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.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
Activity ID
Country ISO Code

I.17. Accompanying documents
Commercial document reference
Date of issue
Country
Place of issue

I.18. Transport conditions
Ambient ☐ Chilled ☐ 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.25. Journey Log

I.26. Total number of packages

I.27. Total quantity

I.28. Total gross weight

I.30. Description of consignment

Commodity	Species	Identification Number	Quantity	Nature of commodity

Identification Mark	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 germinal product processing establishment(1) described in Box I.11. at which the semen(2)/ oocytes(2)/ in vivo derived embryos(2)/ in vitro produced embryos(2)/ micromanipluated embryos(2) was/were processed and 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 4 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)/ micromanipluated embryos(2) described in Part I is/are intended for artificial reproduction and

(2) ☐ either [II.2.1. has/have been collected or produced, processed and stored in a semen collection centre(2)(3)/ by an embryo collection team(2)(3)/ by an embryo production team(2)(3), and/or processed and stored in a germinal product processing establishment(2)(3), and/or stored in a germinal product storage centre(2)(3) 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 processing establishment 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 EQUI-SEM-A-INTRA(4);]
(2) ☐ and/or [Model EQUI-SEM-B-INTRA(4);]
(2) ☐ and/or [Model EQUI-SEM-C-INTRA(4);]
(2) ☐ and/or [Model EQUI-SEM-D-INTRA(4);]
(2) ☐ and/or [Model EQUI-OOCTYES-EMB-A-INTRA(4);]
(2) ☐ and/or [Model EQUI-OOCTYES-EMB-B-INTRA(4);]
(2) ☐ and/or [Model EQUI-OOCTYES-EMB-C-INTRA(4);]
(2) ☐ and/or [Model EQUI-OOCTYES-EMB-D-INTRA(4);]
(2) ☐ and/or [Model EQUI-GP-PROCESSING-INTRA(4);]
(2) ☐ and/or [Model EQUI-GP-STORAGE-INTRA(4);]]

(2) ☐ and/or [II.2.1. has/have been collected or produced, processed and stored in a semen collection centre(2)(3)/ by an embryo collection team(2)(3)/ by an embryo production team(2)(3), and/or processed and stored in a germinal product processing establishment(2)(3), and/or stored in a germinal product storage centre(2)(3) 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 processing establishment indicated in Box I.11. situated in another Member State accompanied by certificate(s) in accordance with:

(2) ☐ either [Model EQUI-SEM-A-INTRA(4);]
(2) ☐ and/or [Model EQUI-SEM-B-INTRA(4);]
(2) ☐ and/or [Model EQUI-SEM-C-INTRA(4);]
(2) ☐ and/or [Model EQUI-SEM-D-INTRA(4);]
(2) ☐ and/or [Model EQUI-OOCTYES-EMB-A-INTRA(4);]
(2) ☐ and/or [Model EQUI-OOCTYES-EMB-B-INTRA(4);]
(2) ☐ and/or [Model EQUI-OOCTYES-EMB-C-INTRA(4);]
(2) ☐ and/or [Model EQUI-OOCTYES-EMB-D-INTRA(4);]
(2) ☐ and/or [Model EQUI-GP-PROCESSING-INTRA(4);]
(2) ☐ and/or [Model EQUI-GP-STORAGE-INTRA(4);]]

Part II: Certification

II. Health information

(2) ☐ and/or [II.2.1. has/have been collected or produced, processed and stored in a semen collection centre(2)(3)/ by an embryo collection team(2)(3)/ by an embryo production team(2)(3), and/or processed and stored in a germinal product processing establishment(2)(3), and/or stored in a germinal product storage centre(2)(3) situated in a third country, territory or zone thereof listed in Annex XII to Commission Implementing Regulation (EU) [C(2021)1800] 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 certificate(s) in accordance with:

(2) ☐ either [Model EQUI-SEM-A-ENTRY(4);]

(2) ☐ and/or [Model EQUI-SEM-B-ENTRY(4);]

(2) ☐ and/or [Model EQUI-SEM-C-ENTRY(4);]

(2) ☐ and/or [Model EQUI-SEM-D-ENTRY(4);]

(2) ☐ and/or [Model EQUI-OOCYTES-EMB-A-ENTRY(4);]

(2) ☐ and/or [Model EQUI-OOCYTES-EMB-B-ENTRY(4);]

(2) ☐ and/or [Model EQUI-OOCYTES-EMB-C-ENTRY(4);]

(2) ☐ and/or [Model EQUI-GP-PROCESSING-ENTRY(4);]

(2) ☐ and/or [Model EQUI-GP-STORAGE-ENTRY(4);]

II.2.2. 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.2.3. 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(a) of Commission Delegated Regulation (EU) 2020/692 and that mark is indicated in Box I.30;

II.2.4. is/are transported in a container which:

II.2.4.1. was sealed and numbered prior to the 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.2.4.2. has been cleaned and either disinfected or sterilised before use, or is a single-use container;

(2)(5) ☐ [II.2.4.3. has been filled in with the cryogenic agent which not have been previously used for other products;]

(2)(6) ☐ [II.2.5. is/are placed in straws or other packages which are securely and hermetically sealed;

II.2.6. is/are transported in a container where they are separated from each other by physical compartments or by being placed in secondary protective bags.]

Part II: Certification

II. Health information

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 germinal product processing establishment of dispatch of the consignment of semen, oocytes, and/or embryos. Only germinal product processing establishments 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.
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 certificate(s) shall correspond to the serial number of the individual official document(s) or health certificate(s) that accompanied the semen, oocytes and/or embryos described in Part I from the semen collection centre where the semen was collected, and/or the embryo collection and/or production team by which the oocytes and/or embryos were collected or produced, and/or the germinal product processing establishment where the semen, oocytes or embryos were processed and stored, and/or the germinal product storage centre where the semen, oocytes or embryos were stored to the germinal product processing establishment described in Box I.11. The original(s) of those document(s) or those certificate(s) or the officially endorsed copies thereof must be attached to this 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.
	"Identification number": Indicate identification number of each donor animal.
	"Identification mark": Indicate mark on the straw or other packages where semen, oocytes and/or embryos of the consignment are placed.
	"Date of collection/production": Indicate the date on which 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 was collected, and/or of the embryo collection and/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)	Only germinal product processing establishments 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.
(2)	Delete if not applicable.
(3)	Only germinal product establishments 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.

Part II: Certification

II. Health information

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

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

(6) Applicable for the consignment where in one container semen, oocytes, in vivo derived embryos, in vitro produced embryos and micromanipulated embryos of equine animals are placed and transported.

Certifying Officer/Official veterinarian

Name (in capital letters)

Authority name

Date of signature

Signature

Stamp