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:

II.1. The semen collection centre (1), in which the semen described in Part I was collected, processed and stored for trade:

II.1.1. was approved and supervised by the competent authority according to the conditions of Chapter I of Annex D to Directive 92/65/EEC;

II.1.2. was situated on the territory or in the case of regionalisation in a part of the territory (2) of a Member State which was on the day semen was collected until the date the semen was dispatched as ☐ [fresh] (2) ☐ [chilled] (2) semen or until the 30 days mandatory storage period for frozen semen elapsed (2) 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.1.3. fulfilled during the period commencing 30 days prior to the date of semen collection until the date the semen was dispatched as ☐ [fresh] (2) ☐ [chilled] (2) semen or until the 30 days mandatory storage period for frozen semen elapsed(2), the conditions of Article 4 of Directive 2009/156/EC;

II.1.4. contained during the period commencing 30 days prior to the date of semen collection until the date the semen was dispatched as ☐ [fresh] (2) ☐ [chilled] (2) semen or until the 30 days mandatory storage period for frozen semen elapsed (2) only equidae which were free of clinical signs of equine viral arteritis and contagious equine metritis;

II.2. All equidae have been admitted onto the centre under the provisions of Article 4 and 5 of Directive 2009/156/EC (3);

II.3. The semen described in Part I was collected from donor stallions, which:

II.3.1. on the day the semen was collected have not shown clinical signs of an infectious or contagious disease,

II.3.2. during at least 30 days prior to collection of the semen have not been used for natural service,

II.3.3. during the 30 day period prior to collection of the semen have been kept on holdings where no equidae showed clinical signs of equine viral arteritis,

II.3.4. during the 60 days period prior to collection of the semen have been kept on holdings where no equidae showed clinical signs of contagious equine metritis,

II.3.5. 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 15 days immediately preceding collection of the semen,

II.3.6. have undergone the following animal health tests, carried out in a laboratory recognised by the competent authority, in accordance with a test programme as specified in point II.3.7.

II.3.6.1. an agar-gel immuno-diffusion test (Coggins test) for equine infectious anaemia with negative result;

and (2) ○ either [II.3.6.2. a serum neutralisation test for equine viral arteritis with negative result at a serum dilution of one in four; and]

(2) ○ or [II.3.6.2. a virus isolation test for equine viral arteritis carried out with negative result on an aliquot of the entire semen of the donor stallion;]

and II.3.6.3. an agent identification test for contagious equine metritis carried out on two occasions on samples collected from the donor stallion with an interval of 7 days by isolation of Taylorella equigenitalis from pre-ejaculatory fluid or a semen sample and from genital swabs taken at least from the penile sheath, urethra and urethral fossa with negative result in each case;

II.3.7. have been subject to the one of the following test programmes (4):

II.3.7.1. The donor stallion was continuously resident on the collection centre for at least 30 days prior to the semen collection, and during the collection period, and no equidae on the collection centre came during that time into direct contact with equidae of lower health status than the donor stallions.

The tests described in point II.3.6 have been carried out on samples taken on (5) and in the case of contagious equine metritis on a second sample taken on (5), being at least 14 days after the commencement of the above residence period and at least at the beginning of the breeding season;

II.3.7.2. The donor stallion was not continuously resident on the collection centre or other equidae on the collection centre came into contact with equidae of lower health status than the donor stallion.

Part II: Certification

II. Health information		

The tests described in point II.3.6 have been carried out on samples taken on (5) and in the case of contagious equine metritis on a second sample taken on (5), being within the 14 days period before the first semen collection and at least at the beginning of the breeding season,

and the test described in point II.3.6.1 for equine infectious anaemia was last carried out on a sample of blood taken on (5), being not more than 120 days before the semen described in Part I was collected;

and (2) ○ either [one of the tests described in point II.3.6.2 for equine viral arteritis was last carried out on a sample collected on (5), being not more than 30 days before the semen described in Part I was collected,]

(2) ○ or [the non-shedder state of the seropositive stallion for equine viral arteritis was confirmed by a virus isolation test which was carried out on an aliquot of the entire semen of the donor stallion collected on (5), being not more than one year before the semen described in Part I was collected;]

II.3.7.3. The tests described in point II.3.6 have been carried out during the 30 days mandatory storage period of frozen semen and not less than 14 days after the collection of the semen on samples taken on (5) and in the case of contagious equine metritis on a second sample taken on (5);

II.4. The semen described in Part I:

II. 4.1. was collected, processed, stored and transported under conditions which comply with the requirements of Chapters II and III of Annex D to Directive 92/65/EEC;

II.4.2. is dispatched from:

(2) ○ either [a semen collection centre or 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 restrictions do not apply to this semen because it was collected before the restrictions were established, and the semen has not been in contact with other semen of a lower health status for an adequate period.]

(2) ○ or [a semen collection centre or a zone subject to movement restrictions affecting equine animals and established for (6), but derogations from movement restrictions have been granted, and:

(2) ☐ [it complies with the requirements set out in (7);]]

(2) ☐ [and, in particular, it is (8).]]

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 semen collection centre of origin of the semen.

Box reference I.12: Place of destination shall correspond to the semen collection or storage centre or to the holding of semen destination.

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

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

Box reference Donor identity shall correspond to the official identification of the animal.

Part II: Certification

II. Health information		

I.30:

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 if not applicable.

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

(4) Cross out the programme(s) that do(es) not apply to the consignment.

(5) Insert date.

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

(7) 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.

(8) 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	