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:

(1) ○ either [II.1. the in vivo derived embryos/in vivo derived ova(1) described in Part I were collected, processed and stored by an embryo collection team(2) approved and supervised in accordance with Chapter I(III)(1) of Annex D to Directive 92/65/EEC;]

(1) ○ or [II.1. the in vitro produced embryos/micromanipulated embryos(1) described in Part I were produced, processed and stored by an embryo production team(2), approved and supervised in accordance with Chapter I(III) (1) and (2) of Annex D to Directive 92/65/EEC;]

(1) ○ either [II.2. the in vivo derived embryos described in Part I meet the requirements of Chapter III(II)(1) of Annex D to Directive 92/65/EEC;]

(1) ○ or [II.2. the in vivo derived ova described in Part I meet the requirements of Chapter III(II)(2) of Annex D to Directive 92/65/EEC;]

(1) ○ or [II.2. the in vitro produced embryos described in Part I meet the requirements of Chapter III(II)(3) of Annex D to Directive 92/65/EEC;]

(1) ○ or [II.2. the micromanipulated embryos described in Part I meet the requirements of Chapter III(II)(4) of Annex D to Directive 92/65/EEC;]

II.3. the ova or embryos described in Part I come from donor mares which:

II.3.1. coming from holdings fulfilling the conditions laid down in Article 4(5) of Directive 2009/156/EC(4) onto which only equidae satisfying the conditions laid down in Articles 4 and 5 or Articles 12 to 16 of Directive 2009/156/EC have been admitted;

II.3.2. meet the additional requirements of Chapter IV(4) of Annex D to Directive 92/65/EEC;

II.3.3. have not been used for natural breeding during at least 30 days prior to the date of collection of ova or embryos and between the date of the first sample referred to in points II.3.4. and II.3.5. and the date of the collection of ova and embryos;

II.3.4. have been subjected with negative result to an agar-gel immuno-diffusion test (Coggins test) or an ELISA for equine infectious anaemia carried out on a blood samples taken on _____ (3), being during the past 30 days prior to the date of the first collection of ova or embryos and the last test was carried out on a sample of blood taken on _____ (3); being not more than 90 days before the ova and embryos were collected;

II.3.5. have been subjected to an agent identification test for contagious equine metritis by isolation of Taylorella equigenitalis after a cultivation of 7 to 14 days carried out with negative results in each case on samples taken during the past 30 days prior to the date of the first collection of ova or embryos from mucosal surfaces of the clitoral fossa and clitoral sinuses on two consecutives oestrus periods on _____ (3) and on _____ (3), and on an additional culture specimen taken during one of the oestrus periods from the endometrial cervix on _____ (3);

(1) ○ either [II.4. the embryos described in Part I were conceived as a result of artificial insemination of the donor mares with semen which was collected, processed, stored and transported under conditions which comply with the requirements of Chapters I(I), II(I) and III(I) of Annex D to Directive 92/65/EEC;]

(1) ○ or [II.4. the embryos described in Part I were conceived as a result of in vitro fertilisation of ova complying with the conditions in point 2 of Chapter III(II) of Annex D to Directive 92/65/EEC with semen which was collected, processed, stored and transported under conditions which comply with the requirements of Chapters I(I), II(I) and III(I) of Annex D to Directive 92/65/EEC;]

(1) ○ or [II.4. the ova have not been in contact with semen of the equine species;]

II.5. the ova or embryos described in Part I were sent to the place of loading in a sealed container in accordance with point 6 of Chapter III(II) of Annex D to Directive 92/65/EEC and bearing the number detailed in Box I.19.

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 I.11: Place of dispatch shall correspond to the embryo collection team or embryo production team of ova/embryos collection/production.

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 or embryo production team of ova/embryos collection/production.

Part II:

(1) Delete as appropriate.

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

(3) Insert date.

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

Certifying Officer/Official veterinarian

Name (in capital letters) Qualification and title

Date of signature Signature

Stamp