**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 the ova/embryos(1) described in Part I:

II.1. were collected, processed and stored under conditions which meet the requirements of Directive 92/65/EEC;

II.2. come from female donors of the ovine/caprine species(1) which meet the requirements of Chapter IV of Annex D to Directive 92/65/EEC;

II.3. are embryos of the ovine or caprine species which comply with the following conditions as regards classical scrapie:

(1) ○ either [II.3.1. they meet the requirements of Chapter III of Annex D to Directive 92/65/EEC and of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001, as applicable on 31 August 2010;]

(1) ○ or [II.3.1. they meet the requirements of Chapter III of Annex D to Directive 92/65/EEC and of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001, as applicable on 31 August 2010, and are destined for a Member State which benefits, for all or part of its territory, from the provisions laid down in point (b) or (c) of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001, as applicable on 31 August 2010 and the donor animals comply regarding scrapie with the guarantees provided for by the programmes referred to in that point and with the guarantees(2) requested by the Member State of destination;]

(1) ○ either [II.3.2. the semen used for fertilisation meets the requirements of Directive 92/65/EEC and the requirements of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001, as applicable on 31 August 2010.]

(1) ○ or [II.3.2. the semen used for fertilisation meets the requirements of Directive 92/65/EEC and the requirements of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001, as applicable on 31 August 2010, and is destined for a Member State which benefits, for all or part of its territory, from the provisions laid down in point (b) or (c) of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001, as applicable on 31 August 2010, and the donor animals comply regarding scrapie with the guarantees provided for by the programmes referred to in that point and with the guarantees(2) requested by the Member State of destination.]

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 embryo collection team of ova/embryos collection.

Box I.12: Place of destination shall correspond to the embryo collection team, embryo production team, germinal product processing establishment, germinal product storage centre or to the establishment of ova/embryos destination.

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

Box I.30: ”Type”: specify if: in vivo derived embryos, in vivo derived oocytes, in vitro produced embryos or micromanipulated embryos.

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 team shall correspond to the embryo collection team of ova/embryos collection indicated in Box I.11.

Part II:

(1) Delete as appropriate.

(2) Additional guarantees as laid down in Article 2 of Regulation (EC) No 546/2006 [OJ L 94, 1.4.2006, p. 28].

Certifying Officer/Official veterinarian

Name (in capital letters) Authority name

Date of signature Signature

Stamp