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
Ambient ☐ Chilled ☐ Frozen ☐

I.19. Container Number / Seal Number

I.20. Certified as
Further keeping ☐ Exhibition ☐ Confined establishment ☐ Registered equine animal ☐
Event or activity near borders ☐

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.27. Total quantity

I.28. Total gross weight

I.30. Description of consignment

1. 01 LIVE ANIMALS

0101 Live horses, asses, mules and hinnies
Registered equine animals

Part I: Description of consignment

#1.	Commodity	Subcategory	Sex	Identification system
Species		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 Article 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), point (i)(i), of Delegated Regulation (EU) 2019/2035.]

(1) ☐ [The single lifetime identification document includes a valid license in accordance with Article 65(1), point (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 last 48 hours prior to the time of its departure, or on the last working day prior to the date of its departure, 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:

(1) ○ either [II.2.1. The animal comes from an establishment or a zone not subject to movement restrictions affecting equine animals and established for reasons of diseases listed for those species or diseases subject to emergency measures relevant for those species, and the animal has not been in contact with kept animals of a listed species of a lower health status for an adequate period.]

(1) or ○ [II.2.1. The animal comes from an establishment or zone subject to movement restrictions affecting equine animals and established for (2), but derogations from movement restrictions have been granted, and:

(1) ☐ [it complies with the requirements set out in (3);]]

(1) ☐ [and in particular, it is (4).]]

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

(1) either ○ [surra has not been reported in the establishment during the last 2 years prior to the date of its departure.]

(1) or ○ [surra has been reported in the establishment during the last 2 years prior to the date of its departure and following the date of the last outbreak, the affected establishment has remained under movement restrictions:

(1) either ○ [until the date on which 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 following the date on which the last infected animal has been removed from the establishment.]]

(1) or ○ [for at least 30 days following the date on which the last animal of listed species in the establishment was either killed and destroyed, or slaughtered, and the premises were cleaned and disinfected.]]

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

(1) either ○ [dourine has not been reported in the establishment during the last 2 years prior to the date of its departure.]

(1) or ○ [dourine has been reported in the establishment during the last 2 years prior to the date of its departure and following the date of the last outbreak the affected establishment has remained under movement restrictions:

Part II: Certification

II. Health information

(1) either ○ [until the date on which 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 following the date on which 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 after the date on which the last equine animal in the establishment was either killed and destroyed, or slaughtered, and the premises were cleaned and disinfected.]]

II.2.4. The animal comes from an establishment in which equine infectious anaemia has not been reported during the last 90 days prior to the date of its departure, and:

(1) either ○ [equine infectious anaemia has not been reported in the establishment during the last 12 months prior to the date of its departure.]

(1) or ○ [equine infectious anaemia has been reported in the establishment during the last 12 months prior to the date of its departure and following the date of the last outbreak the affected establishment has remained under movement restrictions:

(1) either ○ [until the date on which 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 the date on which the infected animals have been killed and destroyed, or slaughtered, and the establishment was cleaned and disinfected.]]

(1) or ○ [for at least 30 days following the date on which the last equine animal in the establishment was either killed and destroyed, or slaughtered, and the premises were cleaned and disinfected.]]

II.2.5. The animal comes from an establishment in which Venezuelan equine encephalomyelitis has not been reported during the last 6 months prior to the date of its departure, and:

(1) either ○ [during the last 2 years prior to the date of 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 last 2 years prior to the date of its departure, Venezuelan equine encephalomyelitis has been reported in the Member State or zone thereof in which the establishment is situated, and during the last 21 days prior to the date of 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 in quarantine protected from attacks by insect vectors, and 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, point 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, point 1(b), of Annex I to Delegated Regulation (EU) 2020/688, carried out, with negative

Part II: Certification

II. Health information

results, on a sample taken not less than 14 days after the date of its entry into quarantine.]]]

(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, point 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, point 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 last 10 days prior to the date of its departure, and
- Part 10, point 2, of Annex I to Delegated Regulation (EU) 2020/688, with negative result, carried out on a sample taken within the last 48 hours prior to the time of its departure, and the animal has been protected from attacks by insect vectors after sampling until the date of 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 last 30 days prior to the date of its departure.

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

II.2.8. The animal comes from an establishment in which, to the best of my knowledge, after due inquiry, and as declared by the operator, 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 last 30 days prior to the date of its departure, and with the requirement referred to in point II.2.7 during the last 15 days prior to the date of its departure.

II.3. 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.4. 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 a valid 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 animal 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).

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

Part II: Certification

II. Health information		

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 of the European Parliament and of the Council.
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, or a veterinary clinic.
Box reference I.22:	“For transit through Member State(s)”: Indicate each Member State of transit. In the case of an animal health certificate that is valid for 30 days, indicate all Member States that the equine animal has transited while returning to the establishment of its departure.
Box reference I.30:	“Identification number”: Indicate the unique code of the equine animal referred to in Article 65(1), point (b), of Delegated Regulation (EU) 2019/2035.

Part II:

(1)	Delete if not applicable.
(2)	Insert the name of the disease(s).
(3)	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.
(4)	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 126(1), points (b)(ii) and (iii), of Regulation (EU) 2016/429.

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