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
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 ☐ Frozen ☐ Ambient ☐

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:

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

II.1.1. 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.1.2. are

(1) ○ either [marked by the implantation of a transponder in accordance with Article 17(1) of Regulation (EU) No 576/2013;]

(1) ○ or [marked by a clearly readable tattoo in accordance with Article 17(1) of Regulation (EU) No 576/2013;]

(1) ○ or [identified in accordance with Article 70 of Commission Delegated Regulation (EU) 2019/2035;]

II.1.3. have received an anti-rabies vaccination that complies with the validity requirements set out in Part 1 of Annex VII to Commission Delegated Regulation (EU) 2020/688.

II.2. The semen(1)/ oocytes(1)/ embryos(1) described in Part I comes/come from a registered establishment assigned by the competent authority with a unique registration number as indicated in Box I.11.

II.3. According to official information, the semen(1)/ oocytes(1)/ embryos(1) described in Part I was/were obtained from donor animals which

II.3.1. come from establishments in which infection with rabies virus has not been confirmed for a period of at least 30 days prior to the date of collection of the semen(1)/ oocytes(1)/ embryos(1);

II.3.2. comply with any preventive health measure for diseases or infections other than rabies set out in Part 2 of Annex VII to Delegated Regulation (EU) 2020/688.

II.4. To the best of my knowledge and as declared by the operator, the semen(1)/ oocytes(1)/ embryos(1) was/were obtained from donor animals which

II.4.1. showed no disease symptoms on the day of collection of the semen(1)/ oocytes(1)/ embryos(1);

II.4.2. were not used for natural breeding during a period of at least 30 days prior to the date of collection of the semen(1)/ oocytes(1)/ embryos(1) and during the collection period.

II.5. The semen(1)/ oocytes(1)/ embryos(1) described in Part I is/are placed in a sealed transport container and the seal bears the number as indicated in Box I.19.

II.6. To the best of my knowledge and based on the documentary check of the data submitted by the operator, 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 Article 11 of Commission Delegated Regulation (EU) 2020/686 and that mark is indicated in Box I.30.

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 address and the unique registration number of the establishment of dispatch of the consignment of semen, oocytes or embryos.

Box reference I.12: "Place of destination": Indicate the address and the unique registration number, if assigned by the competent authority, of the 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.

"Species": Indicate "Canis lupus familiaris" or "Felis silvestris catus" as appropriate.

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

"Identification mark": Indicate mark on the straw or other packages where semen, oocytes or embryos of the consignment are placed.

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

"Approval or registration number of plant/establishment/centre": Indicate the unique registration number of the establishment of the collection or production of 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.

Certifying Officer/Official veterinarian

Name (in capital letters) Qualification and title

Date of signature Signature

Stamp