐ [stored] (2) ☐ [in a semen collection centre] (2)(6) ☐ [by an embryo collection team] (2)(6) ☐ [by an embryo production team] (2)(6), ☐ [and] (2) ☐ [processed] (2) ☐ [stored] (2) ☐ [in a germinal product processing establishment] (2)(6) ☐ [and stored in a germinal product storage centre] (2)(6) situated in the Member State of its/their collection or production and complying with requirements as regards responsibilities, operational procedures, facilities and equipment set out in ☐ [Part 1] (2) ☐ [Part 2] (2) ☐ [Part 3] (2) ☐ [Part 4] (2) ☐ [Part 5] (2)of Annex I to Delegated Regulation (EU) 2020/686, and was/were moved to the germinal product storage centre indicated in box I.11 situated in the Member State of its/their collection or production under animal health certification requirements at least as strict as those provided for in:

(2) ☐ either [Model OV/CAP-SEM-A-INTRA (7);]

(2) ☐ and/or [Model OV/CAP-SEM-B-INTRA (7);]

(2) ☐ and/or [Model OV/CAP-SEM-C-INTRA (7);]

(2) ☐ and/or [Model in Part A of Annex III to Commission Decision 2010/470/EU (7);]

(2) ☐ and/or [Model in Part B of Annex III to Decision 2010/470/EU (7);]

(2) ☐ and/or [Model in Part C of Annex III to Decision 2010/470/EU (7);]

(2) ☐ and/or [Model in Commission Decision 95/388/EC (7);]

(2) ☐ and/or [Model OV/CAP-OOCYTES-EMB-A-INTRA (7);]

(2) ☐ [Model OV/CAP-OOCYTES-EMB-B-INTRA (7);]

Part II: Certification

II. Health information		

and/or

(2) ☐ [Model OV/CAP-OOCYTES-EMB-C-INTRA (7);]
and/or

(2) ☐ [Model OV/CAP-GP-PROCESSING-INTRA (7);]
and/or

(2) ☐ [Model OV/CAP-GP-STORAGE-INTRA (7);]]
and/or

(2) ☐ [II.3.1. has/have been ☐ [collected] (2) ☐ [produced] (2) ☐ [processed] (2) ☐ [stored] (2) ☐ [in a semen collection centre] (2)(6) ☐ [by an embryo collection team] (2)(6) ☐ [by an embryo production team] (2)(6) ☐ [and] (2) ☐ [processed] (2) ☐ [stored] (2) ☐ [in a germinal product processing establishment] (2)(6) ☐ [and stored in a germinal product storage centre] (2)(6) situated in the Member State of its/their collection or production and complying with requirements as regards responsibilities, operational procedures, facilities and equipment set out in ☐ [Part 1] (2) ☐ [Part 2] (2) ☐ [Part 3] (2) ☐ [Part 4] (2) ☐ [Part 5] (2) of Annex I to Delegated Regulation (EU) 2020/686, and was/were moved to the germinal product storage centre indicated in box I.11 situated in another Member State accompanied by animal health certificate(s) in accordance with:
and/or

(2) ☐ [Model OV/CAP-SEM-A-INTRA (7);]
either

(2) ☐ [Model OV/CAP-SEM-B-INTRA (7);]
and/or

(2) ☐ [Model OV/CAP-SEM-C-INTRA (7);]
and/or

(2) ☐ [Model in Part A of Annex III to Decision 2010/470/EU (7);]
and/or

(2) ☐ [Model in Part B of Annex III to Decision 2010/470/EU (7);]
and/or

(2) ☐ [Model in Part C of Annex III to Decision 2010/470/EU (7);]
and/or

(2) ☐ [Model in Decision 95/388/EC (7);]
and/or

(2) ☐ [Model OV/CAP-OOCYTES-EMB-A-INTRA (7);]
and/or

(2) ☐ [Model OV/CAP-OOCYTES-EMB-B-INTRA (7);]
and/or

(2) ☐ [Model OV/CAP-OOCYTES-EMB-C-INTRA (7);]
and/or

(2) ☐ [Model OV/CAP-GP-PROCESSING-INTRA (7);]
and/or

(2) ☐ [Model OV/CAP-GP-STORAGE-INTRA (7);]]
and/or

(2) ☐ [II.3.1. has/have been ☐ [collected] (2) ☐ [produced] (2) ☐ [processed] (2) ☐ [stored] (2) ☐ [in a semen collection centre] (2)(6) ☐ [by an embryo collection team] (2)(6) ☐ [by an embryo production team] (2)(6), ☐ [and] (2) ☐ [processed] (2) ☐ [stored] (2) ☐ [in a germinal product processing establishment] (2)(6) ☐ [and stored in a germinal product storage centre] (2)(6) situated in a third country or territory, or zone thereof listed in Annex X to Commission Implementing Regulation (EU) 2021/404 and complying with requirements as regards responsibilities, operational procedures, facilities and equipment set out in ☐ [Part 1] (2) ☐ [Part 2] (2) ☐ [Part 3] (2) ☐ [Part 4] (2) ☐ [Part 5] (2) of Annex I to Delegated Regulation (EU) 2020/686, and entered the Union accompanied by animal health certificate(s) in accordance with:
and/or

(2) ☐ [Model OV/CAP-SEM-A-ENTRY (7);]
either

Part II: Certification

II. Health information		

(2) ☐ and/or [Model OV/CAP-SEM-B-ENTRY (7);]

(2) ☐ and/or [Model 1 in Part 2, Section A, of Annex II to Commission Decision 2010/472/EU (7);]

(2) ☐ and/or [Model 2 in Part 2, Section B, of Annex II to Decision 2010/472/EU (7);]

(2) ☐ and/or [Model in Annex II to Decision 2008/635/EC (7);]

(2) ☐ and/or [Model OV/CAP-OOCYTES-EMB-A-ENTRY (7);]

(2) ☐ and/or [Model OV/CAP-OOCYTES-EMB-B-ENTRY (7);]

(2) ☐ and/or [Model OV/CAP-GP-PROCESSING-ENTRY (7);]

(2) ☐ and/or [Model OV/CAP-GP-STORAGE-ENTRY (7);]]

II.3.2. has/have been collected, processed and stored in accordance with animal health requirements set out in Annex III to Delegated Regulation (EU) 2020/686;

II.3.3. is/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/or Article 83, point (a), of Commission Delegated Regulation (EU) 2020/692 and that mark is indicated in box I.30;

II.3.4. is/are transported in a container which:

II.3.4.1. was sealed and numbered prior to the date of dispatch from the germinal product storage centre under responsibility of the centre veterinarian, or by an official veterinarian, and the seal bears the number as indicated in box I.19;

II.3.4.2. has been cleaned and either disinfected or sterilised before use, or is single-use container;

(2) (8) ☐ [II.3.4.3. has been filled in with a cryogenic agent which has not been previously used for other products;]

(2) (9) ☐ [II.3.5. is/are placed in straws or other packages which are securely and hermetically sealed;

II.3.6. 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 unique approval number and the name and address of the germinal product storage centre of dispatch of the consignment of semen, oocytes, and/or embryos. Only germinal product storage centres approved by the competent authority and included in the register referred to in Article 101(1), point (b), of Regulation (EU) 2016/429 of the European Parliament and of the Council and Article 7 of Delegated Regulation (EU) 2020/686.

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 semen, oocytes, and/or embryos.

Box reference I.17: “Accompanying documents”: Number(s) of related original animal health certificate(s) shall correspond to the serial number of the individual official document(s) or animal health certificate(s) that accompanied the semen, oocytes and/or embryos described in Part I from the semen collection centre

Part II: Certification

II. Health information		

where the semen was collected, and/or from the embryo collection or from the embryo production team by which the oocytes and/or embryos were collected or produced, and/or from the germinal product processing establishment where the semen, oocytes or embryos were processed and stored, and/or from the germinal product storage centre where the semen, oocytes or embryos were stored, to the germinal product storage centre described in box I.11. The original(s) of those document(s) or those animal health certificate(s) or the officially endorsed copies thereof shall be attached to this animal helath certificate.

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 semen, in vivo derived embryos, in vivo derived oocytes, in vitro produced embryos or micromanipulated embryos.

“Species”: indicate “Ovis aries” and/or “Capra hircus” as appropriate.

“Identification number”: Indicate identification number of each donor animal.

“Identification mark”: Indicate mark on the straw or other packages where the semen, oocytes and/or embryos of the consignment is/are placed.

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

“Approval or registration number of plant/establishment/centre”: Indicate the unique approval number of the semen collection centre where the semen of the consignment was collected, and/or of the embryo collection team and/or the embryo production team by which the oocytes or embryos of the consignment were collected or produced.

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

Part II:

(1) Only germinal product storage centres approved by the competent authority and included in the register referred to in Article 101(1), point (b), of Regulation (EU) 2016/429 and Article 7 of Delegated Regulation (EU) 2020/686.

(2) Delete if not applicable.

(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) Only germinal product establishments approved by the competent authority and included in the register referred to in Article 101(1), point (b), of Regulation (EU) 2016/429 and Article 7 of Delegated Regulation (EU) 2020/686.

(7) The original(s) of the document(s) or the animal health certificate(s) or the officially endorsed copies thereof that accompanied the semen, oocytes or embryos described in Part I from the semen collection centre where the semen was collected, and/or from the embryo collection team or the embryo production team by which the oocytes and/or embryos were collected or produced, and/or from the germinal product processing establishment where the semen, oocytes or embryos were processed and stored, and/or from the germinal product storage centre where the semen, oocytes or embryos were stored, to the germinal product storage centre of the semen, oocytes and/or embryos dispatch described in box I.11 shall be attached to this animal health certificate.

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

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

Certifying Officer/Official veterinarian

Part II: Certification

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