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~~

I.19. Container No / Seal No

I.20. Certified as
Registered equidae ☐

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.27. Total quantity

I.28. Total gross weight

I.30. Description of consignment

Commodity	Species	Subcategory	Sex	Identification system

Identification Number	Age	Quantity

Part II: Certification

II. Health information

I, the undersigned official veterinarian, hereby certify that:

II.1. The equine animal described in Part I meets the following requirements:

II.1.1. The animal is accompanied by its single lifetime identification document as provided for in Article 65, 67 or 68 of Commission Delegated Regulation (EU) 2019/2035, or by a temporary document issued in accordance with Article 61(2) thereof.

(1) ☐ [The single lifetime identification document was issued in accordance with Article 65(2) or 67(1) of Delegated Regulation (EU) 2019/2035, or the temporary document was issued in accordance with Article 61(2) of that Regulation, for a registered equine animal as defined in Article 2(30) of that Delegated Regulation.]

(1) ☐ [The single lifetime identification document includes a valid validation mark in accordance with Article 65(1)(i)(i) of Delegated Regulation (EU) 2019/2035.]

(1) ☐ [The single lifetime identification document includes a valid license in accordance with Article 65(1)(i)(ii) of Delegated Regulation (EU) 2019/2035.]

II.1.2. The animal has not shown signs or symptoms of diseases listed for equine animals during the clinical examination, which was carried out within the 48 hour period prior to its departure, or on the last working day prior to its departure(2), from the registered establishment, on (insert date dd/mm/yyyy).

II.2. According to official information, the animal described in Part I meets the following health requirements:

II.2.1. The animal does not come from an establishment subject to movement restrictions or situated in a restricted zone established for reasons of diseases listed for equine animals, including African horse sickness and infection with Burkholderia mallei (glanders).

II.2.2. The animal comes from an establishment in which surra (Trypanosoma evansi) has not been reported during the 30 day period prior to its departure, and

(1) either ○ [surra has not been reported in the establishment during the 2 year period prior to departure.]

(1) or ○ [surra has been reported in the establishment during the 2 year period prior to its departure and following the last outbreak the establishment has remained under movement restrictions

(1) either ○ [until the remaining animals in the establishment have been subjected to a test for surra with one of the diagnostic methods provided for in Part 3 of Annex I to Commission Delegated Regulation (EU) 2020/688, carried out, with negative results, on samples taken at least 6 months after the last infected animal has been removed from the establishment.]]

(1) or ○ [for at least 30 days from the date of cleaning and disinfection of the establishment, after the last animal of listed species on the establishment was either killed and destroyed or slaughtered.]]

II.2.3. The animal comes from an establishment in which dourine has not been reported during the last 6 months prior to its departure, and

(1) either ○ [dourine has not been reported in the establishment during the 2 year period prior to its departure.]

(1) or ○ [dourine has been reported in the establishment during the 2 year period prior to its departure and following the last outbreak, the establishment has remained under movement restrictions

(1) either ○ [until the remaining equine animals in the establishment, except castrated male equine animals, have been subjected to a test for dourine with the diagnostic method provided for in Part 8 of Annex I to Delegated Regulation (EU) 2020/688, carried out, with negative results, on samples taken at least 6 months after the infected animals have been killed and destroyed or slaughtered, or the infected entire male equine animals have been castrated.]]

(1) or ○ [for at least 30 days from the date of cleaning and disinfection after the last equine animal on the establishment was either killed and destroyed or slaughtered.]]

Part II: Certification

II. Health information

II.2.4. The animal comes from an establishment in which equine infectious anaemia has not been reported during the 90 day period prior to its departure, and

(1) either ○ [equine infectious anaemia has not been reported on the establishment during the 12 month period prior to its departure.]

(1) or ○ [equine infectious anaemia has been reported on the establishment during the 12 month period prior to its departure and following the last outbreak the establishment has remained under movement restrictions

(1) either ○ [until the remaining equine animals in the establishment have been subjected to a test for equine infectious anaemia with the diagnostic method provided for in Part 9 of Annex I to Delegated Regulation (EU) 2020/688, carried out, with negative results, on samples taken on two occasions with a minimum interval of 90 days following cleaning and disinfection of the establishment after the infected animals have been killed and destroyed or slaughtered.]]

(1) or ○ [for at least 30 days from the date of cleaning and disinfection of the establishment, after the last equine animal on the establishment was either killed and destroyed or slaughtered.]]

II.2.5. The animal comes from an establishment in which Venezuelan equine encephalomyelitis has not been reported during the 6 month period prior to its departure, and

(1) either ○ [during the 2 year period prior to its departure, Venezuelan equine encephalomyelitis has not been reported in the Member State or zone thereof in which the establishment is situated.]

(1) or ○ [during the 2 year period prior to its departure, Venezuelan equine encephalomyelitis has been reported in the Member State or zone thereof in which the establishment is situated, and during the 21 day period prior to departure of the animal referred to in point II.1 all equine animals in the establishment have remained clinically healthy, and

(1) either ○ [the animal referred to in point II.1 was kept protected from attacks by insect vectors in a quarantine station, in which any equine animal that showed a rise in daily taken body temperature has been subjected with negative result to a diagnostic test for Venezuelan equine encephalomyelitis with the diagnostic method provided for in Part 10(1)(a) of Annex I to Delegated Regulation (EU) 2020/688, and the animal referred to in point II.1 has been

(1) either ○ [vaccinated against Venezuelan equine encephalomyelitis with a complete primary course and revaccinated according to manufacturer's recommendations not less than 60 days and not more than 12 months prior to the date of its departure.]]]

(1) or ○ [subjected to a serological test for Venezuelan equine encephalomyelitis with the diagnostic method provided for in Part 10(1)(b) of Annex I to Delegated Regulation (EU) 2020/688, carried out, with negative results, on a sample taken not less than 14 days after the date of its entry into the quarantine station.]]]

(1) or ○ [the body temperature of the animal referred to in point II.1 has been taken daily, either without a rise or the animal has been subjected to a diagnostic test for Venezuelan equine encephalomyelitis with the diagnostic method provided for in Part 10(1)(a) of Annex I to Delegated Regulation (EU) 2020/688, with negative results, and the animal referred to in point II.1 has been subjected to tests for Venezuelan equine encephalomyelitis with the diagnostic methods provided for in:

- Part 10(1)(b) of Annex I to Delegated Regulation (EU) 2020/688, without an increase in antibody titre, carried out on paired samples taken on two occasions with an interval of 21 days, the second of which was taken during the 10 day period prior to the date of its departure, and

Part II: Certification

II. Health information

- Part 10(2) of Annex I to Delegated Regulation (EU) 2020/688, with negative result, carried out on a sample taken within the 48 hour period prior to its departure, and the animal has been protected from attacks by insect vectors after sampling until its departure.]]

II.2.6. The animal comes from an establishment in which infection with rabies virus in kept terrestrial animals has not been reported during the 30 day period prior to its departure.

II.2.7. The animal comes from an establishment in which anthrax in ungulates has not been reported during the 15 day period prior to its departure.

II.3. To the best of my knowledge, after due inquiry, and as declared by the operator, the animal comes from an establishment where there were no abnormal mortalities with an undetermined cause and the animal has not been in contact with kept animals of listed species which did not comply with the requirements referred to in points II.2.1. to II.2.6. during the 30 day period prior to its departure, and with the requirement referred to in point II.2.7. during the 15 day period prior to its departure.

(1) ☐ [II.4. According to official information and as declared by the operator, it is a semen donor animal subjected to the testing programme as referred to in point 1(b)(i) of Chapter I of Part 4 of Annex II to Commission Delegated Regulation (EU) 2020/686, and

II.4.1. it comes from a semen collection centre and will be transported directly to another semen collection centre in accordance with Article 19 of Delegated Regulation (EU) 2020/686; and

II.4.2. since the date of its admission, it was continuously resident at the semen collection centre and was subjected, with negative results, to all compulsory routine tests referred to in point 1(a) of Chapter I of Part 4 of Annex II to Delegated Regulation (EU) 2020/686 during the 12 month period prior to the date of its departure; and

II.4.3. the prior consent of the centre veterinarian of the semen collection centre of destination has been obtained by the operator; and

II.4.4. the means of transport used have been cleansed and disinfected before use.]

II.5. Arrangements are made to

(1) either ○ [transport the animal in accordance with Article 4 of Delegated Regulation (EU) 2020/688.]

(1) or ○ [move the animal on foot.]

II.6. This animal health certificate is valid for

(1) either ○ [10 days from the date of issuing, and]

(1) or ○ [30 days from the date of issuing, and a valid validation mark or license is attested in point II.1.1, and]

in the case of transport by waterway/sea of the animal, the period of validity of the animal health certificate may be extended by the duration of the journey by waterway/sea.

Animal welfare attestation

At the time of inspection, the animal covered by this health certificate was fit to be transported in accordance with the provisions of Council Regulation (EC) No 1/2005 on the intended journey due to start on (insert date).

Part II: Certification

II. Health information

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 European Union in this certificate include the United Kingdom in respect of Northern Ireland.

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 reference I.11: "Place of dispatch": Indicate a registered establishment of dispatch of the equine animal or, provided the animal is transported, an establishment approved for assembly operations in accordance with Articles 97 and 99 of Regulation (EU) 2016/429.

Box reference I.12: "Place of destination": Indicate a registered establishment of destination or, provided the animal is transported, an establishment approved for assembly operations in accordance with Articles 97 and 99 of Regulation (EU) 2016/429.

Box reference I.30: "Identification number": Indicate the unique code of the equine animal referred to in Article 65(1)(b) of Delegated Regulation (EU) 2019/2035, or the code displayed by the means of identification defined in point (a), (c) or (e) of Annex III to Delegated Regulation (EU) 2019/2035, if the animal is unweaned and accompanies its dam or foster mare.

Part II:

(1) Delete if not applicable.

(2) Option only available in the case of either:

(a) an equine animals accompanied by its single lifetime identification document as provided for in Article 114(1), point (c), of Regulation (EU) 2016/429 which includes a valid validation mark referred to in Article 92(2), point (a), of Delegated Regulation (EU) 2020/688; or

(b) a registered equine animal accompanied by its single lifetime identification document as provided for in Article 114(1), point (c), of Regulation (EU) 2016/429 which includes a valid license referred to in Article 92(2), point (b), of Delegated Regulation (EU) 2020/688, or by its single lifetime identification document accompanied by the FEI Recognition Card together with the validation sticker.

Certifying Officer/Official veterinarian

Name (in capital letters)
Authority name
Date of signature
Signature
Stamp