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
Exhibition ☐ Slaughter ☐ Further keeping ☐ Event or activity near borders ☐
Confined establishment ☐

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
Non-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 animals (1) of the consignment described in Part I meet the following requirements:

(2) either ☐ [II.1.1. They are accompanied by their single lifetime identification documents as provided for in:

(2) either ○ [Article 65, 67 or 68 of Commission Delegated Regulation (EU) 2019/2035, and are not intended for slaughter for human consumption.]

(2) or ○ [Article 65 or Article 67(1) of Delegated Regulation (EU) 2019/2035, and are intended for slaughter for human consumption.]

(2) ☐ [Their single lifetime identification documents were issued in accordance with Article 65(2) or Article 67(1) of Delegated Regulation (EU) 2019/2035 for registered equine animals as defined in Article 2(30) of that Delegated Regulation.]

(2) ☐ [Their single lifetime identification documents include a valid validation mark in accordance with Article 65(1), point (i)(i), of Delegated Regulation (EU) 2019/2035.]

(2) ☐ [Their single lifetime identification documents include a valid license in accordance with Article 65(1), point (i)(ii) of Delegated Regulation (EU) 2019/2035.]]

(2) and/or ☐ [II.1.2. They are not accompanied by single lifetime identification documents and they are unweaned and accompany their dam or foster mare as provided for in Article 58(2), point (a), and Article 66(2), point (c), of Delegated Regulation (EU) 2019/2035.]

II.1.3. They have 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 departure of the consignment, or on the last working day prior to departure of the consignment, from the registered establishment, on (insert date dd/mm/yyyy).

(2) ☐ [II.1.4. They are intended to be slaughtered for disease eradication purposes as part of an eradication programme, as provided for in Article 31(1) or (2) of Regulation (EU) 2016/429 of the European Parliament and of the Council, and the Member State of destination and, where applicable, the Member State of passage authorised the movement in advance.]

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

(2) either ○ [II.2.1. They come from establishments or zones 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 animals have not been in contact with kept animals of a listed species of a lower health status for an adequate period.]

(2) or ○ [II.2.1. They come from establishments or zones subject to movement restrictions affecting equine animals and established for (3), but derogations from movement restrictions have been granted, and:

(2) ☐ [they comply with the requirements set out in (4);]]

(2) ☐ [and in particular, they are (5).]]

II.2.2. They come from establishments in which surra (Trypanosoma evansi) has not been reported during the last 30 days prior to the date of departure of the consignment, and:

(2) either ○ [surra has not been reported in the establishments during the last 2 years prior to the date of departure of the consignment;]

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

(2) 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

Part II: Certification

II. Health information

results, on samples taken at least 6 months following the date on which the last infected animal has been removed from the establishment.]]

(2) 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. They come from establishments in which dourine has not been reported during the last 6 months prior to the date of departure of the consignment, and:

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

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

(2) 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 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 date on which the infected entire male equine animals have been castrated.]]

(2) 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.2.4. They come from establishments in which equine infectious anaemia has not been reported during the last 90 days prior to the date of departure of the consignment, and:

(2) either ○ [equine infectious anaemia has not been reported in the establishments during the last 12 months prior to the date of departure of the consignment;]

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

(2) either ○ [until the 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.]]

(2) 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.2.5. They come from establishments in which Venezuelan equine encephalomyelitis has not been reported during the last 6 months prior to the date of departure of the consignment, and:

(2) either ○ [during the last 2 years prior to the date of departure of the consignment, Venezuelan equine encephalomyelitis has not been reported in the Member State or zone thereof in which the establishments are situated.]

(2) or ○ [during the last 2 years prior to the date of departure of the consignment, Venezuelan equine encephalomyelitis has been reported in the Member State or zone thereof in which the establishments are situated, and during the last 21 days prior to the date of departure of the consignment all equine animals in the

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establishments have remained clinically healthy, and:

(2) either ○ [the animals of the consignment were kept in quarantine protected from attacks by insect vectors, and any equine animals 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 animals of the consignment have been:

(2) 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 the departure of the consignment;]]]

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

(2) or ○ [the body temperature of the animals of the consignment has been taken daily, either without a rise or the animals have 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 animals of the consignment have 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 departure of the consignment, 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 departure of the consignment, and the animals have been protected from attacks by insect vectors after the date of sampling until the date of departure of the consignment;]]

II.2.6. They come from establishments in which infection with rabies virus in kept terrestrial animals has not been reported during the last 30 days prior to the date of departure of the consignment;

II.2.7. They come from establishments in which anthrax in ungulates has not been reported during the last 15 days prior to the date of departure of the consignment;

II.2.8. They come from establishments in which, to the best of my knowledge and as declared by the operator, there were no abnormal mortalities with an undetermined cause and they have 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 departure of the consignment, and with the requirement referred to in point II.2.7 during the last 15 days prior to the date of departure of the consignment.

II.3. Arrangements are made to transport the consignment in accordance with Article 4 of Delegated Regulation (EU) 2020/688.

II.4. This animal health certificate is valid for 10 days from the date of issuing. In the case of transport by

Part II: Certification

II. Health information

waterway/sea of animals, the period of validity of the certificate may be extended by the duration of the journey by waterway/sea.

(2)(6) ☐ [II.5. Since the date of departure from their registered establishments of origin and prior to the date of arrival to this establishment approved for assembly operations, none of the animals of the consignment has undergone more than two assembly operations, and:

(2) either ○ [they come from registered establishments of departure.]]

(2) or ○ [at least one of the animals of the consignment has undergone one assembly operation in an approved establishment.]]

(2) or ○ [at least one of the animals of the consignment has undergone two assembly operations in the approved establishments.]]

Animal welfare attestation

At the time of inspection, the animals covered by this animal health certificate were 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) (7) (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": Indicate a registered establishment of dispatch of the equine animals or 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 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.17: "Accompanying documents": In case the animals are dispatched from an establishment approved for assembly operations in the Member State of origin, the reference number(s) of the official document(s), based on which the animal health certificate for this consignment is issued in this establishment approved for assembly operations, may be indicated.

In case the animals are dispatched from an establishment approved for assembly operations in the Member State of passage, the reference number(s) of the animal health certificate(s), based on which the animal health certificate for this consignment is issued in this establishment approved for assembly operations, shall be indicated.

Box reference I.22: "For transit through Member State(s)": Indicate each Member State of transit.

Box reference I.30: "Identification number": Indicate for each animal of the consignment the unique code referred to in Article 65(1), point (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) There may be one or more animals in the consignment.

(2) Delete if not applicable.

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

(4) 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.

(5) Insert the specific attestation(s) provided for in and required by the relevant legal act(s) adopted by the

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Commission, as referred to in Article 126(1), points (b)(ii) and (iii), of Regulation (EU) 2016/429.

(6) Applicable in case the consignment is dispatched from the establishment approved for assembly operations.

(7) In the case where a consignment is grouped in an establishment approved for assembly operations and comprises animals that were loaded on different dates, the date which the journey commenced for the whole consignment is considered to be the earliest date when any part of the consignment left the establishment of origin.

(8) This statement does not exempt transporters from their obligation in accordance with Union rules in force in particular regarding the fitness to be transported.

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