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.8. Region of origin Code

I.9. Country of destination ISO 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
Document Type
Accompanying document reference
Date of Issue
Country
Place of issue

I.18. Transport conditions
Chilled ☐ Ambient ☐ 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.24. Estimated journey time

I.25. Journey Log

I.26. Total number of packages	I.27. Total quantity	I.28. Total gross weight

I.30. Description of consignment

1. 05 PRODUCTS OF ANIMAL ORIGIN, NOT ELSEWHERE SPECIFIED OR INCLUDED

0511 Animal products not elsewhere specified or included; dead animals of Chapter 1 or 3, unfit for human consumption

051199 Other

05119985 Other

Part I: Description of consignment

#1.	Commodity	Identification Number	Quantity	Nature of commodity	Identification Mark
Species		Package count	Date of collection / production	Plant / Establishment / Centre	Type

Part II: Certification

II. Health information

I, the undersigned official veterinarian, hereby certify that:

(1) ○ either [II.1. the ☐ [in vivo derived embryos] (1) ☐ [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] (1) ☐ [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] (1) ☐ [embryos] (1) described in Part I come from donor mares which:

II.3.1. come from the 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 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 sample 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] (1) ☐ [embryos] (1) described in Part I:

II.5.1. 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;

II.5.2. are dispatched:

(1) ○ either [by an ☐ [embryo collection team] (1) ☐ [embryo production team] (1) or from a zone not subject to movement restrictions affecting equine animals and established for reasons of listed diseases relevant for those species or diseases subject to emergency measures relevant for those species, or those

Part II: Certification

II. Health information		

restrictions do not apply to these ova or embryos because they were collected before the restrictions were established, and they have not been in contact with other ova or embryos of a lower health status for an adequate period.]

(1) ○ or [by an ☐ [embryo collection team] (1) ☐ [embryo production team] (1) or from a zone subject to movement restrictions affecting equine animals and established for (5), but derogations from movement restrictions have been granted, and:

(1) ☐ [they comply with the requirements set out in (6);]]

(1) ☐ [and in particular, they are (7).]]

Notes

In accordance with the Agreement on the withdrawal of the United Kingdom of Great Britain and Northern Ireland from the European Union and the European Atomic Energy Community, and in particular Article 5(4) of the Protocol on Ireland/Northern Ireland in conjunction with Annex 2 to that Protocol, references to the Union in this animal health certificate include the United Kingdom in respect of Northern Ireland.

This animal health certificate shall be completed in accordance with 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 shall correspond to the embryo collection team or embryo production team of ova/embryos collection/production.

Box reference I.12: Place of destination shall correspond to the embryo collection team, embryo production team or to the holding of ova/embryos destination.

Box reference I.26: Total number of packages shall correspond to the number of containers.

Box reference I.19: Seal number shall be indicated.

Box reference 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 if not applicable.

(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.

(5) Insert the name of the disease(s).

(6) Insert the specific reference to the article(s), title, and number of the relevant legal act(s) adopted by the Commission providing for those requirements.

(7) Insert the specific attestation(s) provided for in and required by the relevant legal act(s) adopted by the Commission, as referred to in Article 159(2), points (a), (b) and (c), of Regulation (EU) 2016/429 of the European Parliament and of the Council.

Certifying Officer/Official veterinarian	
Name (in capital letters)	Qualification and title
Date of declaration	Signature
Stamp	