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. Ova/embryos(1) described in Part I were collected by a collection team(2) approved by the competent authority and processed in an appropriate laboratory;

II.2. Ova/embryos(1) were collected from donor mares which:

II.2.1. on the day of collection have been located in premises situated on the territory or in the case of regionalisation in a part of the territory of a Member State which is not considered to be infected with African horse sickness in accordance with Article 5(2)(a) and (b) of Directive 2009/156/EC(3),

II.2.2. have been located in holdings under veterinary supervision which on the day of collection fulfilled the conditions of Article 4 of Directive 2009/156/EC,

II.2.3. have been kept prior to the collection in holdings free from clinical signs of contagious equine metritis for 60 days,

II.2.4. have not been used for natural breeding during the period of 30 days prior to the collection of ova/embryos(1),

II.2.5. to the best of my knowledge and as far as I could ascertain, have not been in contact with equidae suffering from an infectious or contagious disease during the 15 days immediately preceding the collection of ova/embryos(1),

II.2.6. have on the day of collection not shown clinical signs of an infectious or contagious disease;

II.3. Ova/embryos(1) were collected, processed, stored and transported under conditions which comply with the requirements of Annex D of Directive 92/65/EEC;

II.4. The semen used for the artificial insemination of the donor mares complies with the requirements of Directive 92/65/EEC(4)(1);

II.5. The ova used for the in vitro production of embryos comply with the requirements of Directive 92/65/EEC(1).

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 or to the holding 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.

Donor identity 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.

Part II:

(1) Delete as appropriate.

(2) Only embryo collection teams approved by the competent authority and listed in accordance with Article 11(4) of Council Directive 92/65/EEC.

(3) OJ L 192, 23.7.2010, p. 1.

(4) Does not apply to ova.

Certifying Officer/Official veterinarian

Name (in capital letters) Authority name

Date of signature Signature

Stamp