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. 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/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/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/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 seven 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.

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 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.

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 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.

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 semen collection centre of origin of the semen.

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

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

Box I.30: 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 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 Council Directive 92/65/EEC.

(2) Delete as appropriate.

(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.

Certifying Officer/Official veterinarian

Part II: Certification

II. Health information		
Name (in capital letters) Date of signature Stamp	Authority name Signature	