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
Frozen ☐ Chilled ☐ 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.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 semen described in Part I:

II.1.1. was collected, processed and stored in a semen collection centre(1) approved and supervised by the competent authority in accordance with Chapter I(I)(1) and Chapter I(II)(1) of Annex D to Directive 92/65/EEC;

II.1.2. comes from donor animals which meet the requirements of Chapter II(II) of Annex D to Directive 92/65/EEC;

II.1.3. was collected, processed, stored and transported under conditions which comply with the requirements of Chapters II(II) and III(I) of Annex D to Directive 92/65/EEC;

(2) ○ either [II.1.4. was collected from animals which have been kept continuously since birth on a holding or holdings recognised as having a negligible or controlled risk of classical scrapie according to point 1 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001;]

(2) ○ or [II.1.4. was collected from animals which have been kept continuously for the last three years before the collection on a holding or holdings which has/have complied for the last three years before the collection with the requirements laid down in points 1.3. (a) to (f) of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001;]

(2) ○ or [II.1.4. was collected from animals which have been kept continuously since birth in a Member State or zone of a Member State with a negligible risk status for classical scrapie approved in accordance with point 2.2 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001.;]

(2) ○ or [II.1.4. was collected from ovine animals of the ARR/ARR prion protein genotype;]

II.1.5. was sent to the place of loading in a sealed container in accordance with point 1.4. of Chapter III(I) of Annex D to Directive 92/65/EEC and bearing the number detailed in Box I.23.

(2) ○ either [II.2. No antibiotics or no mixture of antibiotics were added to the semen.]

(2) ○ or [II.2. The following antibiotic or combination of antibiotics was added to produce a concentration in the final diluted semen of not less than(3):

.]

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 I.11: Place of dispatch shall correspond to the semen collection centre of origin of the semen.

Box I.12: Place of destination shall correspond to the semen collection centre, germinal product processing establishment, germinal product storage centre or to the establishment of semen destination.

Box I.19: Identification of container and Seal number shall be indicated.

Box I.30: Identification number shall correspond to the official identification of the animal.

Date of collection shall be indicated in the following format: dd/mm/yyyy.

Approval number of the centre shall correspond to the approval number of the semen centre indicated in Box I.11. where the semen was collected.

Part II:

(1) Only semen collection centres approved by the competent authority and listed in accordance with Article 11(4) of Directive 92/65/EEC.

(2) Delete as appropriate.

(3) Insert names and concentrations.

Certifying Officer/Official veterinarian

Name (in capital letters) Authority name

Date of signature Signature

Stamp