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$ightharpoonup \underline{B}$ COMMISSION IMPLEMENTING REGULATION (EU) 2018/659

of 12 April 2018

on the conditions for the entry into the Union of live equidae and of semen, ova and embryos of equidae

(Text with EEA relevance)

(OJ L 110, 30.4.2018, p. 1)

Corrected by:

►C1 Corrigendum, OJ L 228, 11.9.2018, p. 2 (2018/659)

►<u>C2</u> Corrigendum, OJ L 237, 20.9.2018, p. 13 (2018/659)

COMMISSION IMPLEMENTING REGULATION (EU) 2018/659

of 12 April 2018

on the conditions for the entry into the Union of live equidae and of semen, ova and embryos of equidae

(Text with EEA relevance)

SECTION 1

Subject matter, scope and definitions

Article 1

Subject matter and scope

This Regulation establishes the list of third countries and parts of the territory of third countries from which the entry into the Union of consignments of equidae and of their semen, ova and embryos is authorised.

It also lays down the animal health and veterinary certification requirements applicable to those consignments.

Article 2

Definitions

For the purposes of this Regulation, the following definitions shall apply:

(a) 'regionalisation'

means the official recognition of a part of the territory of a third country with precise geographical delimitations, containing an equidae subpopulation with a distinct health status with respect to one or more specific diseases and subject to appropriate surveillance, disease control and biosecurity measures;

(b) 'identification document'

means any document which may be used to prove the identity of an equine animal and which includes at least the following information:

- (i) a narrative describing the animal and recording its marks depicted in a completed outline diagram;
- (ii) a reference to specific marks, characteristics or identifiers which establish an unambiguous link between the animal and the document;
- (iii) the information set out in points 1, 2, 3 and 6 to 10 of Part A and in points 12 to 18 in Part B of Section 1 of Annex I to Commission Implementing Regulation (EU) 2015/262 (¹);

⁽¹) Commission Implementing Regulation (EU) 2015/262 of 17 February 2015 laying down rules pursuant to Council Directives 90/427/EEC and 2009/156/EC as regards the methods for the identification of equidae (Equine Passport Regulation) (OJ L 59, 3.3.2015, p. 1).

(c) 'registered horse'

means an animal of the species *Equus caballus* registered as defined in Council Directive 90/427/EEC (1), identified by means of an identification document issued by:

- (i) the breeding authority or any other competent authority of the country where the animal originated which manages the studbook or register for that breed of animal; or
- (ii) any international association or organisation which manages horses for competition or racing;
- (d) 'entry'

means the action of moving equidae or their semen, ova or embryos into one of the territories listed in Annex I to Council Directive 97/78/EC (2);

(e) 'type of entry'

means respectively the temporary admission, the re-entry after temporary export, imports and transit;

(f) 'temporary admission'

means the status of a registered horse originating in a third country and moved into the Union territory for a period of less than 90 days;

(g) 'temporary export'

means the movement of a registered horse out of the Union for a period of less than 90 days;

(h) 're-entry'

means the movement of a registered horse from a third country into the Union after temporary export from the Union;

(i) 'imports'

means the movement of a consignment of equidae or their semen, ova or embryos into the Union for an undetermined period;

(j) 'transit'

means the movement of a consignment of equidae across Union territory by road, rail or waterway transport from one third country to another or from one part of the territory of a third country to another part of the territory of the same third country;

(k) 'border inspection post'

means any inspection post as defined in Article 2(2)(f) of Directive 91/496/EEC and Article 2(2)(g) of Directive 97/78/EC and approved for the commodity concerned in accordance with Decision 2009/821/EC;

(l) 'category of equidae'

means respectively registered equidae, equidae for slaughter and equidae for breeding and production as defined in Article 2 of Directive 2009/156/EC, and registered horses;

⁽¹⁾ Council Directive 90/427/EEC of 26 June 1990 on the zootechnical and genealogical conditions governing intra-Community trade in equidae (OJ L 224, 18.8.1990, p. 55).

⁽OJ L 224, 18.8.1990, p. 55).
(2) Council Directive 97/78/EC of 18 December 1997 laying down the principles governing the organisation of veterinary checks on products entering the Community from third countries (OJ L 24, 30.1.1998, p. 9).

(m) 'ova' means the haploid stages of the ootidogenesis including secondary oocytes and ova;

(n) 'operator' means any natural or legal person subject to one or more of the rules provided for in this Regulation who has equidae or their germinal products under its responsibility;

(o) 'isolation' means the separation for a specified period of equidae from other animals to prevent the transmission through direct or indirect contact of specified pathogen(s), while the equidae are undergoing observation and, if appropriate, testing and treatment under the supervision of the veterinary authority;

(p) 'quarantine' means the isolation of equidae on premises operated in accordance with specific biosecurity rules under the control of the veterinary authority;

(q) 'vector-protected means the quarantine of equidae which quarantine'

- (i) is carried out on dedicated premises that are:
 - screened against the intrusion of relevant vectors,
 - included in a system of vector surveillance within the premises and of measures to limit the presence of relevant vectors around the premises;
- (ii) may include exercise of the quarantined animal under official supervision during the vector-low period of the day and subject to application of insecticides and insect repellents and where possible bodycoverage;
- (r) 'vector-proof means the quarantine of equidae in a sealed building which is:
 - furnished with positive pressure ventilation and filtered air inlets,
 - is only accessible through a double door entry-exit system (1),
 - in which a vector surveillance system is operated,
 - where Standard Operating Procedures, including description of back-up and alarm systems, are implemented for the operation of the quarantine and the transport of equidae to the place of loading,
- (s) 'TRACES' means the integrated computerised veterinary system provided for in Decisions 2003/24/EC and 2004/292/EC.

⁽¹⁾ https://ec.europa.eu/food/sites/food/files/animals/docs/ad_control-measures_bt_guidance_vpe_7068_2012.pdf

SECTION 2

List of third countries and parts thereof for the entry into the Union of equidae and semen, ova and embryos of equidae

Article 3

List of third countries and parts of the territory of third countries from which the entry of equidae into the Union is authorised

- 1. Member States shall authorise the entry into the Union of consignments of equidae from the third countries or, where the Union applies regionalisation, parts of the territory of third countries, listed in columns 2 and 4 of the table in Annex I in accordance with the indications set out in that Annex, as follows:
- (a) the temporary admission of registered horses as indicated in column 6 of the table set out in Annex I, and accompanied by an individual health certificate drawn up in accordance with the model health certificate set out in Section A of Part 1 of Annex II;
- (b) the transit of equidae as indicated in column 15 of the table set out in Annex I, and accompanied by an individual health certificate drawn up in accordance with the model health certificate set out in Section B of Part 1 of Annex II;
- (c) the re-entry of registered horses for racing, competition and cultural events after temporary export as indicated in column 7 of the table set out in Annex I, and accompanied by an individual health certificate drawn up in accordance with the appropriate model health certificate set out in Section A or B of Part 2 of Annex II;
- (d) the import of registered horses as indicated in column 8 of the table set out in Annex I, and accompanied by an individual health certificate drawn up in accordance with the model health certificate set out in Section A of Part 3 of Annex II;
- (e) the import of a consignment of equidae for slaughter as indicated in column 9 of the table set out in Annex I, and accompanied by a health certificate drawn up in accordance with the model health certificate set out in Section B of Part 3 of Annex II;
- (f) the import of registered equidae and equidae for breeding and production as indicated in column 10 of the table set out in Annex I, and accompanied by an individual health certificate drawn up in accordance with the model health certificate set out in Section A of Part 3 of Annex II.
- 2. The competent authority of the third country of dispatch shall apply the measures necessary in order to comply with the specific conditions or temporal limitations indicated for that country in column 16 of the table in Annex I.

Third countries and parts of the territory of third countries from which the entry into the Union of semen of equidae is authorised

Member States shall authorise the entry into the Union of consignments of semen of equidae from the third countries, or, where the Union applies regionalisation, parts of the territory of third countries, listed in columns 2 and 4 of the table in Annex I, as indicated in columns 11, 12 and 13 of that table, and provided that the consignment complies with the following conditions:

- (a) the consignment is dispatched from a semen collection or storage centre listed in accordance with Article 17(3)(b) of Directive 92/65/EEC;
- (b) the consignment is accompanied by a health certificate drawn up in accordance with the appropriate model health certificate set out in Part 1 of Annex III.

Article 5

Third countries and parts of the territory of third countries from which the entry into the Union of ova and embryos of equidae is authorised

Member States shall authorise the entry into the Union of consignments of ova and embryos of equidae from the third countries, or, where the Union applies regionalisation, parts of the territory of third countries, listed in columns 2 and 4 of the table in Annex I, as indicated in column 14 of that table, and provided that the consignment complies with the following conditions:

- (a) the consignment is dispatched by an embryo collection or production team listed in accordance with Article 17(3)(b) of Directive 92/65/EEC;
- (b) the consignment is accompanied by a health certificate drawn up in accordance with the appropriate model health certificate set out in Part 2 of Annex III.

SECTION 3

General requirements for entry into the Union of consignments of equidae and of semen, ova and embryos of equidae

Article 6

Certification

- 1. The health certificates, as provided for in Articles 3, 4 and 5, shall be drawn up and issued in accordance with:
- (a) the applicable supplementary guarantees or conditions specified in column 16 of Annex I;
- (b) the explanatory notes provided for in Part 4 of Annex II and Part 3 of Annex III respectively.

2. The provisions of paragraph 1 shall not preclude the use of electronic certification or other agreed systems, whenever harmonised procedures at Union level have been established.

Article 7

Period of validity of health certificates

- 1. The operator responsible for a consignment of equidae or of semen, ova or embryos of equidae intended for entry into the Union shall ensure that the consignment is presented to an approved border inspection post authorised for the consignment concerned no later than 10 days from the date of certification of the consignment in the third country of dispatch.
- 2. Where equidae are transported by sea, the period of 10 days referred to in paragraph 1 shall be extended by the time of transport on sea.

SECTION 4

Transport requirements for entry of equidae into the Union

Article 8

General animal health requirements

- 1. The operator responsible for a consignment of equidae intended for entry into the Union shall ensure that those equidae are transported in compliance with the following:
- (a) the equidae are transported by a means of transport carrying only equidae that are destined for the Union or alternatively are accompanied by a health certificate required for transit;
- (b) the equidae are transported by a means of transport carrying only equidae of the same certified health status, except where otherwise authorised in the specific animal health requirements set out in Sections A and B of Part 1 and in Section A of Part 3 of Annex II;
- (c) the equidae are transported by road or railway or moved on foot only in a third country or a part of the territory of a third country that is authorised for at least one type of entry of at least one category of equidae.
- 2. The operator responsible for a consignment of equidae intended for entry into the Union shall ensure compliance with the following:
- (a) the crates, containers, stalls or jet-stalls and the means of transport or the transport compartment of the means of transport in which equidae will be transported are cleansed and disinfected prior to loading of the animals with a disinfectant officially recognised in the country of dispatch;
- (b) the means of transport used for road or railway transport are designed, constructed and operated to prevent the escape of faeces, urine and fodder during the intended journey;
- (c) measures to protect the animals from attacks of insect vectors shall be applied in case of the occurrence of one of the following diseases:

- (i) African horse sickness or Venezuelan equine encephalomyelitis in the third country of dispatch or transit;
- (ii) one or more of the vector-borne diseases listed in Article 11(1), with the exception of equine infectious anaemia, if the equidae are not immune or vaccinated against the pathogen.

In the case of the diseases referred to in point (i) the vector protection shall include measures such as the netting of the crates, containers, stalls or jet-stalls, forced ventilation and keeping the transport compartment closed, except during loading and unloading of the animals or when attending to the animals.

3. The operator responsible for a consignment of equidae intended for entry into the Union shall ensure that, during the journey, the equidae are only unloaded in a third country or a part of the territory of a third country that is authorised for the entry of equidae into the Union in accordance with Annex I.

Article 9

Specific animal health requirements for transport by air

- 1. The operator responsible for a consignment of equidae intended for entry into the Union by air shall ensure compliance with the following:
- (a) the crates, containers or jet-stalls and the surrounding airspace in the transport compartment are sprayed with an appropriate insect repellent in combination with an insecticide immediately after the closing of the doors of the aircraft;
- (b) the captain of the aircraft completes and signs the declaration set out in Part 1 of Annex V.
- 2. By way of derogation from paragraph 3 of Article 8, Member States may, on request of the operator of the consignment, authorise direct transhipment from one aircraft to another aircraft which takes place in a country not listed in Annex I, provided that the following requirements are satisfied:
- (a) the transhipment is carried out in the same airport within the area of the same customs office under direct supervision of an official veterinarian or the responsible customs officer;
- (b) during the transhipment the equidae are protected from attacks by insect vectors of diseases transmissible to equidae;
- (c) the equidae do not come into contact with equidae of a different health status;
- (d) the measures provided for in points (a) and (b) of paragraph 1 are applied in relation to the aircraft to be used for onward travel;
- (e) compliance with the conditions set out in point (a) of paragraph 1 and in points (a), (b) and (c) of this paragraph is certified by the official veterinarian or the responsible customs officer in the Transhipment Manifest drawn up in accordance with the model set out in Part 3 of Annex V.

Specific animal health requirements for transport by sea

- The operator responsible for a consignment of equidae intended for entry into the Union by sea shall ensure compliance with the following:
- (a) the vessel is scheduled to dock directly at a port in the Union without calling into a port of a third country or in a part of the territory of a third country not included in Annex I;
- (b) the crates, containers or stalls and the surrounding airspace in the transport compartment are sprayed with an appropriate insect repellent in combination with an insecticide immediately after the closing of the compartment;
- (c) the captain of the vessel completes and signs the declaration set out in Part 2 of Annex V.
- 2. By way of derogation from point (a) of paragraph 1, Member States may authorise direct transhipment from one vessel to another vessel which takes place in a country not listed in Annex I, provided:
- (a) the transhipment is carried out in the same port within the area of the same customs office under direct supervision of an official veterinarian or the responsible customs officer;
- (b) the equidae are during the transhipment protected from attacks by insect vectors of diseases transmissible to equidae;
- (c) the equidae do not come into contact with equidae of a different health status;
- (d) compliance with the conditions set out in point (b) of paragraph 1 and points (a), (b) and (c) of this paragraph is certified by the official veterinarian or the responsible customs officer in the Transhipment Manifest drawn up in accordance with the model set out in Part 3 of Annex V.

SECTION 5

General requirements for the testing and vaccination of equidae intended for entry into the Union and of donor equidae whose semen, ova or embryos are intended for entry into the Union

Article 11

General requirements for laboratory testing for the certification of consignments of equidae, or their semen, ova or embryos intended for entry into the Union

1. The competent authority of the third country dispatching equidae or semen, ova or embryos of equidae, which are intended for entry into the Union shall ensure that the laboratory tests provided for in the health certificates set out in Annexes II and III for glanders, dourine, equine infectious anaemia, Venezuelan equine encephalomyelitis, Western and

Eastern equine encephalomyelitis, Japanese encephalitis, West Nile Fever, vesicular stomatitis, equine viral arteritis and contagious equine metritis meet at least the sensitivity and specificity requirements laid down for the disease concerned in the respective Chapter of Section 2.5 of the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals, latest edition, of the World Organisation for Animal Health (OIE).

- 2. The competent authority of the third country dispatching equidae which are destined for the Union shall ensure that the laboratory tests provided for in the health certificates set out in Annex II for African horse sickness are carried out in accordance with Annex IV to Directive 2009/156/EC.
- 3. The competent authority of the third country dispatching equidae or semen, ova or embryos of equidae, which are destined for the Union shall ensure compliance with the following:
- (a) the tests referred to in paragraphs 1 and 2 are carried out in a laboratory recognised by the competent authority in the third country of dispatch;
- (b) the details of sampling and the results of the tests are stated as required in the health certificate set out for the consignment concerned in Annex II or III based on the laboratory report made available to the certifying official veterinarian.

Article 12

Testing upon arrival in the Union

- 1. Where a test carried out, in or on behalf of the Member State of entry, on a sample taken in accordance with Article 4 of Decision 97/794/EC does not confirm the result of a laboratory test attested in a health certificate accompanying equidae or semen, ova or embryos of equidae arriving in the Union, as set out in Annex II or III to this Regulation, the competent authority of that Member State of entry shall ensure that the test is repeated in the national reference laboratory designated for the disease concerned in accordance with Article 4(1) of Regulation (EC) No 882/2004 of the European Parliament and of the Council (1).
- 2. Where the measures provided for in paragraph 1 do not result in a conclusive outcome of the checks for compliance carried out in accordance with Article 4 of Decision 97/794/EC, the competent authority referred to in paragraph 1 shall ensure that the sample referred to in that paragraph is subjected to definitive testing as follows:
- (a) for African horse sickness, in the European Union reference laboratory for African horse sickness designated in accordance with Council Directive 92/35/EEC (2);

⁽¹) Regulation (EC) No 882/2004 of the European Parliament and of the Council of 29 April 2004 on official controls performed to ensure the verification of compliance with feed and food law, animal health and animal welfare rules (OJ L 165, 30.4.2004, p. 1).

⁽²⁾ Council Directive 92/35/EEC of 29 April 1992 laying down control rules and measures to combat African horse sickness (OJ L 157, 10.6.1992, p. 19).

(b) for the diseases referred to in Article 11(1), in the European Union reference laboratory for equine diseases other than African horse sickness, designated in accordance with Regulation (EC) No 180/2008.

Article 13

Application of vaccines and recording of vaccination

- 1. The competent authority of the third country dispatching equidae or semen, ova or embryos of equidae, which are destined for the Union, shall ensure that the vaccination attested in any of the certificates set out in Annexes II or III is carried out in compliance with the following:
- (a) the vaccination is carried out in accordance with the manufacturers' instructions or national legislation, whatever is stricter;
- (b) the vaccination is carried out using a licensed vaccine which meets at least the requirements for safety, sterility and efficacy set out for the vaccine concerned in the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals, latest edition, of the World Organisation for Animal Health (OIE).
- 2. Where the competent authority of a third country attests that a positive laboratory finding in a serological test for African horse sickness is related to previous vaccination, the vaccination shall be documented in the identification document accompanying the equine animal, where such identification document is available.

Article 14

Requirements relating to equine viral arteritis

- 1. Uncastrated male equidae intended for entry into the Union, with the exception of those listed in point 1 of Annex IV, shall be subject to tests for equine viral arteritis to ascertain that their semen is free of equine arteritis virus.
- 2. Vaccination against equine viral arteritis, including the testing required in accordance with point 1(a) of Annex IV, shall be carried out under official veterinary supervision.
- 3. Vaccination against equine viral arteritis shall be valid where there is documented proof accompanying the equine animal of an uninterrupted history of a primary course carried out in compliance with one of the vaccination protocols provided for in point 1(a) of Annex IV and regular revaccination according to manufacturers' recommendations and in any event at intervals of not more than 12 months.

SECTION 6

Identification of equidae intended for entry into the Union

Article 15

Identification of equidae intended for entry into the Union

1. Equidae destined for entry into the Union shall be individually identified so as to ensure an unequivocal correspondence between the animal and its certified health status.

That identification shall:

- (a) either comply with the requirements of Article 14 of Implementing Regulation (EU) 2015/262; or
- (b) provide at least the information detailed in points 1, 2, 3 and 6 to 10 of Part A and in points 12 to 18 of Part B of Section I of Part 1 of Annex I to that Regulation.
- 2. Equidae for slaughter to be imported into the Union shall be marked individually with an electronic transponder or an ear tag, the number of which shall be recorded in the health certificate accompanying the animals during transport.
- 3. Equidae for slaughter to be imported into the Union shall bear on their left front hoof a clear and indelible hot-branded 'S' of the size not less than half the length of the hoof wall, in the following cases:
- (a) if they are marked individually, by way of derogation from paragraph 2, by an alternative method indicated in the health certificate, in which case the animals must be consigned to the slaughterhouse of destination in accordance with Article 21(a);
- (b) if they are destined to be consigned to the slaughterhouse of destination in accordance with Article 21(b).

SECTION 7

Specific animal health and certification requirements for entry into the Union of consignments of equidae

Article 16

Measures to be taken by the competent authorities to ensure traceability of a registered horse admitted temporarily

- 1. Provided compliance with the entry conditions has been established, the competent authority at the border inspection post of entry shall:
- (a) retain a copy of the health certificate referred to in Article 3(1)(a);
- (b) inform, through TRACES, the relevant competent authority or border inspection post of exit, as appropriate, of the entry of a temporarily admitted registered horse, as follows:
 - (i) the competent authority of the place of destination indicated in Box I.6 of the common veterinary entry document ('CVED') set out in Annex I to Regulation (EC) No 282/2004;
 - (ii) the border inspection post of exit declared in the declaration by the owner or the representative of the owner of the registered horse accompanying the health certificate referred to in Article 3(1)(a) by completing Box I.24 of the CVED;

- (iii) the competent authorities responsible for the places of temporary residence indicated in the declaration by the owner or the representative of the owner of the registered horse accompanying the health certificate referred to in Article 3(1)(a);
- (c) deliver at least one copy of the CVED to the operator identified as 'operator responsible for the consignment' in Box I.7 of the CVED referred to in paragraph 1(b).
- 2. Where a registered horse is to be moved from one Member State to another Member State during its temporary admission, the competent authority of the place of dispatch shall:
- (a) provided the animal health conditions of Articles 4 and 5 of Directive 2009/156/EC are fulfilled, issue a health certificate in accordance with Annex III to Directive 2009/156/EC either for an individual registered horse or for a consignment of registered horses of the same origin and with the same destination, and enter in Box I.6 of that certificate a reference to the health certificate referred to in Article 3(1)(a) of each temporarily admitted registered horse forming the consignment and a reference to the CVED referred to in point (i) of paragraph 1(b);
- (b) inform, through TRACES, the competent authority at the place of destination, of the movement of a registered horse to that Member State, and request the verification of arrival by completing a further Part III of the CVED referred to in point (i) of paragraph 1(b);
- (c) deliver to the operator, as identified in Box I.7 of the CVED referred to in point (i) of paragraph 1(b), a new print of the CVED displaying the Part III added in accordance with point (b) of this paragraph;
- (d) invalidate or withdraw any print of the CVED delivered to the operator in accordance with paragraph 1(c), or, if there had been a previous movement to another Member State, in accordance with point (c) of this paragraph.
- 3. The competent authority of the place of destination referred to in point (i) of paragraph 1(b) and in paragraph 2(b) shall acknowledge through TRACES the arrival of the registered horse and document the checks carried out by completing Part III of the CVED.
- 4. At the end of the temporary admission, the competent authority referred to in points (i) or (iii) of paragraph 1(b) which certifies the temporarily admitted registered horse to the third country of origin or to another third country, shall:
- (a) inform, through TRACES, the border inspection post of exit of the departure of the temporarily admitted registered horse from the Union, by completing a further Part III of the CVED referred to in point (i) of paragraph 1(b);
- (b) deliver to the operator, as identified in Box I.7 of the CVED referred to in point (i) of paragraph 1(b), a new print of the CVED displaying the Part III added in accordance with point (a) of this paragraph;

- (c) where the border inspection post of exit is situated in another Member State.
 - (i) issue, in accordance with Decision 93/444/EEC, a certificate in accordance with Annex III to Directive 2009/156/EC either for an individual registered horse or for a consignment of registered horses of the same origin and with the same destination;
 - (ii) enter in Box I.6 of the certificate referred to in point (i) a reference to the health certificate referred to in Article 3(1)(a) of each temporarily admitted registered horse forming the consignment and a reference to the CVED referred to in point (i) of paragraph 1(b).
- 5. The border inspection post of exit referred to in point (a) of paragraph 4 shall document the termination of the temporary admission of the registered horse by completing Part III of the CVED accordingly.
- 6. Where the temporary admission of a registered horse has not been terminated in accordance with paragraph 5 within a period of less than 90 days following the date of issue of the CVED referred to in point (i) of paragraph 1(b), an alert is sent automatically through TRACES to the border inspection post of entry and the competent authorities referred to in this Article until those competent authorities have determined the status of the registered horse.

Operator responsibilities for temporarily admitted registered horses

- 1. The operator responsible for a registered horse temporarily admitted into the Union, as identified in Box I.7 of the CVED referred to in point (i) of Article 16(1)(b), shall ensure that the following conditions are met:
- (a) the registered horse shall at all times during its temporary admission be accompanied by its original health certificate referred to in Article 3(1)(a) and by the CVED issued by the border inspection post of entry into the Union;
- (b) the registered horse shall remain in the respective Member State and on the premises stated in the declaration accompanying the health certificate referred to in Article 3(1)(a);
- (c) where the registered horse is to be moved to another Member State, it shall be accompanied by a health certificate in accordance with Annex III to Directive 2009/156/EC and the modified CVED delivered by the competent authority in accordance with Article 16(2);
- (d) any former prints of the CVED are surrendered to the competent authority for invalidation or withdrawal;
- (e) the registered horse shall leave the Union through a border inspection post indicated in the health certificate referred to in in Article 3(1)(a) not later than 89 days following the date of entry into the Union indicated on the corresponding CVED.

- 2. The operator referred to in paragraph 1 shall remain responsible for the movement of the registered horse during its temporary admission in the Union, and in particular shall inform:
- (a) the competent authority referred to in points (i) and (iii) of Article 16(1)(b) regarding any changes to be made to the movements stated in the declaration accompanying the health certificate referred to in Article 3(1)(a);
- (b) the border inspection post of exit regarding the date when the temporarily admitted registered horse is to depart from the Union;
- (c) the competent authority referred to in points (i) and (iii) of Article 16(1)(b) responsible for the holding regarding the death or loss of the registered horse or any emergency, such as health conditions, requiring veterinary attention beyond the 89 days of temporary admission.

Re-entry after temporary export of registered horses temporarily admitted into the Union

- 1. Registered horses temporarily admitted into the Union may be authorised for re-entry after temporary export to a third country or part of the territory of a third country authorised for the re-entry of registered horses to take part in specific races, competitions or cultural events for which model health certificates for re-entry into the Union are laid down in accordance with Article 20(3), provided that the re-entry into the Union takes place within a period of less than 90 days following the date of issuing of the CVED referred to in point (i) of Article 16(1)(b).
- 2. In order to allow the re-entry of a registered horse referred to in paragraph 1, the competent authority referred to in points (i) and (iii) of Article 16(1)(b) issuing the certificate for the temporary export shall:
- (a) apply the measures provided for in points (a), (b) and, where applicable, (c) of Article 16(4);
- (b) inform, through TRACES, the border inspection post of scheduled re-entry by completing Part III of the CVED;
- (c) deliver to the operator as identified in Box I.7 of the CVED referred to in point (i) of Article 16(1)(b), a new print of the CVED displaying the Part III added in accordance with point (b) of this paragraph;
- (d) invalidate or withdraw any print of the CVED delivered in accordance with Article 16(1)(c) or, if there had been a previous movement to another Member State, in accordance with Article 16(2)(c).
- 3. The border inspection post of re-entry shall:
- (a) retain the original of the health certificate referred to in Article 3(1)(c);

- (b) inform, through TRACES, of the re-entry of the registered horse:
 - (i) the competent authority of the place of destination, as declared in the declaration accompanying the health certificate referred to in Article 16(1)(a), or as modified in accordance with Article 17(2)(a);
 - (ii) the border inspection post of exit, as declared in the declaration accompanying the health certificate referred to in Article 16(1)(a), or as modified in accordance with Article 17(2)(a), by completing Box I.24 of the CVED referred to in point (d);
- (c) request the competent authority of the place of destination to verify and, where appropriate, to confirm the arrival of the registered horse by completing Box I.6 of the CVED referred to in point (d);
- (d) deliver to the operator a print of a new CVED in which Box II.1 is completed with a reference to the number of the CVED delivered previously in accordance with Article 16(1)(c) or, if there had been a previous movement to another Member State, in accordance with Article 16(2)(c), and in which Box II.14 is completed within the deadline for leaving the Union indicated in the CVED referred to in point (i) of Article 16(1)(b);
- (e) invalidate or withdraw any print of the CVED delivered to the operator in accordance with Article 16(1)(c) or, if there had been a previous movement to another Member State, in accordance with Article 16(2)(c).
- 4. Following the re-entry after temporary export of a temporarily admitted registered horse in accordance with paragraph 1, the rules laid down in Article 16 apply for the remaining period of less than 90 days following the date of issuing of the CVED referred to in point (i) of Article 16(1)(b).

Conversion of temporary admission into permanent entry and death or loss of a registered horse

- 1. Where the operator, as identified in Box I.7 of the CVED referred to in point (i) of Article 16(1)(b), submits an application to the competent authority referred to in point (i) or (iii) of Article 16(1)(b) or in Article 16(2)(b), to convert the temporary admission of a registered horse into a permanent entry, a Member State may authorise that conversion provided that the following requirements are met:
- (a) in accordance with Annex I, imports of registered horses are authorised from the third country or part of the territory of the third country concerned;
- (b) the competent authority responsible for the place of temporary residence has complied with the following conditions:
 - (i) that competent authority has carried out with satisfactory results the checks necessary to verify compliance with the test and vaccination requirements for imports of registered horses from the third country or part of the territory of the third country concerned set out in Part 3 of Annex II;

- (ii) that competent authority has ensured that the registered horse remained under official veterinary supervision in that Member State until 3 months have elapsed from the date of its entry into the Union indicated on the CVED referred to in point (i) of Article 16(1)(b).
- 2. The competent authority referred to in paragraph 1, or a border inspection post designated for this task by the Member State, shall:
- (a) terminate the temporary admission in TRACES by choosing 'Conversion into permanent entry' in Part III of the CVED delivered to the operator in accordance with either Article 16(1)(c), or, if there had been a previous movement to another Member State, with Article 16(2)(c) or, if there had been a previous re-entry after temporary export, with Article 18(3)(c);
- (b) deliver to the operator identified in Box I.7 of the CVED referred to in point (i) of Article 16(1)(b), a new print of the CVED referred to in point (a), or a new CVED, in which 'For internal market' is checked in Box I.21;
- (c) invalidate or withdraw any print of the CVED delivered to the operator in accordance with either Article 16(1)(c), or, if there had been a previous movement to another Member State, with Article 16(2)(c) or, if there had been a previous re-entry after temporary export, with Article 18(3)(c);
- (d) invalidate or withdraw the original of the health certificate referred to in Article 3(1)(a).
- 3. During the period of conversion, the operator, as identified in Box I.7 of the CVED issued in accordance with point (i) of Article 16(1)(b) or Article 18(3)(b) of the registered horse shall take the following measures:
- (a) arrange regular visits carried out and recorded by a veterinarian to check the registered horse for clinical signs of possible infectious diseases;
- (b) keep records on the movement of the registered horse and on movements of equidae on and off the holding where it is kept;
- (c) complete the customs procedures, as referred to in Article 15 of Implementing Regulation (EU) 2015/262;
- (d) make an application in accordance with Article 15(1) of Implementing Regulation (EU) 2015/262 for the issuing of an identification document or the adaptation of an existing identification document.
- 4. In the case of death or loss of a registered horse temporarily admitted into the Union, the competent authority of the place of death or loss, where required by the Member State concerned in close collaboration with a border inspection post, shall:
- (a) terminate the temporary admission in TRACES by choosing 'Death/Loss' in Part III of the CVED referred to in point (i) of Article 16(1)(b) or Article 18(3)(b);

(b) invalidate or withdraw any print of the CVED delivered to the operator in accordance with either Article 16(1)(c), or, if there had been a previous movement to another Member State, with Article 16(2)(c) or, if there had been a previous re-entry after temporary export, with Article 18(3)(c).

Article 20

Specific animal health conditions regarding the re-entry of registered horses after temporary export for races, competition and cultural events

- 1. Member States shall authorise the re-entry of registered horses subject to compliance with the following conditions:
- (a) the registered horse has remained outside the Union for not more than 30 days, unless specifically provided for in paragraph 3;
- (b) the registered horse has neither been resident in nor transited on land through any third country or part of the territory of a third country that is not assigned to the same sanitary group as the third country or part of the territory of a third country in which the health certificate in accordance with Section A of Part 2 of Annex II has been signed by the official veterinarian;
- (c) the health certificate for temporary export signed by the official veterinarian in the Member State of origin, or an authorised copy thereof, is presented on request of the border inspection post of re-entry into the Union.
- 2. The competent authority certifying a registered horse for temporary export to a third country shall ensure that in application of Article 2(1) of Decision 93/444/EEC the registered horse is accompanied until the exit point in another Member State by a health certificate in accordance with Annex III to Directive 2009/156/EC.
- 3. The re-entry after temporary export for a period of more than 30 days of registered horses taking part in specific races, competitions or cultural events is subject to specific animal health requirements as contained in the corresponding model health certificates provided for Section B of Part 2 of Annex II in respect of the relevant event.
- 4. The operator, as identified in Box I.7 of the CVED, responsible for the consignment shall ensure that during the temporary export the registered horse neither has been resident in nor has transited on land through any third country or part of the territory of a third country that is not assigned to the same sanitary group as the third country or part of the territory of a third country in which the health certificate in accordance with Section A of Part 2 of Annex II has been signed by the official veterinarian.

Article 21

Specific animal health conditions regarding imports of equidae for slaughter

The operator, as identified in Box I.7 of the CVED referred to in point (i) of Article 16(1)(b), of a consignment of equidae for slaughter shall ensure that after the checks carried out at the border inspection post of entry into the Union, the animals:

- (a) are either conveyed directly, without delay and without coming into contact with equidae of a different health status, to the slaughter-house of destination where they shall be slaughtered within 72 hours of arrival at the slaughterhouse; or
- (b) pass through a single approved market or marshalling centre referred to in Article 7(1) of Directive 2009/156/EC as indicated in the health certificate referred to in Article 3(1)(e) of this Regulation, from where they are to be removed after the market under national rules ensuring traceability directly to a slaughterhouse to be slaughtered, as soon as possible, but at the latest within 5 working days of arrival in the Union without coming into contact with equidae of a different health status.

SECTION 8

Transitional and final provisions

Article 22

Transitional provisions

For a transitional period until 31 December 2018, Member States shall authorise the entry into the Union of consignments of equidae and of semen, ova and embryos of equidae accompanied by health certificates drawn up in accordance with the model health certificates applicable before the date of application of this Regulation specified in the second subparagraph of Article 24.

Article 23

Repeals

Decisions 92/260/EEC, 93/195/EEC, 93/196/EEC, 93/197/EEC, 94/699/EC, 95/329/EC, 2003/13/EC, 2004/177/EC, 2004/211/EC, 2010/57/EU and 2010/471/EU are repealed.

Any reference to those Decisions shall be construed as a reference to this Regulation.

Article 24

Entry into force and applicability

This Regulation shall enter into force on the twentieth day following that of its publication in the Official Journal of the European Union.

It shall apply from 1 October 2018.

However, Article 16(1)(b)(iii), Article 16(2)(b), (c) and (d), Article 16(3), Article 16(4)(a) and (b), Article 16(5) and Article 17(1)(d) shall apply from 14 December 2019.

This Regulation shall be binding in its entirety and directly applicable in all Member States.

ANNEX I

LIST OF THIRD COUNTRIES (¹) AND PARTS OF THE TERRITORY OF THIRD COUNTRIES (²) FOR THE ENTRY INTO THE UNION OF CONSIGNMENTS OF EQUIDAE AND OF SEMEN, OVA AND EMBRYOS OF EQUIDAE

		Code of the			TA	Re-En		Imports			Imp	orts		Transit	Specific conditions
ISO-Code	Third country	part of the territory of the third	Description of the part of the territory of the third country	SG	RH	RH	RH	ES	RE +		SEMEN		O/E	Equidae	
		country			KII	KII	KΠ	ES	EBP	RH	RE	EBP			
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16
AE	United Arab Emirates	AE-0	Whole country	Е	X	X	X	_	_	X	_	_	X	X	
AR	Argentina	AR-0	Whole country	D	X	X	X	X	X	X	X	X	X	X	
AU	Australia	AU-0	Whole country	A	Х	Х	X	X	X	X	X	X	X	X	
ВВ	Barbados	BB-0	Whole country	D	X	X	X	_	_	X	_	_	_	X	
ВН	Bahrain	BH-0	Whole country	Е	Х	X	X	_	_	_	_	_	_	X	
BM	Bermuda	BM-0	Whole country	D	X	X	X	_	_	X	_	_	_	X	
ВО	Bolivia	BO-0	Whole country	D	X	X	X	_	_	X	_	_	_	X	
BR	Brazil	BR-0	Whole country	_	_	_	_	_	_	_	_	_	_		
		BR-1	The states of: Rio Grande do Sul, Santa Catarina, Mato Grosso do Sul, Distrito Federal and Rio de Janeiro	D	Х	Х	Х	_	_	Х	_	_	_	X	
BY	Belarus	BY-0	Whole country	В	X	X	X	X	X					X	
CA	Canada	CA-0	Whole country	С	Х	Х	Х	X	X	Х	X	X	X	X	

		Code of the			TA	Re-En		Imports			Imp	orts		Transit	Specific conditions
ISO-Code	Third country	part of the territory of the third	Description of the part of the territory of the third country	SG	RH	RH	RH	ES	RE +		SEMEN		O/E	Equidae	
		country			KII	KII	KII	ES	EBP	RH	RE	EBP			
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16
СН	Switzerland (1)	СН-0	Whole country	A	X	X	X	X	X	X	X	X	X	X	
CL	Chile	CL-0	Whole country	D	X	X	X	X	X	_	_	_	_	X	
CN	China	CN-0	Whole country	_	_	_	_	_	_	_	_	_	_		
		CN-1	The equine disease-free zone in Conghua City, Guangzhou Municipality, Guangdong Province including the Biosecurity Highway Passage from and to the airport in Guangzhou and Hong Kong (see BOX 1 for details)	G	X	X	X	_	_	_	_	_	_	X	
		CN-2	The venue for the Global Champions Tour at the Expo 2010 No 15 Parking Lot and the passage to the Shanghai Pudong International Airport in the northern part of the Pudong New area and the Eastern part of the Minhang District of the Metropolitan area of Shanghai (see BOX 1 for details)	G	_	х	_	_	_	_	_	_	_		Only if certified in accordance with Chapter 1 of Section B of Part 2 of Annex II

		Code of the			TA	Re-En		Imports			Imp	orts		Transit	Specific conditions
ISO-Code	Third country	part of the territory of the third	Description of the part of the territory of the third country	SG	RH	RH	RH	ES	RE +		SEMEN		O/E	Equidae	
		country			KII	KII	KII	ES	EBP	RH	RE	EBP			
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16
CR	Costa Rica	CR-0	Whole country	_	_	_		_	_	_	_	_	_		
		CR-1	Metropolitan area of San José	D	_	X	_		_		_	_	_		
CU	Cuba	CU-0	Whole country	D	X	X	X	_	_		_	_	_	X	
DZ	Algeria	DZ-0	Whole country	Е	X	X	X	X	X					X	
EG	Egypt	EG-0	Whole country		_	_	_	_	_	_	_	_	_		
		EG-1	The Equine Disease Free Zone established at the Egyptian Armed Forces Veterinary Hospital at El Nasr road, across Al Ahly Club, Cairo, and the highway passage to Cairo International Airport (see BOX 2 for details)	Е	X	_	X	_	_	_	_	_	_	X	
FK	Falkland Islands	FK-0	Whole country	A	X	X	X	_	X					X	
GL	Greenland	GL-0	Whole country	A	Х	X	X	X	X					X	
НК	Hong Kong	HK-0	Whole country	G	Х	X	X	_	_		_	_	_	X	
IL	Israel (3)	IL-0	Whole country	Е	Х	X	X	X	X	X	X			X	
IS	Iceland (5)	IS-0	Whole country	A	Х	Х	X	Х	Х	Х	X	X		Х	

_		Code of the			TA	Re-En		Imports			Imp	orts		Transit	Specific conditions
ISO-Code	Third country	part of the territory of the third	Description of the part of the territory of the third country	SG	RH	RH	RH	ES	RE +		SEMEN		O/E	Equidae	
		country			KII	KII	KII	LS	EBP	RH	RE	EBP			
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16
JM	Jamaica	JM-0	Whole country	D	X	X	X	_	_		_		_	X	
JO	Jordan	JO-0	Whole country	Е	X	X	X	-	_		_	1	_	X	
JP	Japan	JP-0	Whole country	G	X	X	X	_	_		_		_	X	
KG	Kyrgyzstan	KG-0	Whole country	_	_	_	_	_	_	_	_	_	_		
		KG-1	Region of Issyk-Kul	В	_	_	X	_	_		_	_	_	X	
KR	Korea Republic	KR-0	Whole country	G	X	X	X	_	_		_	_	_	X	
KW	Kuwait	KW-0	Whole country	Е	X	X	X	_	_	_	_	_	_	X	
LB	Lebanon	LB-0	Whole country	Е	X	X	X	_	_	_	_	_	_	X	
MA	Morocco	MA-0	Whole country	Е	X	X	X	X	X	X	X	X		X	
ME	Montenegro	ME-0	Whole country	В	X	X	X	X	X					X	
MK	fYROM (4)	MK-0	Whole country	В	X	X	X	X	X					X	
МО	Macao	МО-0	Whole country	G	X	X	X	_	_		_	_	_	X	
MY	Malaysia	MY-0	Whole country	_	_	_	_	_	_	_	_	_	_		
		MY-1	Peninsula	G	X	X	X	_	_		_	_	_	X	
MU	Mauritius	MU-0	Whole country	Е	_	_	X	_	_		_		_	X	

		Code of the			TA	Re-En		Imports			Imp	orts		Transit	Specific conditions
ISO-Code	Third country	part of the territory of the third	Description of the part of the territory of the third country	SG	RH	RH	RH	ES	RE +		SEMEN		O/E	Equidae	
		country			KII	KII	KII	LS	EBP	RH	RE	EBP			
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16
MX	Mexico	MX-0	Whole country	С	_	_	_	_	_	_	_	_	_		
		MX-1	Metropolitan area of Mexico-City	С		X									Only if certified in accordance with Chapter 1 of Section B of Part 2 of Annex II
NO	Norway (5)	NO-1	Whole country	A	X	X	X	X	X	X	X	X	X	X	
NZ	New Zealand	NZ-0	Whole country	A	X	X	X	X	X					X	
OM	Oman	OM-0	Whole country	Е	X	X	X				_	_		X	
PE	Peru	PE-0	Whole country	_	_	_	_	_	_	_	_	_	_		
		PE-1	Region of Lima	D	X	X	X	_	_		_	_		X	
PM	St Pierre & Miquelon	PM-0	Whole country	A			X	—	X					X	
PY	Paraguay	PY-0	Whole country	D	X	X	X	X	X					X	
QA	Qatar	QA-0	Whole country	Е	X	X	X	_	_		_	_	_	X	
RS	Serbia (6)	RS-0	Whole country	В	X	X	X	X	X					X	

		Code of the			TA	Re-En		Imports			Imp	orts		Transit	Specific conditions
ISO-Code	Third country	part of the territory of the third	Description of the part of the territory of the third country	SG	RH	RH	RH	ES	RE +		SEMEN		O/E	Equidae	
		country			KII	KII	KII	ES	EBP	RH	RE	EBP			
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16
RU	Russia	RU-0	Whole country	_	_	_	_	_	_	_	_	_	_		
		RU-1	Provinces of Kaliningrad, Arkhangelsk, Vologda, Murmansk, Leningrad, Novgorod, Pskov, Briansk, Vladimir, Ivanovo, Tver, Kaluga, Kostroma, Moskva, Orjol, Riasan, Smolensk, Tula, Jaroslavl, Nijninov- gorod, Kirov, Belgorod, Voronesh, Kursk, Lipezk, Tambov, Astrahan, Volgograd, Penza, Saratov, Uljanovsk, Rostov, Orenburg, Perm and Kurgan	В	X	X	X	X	X					X	
		RU-2	Regions of Stavropol and Krasnodar	В	X	X	X	X	X					X	
		RU-3	Republics of Karelia, Marij-El, Mordovia, Chuvachia, Kalmykia, Tatarstan, Dagestan, Kabardino-Balkaria, Severnaya Osetia, Ingushetia and Karachaevo-Cherkesia	В	X	Х	Х	Х	Х					Х	
SA	Saudi Arabia	SA-0	Whole country	_	_	_	_	_	_	_	_	_	_		
		SA-1	Whole country, except SA-2	Е	X	X	X	_	_	X	_	_	_	X	

		Code of the			TA	Re-En		Imports			Imp	oorts		Transit	Specific conditions
ISO-Code	Third country	part of the territory of the third	Description of the part of the territory of the third country	SG	DVV	DVV	DVV	FG	RE +		SEMEN		O/E	Equidae	
		country			RH	RH	RH	ES	EBP	RH	RE	EBP			
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16
		SA-2	Protection and surveillance zones in the provinces of Jizan, Asir and Najran as described in BOX 3	_	_	_	_			_					
SG	Singapore	SG-0	Whole country	G	Х	Х	X	_	_		_	_	_	X	
TH	Thailand	TH-0	Whole country	G	X	X	X	_	_		_	_	_	X	
TN	Tunisia	TN-0	Whole country	Е	X	X	X	X	X					X	
TR	Turkey	TR-0	Whole country	_	_	_	_	_	_	_	_	_	_		
		TR-1	Provinces of Ankara, Edirne, Istanbul, Izmir, Kirklareli and Tekirdag	Е	_	_	_	_	_		_	_	_	_	
UA	Ukraine	UA-0	Whole country	В	X	X	X	X	X	X	X	Х		X	
US	United States of America	US-0	Whole country	С	X	X	X	X	X	X	X	X	X	X	
UY	Uruguay	UY-0	Whole country	D	X	X	X	X	X	X	X	X		X	
ZA	South Africa	ZA-0	Whole country	_	_	_	_	_	_	_	_	_	_		
		ZA-1	Metropolitan area of Cape-Town (see BOX 4 for details)	F	_	_	_	_	_			_	_		Commission Decision 2008/698/EC

⁽¹⁾ Without prejudice to specific certification requirements provided for in Decision 2002/309/EC, Euratom of the Council, and of the Commission.

⁽²⁾ Where official regionalisation applies in accordance with Article 13(2)(a) of Directive 2009/156/EC.

⁽³⁾ Hereinafter understood as the State of Israel, excluding the territories under Israeli administration since June 1967, namely the Golan Heights, the Gaza Strip, East Jerusalem and the rest of the West Bank.

⁽⁴⁾ The former Yugoslav Republic of Macedonia - the definitive nomenclature for this country will be agreed following current negotiations at UN level.

⁽⁵⁾ Without prejudice to specific certification requirements provided for in Article 17 of the Agreement on the European Economic Area (OJ L 1, 3.1.1994, p. 3).

⁽⁶⁾ Excluding Kosovo as defined by United Nations Security Council Resolution 1244 of 10 June 1999.

LEGEND TO ANNEX I:

Animal/Product	Categories/conditions
RH	Registered horses as defined in Article 2(c) of this Regulation.
ES	Equidae for slaughter as defined in Article 2(d) of Directive 2009/156/EC.
RE	Registered equidae as defined in Article 2(c) of Directive 2009/156/EC.
EBP	Equidae for breeding and production as defined in Article 2(e) of Directive 2009/156/EC.
SEMEN	Semen of the equine species collected in accordance with Article 17(2)(b)(ii) of Directive 92/65/EEC.
O/E	Ova and embryos of the equine species collected or produced in accordance with Article 17(2)(b)(ii) of Directive 92/65/EEC.

Columns	Information / Description of commodity	Required Health Certificate
1-4	Territorial description	NA
5	Sanitary Group	NA
6	Temporary admission of registered horses	ANNEX II Part 1 Section A
7	Re-entry of registered horses after temporary export for racing, competition and cultural events	ANNEX II Part 2 Section A ANNEX II Part 2 Section B Chapter 1 ANNEX II Part 2 Section B Chapter 2
8	Imports of registered horses	ANNEX II Part 3 Section A
9	Imports of equidae for slaughter	ANNEX II Part 3 Section B
10	Imports of registered equidae and equidae for breeding and production	ANNEX II Part 3 Section A
11	Imports of semen collected from registered horses	Annex III Part 1 Section A Annex III Part 1 Section B Annex III Part 1 Section C Annex III Part 1 Section D

▼<u>B</u>

Columns	Information / Description of commodity	Required Health Certificate
12	Imports of semen collected from registered equidae	Annex III Part 1 Section A Annex III Part 1 Section B Annex III Part 1 Section C Annex III Part 1 Section D
13	Imports of semen collected from equidae for breeding and production	Annex III Part 1 Section A Annex III Part 1 Section B Annex III Part 1 Section C Annex III Part 1 Section D
14	Imports of ova and embryos of the equine species	Annex III Part 2 Section A Annex III Part 2 Section B
15	Equidae in Transit	ANNEX II Part 1 Section B
16	Reference to specific conditions/ additional guarantees	NA

Boxes

- X Entry authorised
- Entry not authorised

Sanitary Groups

Specific animal health guarantees required for entry of equidae into the Union
equine infectious anaemia, equine viral arteritis
equine infectious anaemia, equine viral arteritis, glanders, dourine
equine infectious anaemia, equine viral arteritis, Eastern and Western equine encephalomyelitis, vesicular stomatitis
equine infectious anaemia, equine viral arteritis, glanders, dourine, Eastern and Western equine encephalomyelitis, Venezuelan equine encephalomyelitis, vesicular stomatitis
equine infectious anaemia, equine viral arteritis, glanders, dourine, African horse sickness
equine infectious anaemia, dourine, African horse sickness
equine infectious anaemia, equine viral arteritis, glanders, dourine, Japanese encephalitis

▼<u>B</u>

CN	China	CN-1	The specific equine following delimitation	disease-free zone in the Guangdong Province with the on:
			Core zone:	equestrian site in Reshui Village, Lingkou Town of Conghua City with the surrounding area within a five km radius controlled by the road control post at State Highway 105;
			Surveillance zone:	all administrative divisions in Conghua City surrounding the core zone covering an area of 2 009 km ² ;
			Protection zone:	outwards boundaries of the following contiguous administrative divisions surrounding the surveillance zone:
				 Baiyun District, Luogang District of Conghua City,
				- Huadu District of Guangzhou City,
				— Zengcheng City,
				 administrative divisions in Qingcheng District of Qingyuan City,
				— Fogang County,
				— Xinfeng County,
				— Longmen County
			Biosecurity highway passage:	— from the equestrian site in the core zone to Guangzhou Baiyun International Airport through to the State Highway 105, Jiebei Highway, airport expressway, including the equine exclusion zone of one km around Baiyun Inter- national Airport in Guangzhou City;
				— from the equestrian site in the core zone to Shenzhen Huanggang Port at the border of China with Hong Kong through State Highway 105, Jiebei highway, No. 2 north ring expressway and Guang-Shen highway with the equine exclusion zone on both sides of that highway of at least one km width;
			Pre-entry quarantine:	the quarantine facilities in the protection zone designated by the competent authority for the preparation of equidae from other parts of China for entry into the equine disease free zone.
CN	China	CN-2	Delimitation of the z	zone in the Metropolitan area of Shanghai:
			Western boundary:	Huangpu River from its estuary in the North to the bifurcation of the Dazhi River,
			Southern boundary:	from the bifurcation of the Huangpu River to the estuary of the Dazhi River in the East,
			Northern and Eastern boundaries:	coast line.

▼<u>B</u>

BOX	2		
EG	Egypt	EG-1	The Equine Disease Free Zone (EDFZ) of about 0,1 km² size, established around the Egyptian Armed Forces Veterinary Hospital at El-Nasr Road, across Al Ahly Club, on the Eastern outskirts of Cairo, (localised at 30°04′19.6″N 31°21′16.5″E) and the passage of 10 km on the El-Nasr Road and the Airport Road to Cairo International Airport.
			(a) Delineation of the boundaries of the EDFZ:
			From the crossing of El-Nasr Road with El-Shaheed Ibrahim El-Shaikh Road (at 30°04′13.6″N 31°21′04.3″E) along the El-Shaheed Ibrahim El-Shaikh Road for about 500 m to the North until the first junction with the Passage Inside Armed Forces, turning right and following the Passage for about 100 m to the East, turning right again and following the Passage for 150 m to the South, turning left and following the Passage for 300 m to the East, turning right and following the Passage for 100 m to the South until El-Nasr Road, turning right and following El-Nasr Road for 300 m to South-West until opposite of the junction of El-Nasr Road with Hassan Ma'moon Road, turning right and following the Passage for 100 m to the North, turning left and following the Passage for 120 m to the West, turning left and following the Passage for 200 m to the South, turning right and following El-Nasr Road for 100 m to the West until the crossing of El-Nasr Road with El-Shaheed Ibrahim El-Shaikh Road.
			(b) Delineation of the boundaries of the pre-export quarantine area within the EDFZ:
			From the point opposite of the junction of El-Nasr Road with Hassan Ma'moon Road following the Passage for 100 m to the North, turning right and following the Passage for 250 m to the East, turning right and following the Passage for 50 m to the South until El-Nasr Road, turning right and following El-Nasr Road for 300 m to South-West until opposite of the junction of El-Nasr Road with Hassan Ma'moon Road.
ВОХ	3		
SA	Saudi Arabia	SA-1	Approved Quarantine stations:
			1. Riyadh Airport
			2. King Abdulaziz Race Track (Janadrijah)
		SA-2	Delimitation of the protection and surveillance zones established in accordance with points (a) and (b) of the second paragraph of Article 5(2) of Directive 2009/156/EC:
			1. Province of Jizan
			 Protection zone: the whole province, except the part north of the road control post at Ash-Shuqaiq at road No 5 and north of road No 10;
			 Surveillance zone: the part of the province north of the road control post at Ash-Shuqaiq at road No 5, controlled by the road control post at Al Qahmah, and north of road No 10.
			2. Province of Asir
			— Protection zone: the part of the province delineated by road No 10, between Ad Darb, Abha and Khamis-Mushayt to the north, except the equestrian clubs at their air and military bases, and the part of the province delineated to the north by road No 15 leading from Khamis-Mushayt through Jarash, Al Utfah and Dhahran Al Janoub to the border with the province of Najran, and, the part of the province delineated to the north by the road leading from Al Utfah through Al Fayd to Badr Al Janoub (Province of Najran);

▼B

— Surveillance zone: the equestrian clubs at their air and military bases, the part of the province between the border of the protection zone and road No 209 from Ash-Shuqaiq to the road control post Muhayil on road No 211, the part of the province between the control post on road No 10 south of Abha, the city of Abha and the road control post Ballasmer 65 km from Abha on road No 15 leading north, the part of the province between Khamis-Mushayt and the road control post 90 km from Abha on road No 255 to Samakh and the road control post at Yarah, 90 km from Abha, on road No 10 leading to Riyadh, and, the part of the province south of a virtual line between the road control post at Yarah on road No 10 and Khashm-Ghurab on road No 177 up to the border of the province of Najran.

3. Province of Najran

- Protection zone: the part of the province delineated by the road from Al Utfah (province of Asir) to Badr Al Janoub and to As Sebt and from As Sebt along Wadi Habunah to the conjunction with road No 177 between Najran and Riyadh to the north and from this conjunction by road No 177 leading south to the conjunction with road No 15 from Najran to Sharourah, and, the part of the province south of road No 15 between Najran and Sharourah and the border with the Yemen.
- Surveillance zone: the part of the province south of a line between the road control post at Yarah, on road No 10, and Khashm-Ghurab, on road No 177, from the border of the province of Najran until the road control post at Khashm-Ghurab, 80 km from Najran, and west of road No 175 leading to Sharourah.

BOX 4

ZA	South Africa	ZA-1	Approved Quarantine s	Quarantine stations:		
			1. Kenilworth Quarantine Station			
			Delimitation of the Metropolitan area of Cape-Town (ZA-1):			
			Northern boundary:	Blaauwberg Road (M14);		
			Eastern boundary:	Koeberg Road (M14), Plattekloof Road (M14), N7 Highway, N1 Highway and M5 Highway,		
			Southern boundary:	Ottery Road, Prince George's Drive, Wetton Road, Riverstone Road, Tennant Road, Newlands Drive, Paradise Road, Union Drive, Rhodes Drive up to the Newslands Forestry station and across Echo Gorge of Table Mountain to Camps Bay;		
			Western boundary:	Coastline from Camps Bay to Blaauwberg Road.		

ANNEX II

MODEL HEALTH CERTIFICATES AND MODEL DECLARATIONS FOR THE ENTRY INTO THE UNION OF LIVE EQUIDAE

PART 1

Temporary admission and transit

$Section \ A$

Model health certificate and model declaration for the temporary admission of registered horses into the Union for a period of less than 90 days

NTRY:									V	eterinary certific	ate to E
l.1.	Consignor Name					1.2.	Certificate referer	nce No	1.2	2.a.	
	Address					1.3.	Central competer	nt authority			
	Tel.						Local competent	authority			
1.5.	Consignee Name Address					1.6.					
	Postcode Tel.										
I.5.	Country of origin	ISO code	I.8.	Region of origin	Code	1.9.	Country of destination	ISO code	I.10.	Region of destination	Code
l.11.	Place of origin					I.12.	Place of destinati	on			ı
	Name Approval number Address						Name Address				
							Postcode				
I.13.	Place of loading					I.14. Date of departure					
I.15.	Means of transport					I.16.	Entry BIP in EU				
	Aeroplane ☐ _ Ship ☐ Railway wagon ☐										
	Road vehicle Other Identification Documentary references						No(s) of CITES				
I.18.	18. Description of animal						I.19. Comm	odity	code (HS code) 01 01		
									1.20	. Quantity 1	
I.21.									1.22	. Number of pac	kages
1.23.	I.23. Seal/Container No						1.24				
1.25.	I.25. Animal certified for:										
	Registered horse										
1.26.	1.26.						I.27. For import or	admission in	ito EU		
1.28.	. Identification of the animal										
	Species (Scientific Identification system Ident name) Equus caballus				ificatio	on number	Age		Sex		

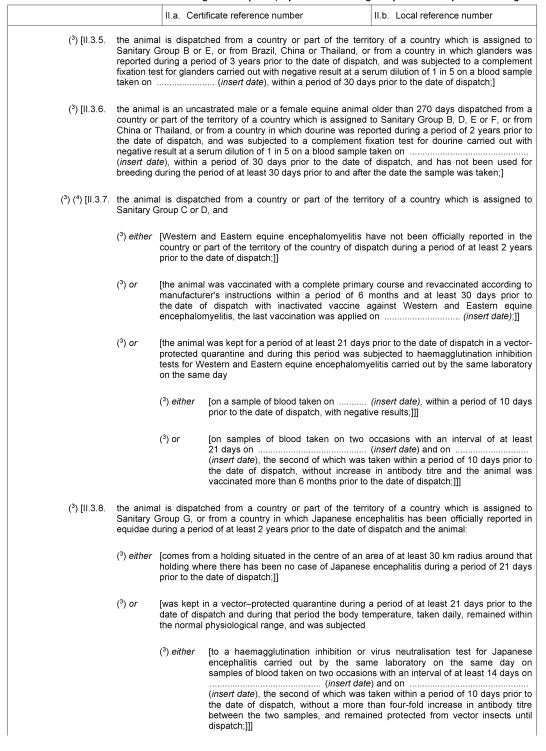
			II.a. Certificate reference number	II.b. Local reference number							
	II.	Attestation of anima	l health and welfare								
	I, the undersigned official veterinarian, hereby certify, that the animal described in Box I.28:										
	_	is a registered horse as defined in Article 2(c) of Commission Implementing Regulation (EU) 2018/659;									
	_	was examined today infestation;	y (1) and found free of clinical signs of disc	eases and of obvious signs of ectoparasite							
	_	is not intended for sla	nughter under a national programme of infectiou	is or contagious disease eradication;							
	_	meets the requirements attested in points II.1 to II.5 of this certificate;									
	_	is accompanied by the written declaration, signed by the owner of the animal or the representative of the owner.									
	II.1.	Attestation on third country or part of the territory of third country and holding of dispatch									
ation	II.1.1.	The animal is dispatched from									
Part II: Certification	II.1.2.	in the country of dispatch the following diseases are compulsorily notifiable: African horse sickness, dourine (<i>Trypanosoma equiperdum</i>), glanders (<i>Burkholderia mallei</i>), equine encephalomyelitis (of all types including Venezuelan equine encephalomyelitis), equine infectious anaemia, vesicular stomatitis, rabies and anthrax;									
Par	II.1.3.	the animal is dispatched from a country or part of the territory of a country:									
		in which to	considered free from African horse sickness in there has been no clinical, serological (in unvact horse sickness during the period of 2 years per no vaccinations against the disease during	ccinated equidae) or epidemiological evidence rior to the date of dispatch and in which there							
			Venezuelan equine encephalomyelitis has not of dispatch;	occurred during the period of 2 years prior to							
c) in which dourine has not occurred during the period of 6				months prior to the date of dispatch;							
		d) in which (glanders has not occurred during the period of 6	6 months prior to the date of dispatch;							
	(³) either	(3) either [e) in which vesicular stomatitis has not occurred during the period of 6 months prior dispatch;]									
	(³) or	and a blo	vesicular stomatitis has occurred during the pe od sample taken from the animal on prior to the date of dispatch, was tested with virus	(insert date), within a period of							
		(³) either	[in a virus neutralisation test at a serum dilution	n of 1 in 32;]]							
		(³) or	[in an ELISA in accordance with the relevan and Vaccines for Terrestrial Animals of the OI								
	II.1.4.	points II.1.4.1 to II.1.	come from a holding and to the best of my k 4.7 was not in contact with animals from holdi ed to in points II.1.4.1 to II.1.4.7 and which last	ngs, which were subject to prohibition orders							

				II.b. Local reference number
			II.a. Certificate reference number	II.b. Local reference number
	II.1.4.1.	in the cas	se of equidae suspected of having contracted dour	ine,
		(³) either	[6 months beginning on the date of the last a suspected of having contracted dourine or infected	
		(³) or	[in the case of a stallion, until the animal is castra	ated;]
		(³) or	[30 days following the date of completion of the after all animals of susceptible species have bee	
	II.1.4.2.	in the cas	se of glanders,	
		(³) either	[6 months beginning on the day on which th subjected with positive results to a test for Burkholderia mallei or antibodies to that pathoge	the detection of the causative pathogen
		(³) or	[30 days following the date of completion of the after all animals of susceptible species have bee	
	II.1.4.3.	in the cas	se of equine encephalomyelitis of any type,	
		(³) either	[6 months beginning on the day on which the equal slaughtered;]	uidae suffering from the disease have been
		(³) or	[6 months beginning on the day on which the eq Nile Fever, Eastern equine encephalomyelitis of died, been removed from the holding or fully reco	or Western equine encephalomyelitis have
		(³) or	[30 days following the date of completion of the after all animals of susceptible species have bee	
	II.1.4.4.	slaughter gel immi	se of equine infectious anaemia, until the date or led, the remaining equine animals on the holding lunodiffusion test (AGID or Coggins test) carrisions 3 months apart;	have shown a negative reaction in an agar
	II.1.4.5.	in the cas	se of vesicular stomatitis,	
		(³) either	[6 months following the last case;]	
		(³) or	[30 days following the date of completion of the after all animals of susceptible species have bee	
	II.1.4.6.		se of rabies, 30 days following the last case and to on of the premises;	he date of completion of the cleansing and
	II.1.4.7.		se of anthrax, 15 days following the last case and ton of the premises;	the date of completion of the cleansing and
II.1.5.			wledge, during the period of 15 days prior to the c infected or suspected of an infectious or contagion	
II.2.	Attestation	n of residen	ce and pre-export isolation	
(³) either	[II.2.1.	holdings	period of at least 40 days prior to the date of c under veterinary supervision situated in the coun which is assigned to Sanitary Group A, B, C, D, E	try or part of the territory of the country of

		rogiciorea equidad, equidad for steeding and production equidad for claugh
		II.a. Certificate reference number II.b. Local reference number
	(³) either	[in a Member State of the Union;]]
	(³) or	(in a country or part of the territory of a country with Code:
		(3) either [assigned to the same Sanitary Group
		(³) and/or [assigned to Sanitary Group A, B or C;]]]
		(³) andlor [China (⁵), Hong Kong, Japan, Korea, Macao, Malaysia (Peninsula), Singapore, Thailand of the United Arab Emirates;]]]
(³) (⁴) or	[II.2.1.	During a period of at least 60 days prior to the date of dispatch, the animal has been resident of holdings under veterinary supervision situated in the country or part of the territory of the country dispatch which is assigned to Sanitary Group F, or was imported during the 60 days prior to the day of dispatch from a Member State of the Union before entering the vector–protected or vector programatine station in accordance with point II.2.2;]
(³) (⁴) either	[II.2.2.	the animal is dispatched from a country or part of the territory of a country which is assigned to Sanitary Group E and
	(³) either	[has been kept in isolation in the country or part of the territory of the country of dispatch protecte from vector insects for a period of at least 40 days prior to the date of dispatch, or since entry into the country or part of the territory of the country of dispatch, if it was imported in accordance will point II.2.1 from a Member State of the Union or a country or part of the territory of a country which assigned to Sanitary Group A, B, C, D, E or G;]]
	(³) or	[has been kept in designated premises under official veterinary supervision for a period of at lea 40 days prior to the date of dispatch, or since entry into the country or part of the territory of the country of dispatch, if it was imported in accordance with point II.2.1 from a Member State of the Union or a country or part of the territory of a country which is assigned to Sanitary Group A, B, C, I E or G, and the country or part of the territory of the country of dispatch is recognised by the OIE a officially free of African horse sickness and is not adjacent to a country in which African horse sickness has occurred during the period of 2 years prior to the date of dispatch;]]
(³) (4) or	[II.2.2.	the animal is dispatched from a country or part of the territory of country which is assigned to Sanital Group F and was kept:
	(³) either	(insert name of quarantine station) during at least the last 40 days prior to the date of dispate from
	(³) or	[permanently confined in the approved vector-proof quarantine station of
II.3.	Attestation	of vaccination and health tests
(³) either	[II.3.1.	The animal was not vaccinated against African horse sickness in the country of dispatch and there no information suggesting previous vaccination;]

EUROPEAN UNION	Registered equidae, equidae for breeding and production equidae for slaughte									
	II.a. Certificate reference number II.b. Local reference number									
(³) or [II.3.1.	The animal was vaccinated against African horse sickness, and this vaccination was carried ou									
(²) either	[more than 12 months prior to the date of dispatch;]]									
(²) or	[more than 60 days and less than 12 months prior to the date of admission into the part of the territory of the country referred to in point II.1.3.(a), from where it is dispatched;]]									
(³) (⁴) or [II.3.1.	The animal is dispatched from a country or part of the territory of a country which is assigned to Sanitary Group F and was vaccinated against African horse sickness on									
II.3.2.	the animal was not vaccinated against Venezuelan equine encephalomyelitis during the period of 60 days prior to the date of dispatch from									
(³) either	[a country of which all parts of the territory are free of Venezuelan equine encephalomyelitis for a period of at least 2 years prior to the date of dispatch;]									
(³) (⁴) or	[a part of the territory of a country which is assigned to Sanitary Group C or D, which is free of Venezuelan equine encephalomyelitis for a period of at least 2 years prior to the date of dispatch and Venezuelan equine encephalomyelitis occurs in the remaining parts of the territory of the country of dispatch, and									
	(3) either [is vaccinated against Venezuelan equine encephalomyelitis with a complete primary course and revaccinated according to manufacturer's recommendations not less than 60 days and no more than 12 months prior to the date of dispatch, and was kept in vector-protected quarantine for a period of at least 21 days prior to the date of dispatch, and during that period remained clinically healthy, and its body temperature, taken daily, remained within the normal physiological range, and any equine animal on the same holding which showed a rise in body temperature, taken daily, was subjected to a blood test for virus isolation for Venezuelan equine encephalomyelitis with negative results;]]									
	(3) or [is not vaccinated against Venezuelan equine encephalomyelitis and was kept in vector-protected quarantine for a period of at least 21 days, and during that period remained clinically healthy, and its body temperature, taken daily, remained within the normal physiological range, and any equine animal on the same holding which showed a rise in body temperature, taken daily, was subjected to a blood test for virus isolation for Venezuelan equine encephalomyelitis with negative results, and the animal to be dispatched was subjected to a diagnostic test for Venezuelan equine encephalomyelitis with negative result conducted on a sample taken not less than 14 days after the date of entry into of the vector-protected quarantine and remained protected from vector insects until dispatch;]]									
	(3) or [was subjected to a haemagglutination inhibition test for Venezuelan equine encephalomyelitis carried out by the same laboratory on the same day on samples taken on two occasions with an interval of 21 days on									
(³) [II.3.3.	the animal is an uncastrated male equine animal older than 180 days, and									
(³) either	[is dispatched from a country in which equine viral arteritis (EVA) is a compulsorily notifiable disease and has not been officially reported during the period of 6 months prior to the date of dispatch;]]									

			ragiotoroa oquidae, equidae for bree	aning and production equidae for slaught
			II.a. Certificate reference number	II.b. Local reference number
	(³) or		ed on a blood sample taken on(insert spatch, by virus neutralisation test for EVA with r	
	(³) or	of 21 day	ed on an aliquot of its entire semen taken on s prior to the date of dispatch, by virus isolatio PCR for EVA with negative result.]]	
	(³) or	and re-va	inated against EVA on(insert of circlinated at regular intervals according to the by the competent authority, and the initial vaccin	manufacturer's instructions, with a vaccin
		(³) either	[before 31 December 2017, on the day a blootested in a virus neutralisation test for EVA of 1 in 4;]]]	·
		(³) or	[before 31 December 2017, during a period of official veterinary supervision, commencing or was tested during that isolation period in a vir result at a serum dilution of 1 in 4;]]]	n the day a blood sample was taken whic
		(³) or	[at the age of 180 to 270 days, during a part supervision, during which the animal was subcarried out with negative result at a serum dilute by the same laboratory with stable or declining 10 days apart; []]	jected to a virus neutralisation test for EV ion of 1 in 4, or carried out on the same da
		(³) or	[after the animal was subjected to a virus neutral a serum dilution of 1 in 4, carried out on a bloc commencing a period of uninterrupted isolal vaccination;]]]	od sample taken not earlier than 7 days afte
		(³) or	[at the age of 180 to 250 days, after the anima for EVA carried out with negative result at a sr same day by the same laboratory with stable or at least 14 days apart;]]]	erum dilution of 1 in 4, or carried out on th
	(³) or	carried of sample of	ected to a virus isolation test, polymerase chai ut with negative result on an aliquot of its ent that animal taken on	ire semen collected after the date a bloc within a period of 6 months prior to the da
	(³) or	legislation animal is legal act a that any	irements for testing for EVA or vaccination in	cable Union legal act) on the ground that the truction in the equestrian event specified in the equidae not participating in such event are
(³) (⁴) either	[11.3.4.	anaemia,	al is dispatched from Iceland, which is certific where it was continuously resident since birth, we entered Iceland from other countries;]	
(³) or	[II.3.4.	Coggins t	al was subjected with negative result to an est) or to an ELISA for equine infectious anaer (insert date), this being within	
		(³) either	[a period of 90 days prior to the date of dispatch	n;]]
		(³) or		ch from a country or part of the territory of



		· g 1 - 1 - 1	andac, equidae for breeding and production equidae for staughte			
	II.a. Cert	ificate refere	ence number II.b. Local reference number			
	(³) or	encephali not earlie	capture ELISA test for the detection of antibodies against Japanese tis virus with negative result, carried out on a blood sample taken than 7 days after the date the isolation commenced on			
	(³) or	and reva	cinated against Japanese encephalitis with a complete primary course ccinated according to manufacturer's recommendations during a not less than 21 days and not more than 12 months prior to the date h;]]			
	Sanitary Group E, a	al is dispatched from a country or part of the territory of a country which is assigned to Group E, and was subjected to a serological test for African horse sickness as described it to Directive 2009/156/EC, which was carried out by the same laboratory on the same day				
(1	(3) either [on blood	samples tal	ken on two occasions with an interval of between 21 and 30 days, on			
			(insert date) and on			
	(³) either	[with nega	ative results in each case.]]]			
	(³) or	[with a po	sitive result in the first sample, and			
		(³) either	[the second sample was subsequently tested with negative result in an agent identification test as described in Annex IV to Directive 2009/156/EC.]]]]			
		(³) or	[the two samples were tested without more than a two-fold increase in antibody titre in a virus neutralisation test as described in point 2.4 of Chapter 2.5.1 of the OIE Terrestrial Manual for Diagnostic Tests and Vaccines.]]]]			
(*	prior to t dispatch adjacent	ne date of oils is recognise to a country	ken on			
	he animal is dispa Sanitary Group F, ai		a country or part of the territory of a country which is assigned to			
(*	Directive day on bloonnot taker	2009/156/E ood samples (in less than	erological test for African horse sickness as described in Annex IV to C, which was carried out by the same laboratory on the same is taken on two occasions with an interval of between 21 and 30 days, sert date) and on			
	(³) either	[with nega	ative results in each case.]]]			
	(³) or	[with a po	sitive result in the first sample, and			
		(³) either	[the second sample was subsequently tested with negative result in an agent identification test as described in Annex IV to Directive 2009/156/EC.]]]]			
		(³) or	[the two samples were tested without more than a two-fold increase in antibody titre in a virus neutralisation test as described in point 2.4 of Chapter 2.5.1 of the OIE Terrestrial Manual for Diagnostic Tests and Vaccines.]]]]			

Registered equidae, equidae for breeding and production equidae for slaughter

EUROPEAN U	NION		Registered equidae, equidae for breeding	g and	production equidae for slaughter
			II.a. Certificate reference number	II.b.	Local reference number
		(³) or	[was subjected to a serological and an agent identif described in Annex IV to Directive 2009/156/EC, case on a blood sample taken on	carriec	d out with negative result in each (insert date) not less than
		(³) or	[was subjected to an agent identification test for Annex IV to Directive 2009/156/EC, carried out v taken on	vith ne t less	egative result on a blood sample than 14 days after the date of
II.4.	Attestation	n of the trar	sport conditions		
(³) (⁴) either	[II.4.1.	Sanitary Union, w	nal is dispatched from a country or part of the terr Group A, B, C, D, E or G and arrangements have lithout passing through a market, marshalling or as irith other equidae of a different health status.]	oeen n	nade to transport it directly to the
(³) (⁴) or	[II.4.1.	Sanitary quarantin	nal is dispatched from a country or part of the terr Group F and arrangements have been made to trans- ie station without coming into contact with other either for imports or for temporary admission into the	sport it equida	directly from the vector-protected e not accompanied by a health
		(²) either	[to the airport under vector-protected conditions a the aircraft being cleansed and disinfected in recognised in the third country of dispatch, and spi take off.]]	advar	nce with a disinfectant officially
		(³) or	[to a sea port in that country or part of the territor conditions and arrangements have been made scheduled directly to a port in the Union without creat of the territory of a country not approved for stalls which were cleansed and disinfected in recognised in the third country of dispatch and sprideparture.]]	to tra alling in the er advar	insport it on a vessel which is nto a port situated in a country or ntry into the Union of equidae, in nce with a disinfectant officially
	II.4.2.	with at le	nents have been made and verified to prevent any cleast the same health requirements as described in fication until dispatch to the Union.		
	II.4.3.	disinfecte	sport vehicles or containers in which the animal is to before loading with a disinfectant officially recogn so constructed that faeces, urine, litter or fodder cann	ised in	the third country of dispatch and
II.5.	Attestation	n of animal	welfare		
			d in Box I.28 was examined today (1) and found fit to ere made to protect its health and well-being effective		
Notes:					
Part I:					
Box I.8.:			the country or part of the territory of the country on Implementing Regulation (EU) 2018/659.	f dispa	atch as appearing in column 3 of
Box I.15.:	informatio		(railway wagons or container and lorries), flight rovided. In case of unloading and reloading, the cons Union.		

Box I.23.: The container number and the seal number (if applicable) should be included.

II.b. Local reference number

EUROPEAN UNION

Registered equidae, equidae for breeding and production equidae for slaughter

Identification system: The animal must bear an individual identifier which permits to link the animal to the

Box I.28.: identification document as defined in Article 2(b) of Commission Implementing Regulation (EU) 2018/659. Specify the identification system (such as ear tag, tattoo, brand, transponder etc.) and the anatomic place used on the

II.a. Certificate reference number

If a passport accompanies the animal, its number should be stated and the name of the competent authority which validated it.

Age: Date of birth (dd/mm/yyyy).

Sex (M = male, F = female, C = castrated).

Part II:

 $(^{1})$ The certificate must be issued on the day of loading or on the last working day before loading of the animal for dispatch to the Member State of destination in the Union.

The temporary admission of this registered horse shall not be allowed when the animal was loaded either prior to the date of authorisation for temporary admission into the Union from the respective country or part of the territory of the country referred to in point II.1.1, or during a period where restrictive measures have been adopted by the Union against the entry of equidae from this country or this part of the territory of the country of dispatch.

- Code of the country or part of the territory of the country and the Sanitary Group as appearing in columns 3 and 5 $(^{2})$ respectively of Annex I to Commission Implementing Regulation (EU) 2018/659.
- $(^{3})$ Delete as appropriate.
- Statements that relate entirely and exclusively to a Sanitary Group different from the Sanitary Group to which the country $(^{4})$ of dispatch, or part of its territory, is assigned, may be left out, provided that the numbering of the subsequent statements is maintained.
- Part of the territory of country authorised for temporary admission as appearing in columns 3 and 6 respectively of $(^{5})$ Annex I to Commission Implementing Regulation (EU) 2018/659.

This health certificate shall:

- be drawn up in at least a language understood by the certifying officer and one of the official languages of the Member State of destination and of the Member State where the registered horse will enter Union territory and undergo the veterinary border checks;
- be made out to a single consignee;
- accompany the registered horse in the original throughout its temporary admission in the Union: (c)
- (d) be signed and stamped in a colour different to the colour of the printing;
- consist of a single sheet of paper or all sheets of paper required are part of an integrated whole and indivisible by (e) inserting page numbers and total number of pages, and each page shall bear the certificate reference number at the top of the page and those pages are stapled and stamped.

Official veterinarian	
Name (in capital letters):	Qualification and title:
Date:	Signature:
Stamp:	

						ner or representativ dmission of a regis						
Iden	tification o	f the	animal (1)									
Spe	ecies (Scie	ntific	name)	Identification	system	Identification numb	er	Ą	ge		Se	×
	Equus ca	aball	us									
I, the	e undersigr	ned c	owner (²) or	representative	of the owner (2) of the registered ho	rse describe	ed above, I	nereby de	clare, t	that:	
_	the horse											
	(²) either				(insert name of ate of dispatch;]	country or part of the	territory of	a country	of dispatci	h) duri	ng a pe	riod of
	(²) or					f country or part of prior to the date of d		y of a co	untry of a	lispatc	h) durii	ng the
		(a)				mf country of dispatch)		name of	country	from	where	horse
		(b)				mf country of dispatch)		name of	country	from	where	horse
		(c)	on		(<i>insert date</i>) fror	mf country of dispatch)	(insert	name of	country	from	where	horse
_			iod of 15 da		date of dispatch	n the horse has not b	· -	act with an	imals suffe	ering f	rom infe	ectious
_	•	orta	tion will be		•	alth and well-being o	of the horse of	can be pro	tected effe	ectivel	y at all	stages
_	the condi	tions	for reside			as applicable in acco		point II.2	of the ac	ccomp	anying	health
_	the condi	tions	for the trai	nsport as applic	cable in accorda	ance with point II.4 of		panying he	ealth certif	icate f	or the c	ountry
_	during its	resi		•	ispatch are fulfil for a period of	less than 90 days	the horse w	vill be acc	commodate	ed on	the fol	lowing
	premises											
	` '		•	,	, ,	in	· ·	•		•		,
	(b) from		(da	ate) to	(date)	in	(place of h	olding) in		(Me	ember S	State)
	(c) from		(da	ate) to	(date)	in	(place of h	o <i>lding</i>) in		(Ме	ember S	State)
	(d) from		(da	ate) to	(date)	in	(place of ho	o <i>lding</i>) in		(Ме	ember S	State);
_	this decla	ratio	n, it must b	e accompanied	d by a health cei	one Member State on tificate issued by an State of destination;	official veter					
_				leave the Unio		(<i>date</i>) at	the border p	oost of				
				er (²) or represo (dd/mm/y								
(¹) (²)	Article 2(b transponde If a passpo Age: Date	o) of er) and ert acc of birt nale, l	Commission d the anaton companies th th (dd/mm/yy F = female,	n Implementing nic place used on ne animal, its nun	Regulation (EU) the animal.	ntifier which permits to I 2018/659. Specify the ated and the name of the	e identification	n system (such as e	ar tag,		

Section B

Model health certificate and model declaration for the transit of live equidae through the Union from one third country or part of the territory of a third country to another third country or another part of the territory of the same third country

COUN	TRY:								Veterinary	certificate to EU
	l.1.	Consignor Name				1.2.	Certificate reference No I.2.a.			
		Address				1.3.	Central competen	t authority		
		Tel.				1.4.	Local competent a	authority		
ment	1.5.	Consignee				1.6.	Person responsible	e for the load	in EU	
nsign		Name Address					Name Address			
100 p										
atche		Postcode Tel.					Postcode Tel.			
Part I : Details of dispatched consignment	1.7.	Country of ISO o	ode I.	8. Region of origin	Code	1.9.	Country of destination	ISO code	I.10. Region of destination	
: Deta	l.11.	Place of origin				1.12.				
Part		Name	Apr	proval number						
		Address								
	I.13.	Place of loading				I.14. Date of departure				
	I.15.	Means of transpor	t			I.16.	Entry BIP in EU			
		Aeroplane \square	Ship	☐ Railway w	agon 🗖					
		Road vehicle Identification	Other	· 🗆		I.17. No(s) of CITES				
		Documentary refe	rences							
	I.18.	Description of anir	nals					I.19. Commo	dity code (HS o	ode)
							l		I.20. Quantity	
	I.21.								I.22. Number	of packages
	1.23.	Seal/Container No	ı						1.24.	
	1.25.	Animals certified for	or:							
		Registered equida	е С] bree	ding and	prod	uction \square		slaughter	
	1.26.	For transit through	EU to th	ird country	Х		1.27.			
		Third country		ISO code						
	1.28.	Identification of the	e animals	i						
	s	pecies (Scientific name)	lden	tification system	Identi	fication	on number	Age		Sex

			II.a. Certificate reference number	II.b. Local reference number									
	II.	Attestation of anim	al health and welfare										
	I, the under	signed official veterina	rian, hereby certify, that the equine animal described in	Box I.28:									
	_	was examined toda infestation;	y $(^1)$ and found free of clinical signs of diseases a	and of obvious signs of ectoparasite									
ation	_	- is not intended for slaughter under a national programme of infectious or contagious disease eradication;											
ertific	_	meets the requirements attested in points II.1 to II.5 of this certificate;											
Part II: Certification	_	is accompanied by the	ne written declaration, signed by the owner of the anima	al or the representative of the owner.									
ă	II.1.	Attestation on third o	country or part of the territory of third country and holdin	g of dispatch									
	II.1.1.	1. The animal is dispatched from											
	II.1.2.	(Trypanosoma equi	spatch the following diseases are compulsorily notificoerdum), glanders (Burkholderia mallei), equine encencephalomyelitis), equine infectious anaemia, vesicula	ephalomyelitis (of all types including									
	II.1.3.	the animal is dispato	hed from a country or part of the territory of a country										
		in which of Africa	considered free from African horse sickness in according there has been no clinical, serological (in unvaccinated in horse sickness during the period of 2 years prior to the no vaccinations against the disease during the period of 2 years prior to the no vaccinations against the disease during the period.	d equidae) or epidemiological evidence he date of dispatch and in which there									
		,	Venezuelan equine encephalomyelitis has not occurre of dispatch;	ed during the period of 2 years prior to									
		c) in which	dourine has not occurred during the period of 6 months	s prior to the date of dispatch;									
		d) in which	glanders has not occurred during the period of 6 month	ns prior to the date of dispatch;									
	(3) either	[e) in which dispatch	vesicular stomatitis has not occurred during the per:]	riod of 6 months prior to the date of									
	(³) or	and a blo	vesicular stomatitis has occurred during the period of bod sample taken from the animal on	(insert date), within a period of									
		(³) either	[in a virus neutralisation test at a serum dilution of 1 i	n 32;]]									
		(³) or	[in an ELISA in accordance with the relevant Chap and Vaccines for Terrestrial Animals of the OIE;]]	ter of the Manual of Diagnostic Tests									
	II.1.4.	points II.1.4.1 to II.1	come from a holding, and to the best of my knowled. 4.7 was not in contact with animals from holdings, where to in points II.1.4.1 to II.1.4.7 and which last for:										

			registered equidae, equidae for breeding e	p				
			II.a. Certificate reference number	II.b. Local reference number				
	II.1.4.1.	in the cas	e of equidae suspected of having contracted dourine,					
		(³) either	either [6 months beginning on the date of the last actual or possible contact with an anin suspected of having contracted dourine or infected with Trypanosoma equiperdum;]					
		(³) or	[in the case of a stallion, until the animal is castrated;]					
		(³) or	[30 days following the date of completion of the clear after all animals of susceptible species have been sla					
	II.1.4.2.	in the cas	e of glanders,					
		(³) either	[6 months beginning on the day on which the eq subjected with positive result to a test for the of Burkholderia mallei or antibodies to that pathogen, we	detection of the causative pathogen				
		(³) or	[30 days following the date of completion of the clear after all animals of susceptible species have been killed					
	II.1.4.3.	in the cas	e of equine encephalomyelitis of any type,					
		(³) either	[6 months beginning on the day on which the equidae slaughtered;]	e suffering from the disease have been				
		(³) or	[6 months beginning on the day on which the equidae infected with the virus causi Nile Fever, Eastern equine encephalomyelitis or Western equine encephalomyel died, been removed from the holding or fully recovered;]					
		(³) or	[30 days following the date of completion of the clear after all animals of susceptible species have been sla					
	II.1.4.4.	slaughter	se of equine infectious anaemia, until the date on whed, the remaining animals on the holding have show ffusion test (AGID or Coggins test) carried out on bloo apart;	n a negative reaction in an agar gel				
	II.1.4.5.	in the cas	e of vesicular stomatitis,					
		(3) either	[6 months following the last case;]					
		(³) or	[30 days following the date of completion of the clear after all animals of susceptible species have been sla					
	II.1.4.6.		e of rabies, 30 days following the last case and the do on of the premises;	ate of completion of the cleansing and				
	II.1.4.7.		e of anthrax, 15 days following the last case and the d on of the premises;	ate of completion of the cleansing and				
II.1.5.			wledge, during the period of 15 days prior to the date infected or suspected of an infectious or contagious dis					
II.2.	Attestation	of residen	ce and pre-export isolation					
(³) either	[II.2.1.	holdings	period of at least 40 days prior to the date of disparunder veterinary supervision situated in a country owhich is assigned to Sanitary Group A, B, C, D, E or G	r part of the territory of a country of				

			II.a. Certificate reference number	II.b. Local reference number					
	(³) either	[in a Member State of the Union;]]							
	(³) and/or	for tempora country or required in	y or part of the territory of country with Code:	d from which it was imported into the conditions at least as strict as those orary admission of registered horses					
			assigned to the same Sanitary Grouphe territory of the country of dispatch;]]]	(2) as the country or part of					
		(3) and/or [assigned to Sanitary Group A, B or C;]]]						
			assigned to Sanitary Group D, E or G and the anim Article 2(c) of Commission Implementing Regulation (I						
(³) (⁴) or	[II.2.1.	holdings undispatch wordispatch	During a period of at least 60 days prior to the date of dispatch, the animal has been resident on noldings under veterinary supervision situated in a country or part of the territory of a country of dispatch which is assigned to Sanitary Group F, or was imported during the 60 days prior to the date of dispatch from a Member State of the Union before entering the vector-protected or vector proof quarantine station in accordance with point II.2.2;]						
(³) (⁴) either	[II.2.2.	the animal Sanitary G	is dispatched from a country or part of the territor roup E and	y of a country which is assigned to					
	(³) either	from vector country or point II.2.1	kept in isolation in the country or part of the territory insects for a period of at least 40 days prior to the d part of the territory of the country of dispatch, if from a Member State of the Union or a country or pa	ate of dispatch, or since entry into the it was imported in accordance with					
	(³) or	40 days procountry of Union or a E or G, and officially from	kept in designated premises under official veterinar ior to the date of dispatch, or since entry into the dispatch, if it was imported in accordance with poir country or part of the territory of a country which is a d the country or part of the territory of the country of se of African horse sickness and is not adjacent as occurred during the period of 2 years prior to the data.	country or part of the territory of the tt II.2.1 from a Member State of the seigned to Sanitary Group A, B, C, D, dispatch is recognised by the OIE as to a country in which African horse					
(³) (⁴) or	[II.2.2.		is dispatched from a country or part of the territor oup F and was kept	y of a country which is assigned to					
	(³) either	to sunset ufollowing the Culicoides prepared for	roved vector-protected quarantine station of	th from(insert date) to oremises at least from two hours prior d under official veterinary supervision, with an insecticide effective against ict isolation from equidae not being					
	(³) or	of quaranti monitoring	tly confined in the approved vector-proof quarantine sine station) during the period of at least 14 days prior of the vector protection has proven absence of vector station.]	to the date of dispatch and constant					
II.3.	Attestation	of vaccinati	on and health tests						
(³) either	[II.3.1.		was not vaccinated against African horse sickness i ion suggesting previous vaccination;]	n the country of dispatch and there is					

			Trogistorea equidae, equidae for brocaring t	The production equidue for slaughter					
			II.a. Certificate reference number	II.b. Local reference number					
(³) or	[II.3.1.	The anima	ne animal was vaccinated against African horse sickness, and this vaccination was carried out						
	(³) either	[more thar	n 12 months prior to the date of dispatch;]]						
	(³) or		n 60 days and less than 12 months prior to the date of ntry referred to in point II.1.3.(a), from where it is dispa						
(³) (⁴) or	[II.3.1.	Sanitary G (insert dat protected	The animal is dispatched from a country or part of the territory of a country which is assigned to Sanitary Group F and was vaccinated against African horse sickness on						
	II.3.2.		I was not vaccinated against Venezuelan equine e ior to the date of dispatch from	ncephalomyelitis during the period of					
	(³) either		of which all parts of the territory are free of Venez it least 2 years prior to the date of dispatch;]	zuelan equine encephalomyelitis for a					
	(³) (⁴) or	Venezuela	the territory of a country which is assigned to Sar in equine encephalomyelitis for a period of at least 2 in equine encephalomyelitis occurs in the remaining and	years prior to the date of dispatch and					
			[is vaccinated against Venezuelan equine encept course and revaccinated according to manufactur 60 days and no more than 12 months prior to the da protected quarantine for a period of at least 21 daduring that period remained clinically healthy, and remained within the normal physiological range, a holding which showed a rise in body temperature, tak for virus isolation for Venezuelan equine encephalom	rer's recommendations not less than te of dispatch, and was kept in vectorays prior to the date of dispatch, and its body temperature, taken daily, and any equine animal on the same ten daily, was subjected to a blood test					
			[is not vaccinated against Venezuelan equine ence protected quarantine for a period of at least 21 da clinically healthy, and its body temperature, taken physiological range, and any equine animal on the body temperature, taken daily, was subjected to Venezuelan equine encephalomyelitis with negatispatched was subjected to a diagnostic test for with negative result conducted on a sample taken nentry into vector-protected quarantine and remaine dispatch;]]	ays, and during that period remained in daily, remained within the normal same holding which showed a rise in a blood test for virus isolation for tive results, and the animal to be Venezuelan equine encephalomyelitis not less than 14 days after the date of					
		· /	[was subjected to a haemagglutination inhibited encephalomyelitis carried out by the same laborator on two occasions with an interval of 21 days on	y on the same day on samples taken (insert date) and on was taken during a period of 10 days in the antibody titre, and a RT-PCR test for the detection of Venezuelan with negative result on a sample taken (insert date), and noment of the RT-PCR sampling until issect repellents and insecticides on the					
(³) (⁴) either	[11.3.3.	anaemia,	I is dispatched from Iceland, which is certified as where it was continuously resident since birth and de entered Iceland from other countries;]						

				. , , ,	· · · · · · · · · · · · · · · · · · ·					
			II.a. Ce	rtificate reference number	II.b. Local reference number					
(³) or	[II.3.3.	Coggins t	est) or to a	bjected with negative result to an agar an ELISA for equine infectious anaemia ca <i>ert date),</i> this being within						
		(³) either	[a period o	of 90 days prior to the date of dispatch;]]						
		(³) or		a period of 30 days prior to the date of dispatch from a country or part of the territory of a ountry which is assigned to Sanitary Group D, E or F;]]						
	(³) [II.3.4.	Sanitary (reported of fixation te	Group B or during a pe st for glan	is dispatched from a country or part of the territory of a country which is assigned to roup B or E, or from Brazil, China or Thailand, or from a country in which glanders was uring a period of 3 years prior to the date of dispatch, and was subjected to a complement of the forglanders carried out with negative result at a serum dilution of 1 in 5 on a blood en on						
	(³) (⁴) [II.3.5.		al is dispat Group C or	ched from a country or part of the territo D, and	ory of a country which is assigned to					
		(³) either	country or	Nestern and Eastern equine encephalomyelitis have not been officially reported in th ountry or part of the territory of the country of dispatch during a period of at least 2 year rior to the date of dispatch;]]						
		(³) or	[the animal was vaccinated with a complete primary course and revaccinated acc manufacturer's instructions within a period of 6 months and at least 30 days pr date of dispatch with inactivated vaccine against Western and Eastern encephalomyelitis, the last vaccination was applied on (insert of							
		(³) or	[the animal was kept for a period of at least 21 days prior to the date of dispatch in a protected quarantine and during this period was subjected to haemagglutination in tests for Western and Eastern equine encephalomyelitis carried out by the same lab on the same day							
			(³) either	[on a sample of blood taken onperiod of 10 days prior to the date of disp						
			(³) or	[on samples of blood taken on two of 21 days on	e) and on (insert date), period of 10 days prior to the date of e and the animal was vaccinated more					
	(³) [II.3.6.	Sanitary (Group G, o	ched from a country or part of the territo or from a country in which Japanese ence iod of at least 2 years prior to the date of di	phalitis has been officially reported in					
		(³) either	holding wh	om a holding situated in the centre of an are here there has been no case of Japanese e e date of dispatch;]]						
		(³) or	date of dis	in a vector-protected quarantine during a spatch, and during that period the body tem I physiological range, and was subjected						
			(³) either	[to a haemagglutination inhibition Japanese encephalitis carried out by the samples of blood taken on two occasions	with an interval of at least 14 days on (insert date), the eriod of 10 days prior to the date of increase in antibody titre between the					
1										

	,	9.010.00.00	m.a.a., = q.a.a.a	nd production equidae for slaughte	
	II.a. Cer	tificate refe	rence number	II.b. Local reference number	
	(³) or	Japanese sample ta	encephalitis virus with nega ken not earlier than 7 days after	he detection of antibodies against tive result, carried out on a blood the date the isolation commenced on d remained protected from vector	
	revaccinate	ed accordin		th a complete primary course and ations during a period of not less than e of dispatch;]]	
Sanitary G	al is dispatched from a country or part of the territory of a country which is assigned Group E and was subjected to a serological test for African horse sickness as described to Directive 2009/156/EC, which was carried out by the same laboratory on the same day				
		(ir		terval of between 21 and 30 days, on (insert date), the second of which of dispatch	
	(³) either	[with nega	ative results in each case.]]]		
	(³) or	[with a po	sitive result in the first sample, a	nd	
		(³) either		equently tested with negative result in s described in Annex IV to Directive	
		(³) or	in antibody titre in a virus	without more than a two-fold increase neutralisation test as described in of the OIE Terrestrial Manual for s.]]]]	
	prior to the dispatch is adjacent to	e date of or recognise a country of	dispatch, and the country or p d by the OIE as officially free	nsert date), within a period of 21 days lart of the territory of the country of of African horse sickness and is not by in which African horse sickness has of dispatch.]]]	
	I is dispato Group F, and		a country or part of the territor	ry of a country which is assigned to	
,	[was subjected to a serological test for African horse sickness as described in Annex IV to Directive 2009/156/EC, which was carried out by the same laboratory on the same day on blood samples taken on two occasions with an interval of between 21 and 30 days, on				
	(³) either	[with nega	ative results in each case.]]]		
	(³) or	[with a po	sitive result in the first sample, a	nd	
		(³) either		equently tested with negative result in s described in Annex IV to Directive	
		(³) or	in antibody titre in a virus	without more than a two-fold increase neutralisation test as described in of the OIE Terrestrial Manual for §.]]]]	

EUROPEAN U	INION		Registered equidae, equidae for breeding a	ind production equidae for slaughter
			II.a. Certificate reference number	II.b. Local reference number
		(³) or	[was subjected to a serological and an agent identifias described in Annex IV to Directive 2009/156/EC, case on a blood sample taken on	carried out with negative result in each (insert date) not less than 28 days
		(³) or	[was subjected to an agent identification test for Af Annex IV to Directive 2009/156/EC, carried out wit taken on(insert date) not less than into the vector-protected quarantine and not more tha	h negative result on a blood sample 14 days after the date of introduction
II.4.	Attestation	of the trans	sport conditions	
(³) (⁴) either	[II.4.1.	Sanitary C Union, wit	al is dispatched from a country or part of the territo Group A, B, C, D, E or G and arrangements have be shout passing through a market, marshalling or asse th other equidae of a different health status.]	en made to transport it directly to the
(³) (⁴) or	[II.4.1.	Sanitary G quarantine	al is dispatched from a country or part of the territo Group F and arrangements have been made to transpe e station without coming into contact with other eq either for imports or for temporary admission into the l	ort it directly from the vector-protected uidae not accompanied by a health
		(³) either	[to the airport under vector-protected conditions a disinfected in advance with a disinfectant officially dispatch, and sprayed against vector insects just prior	recognised in the third country of
		(³) or	[to a sea port in that country or part of the territory conditions and arrangements have been made to scheduled directly to a port in the Union without callipart of the territory of a country not approved for the stalls which were cleansed and disinfected in a recognised in the third country of dispatch and spray departure.]	o transport it on a vessel which is ing into a port situated in a country or ne entry into the Union of equidae, in dvance with a disinfectant officially
	II.4.2.	with at lea	ents have been made and verified to prevent any con ast the same health requirements as described in th ication until dispatch to the Union.	
	II.4.3.	disinfected	port vehicles or containers in which the animal is g d before loading with a disinfectant officially recognise o constructed that faeces, urine, litter or fodder cannot	ed in the third country of dispatch and
	11.4.4.	outside th	e animal is proceeding to e <i>Union</i>). Arrangements have been made and the e ensure that the animal transits the Union without dela	necessary animal health conditions
II.5.	Attestation	of animal v	velfare	
			in Box I.28 was examined today (1) and found fit to be re made to protect its health and well-being effectively	
Notes:				
Part I:				
Box I.6.:	Person res	sponsible fo	r the load in Union.	

II.b. Local reference number

EUROPEAN UNION

Registered equidae, equidae for breeding and production equidae for slaughter

Provide the code of the country or part of the territory of the country of dispatch as appearing in column 3 of

Box I.8.: Provide the code of the country or part of the territory of the country of dispatch as appearing in column 3 of Annex I to Commission Implementing Regulation (EU) 2018/659.

II.a. Certificate reference number

Box I.15.: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) and information is to be provided. In case of unloading and reloading, the consignor must inform the Border Inspection Post of entry into the Union.

Box I.23.: The container number and the seal number (if applicable) should be included.

Box I.28.: Species: Select amongst: Equus caballus, Equus asinus, Equus africanus, Equus hemionus, Equus kiang, Equus quagga, Equus zebra, Equus grevyi, or indicate any cross between those

Identification system: The animal must bear an individual identifier which permits to link the animal to the identification document as defined in Article 2(b) of Commission Implementing Regulation (EU) 2018/659. Specify the identification system (such as ear tag, tattoo, brand, transponder) and the anatomic place used on the animal.

If a passport accompanies the animal, its number should be stated and the name of the competent authority which validated it.

Age: Date of birth (dd/mm/yyyy).

Sex (M = male, F = female, C = castrated).

Part II:

(¹) The certificate must be issued on the day of loading or in the case of a registered horse on the last working day before loading of the animal for dispatch to the Member State of destination in the Union.

The entry into the Union of these animals shall not be allowed when the animals were loaded either prior to the date of authorisation for transit through the Union from the respective country or part of the territory of the country referred to in point II.1.1, or during a period where restrictive measures have been adopted by the Union against the entry of equidae from this country or this part of the territory of the country of dispatch.

- (2) Code of the country or part of the territory of the country of dispatch, and the Sanitary Group as appearing in columns 3 and 5 of Annex I to Commission Implementing Regulation (EU) 2018/659.
- (3) Delete as appropriate.
- (4) Statements that relate entirely and exclusively to a Sanitary Group different from the Sanitary Group to which the country of dispatch, or part of its territory, is assigned, may be left out, provided that the numbering of the subsequent statements is maintained.

This health certificate shall:

- (a) be drawn up in at least a language understood by the certifying officer and one of the official languages of the Member State of destination and of the Member State where the animal will enter the Union territory and undergo the veterinary border checks:
- (b) be made out to a single consignee;
- (c) be signed and stamped in a colour different to the colour of the printing;
- (d) consist of a single sheet of paper or all sheets of paper required are part of an integrated whole and indivisible by inserting page numbers and total number of pages, and each page shall bear the certificate reference number at the top of the page and those pages are stapled and stamped.

Official veterinarian	
Name (in capital letters):	Qualification and title:
Date:	Signature:
Stamp:	

	Declaration by the owner or representative of the owner for transit through the Union of an equine animal									
Ider	Identification of the animal (¹)									
Sp	Species (Scientific name) Identification system Identification number Age Sex									
I, th	e undersig	ned c	owner (²) o	representative of	the owner (2) of the animal describe	ed above, hereby d	eclare, that:		
_	the anima	al								
	(²) either			in (ins		country or part of the te	erritory of a country	of dispatch) during	a period of	
	(²) or					f country or part of the prior to the date of disp		ountry of dispatch)	during the	
		(a)				mf country of dispatch)	(insert name o	f country from wh	ere animal	
		(b)				mf country of dispatch)	(insert name o	f country from wh	ere animal	
		(c)	on entered	(ins country or part of t	sert date) fror he territory o	mf country of dispatch);]	(insert name o	f country from wh	ere animal	
				days prior to the diseases transmis		patch the animal has lae;	not been in conta	ct with animals sut	ffering from	
						as applicable in accord ountry of dispatch are fu		2 of the accompan	ying health	
_				nsport as applicat he country of disp		ance with point II.4 of the led;	ne accompanying h	nealth certificate for	the country	
_	the trans stages of			e effected in such	ı a way that	health and well-being	of the animal can	be protected effect	ctively at all	
				o leave the Union f border post of ex		(insert da	ate) at the border p	ost of		
Nan	ne and add	Iress	of the own	er (²) or represent	ative (²):					
Date	ə:			(dd/mm/yyyy	")					
(1)				quus caballus, Equu between those.	ıs asinus, Equ	us africanus, Equus hemi	onus, Equus kiang, E	quus quagga, Equus :	zebra, Equus	
	Identification	on sy.	stem: The a	nimal must bear an	gulation (EU)	ntifier which permits to link 2018/659. Specify the i				
	If a passpo Age: Date	ort acc	companies t th (dd/mm/y	ne animal, its numbe		ated and the name of the o	competent authority w	hich validated it.		
(2)	Delete as			o castratou).						

PART 2

Re-entry after temporary export

Section A

Model health certificate and model declaration for the re-entry into the Union of registered horses for racing, competition and cultural events after temporary export for a period of less than 30 days

COUN	OUNTRY: Veterinary certificate to EU								
	l.1.	Consignor Name	Certificate reference No	I.2.a.					
		Address	Central competent author	prity					
ŧ		Tel.	Local competent authori	ty					
gnme	1.5.	Consignee Name							
consi		Address	_						
atched		Postcode Tel.							
Part I : Details of dispatched consignment	1.7.	Country of ISO code I.8. Region of origin Code	Country of ISO destination	code I.10. Region of Code destination					
Detai	l.11.	Place of origin	. Place of destination						
Part I:		Name Approval number Address	Name Address						
			Postcode						
	I.13.	Place of loading	. Date of departure						
	I.15.	Means of transport	. Entry BIP in EU						
		Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification Documentary references	. No(s) of CITES						
	I.18.	Description of animal	I.19. (Commodity code (HS code) 01 01					
				I.20. Quantity					
	I.21.			I.22. Number of packages					
	1.23.	Seal/Container No		1.24.					
	1.25.	Animal certified for:							
		Registered horse	,						
	1.26.		I.27. For import or admis	sion into EU					
	1.28.	Identification of the animal							
	Sı		ion number Aç	ge Sex					

					ina production equidae for claughter				
				II.a. Certificate reference number	II.b. Local reference number				
	II.	Attestation of a	anima	al health and welfare					
	I the under	signed official vete	arinari	an, hereby certify, that the animal described in Box I.28	3.				
	i, the unders	ngried official vete	ziiiaii	an, nereby certify, that the animal described in Box 1.20	J.				
	_	is a registered h	norse	as defined in Article 2(c) of Commission Implementing	Regulation (EU) 2018/659;				
uo	_	was examined today (1) and found free of clinical signs of diseases and of obvious signs of ectoparasite infestation;							
tificat	_	is not intended for slaughter under a national programme of infectious or contagious disease eradication;							
Part II: Certification	_	meets the requir	remer	nts attested in points II.1 to II.3 of this certificate;					
Part	_	is accompanied	by th	e written declaration, signed by the owner of the horse	or the representative of the owner.				
	II.1.	Attestation on th	hird co	ountry or part of the territory of third country and holding	g of dispatch				
	II.1.1.	or part of the ter	rritory	thed from(insert name of country or part of a country which on the date of issuing this certificate ry Group					
	II.1.2.	(Trypanosoma	equip	spatch the following diseases are compulsorily notificated berdum), glanders (Burkholderia mallei), equine enconcephalomyelitis), equine infectious anaemia, vesicula	ephalomyelitis (of all types including				
	II.1.3.	the animal is dis	spatch	ned from a country or part of the territory of a country:					
		, Afric have	ch the	considered free from African horse sickness in accorda ere has been no clinical, serological (in unvaccinated et orse sickness during the period of 2 years prior to the en no vaccinations against the disease during the per	quidae) or epidemiological evidence of e date of dispatch and in which there				
				Venezuelan equine encephalomyelitis has not occurre of dispatch;	d during the period of 2 years prior to				
		c) in w	hich o	dourine has not occurred during the period of 6 months	prior to the date of dispatch;				
		d) in w	hich (glanders has not occurred during the period of 6 month	s prior to the date of dispatch;				
	II.1.4.	points II.1.4.1 to	II.1.4	come from a holding, and to the best of my knowled 4.7 was not in contact with animals from holdings, which to in points II.1.4.1 to II.1.4.7 and which last for:					
		II.1.4.1. in th	ne cas	se of equidae suspected of having contracted dourine,					
		(³) <i>e</i>	either	[6 months beginning on the date of the last actual suspected of having contracted dourine or infected with					
		(³) <i>o</i>	or	[in the case of a stallion, until the animal is castrated;]					
		(³) O	or	[30 days following the date of completion of the clear after all animals of susceptible species have been sla					

EUROPEAN UNION Registered equidae, equidae for breeding and production equidae for slaughter II.a. Certificate reference number II.b. Local reference number II.1.4.2. in the case of glanders. (3) either [6 months beginning on the day on which the equidae suffering from the disease or subjected with positive results to a test for the detection of the causative pathogen Burkholderia mallei or antibodies to that pathogen, were killed and destroyed;] [30 days following the date of completion of the cleansing and disinfection of the premises after all animals of susceptible species have been killed and destroyed;] II.1.4.3. in the case of equine encephalomyelitis of any type, (3) either [6 months beginning on the day on which the equidae suffering from the disease have been slaughtered;] (3) or [6 months beginning on the day on which the equidae infected with the virus causing West Nile Fever, Eastern equine encephalomyelitis or Western equine encephalomyelitis have died, been removed from the holding or fully recovered;] [30 days following the date of completion of the cleansing and disinfection of the premises (3) or after all animals of susceptible species have been slaughtered;] II.1.4.4. in the case of equine infectious anaemia, until the date on which, the infected animals having been slaughtered, the remaining equine animals on the holding have shown a negative reaction in an agar gel immunodiffusion test (AGID or Coggins test) carried out on blood samples collected on two occasions 3 months apart; in the case of vesicular stomatitis, II.1.4.5. (3) either [6 months following the last case:] [30 days following the date of completion of the cleansing and disinfection of the premises (3) or after all animals of susceptible species have been slaughtered;] in the case of rabies, 30 days following the last case and the date of completion of the cleansing and II.1.4.6. disinfection of the premises; in the case of anthrax, 15 days following the last case and the date of completion of the cleansing and II.1.4.7. disinfection of the premises; II.1.5. to the best of my knowledge, during the period of 15 days prior to the date of dispatch the animal has not been in contact with equidae infected or suspected of an infectious or contagious disease. 11.2. Attestation of residence and pre-export isolation II.2.1. The animal was imported on (insert date) (3) either [directly from the EU Member State (insert name of EU Member State);] [from a country or part of the territory of a country (3) or (insert name of country) under conditions at least as strict as those set out in this certificate;] 1122 the animal exited from the Union less than 30 days ago, and since exit from the Union it was never in a country or part of the territory of a country (1) other than those of the same Sanitary Group, and resident on holdings under veterinary supervision, accommodated in separated stables without coming into contact with equidae of lower health status, except during racing, competition or the cultural event.

II.3. Attestation of animal welfare

The animal described in Box I.28 was examined today (¹) and found fit to be transported on the intended journey and arrangements were made to protect its health and well-being effectively at all stages of the journey.

Qualification and title:

Signature:

Name (in capital letters):

Date:

Stamp:

EUROPEAN UNION Registered equidae, equidae for breeding and production equidae for slaughter II.a. Certificate reference number II.b. Local reference number Notes: Part I: Box I.8.: Provide the code of the country or part of the territory of the country as appearing in column 3 of Annex I to Commission Implementing Regulation (EU) 2018/659. Box I.15.: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) and information is to be provided. In case of unloading and reloading, the consignor must inform the Border inspection Post of entry into the Union. Box I.23.: The container number and the seal number (if applicable) should be included. Box I.28.: Identification system: The animal must bear an individual identifier which permits to link the animal to the identification document as defined in Article 2(b) of Commission Implementing Regulation (EU) 2018/659. Specify the identification system (such as ear tag, tattoo, brand, transponder) and the anatomic place used on the animal. The number of the accompanying passport must be stated and the name of the competent authority which validated it. Age: Date of birth (dd/mm/yyyy). Sex (M = male, F = female, C = castrated). Part II: The certificate must be issued on the day of loading or on the last working day before loading of the animal for dispatch to the Member State of destination in the Union. The re-entry after temporary export of this registered horse shall not be allowed when the animal was loaded either prior to the date of authorisation for re-entry into the Union from the respective country or the part of the territory of the country referred to in point II.1.1, or during a period where restrictive measures have been adopted by the Union against the entry of live equidae from this country or this part of the territory of the country of dispatch. Code of the country or part of the territory of the country and the Sanitary Group as appearing in columns 3 and 5 of (²) Annex I to Commission Implementing Regulation (EU) 2018/659. $(^{3})$ Delete as appropriate. This health certificate shall: be drawn up in at least a language understood by the certifying officer and one of the official languages of the Member (a) State of destination and of the Member State where the registered horse will enter the Union territory and undergo the (b) be made out to a single consignee: be signed and stamped in a colour different to the colour of the printing; (c) consist of a single sheet of paper or all sheets of paper required are part of an integrated whole and indivisible by inserting page numbers and total number of pages, and each page shall bear the certificate reference number at the top of the page and those pages are stapled and stamped Official veterinarian

	Declaration by the owner or representative of the owner for the re-entry after temporary export of a registered horse for racing, competition and cultural events								
lder	Identification of the animal (1)								
Sp	ecies (Scie	ntific name)	Identification system	Identification number	Age	Sex			
	Equus c	aballus							
I, th	e undersig	ned owner (²) o	r representative of the owne	r (2) of the registered horse descr	ibed above, hereby dec	clare, that:			
_	the horse								
	(²) either		rily exported from the Union prior to this declaration;]	to the country of dispatch on		(insert date) less			
	(²) or		ountry of dispatch on where horse entered country	of dispatch);]	om	(insert name of			
_			ays prior to the date of disparansmissible to equidae;	atch the horse has not been in co	ntact with animals suffe	ering from infectious			
_	the transpof the jou		effected in such a way that	health and well-being of the hors	se can be protected effe	ectively at all stages			
_	 the conditions for residence and pre-export isolation as applicable in accordance with point II.2 of the accompanying health certificate for the country or part of the territory of the country of dispatch are fulfilled. 								
Nar	ne and add	ress of the own	er (²) or representative (²):						
Dat	e:		(dd/mm/yyyy)						
(1)	Article 2(the transponder of the transponder of the transponder of	o) of Commission er) and the anator ort accompanies the of birth (dd/mm/yy nale, F = female,	n Implementing Regulation (E nic place used on the animal. he animal, its number should be yyy).	dentifier which permits to link the ani LU) 2018/659. Specify the identifical estated and the name of the competer	ition system (such as ea	ar tag, tattoo, brand,			
(2)	Delete as	appropriate.							

Section B

Model health certificates and model declarations applicable to re-entry into the Union of registered horses temporarily exported for specific competitions or races

Chapter 1

Model health certificate and model declaration applicable to re-entry into the Union of registered horses for competition after temporary export for less than 90 days to participate in equestrian events organised under the auspices of the Fédération Equestre Internationale (FEI)

▼<u>B</u>

(Test event in preparation of the Olympic Games, Olympic Games, Paralympics, World Equestrian Games, Asian Equestrian Games, American Equestrian Games, Endurance World Cup in United Arab Emirates)

COUNT	RY:		Veterinary certificate to EU		
	l.1.	Consignor Name	I.2. Certificate reference No I.2.a.		
		Address	I.3. Central competent authority		
_		Tel.	I.4. Local competent authority		
signment	1.5.	Consignee Name Address	1.6.		
ched con		Postcode Tel.			
Part I : Details of dispatched consignment	1.7.	Country of ISO code I.8. Region of code origin Code	I.9. Country of ISO code I.10. Region of Code destination		
: Detai	1.11.	Place of origin	I.12. Place of destination		
Part		Name Approval number Address	Name Address		
			Postcode		
	I.13.	Place of loading	I.14. Date of departure		
	1.15.	Means of transport	I.16. Entry BIP in EU		
		Aeroplane ☐ Ship ☐ Railway wagon ☐			
		Road vehicle ☐ Other ☐ Identification Documentary references	I.17. No(s) of CITES		
	I.18.	Description of animal	I.19. Commodity code (HS code) 01 01		
			I.20. Quantity		
	I.21.		I.22. Number of packages		
	1.23.	Seal/Container No	1.24.		
	1.25.	Animal certified for:			
		Registered horse			
	1.26		I.27. For import or admission into EU		
	1.28.	Identification of the animal			
		Species Identification system Identification Identi	ation number Age Sex		

				II.a. Certificate reference number	II.b. Local reference number				
	II.	Attestation	of anima	l health and welfare					
	I, the under	signed official	veterinari	an, hereby certify, that the animal described in Box I.2	8:				
	_	is a registered horse as defined in Article 2(c) of Commission Implementing Regulation (EU) 2018/659;							
	 was examined today (¹) and found free of clinical signs of diseases and of obvious signs infestation; 								
ation	_	is not intended for slaughter under a national programme of infectious or contagious disease eradication;							
ertifica	_	meets the re	equiremer	ats attested in points II.1 to II.3 of this certificate;					
Part II: Certification	_	is accompa	nied by th	e written declaration, signed by the owner of the horse	e, or the representative of the owner.				
Pa	II.1.	Attestation	on third co	ountry or part of the territory of third country and holding	g of dispatch				
	II.1.1.	The animal is dispatched from (insert name of country or part of the territory country), a country or part of the territory of a country which on the date of issuing this certificate Code:							
	II.1.2. in the country of dispatch the following diseases are compulsorily notifiable: African horse si (<i>Trypanosoma equiperdum</i>), glanders (<i>Burkholderia mallei</i>), equine encephalomyelitis (of all Venezuelan equine encephalomyelitis), equine infectious anaemia, vesicular stomatitis, rabies and								
	II.1.3.	the animal is	s dispatch	ed from a country or part of the territory of a country:	of a country:				
		a) which is considered free from African horse sickness in accordance with Directive 2009/156/EC and in which there has been no clinical, serological (in unvaccinated equidae) or epidemiological evidence of African horse sickness during the period of 2 years prior to the date of dispatch and in which there have been no vaccinations against the disease during the period of 12 months prior to the date of dispatch;							
				/enezuelan equine encephalomyelitis has not occurre f dispatch;	ed during the period of 2 years prior to				
		c)	in which o	lourine has not occurred during the period of 6 months	s prior to the date of dispatch;				
		d)	in which g	planders has not occurred during the period of 6 month	ns prior to the date of dispatch;				
	II.1.4.	the animal does not come from a holding, and to the best of my knowledge for the time periods referred to in points II.1.4.1 to II.1.4.7 was not in contact with animals from holdings, which were subject to prohibition orders for the reasons referred to in points II.1.4.1 to II.1.4.7 and which last for:							
		II.1.4.1.	II.1.4.1. in the case of equidae suspected of having contracted dourine,						
			(³) either	[6 months beginning on the date of the last actual suspected of having contracted dourine or infected w					
			(³) or	[in the case of a stallion, until the animal is castrated;	1				
			(³) or	[30 days following the date of completion of the clea after all animals of susceptible species have been sla					

			II.a. Certificate reference number	II.b. Local reference number	
	II.1.4.2. in the case of glanders,				
		(³) either	[6 months beginning on the day on which the eq subjected with positive results to a test for the a Burkholderia mallei or antibodies to that pathogen, we	detection of the causative pathogen	
		(³) or	[30 days following the date of completion of the clear after all animals of susceptible species have been killed		
	II.1.4.3.	in the case	e of equine encephalomyelitis of any type,		
		(³) either	either [6 months beginning on the day on which the equidae suffering from the disease have been slaughtered;]		
		(³) or	or [6 months beginning on the day on which the equidae infected with the virus causing We Nile Fever, Eastern equine encephalomyelitis or Western equine encephalomyelitis had ided, been removed from the holding or fully recovered;]		
		(³) or	[30 days following the date of completion of the cleansing and disinfection of the premises after all animals of susceptible species have been slaughtered;]		
	II.1.4.4.	slaughtere gel immu	in the case of equine infectious anaemia, until the date on which, the infected animals having bee slaughtered, the remaining equine animals on the holding have shown a negative reaction in an ag- gel immunodiffusion test (AGID or Coggins test) carried out on blood samples collected of two occasions 3 months apart;		
	II.1.4.5.	in the case	in the case of vesicular stomatitis,		
		(³) either	[6 months following the last case;]		
		(³) or	[30 days following the date of completion of the clear after all animals of susceptible species have been sla		
	II.1.4.6.		e of rabies, 30 days following the last case and the dan of the premises;	ate of completion of the cleansing and	
	II.1.4.7.		e of anthrax, 15 days following the last case and the d n of the premises;	ate of completion of the cleansing and	
II.1.5.			wledge, during the period of 15 days prior to the date infected or suspected of an infectious or contagious dis		
II.2.	Attestation	of residenc	ce and pre-export isolation		
II.2.1.	The anima		ted into the country or part of the territory of the count	ry of dispatch on	
	(³) either	[directly fr	om the EU Member State	(insert name of EU Member State);]	
	(³) or		untry or part of the territory of a countrynder conditions at least as strict as those set out in thi		
II.2.2.	the animal	exited from	the Union		

			II.a. Certificate reference number	II.b. Local reference number			
	(³) either	a country supervisio health sta	[less than 30 days ago, and since exit from the Union was never in a country, or pa a country (¹) other than those of the same Sanitary Group, and resident on holding supervision, accommodated in separated stables without coming into contact with health status except during competition and has taken part in or was stabled tog participating in the LG Global Champions Tour				
		(³) either	either [in the Metropolitan area of Mexico City, Mexico;]]				
		(3) and/or	[in Miami, Unites States of America;]				
		(³) or	[in Shanghai, China;]]				
	(³) or	a country supervisio	60 days ago, and since exit from the Union was never (1) other than those of the same Sanitary Group, and n, accommodated in separated stables without comitus except during competition and has taken part in the interval of the competition and the	resident on holdings under veterinary ng into contact with equidae of lower			
		(3) either	[the Asian Games in	(insert place).]]			
		(³) or	[the American Games in	(insert place).]]			
		(³) or	[the Endurance World Cup in United Arab Emirates.]]				
	(³) or	[less than 90 days ago, and since exit from the Union was never in a country, or part of the a country (1) other than those of the same Sanitary Group, and resident on holdings und supervision, accommodated in separated stables without coming into contact with equinealth status except during competition and has taken part in or was stabled together participating in					
		(³) either	[the Test event for the Olympic Games in	(insert place).]]			
		(3) or	[the Olympic Games in	(insert place).]]			
		(3) or	[the Paralympics in	(insert place).]]			
		(³) or	[the World Equestrian Games in	(insert place).]]			
II.3.	Attestation	of animal w	velfare				
	The anima and arrang	The animal described in Box I.28 was examined today (¹) and found fit to be transported on the intended journey and arrangements were made to protect its health and well-being effectively at all stages of the journey.					
Notes:							
Part I:							
Box I.8.:	Provide the code of the country or part of the territory of the country as appearing in column 3 of Annex I to Commission Implementing Regulation (EU) 2018/659.						
Box I.15.:	information		(railway wagons or container and lorries), flight no ovided. In case of unloading and reloading, the consig Jnion.				
Box I.23.:	The contain	ner number	and the seal number (if applicable) should be included	d.			

EURO	PEAN UNION	Registered equida	e, equidae for breeding a	nd production equidae for slaught
		II.a. Certificate reference	number	II.b. Local reference number
Вох	identification docu	ument as defined in Article 2(b) o system (such as ear tag, tattoo, l	of Commission Implementing brand, transponder) and th	ch permits to link the animal to the ng Regulation (EU) 2018/659. Specif e anatomic place used on the anima ne of the competent authority whic
	Age: Date of birth	(dd/mm/yyyy).		
	Sex (M = male, F	= female, C = castrated).		
Part	II:			
(1)	The certificate must be is: the Member State of dest		the last working day befor	e loading of the animal for dispatch t
	to the date of authorisation referred to in point II.1.1,	on for re-entry into the Union fror	n the respective country or ive measures have been a	en the animal was loaded either pric the part of the territory of the countr dopted by the Union against the entr
(2)		part of the territory of the country nplementing Regulation (EU) 20		as appearing in columns 3 and 5 c
(3)	Delete as appropriate.			
This	health certificate shall:			
(a)		of the Member State where the		the official languages of the Membe nter Union territory and undergo th
(b)	be made out to a single c	onsignee;		
(c)	be signed and stamped in	n a colour different to the colour o	of the printing;	
(d)	inserting page numbers a			n integrated whole and indivisible b ertificate reference number at the to
Offic	ial veterinarian			
	Name (in capital letters):			Qualification and title:
	Date:			Signature:
	Stamp:			

	Declaration by the owner or representative of the owner for the re-entry after temporary export of a registered horse for competition									
Ider	Identification of the animal (¹)									
Spe	ecies (Scie	ntific name)	Identification system	Identification number	Age	Sex				
	Equus c	aballus								
I, th	e undersig	ned owner (²)	or representative of the owner	er (2) of the registered horse descri	ibed above, hereby dec	lare, that:				
_	the horse									
	(²) either		rarily exported from the Union 0 days (²) or 90 days (²) prior t	to the country of dispatch on to this declaration;]		(insert date)				
	(²) or		country of dispatch on n where horse entered country	y of dispatch);]		(insert name of				
_	the horse	has been ter	mporarily exported from the U	nion to take part in						
	(²) either	[the Asian G	Sames in	(insert pi	lace);]					
	(²) or	[the America	an Games in	(insert pi	lace);]					
	(²) or	[the Endura	nce World Cup in United Arab	Emirates;]						
	(²) or	[the Test ev	ent for the Olympic Games in	(insert pi	/ace);]					
	(²) or	[the Olympic	Games in	(insert pi	lace);]					
	(²) or	[the Paralyn	npics in	(insert pi	lace);]					
	(²) or	[the World E	Equestrian Games in	(insert pi	lace);]					
	(²) or	[the LG Glo	bal Champions Tour in							
		(²) either [the Metropolitan area of Mexi	ico City, Mexico;]						
		(²) and/or [Miami, Unites States of Amer	rica;]						
		(3) or [Shanghai, China;]							
_			days prior to the date of disp transmissible to equidae;	patch the horse has not been in co	ntact with animals suffe	ring from infectious				
_				on as applicable in accordance we country of dispatch are fulfilled;	vith point II.2 of the acc	companying health				
—	the transpof the jou		be effected in such a way tha	t health and well-being of the hors	e can be protected effe	ctively at all stages				
Nan	ne and add	lress of the o	wner (²) or representative (²):							
Date	ə:		(dd/mm/yyyy)							
(1)	Article 2(b) of Commission Implementing Regulation (EU) 2018/659. Specify the identification system (such as ear tag, tattoo, brand, transponder) and the anatomic place used on the animal.									
	Age: Date	of birth (dd/mm	n/yyyy).	e stated and the name of the competer	n admonty which validated	n.				
(²)	•	nale, F = femal appropriate.	e, C = castrated).							
()	Delete do	appropriate.								

Chapter 2

Model health certificate and model declaration applicable to re-entry into the Union of registered horses for racing after temporary export for less than 90 days to participate in specific race events in Australia, Canada, the United States of America, Hong Kong, Japan, Singapore, the United Arab Emirates or Qatar

 $(International\ Group/Grade\ meetings,\ the\ Japan\ Cup,\ the\ Melbourne\ Cup,\ the\ Dubai\ Racing\ World-Cup,\ the\ Hong\ Kong\ International\ Races)$

COU	OUNTRY: Veterinary certificate to EU											
	l.1.	Consignor Name				1.2.	Certificate refere	ence No		I.2.a.		
	Address						I.3. Central competent authority					
#		Tel.		1.4.	Local competent	authority						
nme	1.5.	Consignee		1.6.								
nsig		Name Address										
ched co		Postcode Tel.										
spat												
s of dis	1.7.	Country of origin	ISO code	I.8. Region origin	of Code	1.9.	Country of destination	ISO code		Region of destination	Code	
Detail	l.11.	Place of origin				1.12	Place of destinat	tion				
Part I : Details of dispatched consignment		Name Approval number Address					Name Address					
					Postcode							
	I.13.	Place of loading				I.14. Date of departure						
	I.15. Means of transport					I.16. Entry BIP in EU						
		Aeroplane ☐ Ship ☐ Railway wagon ☐										
		Road vehicle Identification Documentary references	Other C erences	J		1.17	No(s) of CITES					
	I.18.	Description of an	imal					I.19. Cor	mmodity	code (HS cod 01 01	e)	
									I.	.20. Quantity 1		
	I.21.								1.	.22. Number o	f packages	
	1.23.	Seal/Container N	lo						1.	24.		
	1.25.	Animal certified for	or:						<u> </u>			
Registered horse												
	1.26.						I.27. For import of	or admission	into EU			
	1.28.	Identification of the	ne animal									
		Species (Scientific name) Equus caballus	Identi	fication system	Ident	tificatio	on number	Age		Se.	x	

				a.uuo 101 b1000	anig ania production oquidae for changino						
			II.a. Certificate reference numb	er	II.b. Local reference number						
	II.	Attestation of	al health and welfare								
	I, the under	signed official vet	rian, hereby certify, that the animal	described in Bo	x I.28:						
	 is a registered horse as defined in Article 2(c) of Commission Implementing Regulation (EU) 2018/659; 										
Part II: Certification	_	was examined infestation;	y (1) and found free of clinical	signs of diseas	ses and of obvious signs of ectoparasite						
Certifi	_	is not intended	aughter under a national programn	ighter under a national programme of infectious or contagious disease eradication;							
Part II:	_	meets the requ	ents attested in points II.1 to II.3 of t	his certificate;							
_	_	is accompanied	ne written declaration, signed by th	e owner of the h	horse or the representative of the owner.						
	II.1.	Attestation on o	ry or part of the territory of the cour	try and holding	of dispatch						
	II.1.1.	country), a cou	or part of the territory of a cour	ntry which at th	y or part of the territory of a he date of issuing this certificate has the y Group						
	II.1.2.	(Trypanosoma	ispatch the following diseases are compulsorily notifiable: African horse sickness, dourine iperdum), glanders (Burkholderia mallei), equine encephalomyelitis (of all types including encephalomyelitis), equine infectious anaemia, vesicular stomatitis, rabies and anthrax;								
	II.1.3.	the animal is di	hed from a country or part of the te	ed from a country or part of the territory of a country:							
		whi Afri	ere has been no clinical, serologica norse sickness during the period c en no vaccinations against the dis	al (in unvaccina f 2 years prior	cordance with Directive 2009/156/EC and in ted equidae) or epidemiological evidence of to the date of dispatch and in which there he period of 12 months prior to the date of						
			Venezuelan equine encephalomyof dispatch;	elitis has not oc	ocurred during the period of 2 years prior to						
		c) in v	dourine has not occurred during th	e period of 6 m	onths prior to the date of dispatch;						
		d) in v	glanders has not occurred during t	he period of 6 n	months prior to the date of dispatch;						
	II.1.4.	the animal does not come from a holding, and to the best of my knowledge for the time periods referred to in points II.1.4.1 to II.1.4.7 was not in contact with animals from holdings, which were subject to prohibition orders for the reasons referred to in points II.1.4.1 to II.1.4.7 and which last for:									
	II.1.4.1. in the case of equidae suspected of having contracted dourine,										
		(3)			actual or possible contact with an animal ted with <i>Trypanosoma equiperdum</i> ;]						
		(3)	[in the case of a stallion, until the	animal is castr	rated;]						
		(3)	[30 days following the date of co		e cleansing and disinfection of the premises en slaughtered;]						

			II.a. Certificate reference number						
	II.1.4.2.	in the cas	le of glanders,						
	11.1.4.2.	111 1110 000	o or grandoro,						
		(³) either	[6 months beginning on the day on which the equidae suffering from the disease o subjected with positive results to a test for the detection of the causative pathoger <i>Burkholderia mallei</i> or antibodies to that pathogen, were killed and destroyed;]						
		(³) or	[30 days following the date of completion of the cleansing and disinfection of the premis after all animals of susceptible species have been killed and destroyed;]						
	II.1.4.3.	in the cas	case of equine encephalomyelitis of any type,						
		(³) either	[6 months beginning on the day on which the equidae suffering from the disease have be slaughtered;]						
		(³) or	[6 months beginning on the day on which the equidae infected with the virus causing W Nile Fever, Eastern equine encephalomyelitis or Western equine encephalomyelitis hadied, been removed from the holding or fully recovered;]						
		(³) or	[30 days following the date of completion of the cleansing and disinfection of the premis after all animals of susceptible species have been slaughtered;]						
	II.1.4.4.	slaughtere gel immu	se of equine infectious anaemia, until the date on which, the infected animals having be ed, the remaining equine animals on the holding have shown a negative reaction in an a unodiffusion test (AGID or Coggins test) carried out on blood samples collected sions 3 months apart;						
	II.1.4.5.	in the cas	e of vesicular stomatitis,						
		(³) either	[6 months following the last case;]						
		(³) or	[30 days following the date of completion of the cleansing and disinfection of the premis after all animals of susceptible species have been slaughtered;]						
	II.1.4.6.		se of rabies, 30 days following the last case and the date of completion of the cleansing a on of the premises;						
	II.1.4.7.		ee of anthrax, 15 days following the last case and the date of completion of the cleansing a on of the premises;						
l.1.5.			wledge, during the period of 15 days prior to the date of dispatch the animal has not beer infected or suspected of an infectious or contagious disease.						
1.2.	Attestation	n of residen	ce and pre-export isolation						
I.2.1.	The anim		mported into the country or part of the territory of the country of dispatch (insert date)						
	(³) either	[directly fr the partici	rom the EU Member State (insert name of EU Member State) ipation in						
		(³) either	[The Japan Cup;]						
		(³) or	[The Melbourne Cup;]						
		(³) or	[The Dubai Racing World-Cup;]						
		(³) or							

II.b. Local reference number

EUROPEAN UNION

Registered equidae, equidae for breeding and production equidae for slaughter

(³) or	[from Australia (³), Canada (³), the United States Singapore (³), United Arab Emirates (³) or Qatar (³) fo meetings in the country of dispatch:	

II.2.2. as far as can be ascertained and based on the declaration of the owner of the horse or representative of the owner (3) accompanying this certificate, the animal was:

II.a. Certificate reference number

- not continuously outside the Union for more than 90 days, the date of scheduled return in accordance with this certificate included:
- not outside the country of dispatch or in case of International Group/Grade meetings outside Australia,
 Canada, the United States of America, Hong Kong, Japan, Singapore, the United Arab Emirates or Qatar;
- resident on holdings under veterinary supervision, accommodated in separated stables without coming into contact with equidae of lower health status except during racing;
- II.2.3. the animal entered the country of dispatch under animal health conditions at least as strict as those laid down in this health certificate.
- II.3. Attestation of animal welfare

The animal described in Box I.28 was examined today (¹) and found fit to be transported on the intended journey and arrangements were made to protect its health and well-being effectively at all stages of the journey.

Notes:

Part I:

- Box I.8.: Provide the code of the country or part of the territory of the country as appearing in column 3 of Annex I to Commission Implementing Regulation (EU) 2018/659.
- Box I.15.: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) and information is to be provided. In case of unloading and reloading, the consignor must inform the Border Inspection Post of entry into the Union.
- Box I.23.: The container number and the seal number (if applicable) should be included.
- Box I.28.: Identification system: The animal must bear an individual identifier which permits to link the animal to the identification document as defined in Article 2(b) of Commission Implementing Regulation (EU) 2018/659. Specify the identification system (such as ear tag, tattoo, brand, transponder) and the anatomic place used on the animal. The number of the accompanying passport must be stated and the name of the competent authority which validated it.

Age: Date of birth (dd/mm/yyyy).

Sex (M = male, F = female, C = castrated).

Part II:

(1) The certificate must be issued on the day of loading or on the last working day before loading of the animal for dispatch to the Member State of destination in the Union.

The re-entry after temporary export of this registered horse shall not be allowed when the animal was loaded either prior to the date of authorisation for re-entry into the Union from the respective country or part of the territory of the country referred to in point II.1.1, or during a period where restrictive measures have been adopted by the Union against the entry of live equidae from this country or this part of the territory of the country of dispatch.

- (2) Code of the country or part of the territory of the country, and the Sanitary Group as appearing in columns 3 and 5 of Annex I to Commission Implementing Regulation (EU) 2018/659.
- (3) Delete as appropriate.

EURO	PEAN UNION	Registered equidae, equidae for bree	eding and production equidae for slaughter					
		II.a. Certificate reference number	II.b. Local reference number					
This	health certificate shall:							
(a)	 (a) be drawn up in at least a language understood by the certifying officer and one of the official languages of the Memb State of destination and of the Member State where the registered horse will enter Union territory and undergo to veterinary border checks; 							
(b)	be made out to a single consi	gnee;						
(c)	be signed and stamped in a c	olour different to the colour of the printing;						
(d)	d) consist of a single sheet of paper or all sheets of paper required are part of an integrated whole and indivisible by inserting page numbers and total number of pages, and each page shall bear the certificate reference number at the top of the page and those pages are stapled and stamped.							
Offic	ial veterinarian							
	Name (in capital letters):		Qualification and title:					
	Date:		Signature:					
	Stamp:							

Declaration by the owner or representative of the owner for the re-entry after temporary export of a registered horse for racing										
Identification of the animal (1)										
Spe	ecies (Scie	ntific name)	Identification system	Identification number	Age	Sex				
	Equus c	aballus								
I, th	e undersig	ned owner (²) oı	r representative of the owne	er (²) of the registered horse desc	cribed above, hereby decla	ire, that:				
_	the horse									
	(²) either		rily exported from the Union ays prior to this declaration;	to the country of dispatch on		(insert date)				
	(²) or		ountry of dispatch onry	(insert date) from .v of dispatch);]		(insert name of				
_	the horse	has been temp	orarily exported from the Ur	nion to take part in						
	(²) either	[The Japan Cu	nb:]							
	(²) or	[The Melbourn	ne Cup;]							
	(²) or	[The Dubai Ra	acing World-Cup;]							
	(²) or	[The Hong Ko	ng International Races;]							
	(²) or		Group/Grade meetings in <i>A</i> gapore (²), United Arab Emir	Australia (2), Canada (2), the Urrates (2) or Qatar (2);]	nited States of America (²), Hong Kong (²),				
—			ays prior to the date of disparansmissible to equidae;	atch the horse has not been in c	ontact with animals sufferi	ng from infectious				
_				on as applicable in accordance e country of dispatch are fulfilled;		ompanying health				
_	 the transportation will be effected in such a way that health and well-being of the horse can be protected effectively at all stages of the journey. 									
Name and address of the owner (²) or representative (²):										
Date	Date:(dd/mm/yyyy)									
(¹)	Identificati	on system: The a	nimal must bear an individual i	dentifier which permits to link the a	nimal to the identification doc	ument as defined in				
()	Article 2(b	o) of Commission er) and the anator	n Implementing Regulation (E nic place used on the animal.	EU) 2018/659. Specify the identific	cation system (such as ear	tag, tattoo, brand,				
		ort accompanies to of birth (dd/mm/y		e stated and the name of the compet	ent authority which validated it	i.				
0	Sex (M = r	male, F = female,	• • • •							
(²)	Delete as appropriate.									

PART 3

Imports

Section A

Model health certificates and model declaration for imports into the Union of an individual registered horse, registered equine animal or equine animal for breeding and production

NTRY:				•	Veterinary certificate t			
l.1.	Consignor Name	1.2.	Certificate refere	nce No	I.2.a.			
	Address	1.3.	Central compete	nt authority				
	Tel.	1.4.	Local competent	authority				
1.5.	Consignee	1.6.						
1	Name							
	Address							
	Postcode Tel.							
I.5. I.7.	Country of ISO code I.8. Region of origin Code	1.9.	Country of destination	ISO code	I.10. Region of Co			
l.11.	Place of origin	I.12.	Place of destinat	ion				
	Name Approval number Address		Name Address					
			Postcode					
I.13.	Place of loading	I.14. Date of departure						
I.15.	Means of transport	I.16.	Entry BIP in EU					
	Aeroplane ☐ Ship ☐ Railway wagon ☐							
	Road vehicle Other I Identification Documentary references	l.17.	No(s) of CITES					
I.18.	Description of animal			I.19. Commo	odity code (HS code) 01 01			
					I.20. Quantity 1			
I.21.					I.22. Number of packages			
1.23.	Seal/Container No				1.24.			
1.25.	Animal certified for:							
	Registered horse \Box registered equine animal \Box breeding and production \Box							
I.26.			.27. For import o	r admission in	ito EU 🗖			
1.28.	Identification of the animal							
	pecies (Scientific Identification system Identific			Age	Sex			

				regiotorea equidae, equidae for breed	iiig uii	a production equidae for eladynta							
				II.a. Certificate reference number	II.b.	Local reference number							
	II.	Attestatio	n of anima	al health and welfare									
	I, the unders												
— (¹) either [is a registered equine animal, other than horse, as defined in Article 2(c) of Directive 2009/1													
	(1) or [is a registered horse as defined in Article 2(c) of Commission Implementing Regulation												
	sion implementing Regulation (EO)												
u O		(1) or	or [is an equine animal for breeding and production as defined in Article 2(e) of Directive 2009/156/EC;]										
ertificat	 comes from a country or part of the territory of a country which is authorised for imports into the Union of the category of equidae specified in the first indent above; 												
Рап II: Сеппсатіоп	 was examined today (²) and found free of clinical signs of diseases and of obvious signs of ectoparas infestation; 												
	 is not intended for slaughter under a national programme of infectious or contagious disease eradication; 												
	_	meets the	requireme	nts attested in points II.1 to II.5 of this certificate;									
	_	is accomp	anied by th	e written declaration, signed by the owner of the a	ınimal o	or the representative of the owner.							
	II.1.	Attestation	on third c	ountry or part of the territory of third country and h	olding (of dispatch							
	II.1.1.	.1.1. The animal is dispatched from											
	II.1.2.	(Trypanos	oma equip	spatch the following diseases are compulsorily redum), glanders (<i>Burkholderia mallei</i>), equine ncephalomyelitis), equine infectious anaemia, ves	encep	phalomyelitis (of all types including							
	II.1.3.	the anima	l is dispatch	ned from a country or part of the territory of country	y								
		a)	which the African h	considered free from African horse sickness in acc ore has been no clinical, serological (in unvaccinal orse sickness during the period of 2 years prior on no vaccinations against the disease during th	ed equ to the	iidae) or epidemiological evidence o date of dispatch and in which there							
		b)		Venezuelan equine encephalomyelitis has not oc of dispatch;	curred	during the period of 2 years prior to							
		c)	in which	dourine has not occurred during the period of 6 mo	onths p	rior to the date of dispatch;							
		d)	d) in which glanders has not occurred during the period of 6 months prior to the date of dispatch;										
	(1) either	[e)	in which dispatch;	vesicular stomatitis has not occurred during the	e perio	nd of 6 months prior to the date o							
	(¹) or	[e)	and a blo	vesicular stomatitis has occurred during the perio od sample taken from the animal on		(insert date), within a period							
			(1) either	[in a virus neutralisation test at a serum dilution of	of 1 in 3	32;]]							
			(¹) or	[in an ELISA in accordance with the relevant of and Vaccines for Terrestrial Animals of the OIE;]		r of the Manual of Diagnostic Tests							

		Registered equidae, equidae for breed	ing and production equidae for slaughter						
		II.a. Certificate reference number	II.b. Local reference number						
points II.1	.4.1 to II.1.4	come from a holding, and to the best of my kno 1.7 was not in contact with animals from holdings, to in points II.1.4.1 to II.1.4.7 and which last for:							
II.1.4.1.	in the cas	se of equidae suspected of having contracted dour	ine,						
	(¹) either	[6 months beginning on the date of the last actual or possible contact with an animal suspected of having contracted dourine or infected with <i>Trypanosoma equiperdum</i> ;]							
	(1) or [in the case of a stallion, until the animal is castrated;]								
	(¹) or	[30 days following the date of completion of the after all animals of susceptible species have been							
II.1.4.2.	in the cas	se of glanders,							
	(¹) either	[6 months beginning on the day on which th subjected with positive results to a test for burkholderia mallei or antibodies to that pathoger	he detection of the causative pathogen						
	(¹) or	[30 days following the date of completion of the after all animals of susceptible species have been							
II.1.4.3.	in the cas	se of equine encephalomyelitis of any type,							
	(¹) either	[6 months beginning on the day on which the equal slaughtered;]	uidae suffering from the disease have been						
	(¹) or	[6 months beginning on the day on which the eq Nile Fever, Eastern equine encephalomyelitis o died, been removed from the holding or fully reco	r Western equine encephalomyelitis have						
	(¹) or	[30 days following the date of completion of the after all animals of susceptible species have been							
II.1.4.4.	II.1.4.4. in the case of equine infectious anaemia, until the date on which, the infectious slaughtered, the remaining equine animals on the holding have shown a negel immunodiffusion test (AGID or Coggins test) carried out on block two occasions 3 months apart;								
II.1.4.5.	in the cas	ee of vesicular stomatitis,							
	(¹) either	[6 months following the last case;]							
	(¹) or	[30 days following the date of completion of the after all animals of susceptible species have been							
II.1.4.6.		se of rabies, 30 days following the last case and the premises;	ne date of completion of the cleansing and						
II.1.4.7.		se of anthrax, 15 days following the last case and ton of the premises;	he date of completion of the cleansing and						
		wledge, during the period of 15 days prior to the c infected or suspected of an infectious or contagiou							

			II.a. Certificate reference number II.b. Local reference number					
II.2.	Attestation	of resider	nce and pre-export isolation					
(¹) either	[II.2.1.	than 90 o	period of at least the 90 days prior to the date of di days old, or since entry if the animal was imported prior to the date of dispatch, the animal has bee on situated in a country or part of the territory of a co	directly from the Union during a period of n resident on holdings under veterinary				
	(¹) (⁴) either		to Sanitary Group A, and during the period of at least 30 days prior to the date of dispatch, tapart from equidae not of equivalent health status;]]					
	(¹) (⁴) or	of dispate	It to Sanitary Groups B, C, D or G, and during the period of at least 30 days prior to the date on it was kept in pre-export isolation under veterinary supervision without coming into contact dae not of equivalent health status;]] It to Sanitary Group E, and it was kept in the approved isolation centre described as place of 30x I.11, protected from vector insects [during the period of at least 40 days prior to the date of dispatch;]]					
	(¹) (⁴) or							
		(1) either						
		(¹) or	[during the period of at least 30 days prior to the date of dispatch from the United Emirates;]]					
(¹) (⁴) or	[II.2.1.	[II.2.1. The animal is dispatched from a country of which at least a part of the territory of the cour assigned to Sanitary Group F, and during the period of at least 90 days prior to the date of disports or since birth if the animal is less than 90 days old, it was resident on holdings under vete supervision and was kept during the period of at least 60 days prior to the date of dispatch, or entry if it was imported directly from the Union during the period of 60 days prior to the date of dispatch, in the part of the territory described in point II.1.3 which is considered free of African sickness in accordance with the Union legislation and underwent the pre-export isolation						
	(¹) either	of quare from premises provided combinat in strict i	approved vector-protected quarantine station of					
	(¹) or	(insert na	ently confined in the approved vector-proof quarantiname of quarantine station) during the period of at lestant monitoring of the vector protection has proved part of the quarantine station.]]	east 14 days prior to the date of dispatch				
II.3.	Attestation	of vaccina	ation and health tests					
(¹) either	[II.3.1.		nal was not vaccinated against African horse sickne nation suggesting previous vaccination;]	ss in the country of dispatch and there is				
(¹) or	[II.3.1.	The anim	nal was vaccinated against African horse sickness, a	and this vaccination was carried out:				
	(¹) either	[more that	an 12 months prior to the date of dispatch;]]					
	(¹) or		an 60 days and less than 12 months prior to the data ory of the country referred to in point II.1.3.(a), from v	,				
(¹) (⁴) or	[II.3.1.	Sanitary date) not quarantir	nal is dispatched from a country or part of the ter Group F and was vaccinated against African horse some than 24 months and at least 40 days prior to be by administration of a registered vaccine according a against the circulating serotypes of the African hor	sickness on(insert to the date of entry in the vector-protected and to manufacturer's instructions which is				

EUROPEAN UNION Registered equidae, equidae for breeding and production equidae for slaughter II.a. Certificate reference number II.b. Local reference number II.3.2. the animal was not vaccinated against Venezuelan equine encephalomyelitis during the period of 60 days prior to the date of dispatch from (1) either Ia country of which all parts of the territory are free of Venezuelan equine encephalomyelitis for a period of at least 2 years prior to the date of dispatch;] (1) (4) or [a part of the territory of a country which is assigned to Sanitary Group C or D, which is free of Venezuelan equine encephalomyelitis for a period of at least 2 years prior to the date of dispatch and Venezuelan equine encephalomyelitis occurs in the remaining parts of the territory of the country of dispatch, and (1) either [is vaccinated against Venezuelan equine encephalomyelitis with a complete primary course and revaccinated according to manufacturer's recommendations not less than 60 days and no more than 12 months prior to the date of dispatch, and was kept in vectorprotected quarantine for a period of at least 21 days prior to the date of dispatch, and during that period remained clinically healthy, and its body temperature, taken daily, remained within the normal physiological range, and any equine animal on the same holding which showed a rise in body temperature, taken daily, was subjected to a blood test for virus isolation for Venezuelan equine encephalomyelitis with negative results;]] (1) or [is not vaccinated against Venezuelan equine encephalomyelitis and was kept in vectorprotected quarantine for a period of at least 21 days, and during that period remained clinically healthy, and its body temperature, taken daily, remained within the normal physiological range, and any equine animal on the same holding which showed a rise in body temperature, taken daily, was subjected to a blood test for virus isolation for Venezuelan equine encephalomyelitis with negative results, and the animal to be dispatched was subjected to a diagnostic test for Venezuelan equine encephalomyelitis with negative result conducted on a sample taken not less than 14 days after the date of entry into the vector protected quarantine and remained protected from vector insects until dispatch:11 (1) or [was subjected to a haemagglutination inhibition test for Venezuelan equine encephalomyelitis carried out by the same laboratory on the same day on samples taken on two occasions with an interval of 21 days on (insert date) and on (insert date), the second of which was taken during the period of 10 days prior to the date dispatch, without an increase in antibody titre, and a RT-PCR (reverse transcription-polymerase chain reaction) test for the detection of Venezuelan equine encephalomyelitis virus genome, carried out with negative result on a sampling until loading for dispatch, by combined use of approved insect repellents and insecticides on the animal and disinsectisation of the stable and the means in which it is transported:11 (1) [II.3.3. the animal is an uncastrated male equine animal older than 180 days, and (1) either [is dispatched from a country in which equine viral arteritis (EVA) is a compulsorily notifiable disease and has not been officially reported during the period of 6 months prior to the date of dispatch;]] (insert date), within a period of (1) or [was tested on a blood sample taken on 21 days prior to the date of dispatch, by virus neutralisation test for EVA with negative result at a serum dilution of 1 in 4:11 (1) or a period of 21 days prior to the date of dispatch, by virus isolation test, polymerase chain reaction (PCR) or real-time PCR for EVA with negative result;]] (insert date) under official veterinary (1) or [was vaccinated against EVA on supervision, and re-vaccinated at regular intervals according to the manufacturer's instructions, with a vaccine approved by the competent authority, and the initial vaccination was carried out

			II.a. Certificate reference number	II.b. Local reference number		
		(¹) either	[before 31 December 2017, on the day a blood tested in a virus neutralisation test for EVA v of 1 in 4;]]]			
		(¹) or	[before 31 December 2017, during a period of isolation of not more than 15 days under official veterinary supervision, commencing on the day a blood sample was taken which was tested during that isolation period in a virus neutralisation test for EVA with negative result at a serum dilution of 1 in 4;]]]			
		(¹) or	[at the age of 180 to 270 days, during a period of isolation under official veterinary supervision, during which the animal was subjected to a virus neutralisation test for EVA carried out with negative result at a serum dilution of 1 in 4, or carried out on the same day by the same laboratory with stable or declining titres on two blood samples taken at least 10 days apart;]]]			
		(¹) or	after the animal was subjected to a virus neutralisation test for EVA with negative result at a serum dilution of 1 in 4, carried out on a blood sample taken not earlier than 7 days after commencing a period of uninterrupted isolation which lasted until 21 days following vaccination;]]]			
		(¹) or	[at the age of 180 to 250 days, after the animal w for EVA carried out with negative result at a sert same day by the same laboratory with stable or d at least 14 days apart;]]]	um dilution of 1 in 4 or carried out on the		
	(¹) or	carried of sample of to the da	jected to a virus isolation test, polymerase chain in ut with negative result on an aliquot of its entire f that animal taken on	semen collected after the date a blood rt date), within a period of 6 months prior		
(¹) (⁴) either	[11.3.4.	anaemia,	al is dispatched from Iceland, which is certified where it was continuously resident since birth an we entered Iceland from other countries;]			
(¹) or	[II.3.4.	Coggins	nal was subjected with negative result to an a test) or to an ELISA for equine infectious anaemia (insert date), this being within a perio	a carried out on a blood sample taken on		
	(¹) [II.3.5.	Sanitary during a test for g	imal is dispatched from a country or part of the territory of a country which is assigned ry Group B, D or E, or from China or Thailand, or from a country in which glanders was reporte a period of 3 years prior to the date of dispatch, and was subjected to a complement fixation ry glanders carried out with negative result at a serum dilution of 1 in 5 on a blood sample take			
	(¹) [II.3.6.	country of China or the date negative (insert date	al is an uncastrated male or a female equine animal older than 270 days dispatched from or part of the territory of a country which is assigned to Sanitary Group B, D, E or F, or from Thailand, or from a country in which dourine was reported during a period of 2 years prior to dispatch, and was subjected to a complement fixation test for dourine carried out with result at a serum dilution of 1 in 5 on a blood sample taken on			
	(¹) [II.3.7.		al is dispatched from a country or part of the te Group C or D, and	rritory of a country which is assigned to		
		(¹) either	[Western and Eastern equine encephalomyelitis country or part of the territory of the country of diprior to the date of dispatch;]]			

		II.a. Certif	icate refere	nce number	II.b. Local reference number		
	(¹) or	manufactu date of	rer's instrud dispatch w	ctions within a period of 6 r rith inactivated vaccine a	ary course and revaccinated according to months and at least 30 days prior to the against Western and Eastern equine d on(insert date);]]		
	(¹) or	protected of	quarantine,	and during this period subje	ys prior to the date of dispatch in a vector ected to haemagglutination inhibition tests s carried out by the same laboratory		
		(¹) either	[on a sample of blood taken on				
		(¹) or	21 days o (insert day the date	n(<i>inser</i> e), the second of which was	o occasions with an interval of at least t date) and on		
(¹) [II.3.8.	Sanitary	Group G, o	al is dispatched from a country or part of the territory of a country which is assigned to Group G, or from a country in which Japanese encephalitis has been officially reported in luring the past 2 years, and the animal				
	(¹) either	holding wh	ere there h		area of at least 30 km radius around that se encephalitis during a period of at least		
	(¹) or	date of dis	[was kept in a vector–protected quarantine during a period of at least 21 days prior to the date of dispatch, and during that period the body temperature, taken daily, remained within the normal physiological range, and was subjected				
		(¹) either	Japanese samples of second of dispatch,	encephalitis carried out by if blood taken on two occasi	on or virus neutralisation test for the same laboratory on the same day on ons with an interval of at least 14 days on (insert date), the period of 10 days prior to the date of ld increase in antibody titre between the from vector insects until dispatch;]]]		
		(¹) or	Japanese sample ta	encephalitis virus with not earlier than 7 days a(insert date), and	or the detection of antibodies against egative result, carried out on a blood after the date the isolation commenced on d remained protected from vector insects		
	(¹) or	revaccinat	ed accordin		s with a complete primary course and endations during a period of not less than date of dispatch;]]		
(¹) (⁴) either [II.3.9.	Sanitary	Group E, ar	nd was subj	ected to a serological test for	ritory of a country which is assigned to or African horse sickness as described in the same laboratory on the same day		
	(¹) either	-			n interval of between 21 and 30 days, on(insert date), the second r to the date of dispatch		
		(¹) either	[with nega	tive results in each case;]]]			
		(¹) or	[with posit	ive result in the first sample,	and		
			(¹) either		ubsequently tested with negative result in that as described in Annex IV to Directive		

		-		J	quiduo, equidue for breeding and production equidue for claughter		
			II.a. Certif	icate refere	nce number II.b. Local reference number		
				(¹) or	[the two samples were tested without more than a two-fold increase in antibody titre in a virus neutralisation test as described in point 2.4 of Chapter 2.5.1 of the OIE Terrestrial Manual for Diagnostic Tests and Vaccines;]]]		
	(¹) or	to the dat by the OI	e of dispato E as official	ch, and the ly free of A			
(¹) (⁴) or	[II.3.9.	the animal is dispatched from a country or part of the territory of a country which is Sanitary Group F and					
		(¹) either	r [was subjected to a serological test for African horse sickness as describe Annex IV to Directive 2009/156/EC, which was carried out by the same laboratory of same day on blood samples taken on two occasions with an interval of between 21 30 days, on				
			(1) either	[with nega	ative results in each case;]]]		
			(1) or	[with posi	tive result in the first sample, and		
				(¹) either	[the second sample was subsequently tested with negative result in an agent identification test as described in Annex IV to Directive 2009/156/EC;]]]]		
				(¹) or	[the two samples were tested without more than a two-fold increase in antibody titre in a virus neutralisation test as described in point 2.4 of Chapter 2.5.1 of the OIE Terrestrial Manual for Diagnostic Tests and Vaccines;]]]]		
		(¹) or	as describ case on a 28 days a	ed in Anne: blood samp ifter the da	serological and an agent identification test for African horse sickness x IV to Directive 2009/156/EC, carried out with negative result in each ole taken on		
		(¹) or	Annex IV taken on .	to Directive	agent identification test for African horse sickness as described in a 2009/156/EC, carried out with negative result on a blood sample (insert date) not less than 14 days after the date of introduction ted quarantine and not more than 72 hours before dispatch;]]		
II.4.	Attestatio	on of the tran	sport condi	tions			
(¹) either	[II.4.1.	Sanitary (market, r	e animal is dispatched from a country or part of the territory of a country which is assigned to nitary Group A, B, C, D, E or G and is transported directly to the Union, without passing through a urket, marshalling or assembly centre and without coming into contact with other equidae of a ferent health status.]				
(¹) (⁴) or	[II.4.1.	Sanitary coming in	Group F ai	nd is trans vith other e	a country or part of the territory of a country which is assigned to ported directly from the vector-protected quarantine station without quidae not accompanied by a health certificate either for imports or for nion		
		(¹) either	the aircra	ft being c	vector-protected conditions and arrangements have been made that leansed and disinfected in advance with a disinfectant officially d country of dispatch, and sprayed against vector insects just prior to		

II h I ocal reference number

EUROPEAN UNION

Registered equidae, equidae for breeding and production equidae for slaughter

	ii.a. Certificate reference number	II.b. Local reference number
(¹) or	[to a sea port in that country or part of the territor conditions and arrangements have been made scheduled directly to a port in the Union without opart of the territory of a country not approved fo stalls which were cleansed and disinfected in recognised in the third country of dispatch and space departure.]]	e to transport it on a vessel which is calling into a port situated in a country or r the entry into the Union of equidae, in a advance with a disinfectant officially

Il a Certificate reference number

- II.4.2. Arrangements have been made and verified to prevent any contact with other equidae not complying with at least the same health requirements as described in this health certificate during the period from certification until dispatch to the Union.
- II.4.3. The transport vehicles or containers in which the animal is going to be loaded were cleaned and disinfected before loading with a disinfectant officially recognised in the third country of dispatch and they are so constructed that faeces, urine, litter or fodder cannot escape during transportation.

II.5. Attestation of animal welfare

The animal described in Box I.28 was examined today (²) and found fit to be transported on the intended journey and arrangements were made to protect its health and well-being effectively at all stages of the journey.

Notes:

Part I:

- Box I.8.: Provide the code of the country or the part of the territory of the country as appearing in column 3 of Annex I to Commission Implementing Regulation (EU) 2018/659.
- Box I.15.: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) and information is to be provided. In case of unloading and reloading, the consignor must inform the BIP of entry into the EU.
- Box I.23.: The container number and the seal number (if applicable) should be included.
- Box I.28.: Species: Select amongst: Equus caballus, Equus asinus, Equus africanus, Equus hemionus, Equus kiang, Equus quagga, Equus zebra, Equus grevyi, or indicate any cross between those.

Identification system: The animal must bear an individual identifier which permits to link the animal to the identification document as defined in Article 2(b) of Commission Implementing Regulation (EU) 2018/659. Specify the identification system (such as ear tag, tattoo, brand, transponder) and the anatomic place used on the animal.

If a passport accompanies the animal, its number should be stated and the name of the competent authority which validated it.

Age: Date of birth (dd/mm/yyyy).

Sex (M = male, F = female, C = castrated).

Part II:

- Delete as appropriate.
- (²) The certificate must be issued on the day of loading or in the case of a registered horse on the last working day before loading of the animal for dispatch to the Member State of destination in the Union.

The import of this equine animal shall not be allowed when the animal was loaded either prior to the date of authorisation for imports of an individual registered equine animal or equine animal for breeding and production into the Union from the respective country or part of the territory of the country mentioned under point II.1.1, or during a period where restrictive measures have been adopted by the Union against the entry of live equidae from this country or this part of the territory of the country of dispatch.

		II.a. Certificate reference number	II.b. Local reference number					
(3)	Code of the country or part of the territory of the country and the Sanitary Group as appearing in columns 3 and 5 respectively of Annex I to Commission Implementing Regulation (EU) 2018/659.							
(4)	Statements that relate entirely and exclusively to a Sanitary Group different from the Sanitary Group to which the country of dispatch, or part of its territory, is assigned, may be left out, provided that the numbering of the subsequent statements is maintained.							
This	health certificate shall:							
(a)	be drawn up in at least a language understood by the certifying officer and one of the official languages of the Member State of destination and of the Member State where the animal will enter Union territory and undergo the veterinary border checks;							
(b)	be made out to a single cons	ignee;						
(c)	be signed and stamped in a	colour different to the colour of the printing;						
(d)	consist of a single sheet of paper or all sheets of paper required are part of an integrated whole and indivisible by inserting page numbers and total number of pages, and each page shall bear the certificate reference number at the top of the page and those pages are stapled and stamped.							
Offic	ial veterinarian							
	Name (in capital letters):		Qualification and title:					
	Date:		Signature:					
	Stamp:							

▼<u>B</u>

	Declaration by the owner or representative of the owner for entry into the Union of an equine animal								
Iden	tification o	f the animal (1)							
Species (Scientific name) Identification system Identification number Age Sex									
I, the	e undersigr	ned owner (²) or	representative of the owner	(²) of the animal described abo	ve, hereby declare, that:				
_	the anima	al							
	(²) either			territory of the country of dispa animal is less than 90 days of a		east 90 days prior			
	(²) or			of the country of dispatch duri Member State of the Union;]	ng the required residence	period of at least			
_			days prior to the date of diseases transmissible to equ	lispatch the animal has not be uidae;	een in contact with anima	lls suffering from			
_				n as applicable in accordance country of dispatch are fulfilled;		ompanying health			
_			nsport as applicable in accor ne country of dispatch are fu	dance with point II.4 of the acc lfilled;	ompanying health certifica	te for the country			
_		portation will be the journey;	effected in such a way the	at health and well-being of the	animal can be protected	effectively at all			
Nan	ne and add	ress of the own	er (2) or representative (2):						
Date	e:		(dd/mm/yyyy)						
(¹)	<i>grevyi</i> , or i	ndicate any cross	between those.	quus africanus, Equus hemionus, E					
	Article 2(b) of Commission		lentifier which permits to link the ar J) 2018/659. Specify the identific					
	If a passpo	*	ne animal, its number should be	stated and the name of the compete	ent authority which validated it				
	-	nale, F = female, (
(²)	Delete as a	appropriate.							

Section B

Model health certificate and model declaration for imports into the Union of consignments of domestic equidae for slaughter

UN	TRY	:				Veterinary certificate to E	
	l.1.	Consignor Name	I.2. I.3.	Certificate reference		l.2.a.	
		Address	1.3.	Central Competent	authority		
		Tel.	1.4.	Local competent a	uthority		
ment	1.5.	Consignee	1.6.				
igu		Name					
Sons		Address					
) jed		Postcode					
atcı		Tel.					
Part I : Details of dispatched consignment	I.7.	Country of ISO code I.8. Region of Code origin Code	1.9.	Country of destination	ISO code	I.10. Region of Code destination	
בום	l.11.	. Place of origin	1.12	Place of destinatio	n		
		•					
-		Name Approval number Address		Name Address			
				Postcode			
	I.13.	. Place of loading	I.14. Date of departure				
	I.15.	. Means of transport	1.16	Entry BIP in EU			
		Aeroplane ☐ Ship ☐ Railway wagon ☐					
		Road vehicle Other Ot	117	No(s) of CITES			
		Identification	i. Tr. Tio(o) of office				
		Documentary references					
	I.18.	. Description of animals			I.19. Commo	dity code (HS code) 01 01	
						I.20. Quantity	
	I.21.					I.22. Number of packages	
	1.23.	. Seal/Container No				1.24.	
	1.25.	. Animals certified for:					
		Slaughter					
	1.26. I.27. For import or admission into EU						
}	1.28	. Identification of the animals					
			catio	n number	Δαρ	Sex	
	č	Species (Scientific Identification system Identifi name)	catioi	i ilullibel	Age	Sex	

			II.a. Certificate reference number	II.b. Local reference number						
	II. Attestation of animal health, animal welfare and public health									
	I, the undersigned official veterinarian, hereby certify, that the animals described in Box I.28:									
	_	are equidae for slauç	hter as defined in Article 2(d) of Directive 2009/156/E	EC;						
	_	were examined toda infestation;	ay (1) and found free of clinical signs of diseases	and of obvious signs of ectoparasite						
	_	are not intended for	slaughter under a national programme of infectious or	contagious disease eradication;						
	_	meet the requiremen	ts attested in points II.1 to II.5 of this certificate;							
_	_	are accompanied by owner.	the written declaration, signed by the owner of the	ne animals or the representative of the						
ication	II.1.	Attestation on third c	ountry or part of the territory of third country and hold	ing of dispatch						
Part II: Certification	II.1.1. The animals are dispatched from									
Pa	II.1.2.	in the country of dispatch the following diseases are compulsorily notifiable: African horse sickness, dourine (<i>Trypanosoma equiperdum</i>), glanders (<i>Burkholderia maller</i>), equine encephalomyelitis (of all types including Venezuelan equine encephalomyelitis), equine infectious anaemia, vesicular stomatitis, rabies and anthrax;								
	II.1.3.	the animals are dispa	tched from a country or part of the territory of country							
		which the African h	considered free from African horse sickness in accordance with Directive 2009/156/EC and in the has been no clinical, serological (in unvaccinated equidae) or epidemiological evidence of orse sickness during the period of 2 years prior to the date of dispatch and in which there are no vaccinations against the disease during the period of 12 months prior to the date of							
			n which Venezuelan equine encephalomyelitis has not occurred during the period of 2 years prior to ne date of dispatch;							
		c) in which	dourine has not occurred during the period of 6 mont	hs prior to the date of dispatch;						
		d) in which	glanders has not occurred during the period of 6 mon	ths prior to the date of dispatch;						
	(3) either		in which vesicular stomatitis has not occurred during the period of 6 months prior to the date of dispatch;]							
	(³) or	and a blo period of	in which vesicular stomatitis has occurred during the period of 6 months prior to the date of dispatch, and a blood sample taken from each of the animals on (insert date), within a period of 21 days prior to the date of dispatch, was tested with negative results for antibody to the vesicular stomatitis virus							
	(3) either [in a virus neutralisation test at a serum dilution of 1 in 32;]]									
		(³) or	[in an ELISA in accordance with the relevant Cha and Vaccines for Terrestrial Animals of the OIE;]]	apter of the Manual of Diagnostic Tests						
	II.1.4.	points II.1.4.1 to II.1	come from holdings, and to the best of my knowle 4.7 have not been in contact with animals from hol s referred to in points II.1.4.1 to II.1.4.7 and which las	dings, which were subject to prohibition						

		II.a. Certificate reference number	II.b. Local reference number			
II.1.	4.1. in the cas	se of equidae suspected of having contracted douring	ə ,			
	(³) either	[6 months beginning on the date of the last act suspected of having contracted dourine or infected				
	(³) or	[in the case of a stallion, until the animal is castrate	d;]			
	(³) or	or [30 days following the date of completion of the cleansing and disinfection of the prafter all animals of susceptible species have been slaughtered;]				
II.1.	4.2. in the cas	in the case of glanders,				
	(³) either	[6 months beginning on the day on which the subjected with positive results to a test for the Burkholderia mallei or antibodies to that pathogen,	detection of the causative pathogen			
	(³) or	[30 days following the date of completion of the cleansing and disinfection of the premafter all animals of susceptible species have been killed and destroyed;]				
II.1.	4.3. in the cas	se of equine encephalomyelitis of any type,				
	(³) either	ither [6 months beginning on the day on which the equidae suffering from the disease I slaughtered;]				
	(³) or		on which the equidae infected with the virus causing West ephalomyelitis or Western equine encephalomyelitis have ding or fully recovered;]			
	(³) or	[30 days following the date of completion of the cleafter all animals of susceptible species have been s				
II.1.	slaughter agar gel	se of equine infectious anaemia, until the date on vered, the remaining equine animals on the holding immunodiffusion test (AGID or Coggins test) carrisions 3 months apart;	have shown a negative reaction in an			
II.1.	4.5. in the cas	se of vesicular stomatitis,				
	(³) either	[6 months following the last case;]				
	(³) or	[30 days following the date of completion of the cleafter all animals of susceptible species have been s				
II.1.		se of rabies, 30 days following the last case and the on of the premises;	date of completion of the cleansing and			
II.1.		se of anthrax, 15 days following the last case and the on of the premises;	e date of completion of the cleansing and			
		owledge, during the period of 15 days prior to the dat ae infected or suspected of an infectious or contagiou				
II.2. Atte	estation of resider	nce and pre-export isolation				
of 9	0 days prior to th	een resident in the country or part of the territory of the date of dispatch, or since birth if the animals are lend, and they are dispatched from a country or part of the since the sinc	ess than 90 days old, on holdings under			
(³) <i>e</i>		d to Sanitary Group A and during the period of at lead kept apart from equidae not of equivalent health sta				

			II.a. Certificate reference number	II.b. Local reference number					
	(³) or	dispatch	If to Sanitary Groups B, C or D and during the perior they were kept in pre-export isolation under vete rith equidae not of equivalent health status;]						
	(³) or		ned to Sanitary Group E and for the period of at least 40 days prior to the date of dispatch they kept in the approved isolation centre described in Box I.11, protected from vector insects.]						
II.3.	Attestation	n of vaccina	f vaccination and health tests						
(³) either	[II.3.1.		The animals were not vaccinated against African horse sickness in the country of dispatch and there is no information suggesting previous vaccination;]						
(³) or	[II.3.1.		nals were vaccinated against African horse sicknes n 12 months prior to dispatch;]]	s, and this vaccination was carried out					
	II.3.2.		als were not vaccinated against Venezuelan equindispatch from	e encephalomyelitis during the 60 days					
	(³) either		y of which all parts of the territory are free of Ven at least 2 years prior to the date of dispatch;]	ezuelan equine encephalomyelitis for a					
	(³) (⁴) or	Venezuel	of the territory of a country which is assigned to Sanitary Group C or D, which is free of uelan equine encephalomyelitis for a period of at least 2 years prior to the date of dispatch and uelan equine encephalomyelitis occurs in the remaining parts of the territory of the country of h, and						
		(³) either	[were vaccinated against Venezuelan equine enc course and revaccinated according to manufact 60 days and not more than 12 months prior to the of protected quarantine for a period of at least 21 during that period remained clinically healthy, ar remained within the normal physiological range, holding which showed a rise in body temperature, to for virus isolation for Venezuelan equine encephalogical	urer's recommendations not less than late of dispatch, and were kept in vector-days prior to the date of dispatch, and their body temperature, taken daily, and any equine animal on the same aken daily, was subjected to a blood test					
		(³) or	[were not vaccinated against Venezuelan equinvector-protected quarantine for a period of at leas and during that period remained clinically healthy. remained within the normal physiological range, holding which showed a rise in body temperature. It for virus isolation for Venezuelan equine encephanianist to be dispatched were subjected to a encephalomyelitis with negative result conducted cafter the date of entry into the vector-protected quector insects until dispatch;]]	at 21 days prior to the date of dispatch. and their body temperature, taken daily, and any equine animal on the same aken daily, was subjected to a blood test lomyelitis with negative results, and the diagnostic test for Venezuelan equine on a sample taken not less than 14 days					
(³) (⁴) either	[II.3.3.	the animals are dispatched from Iceland, which is certified as officially free from equine infectious anaemia, where they have been continuously resident since birth and did not come into contact with equidae which have entered Iceland from other countries;]							
(³) or	[II.3.3.	the animals were subjected to an agar gel immunodiffusion test (AGID or Coggins test) or to an ELISA for equine infectious anaemia carried out with negative result in each case on blood samples taken on							
(3) [[II.3.4. the animals are dispatched from a country or part of the territory of Sanitary Group B, D or E, or from a country in which glanders was 3 years prior to the date of dispatch, and were subjected to a compl carried out with negative result in each case at a serum dilution of 1 i				ders was reported during the period of a complement fixation test for glanders ion of 1 in 5 on blood samples taken on					

		II.a. Certifi	cate referer	nce number	II.b.	Local reference number	
c c v a	country or country in vere subj at a serun	als are uncastrated males or female equine animals older than 270 days dispatched from a propert of the territory of a country which is assigned to Sanitary Group B, D or E or from a nowhich dourine was reported during the period of 2 years prior to the date of dispatch, and objected to a complement fixation test for dourine carried out with negative result in each case modification of 1 in 5 on blood samples taken on					
		als are dispatched from a country or part of the territory of a country which is assigned to Group C or D, and					
(*	³) either	[Western and Eastern equine encephalomyelitis have not been officially reported in the country or part of the territory of the country of dispatch during the period of 2 years prior to the date of dispatch;]]					
Ć	³) or	[the animals were vaccinated with a complete primary course and revaccinated according to manufacturer's instructions within the period of 6 months and at least 30 days prior to the date of dispatch with inactivated vaccine against Western and Eastern equine encephalomyelitis, the last vaccination was applied on					
(³) or	period sub	jected to h		tests	om vector insects and during this for Western and Eastern equine carried out on	
		(3) either [a sample of blood taken from each of the animals in the consignment on					
		(³) or	occasions and on period of 1	with an interval of at least 21 (insert date), the	days secon spatch	nimals in the consignment on two on	
S	Sanitary C	Broup E, and	d were subje	ected to a serological test for	r Afric	of a country which is assigned to an horse sickness as described in me laboratory on the same day	
(*	³) either	with an inte	rval of betw) and on	een 21 and 30 days, on	 rt date	e consignment on two occasions), the second of which was taken	
		(³) either	[with negat	ive result in each case;]]]			
		(³) or	[with positive	ve results in the first sample,	and		
			(³) either		entifica	quently tested with negative result tion test as described in Annex IV	
			(³) or	without more than a two-fo	old inc bed in	of the consignment were tested rease in antibody titre in a virus point 2.4 of Chapter 2.5.1 of the stic Tests and Vaccines;]]]]	
(°	³) or	consignment dispatch, a the OIE as	nt onnd nd the coun officially fre	(insert date), within the try or part of the territory of the territ	the pe the co and is	en from each of the animals in the riod of 10 days prior to the date of untry of dispatch is recognised by not adjacent to a country in which 2 years.]]	

Registered equidae, equidae for breeding and production equidae for slaughter

			•	109,010,0	u oqu.uu	-,		.	pioddo		
			II.a. Ce	rtificate re	ference n	umber		II.b.	Local re	eference n	umber
II.4.	Attestation	n of the tran	sport con	ditions							
(³) either	[II.4.1.	slaughterl assembly	nouse on centre re	the terr	itory of t in Article	he Union, 7(1) of Dire	without pas	sing t	hrough	a market	ed directly to , marshalling oming into conta
(³) or	[II.4.1.	Arrangements were made and verified to ensure that before the animals are transported to a slaughterhouse on the territory of the Union they pass only through a single approved market, marshalling or assembly centre referred to in Article 7(1) of Directive 2009/156/EC situated in the same Member State, from where they are transferred directly to the slaughterhouse without coming into contact with other equidae not authorised for the entry into the Union.]									
	II.4.2.		same he	ealth requ	irements	as describ					complying with the period fro
	II.4.3.	disinfecte	d before	loading w	ith a disir	nfectant offi		ised i	n the thi	rd country	ere cleaned ar of dispatch ar portation.
II.5.	Attestation	n of animal	welfare								
	The animals described in Box I.28 were examined today (1) and found fit to be transported on the intended journey and arrangements have been made to protect their health and well-being effectively at all stages of the journey.										
II.6.	Attestation	n of public h	ealth								
	androgeni		ic or beta	a-agonist :	substance	es for purpo					any oestrogeni chnical treatme
		antees cove of Directive				y the resid	ue plan sub	mitted	and app	proved in	accordance wi
Notes:											
Part I:											
Box I.8.:		ne code of on Impleme					he country a	as app	earing i	in column	3 of Annex I
Box I.15.:	informatio		ovided. Ir								ame (ship) ar Border Inspectio
Box I.23.:	The conta	iner numbe	r and the	seal numl	ber (if app	olicable) sho	ould be inclu	ded.			
Box I.28.:	Species: S	Select amor	ıgst: " <i>Equ</i>	ıus caballı	ıs", "Equi	us asinus" c	or "Equus ca	ballus	x Equus	asinus".	
	identificati		nt. Speci	fy the ide							the animal to th ponder) and th
	Age: Date	of birth (dd	/mm/yyyy	').							

Sex (M = male, F = female, C = castrated).

		II.a. Certificate reference number	II.b. Local reference number						
Part	II:								
(¹)	The certificate must be issued Union.	d on the day of loading of the animals for dispatch to	o the Member State of destination in the						
	date of authorisation for impeterritory of a country mention	nimals for slaughter shall not be allowed when the orts of live equidae for slaughter into the Union froed under point II.1.1, or during a period where rest equidae from this country or this part of the territory	om the respective country or part of the rictive measures have been adopted by						
(²)	Code of the country or part of the territory of the country and the Sanitary Group as appearing in columns 3 and 5 respectively of Annex I to Commission Implementing Regulation (EU) 2018/659.								
(³)	Delete as appropriate.								
(⁴)		y and exclusively to a Sanitary Group different from ory, is assigned, may be left out, provided that the n							
This	health certificate shall:								
(a)		guage understood by the certifying officer and one ne Member State where the animals will enter Uni							
(b)	be made out to a single consi	gnee;							
(c)	be signed and stamped in a c	olour different to the colour of the printing;							
(d)		paper or all sheets of paper required are part of otal number of pages, and each page shall bear the are stapled and stamped.							
Offici	ial veterinarian								
	Name (in capital letters):		Qualification and title:						
	Date:		Signature:						
	Stamp:								

				owner or representative of the or consignments of live equidae fo		
Iden	itification c	of the animal (1))			
Spe	ecies (Scie	entific name)	Identification system	Identification number	Age	Sex
I, the	e undersig	ned owner (²) (or representative of the owne	$\mathrm{er}(^2)$ of the animals described above	re, hereby declare, that:	
_	the anima		ned in the country or part of t	the territory of the country of dispat	tch for at least 90 days p	prior to the date o
_			days prior to the date of diseases transmissible to ed	lispatch the animals have not becquidae;	en in contact with anima	als suffering fron
_				on as applicable in accordance wi e country of dispatch are fulfilled;	th point II.2 of the acco	ompanying health
_			ansport as applicable in acco the country of dispatch are f	ordance with point II.4 of the according in the according in the control of the according in the control of the according in	mpanying health certifica	ate for the country
_		sportation will to f the journey;	pe effected in such a way the	nat health and well-being of the a	animal can be protected	d effectively at a
_	the anima	als will be sent				
	(²) either		the premises of dispatch to of the same health status;]	o the slaughterhouse of destination	n without coming into o	contact with othe
	(²) or	marshalling of		slaughterhouse of destination pas to in Article 7(1) of Directive 2009 status;]		
Nan	ne and add	dress of the ow	ner (²) or representative (²):			
Date	ə:		(dd/mm/yyyy)			
(1)	Identificati Article 2(I transpond If a passpo Age: Date	ion system: The b) of Commission er) and the anato	animal must bear an individual in Implementing Regulation (Examic place used on the animal, the animal, its number should be byyyy).	r indicate any cross between those. identifier which permits to link the anin EU) 2018/659. Specify the identificati e stated and the name of the competent	on system (such as ear	tag, tattoo, brand

(2) Delete as appropriate.

PART 4

Explanatory notes for the certification

(a) The health certificates shall be issued by the competent authority of the exporting country, based on the model set out in Part 1, 2 or 3 of Annex II, according to the layout of the model that corresponds to the animals concerned.

They shall contain, in the numbered order that appears in the model, the attestations that are required for any country and, as the case may be, those supplementary guarantees that are required for the exporting country or part of the territory of the country.

- (b) Where the model health certificate states that certain statements shall be kept as appropriate, statements which are not relevant, may be crossed out and initialled and stamped by the certifying officer, or completely deleted from the health certificate.
- (c) A separate and unique health certificate shall be issued for animals that are exported from a single territory appearing in columns 2 and 4 of Annex I which are consigned to the same destination and transported in the same railway wagon, lorry, aircraft or ship.
- (d) The original of the health certificate shall consist of a single sheet of paper, or, where more text is required, it shall be in such a form that all sheets of paper required are part of an integrated whole and indivisible.
- (e) The health certificate shall be drawn up in at least one of the official languages of the EU Member State of the border inspection post of entry of the consignment into the Union and of the EU Member State of destination. However, those EU Member States may authorise the health certificate to be drawn up in the official language of another EU Member State, and accompanied, if necessary, by an official translation.
- (f) If for reasons of identification of the animals of the consignment (schedule in Box 1.28 of the model health certificate), additional sheets of paper are attached to the health certificate, those sheets of paper shall also be considered as forming part of the original of the health certificate by application of the signature and stamp of the certifying official veterinarian, on each of the pages.

- (g) When the health certificate, including additional schedules referred to in (f), comprises more than one page, each page shall be numbered -(page number) of (total number of pages) at the end of the page and shall bear the certificate reference number that has been designated by the competent authority on the top of the pages.
- (h) The original of the health certificate shall be completed and signed by an official veterinarian within 24 hours prior loading the consignment, or in the case of registered horses on the last working day prior to loading, for exportation to the Union. The competent authorities of the exporting country shall ensure that the principles of certification equivalent to those laid down in Council Directive 96/93/EC (*) are followed.

The colour of the signature and the stamp of the official veterinarian shall be different to that of the printing on the health certificate. This requirement also applies to stamps other than those embossed or watermark.

- (i) The original of the health certificate shall accompany the consignment until it reaches the border inspection post of entry into the Union.
- j) The certificate reference number referred to in Box I.2 and Box II.a of the model health certificate shall be provided by the competent authority of the exporting country.

ANNEX III

MODEL HEALTH CERTIFICATES FOR ENTRY INTO THE UNION OF SEMEN, OVA AND EMBRYOS OF EQUIDAE

PART 1

Model health certificate for imports of semen

▼<u>C2</u>

Section A

MODEL 1- Model health certificate for imports of consignments of semen of equidae collected in accordance with Directive 92/65/EEC after 30 September 2014 and dispatched from an approved semen collection centre of origin of the semen

COUN	TRY:									٧	eterinary certifica	te to EU
	l.1.	Consignor Name					1.2.	Certificate referen	ce No	I.	2.a.	
		Address					1.3.	Central competent	t authority			
ent		Tel.					1.4.	Local competent a	uthority			
Part I : Details of dispatched consignment	1.5.	Consignee Name Address					1.6.	Person responsibl Name Address	e for the load	in E	U	
patched o		Postal code Tel.						Postal code Tel.				
s of disp	1.7.	Country of Is origin	SO code		Region of origin	Code	1.9.	Country of destination	ISO code	I.10.	Region of destination	Code
: Detai	l.11.	Place of origin Semen centre					l.12.	Place of destination Semen centre ☐	on	Нс	olding \square	
Part	Name Approval number Address Postal code							Name Address		Appı	roval number	
								Postal code				
	I.13.	I.13. Place of loading						Date of departure				
	I.15. Means of transport						I.16.	Entry BIP in EU				
		Aeroplane	Ship		Railway wag	on 🔲						
		Road vehicle Identification Documentary		r□			l.17.					
	I.18.	Description of	f commodity	,			I.19. Commodity code (HS code) 05 11 99 85					
							I.20. Quantity					
	I.21.										2. Number of packa	iges
	1.23.	Seal/Containe	er No							1.24	1.	
	I.25. Commodities certified for: Artificial reproduction											
	I.26. For transit through EU to third country						I.27. For import or	admission into	o EU			
	Third country ISO code											
	I.28. Identification of the commodities											
	Species (Scientific name) Donor identity						Date of collection Quantity				Quantity	

Equine semen - Section A

	II. Health inf	formation		II.a. Certificate reference No	II.b.						
	I, the undersigr	ned, official veterinar	ian, of the	e exporting country (²)(name of exporting							
				(name of exporting	oodnay)						
	certify that:										
uo	II.1. The semen collection centre (3), in which the semen described above was collected, processed and stored for export to the Union is approved and supervised by the competent authority in accordance with the conditions of Chapters I(I)(1) and I(II)(1) of Annex D to Directive 92/65/EEC (4);										
Part II: Certification	II.2. During the period commencing 30 days prior to the date of first collection of the semen described above until the date the fresh or chilled semen was dispatched or until the 30 days storage period for frozen semen elapsed, the semen collection centre:										
Part II:	II		was situated in the exporting country or, in the case of regionalisation according to Article 1 Directive 2009/156/EC $(^5)$, in that part of the territory of the exporting country which was:								
				ed to be infected with African horse sickness ective 2009/156/EC,	in accordance with Article 5(2)(a)						
		— free	from Ver	nezuelan equine encephalomyelitis for a period	of at least 2 years,						
		— free	from glar	nders and dourine for a period of at least 6 mon	hs;						
	II	I.2.2. fulfilled th	e conditio	ons for a holding laid down in Article 4(5) of Dire	ctive 2009/156/EC and in particular:						
				a case of a disease mentioned below not all the animals of species susceptible to ase located in the holding were slaughtered or killed and the holding has been free:							
	be			from any type of equine encephalomyelitis for a period of at least 6 months beginning on the day on which the equidae suffering from the disease are slaughtered,							
			ne Oi	from equine infectious anaemia (EIA) for at least the period required to obtain a negative result in an agar gel immunodiffusion test (AGID or Coggins test) carried out on samples taken after the infected animals were slaughtered on two occasions 3 months apart from each of the remaining animals,							
				from vesicular stomatitis (VS) for a period of at least 6 months from the last recorded case,							
			— fr	om rabies for a period of at least one month from	n the last recorded case,						
			— fr	om anthrax for a period of at least 15 days from	the last recorded case,]						
	(*		disease and the encepha the case	g a case of a disease mentioned below all the animals of species susceptible to that located in the holding have been slaughtered or killed and the premises disinfected, a holding was free for a period of at least 30 days from any type of equine alomyelitis, equine infectious anaemia, vesicular stomatitis and rabies or 15 days in e of anthrax, beginning on the day on which following the destruction of the animals affection of the premises was satisfactorily completed;]							
	II	I.2.3. contained metritis,	only equ	uidae which were free of clinical signs of equine	viral arteritis and contagious equine						
	II.3. Prior to entering the semen collection centre the donor stallions and any other equidae located in the centre:										

Equine semen - Section A

11 11==10	informatic:		II a Cortificato reference No	II b
II. Health	n information		II.a. Certificate reference No	II.b.
	II.3.1.	a Member State of regionalisation in	resident for a period of 3 months (or since eif the Union during the 3 months period) in the accordance with Article 13 of Directive 2009, try which was during that period:	ne exporting country or, in the case of
			ed to be infected with African horse sickne ective 2009/156/EC,	ess in accordance with Article 5(2)(a)
		— free from Ve	nezuelan equine encephalomyelitis for a perio	d of at least 2 years,
		— free from gla	nders and dourine for a period of at least 6 mo	onths;
(¹) either	[11.3.2.		e country of export which was on the day of (VS) for a period of at least 6 months,]	of admission into the centre free from
(¹) or	[II.3.2.	result at a serum with the relevant	a virus neutralisation test for vesicular stom dilution of 1 in 32 or a VS ELISA carried ou Chapter of the Manual of Diagnostic Tests a I sample taken (⁶) within 14 days prior to ente	t with a negative result in accordance nd Vaccines for Terrestrial Animals of
	II.3.3.	originated from ho point II.2.2;	oldings which on the day of admission onto t	the centre fulfilled the requirements of
II.4.	The seme	n described above v	vas collected from donor stallions which:	
	II.4.1.		clinical sign of an infectious or contagious dis- centre and on the day the semen was collected	
	II.4.2.		eriod of at least 30 days prior to the date of s shown any clinical sign of equine viral arterit	
	II.4.3.	collection and bet	natural mating during a period of at least 30 veen the dates of the first sample referred to if the collection period;	
	II.4.4.	Manual of Diagno which is recognise	owing tests, which meet at least the require stic Tests and Vaccines for Terrestrial Animals d by the competent authority and has the test valent to that provided for in Article 12 of	s of the OIE, carried out in a laboratory ts referred to hereinafter included in its
	(test) or	ne infectious anaemia (EIA), an agar-gel im an enzyme-linked immunosorbent assay (ELI: ve result;]	
		II.4.4.2. for equi	ne viral arteritis (EVA),	
	((¹) either [II.4.4.2	a serum neutralisation test with a nega in four;]	tive result at a serum dilution of one
	((¹) and/or [II.4.4.2	a virus isolation test, polymerase chain renegative result on an aliquot of the entire	
		three sp	stagious equine metritis (CEM), an agen secimens (swabs) taken from the donor stallic than 7 days at least from the penile sheath	on on two occasions with an interval of

cou	NTRY				Equine semen – Section A				
II.	Health information	on	I	I.a. Certificate reference No	II.b.				
			(local trea transport r	aples were in no case taken earlier than 7 days (systemic treatment) or 21 days eatment) after antimicrobial treatment of the donor stallion and were placed in t medium with activated charcoal, such as Amies medium, before dispatch to the ry where they were subjected with a negative result to a test for:					
		(¹) either	[II.4.4.3.1.	the isolation of <i>Taylorella equigenitalis</i> after cultive conditions for a period of at least 7 days, set up wit specimens from the donor animal, or 48 hours who cool during transport;]	hin 24 hours after taking the				
		(¹) and/or	[11.4.4.3.2.	the detection of genome of <i>Taylorella equigenitalis</i> carried out within 48 hours after taking the specime					
				the results specified in point II.4.4 in each case direspectively in points 1.6(a), (b) and (c) of Chaptes:					
		(⁹) [II.4.5.1.	at least 30 the semen	stallion was continuously resident on the semen coll days prior to the date of the first collection and durin described above, and no equidae on the semen co to direct contact with equidae of lower health status t	ng the period of collection of ollection centre came during				
			stallion at collection and not les	described in point II.4.4 were carried out on sample least once a year at the beginning of the breeding of semen intended for imports into the Union of frest than 14 days following the date of the commencer 30 days prior to the first semen collection.]	season or prior to the first sh, chilled or frozen semen				
		(⁹) [II.4.5.2.	30 days posemen des	r stallion was resident on the semen collection cer rior to the date of the first collection and during the scribed above, but left the semen collection centre un erinarian for a continuous period of less than 14 day collection centre came into direct contact with equida-	e period of collection of the nder the responsibility of the ys, and/or other equidae on				
			stallion at the first co semen and	The tests described in point II.4.4 were carried out on samples taken (⁶) from the donor stallion at least once a year at the beginning of the breeding season or prior to the date of the first collection of semen intended for imports into the Union of fresh, chilled or frozen semen and not less than 14 days following the date of the commencement of the residence period of at least 30 days prior to the first semen collection,					
		and	chilled or	period of collection of the semen intended for imperfozen semen the donor stallion was subjected , as follows:					
			(a)	for equine infectious anaemia, one of the tests de last carried out on a sample of blood taken (⁶) not the collection of the semen described above;					
			(b)	for equine viral arteritis, one of the tests described					
			(¹) either	[in point II.4.4.2 was last carried out on a samp 30 days prior to the date of the collection of the sem					
			(¹) or	[in point II.4.4.2.2 was carried out on an aliquot donor stallion taken (6) not more than 6 month collection of the semen described above and a blood donor stallion during the 6 months period reacted serum neutralisation test for equine viral arteritis sthan one in four;]	s prior to the date of the od sample taken (6) from the d with a positive result in a				

Equine semen - Section A

II. Health in	nformation	า		II.a. (Certificate re	ference No			II.b		
			(c)	for contagious equine metritis, the test described in point II.4.4.3 was last carried out on three specimens (swabs) taken (6) not more than 60 days prior to the date of the collection of semen described above							
			(¹) either	[on	two occasio	ns;]					
			(¹) or	[on	a single occ	asion and s	ubjected to a	PCR or real-	time PCR.]]		
				D to D					.6(a) and (b) r imports into		
									e carried out reginning of t		
			the dono from the semen c	r stallic date c collection	on during the	e storage po tion of the not less tha	eriod of the s semen and b ın 14 days a	emen of a mefore the se	on samples ta inimum perioo men is remov e than 90 da	d of 30 days red from the	
and (1) either [the tests for equine viral arteritis described in point II.4.4.2 were carried out on samples taken (6) during the storage period of the semen of a minimum period of 30 days from the date of the collection of the semen and before the semen is removed from the semen collection centre or used, not less than 14 days and not more than 90 days after the date of the collection of the semen described above.]											
			(¹) or	was neg stal	s confirmed pative result lion taken (⁶ lion has rea	by virus iso on sample) twice a ye cted with a	lation test, Po s of an aliquation at an inter	CR or real-timuot of the erwal of at leas It at a serum	re for equine ne PCR carrientire semen of the 4 months are dilution of at ritis.]	ed out with a of the donor and the donor	
	II.4.6.	underwent dates:	the testi	ting provided for in points II.3.2 (1) and II.4.5 on samples taken on the following							
<u></u>		Start	date (6)		Date of sampling for health tests (⁶)						
Identification of semen	Test	Donor	Sem	nen	VS (1)	EIA		A II. I.2.		EM 4.3.	
Identif	T prog	residence	collec		II.3.2	II.4.4.1.	Blood sample	Semen sample	1. sample	2. sample	

▼<u>C2</u>

COUNTRY

Equine semen - Section A

	n informatior		II.a. Certificate reference No	II.b.					
(1) either	[11.5.	No antibiotics were	added to the semen;]						
(¹) or	[II.5.	diluted semen of n	, ,	•					
II.6.	The seme	en described above w	as:						
	II.6.1.		ed, stored and transported under conditions vid III(I) of Annex D to Directive 92/65/EEC;	which comply with the requirements of					
	II.6.2.		of loading in a sealed container in accordar /e 92/65/EEC and bearing the number indicat						
Notes									
Part I:									
Box I.11.:	The place	e of origin shall corres	pond to the semen collection centre of the se	men origin.					
Box I.22.:	The numl	ber of packages shall	correspond to the number of containers.						
Box I.23.:	The ident	The identification of container and seal number shall be indicated.							
Box I.28.:	The dono	The donor identity shall correspond to the official identification of the animal.							
	The date	of collection shall be	indicated in the following format: dd/mm/yyyy.						
Part II:									
Guidance f	or the comp	letion of the table in p	oint II.4.6.						
Abbreviatio	ns:								
VS	Vesic	cular stomatitis (VS) te	esting if required in accordance with point II.3.2	2					
EIA-1	Equir	ne infectious anaemia	(EIA) testing first occasion						
EIA-2	EIA te	esting second occasion	on						
EVA-B	1 Equir	ne viral arteritis (EVA)	testing on blood sample first occasion						
EVA-B	2 EVA	testing on blood samp	ole second occasion						
EVA-S	1 EVA	testing on semen san	ple first occasion						
EVA-S	2 EVA	testing on semen san	nple second occasion						
CEM-1	1 Conta	agious equine metritis	(CEM) testing first occasion first sample						
CEM-1	2 CEM	testing first occasion	second sample taken 7 days after CEM-11						
CEM-2	21 CEM	testing second occas	ion first sample						
	O CEM	testing second occas	ion second sample taken 7 days after CEM-2	1					
CEM-2	Z CEIVI	testing second occas	ion scoond sample taken r days after OLM-2	•					

For each semen identified in column A in correspondence with Box I.28, the test programme (points II.4.5.1, II.4.5.2 and/or II.4.5.3) shall be specified in column B, and columns C and D shall be completed with the dates required.

▼ <u>C2</u>

COUNTRY

Equine semen - Section A

II.	Health information	II.a. Certificate reference No	II.b.

The dates when samples were taken for laboratory testing prior to the first collection of the semen described above as required in points II.4.5.1, II.4.5.2 and II.4.5.3, shall be entered in the upper line of columns 5 to 9 of the table, this being the boxes marked with EIA-1, EVA-B1 or EVA-S1 and CEM-11 and CEM-12 in the example below.

The dates when samples were taken for repeat laboratory testing as required in accordance with point II.4.5.2. or II.4.5.3. shall be entered in the lower line of columns 5 to 9 in table , this being the boxes EIA-2, EVA-B2 or EVA-S2 and CEM-21 and CEM-22 in the example below.

	of	40	Start	date	Date of sampling for health tests						
	Identification semen	Test programme	Donor	Semen	VS II.3.2.	EIA II.4.4.1.	EVA II.4.4.2.		CEM II.4.4.3.		
		brog	residence	collection			Blood sample	Semen sample	1. sample	2. sample	
	Δ.	D 6 D		D	vs	EIA-1	EVA-B1	EVA-S1	CEM-11	CEM-12	
	Α	В	B C	ا ت	VS	EIA-2	EVA-B2	EVA-S2	CEM-21	CEM-22	

- (¹) Delete as necessary.
- (²) Imports of equine semen are authorised from a third country listed in column 2 of Annex I to Commission Implementing Regulation (EU) 2018/657 provided that the semen was collected in the part of the territory of the third country detailed in column 4 of that Annex from a donor stallion of the category of equidae indicated in columns 11, 12 or 13 of that Annex.
- $(^{3})$ Only approved semen collection centres listed in accordance with Article 17(3)(b) of Directive 92/65/EEC on the Commission website: http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm
- Council Directive 92/65/EEC of 13 July 1992 laying down animal health requirements governing trade in and imports into the Community of animals, semen, ova and embryos not subject to animal health requirements laid down in specific Community rules referred to in Annex A(I) to Directive 90/425/EEC (OJ L 268, 14.9.1992, p. 54).
- Council Directive 2009/156/EC of 30 November 2009 on animal health conditions governing the movement and importation from third countries of equidae (OJ L 192, 23.7.2010, p. 1).
- (⁶) Insert date in table in point II.4.6 (follow Guidance in Part II of the Notes).
- (⁷) Regulation (EC) No 882/2004 of the European Parliament and of the Council of 29 April 2004 on official controls performed to ensure the verification of compliance with feed and food law, animal health and animal welfare rules (OJ L 165, 30.4.2004, p. 1).
- The agar gel immunodiffusion test (AGID or Coggins test) or the ELISA for equine infectious anaemia are not required for donor equidae which have continuously resided in Iceland since birth, provided that Iceland has remained officially free of equine infectious anaemia and no equidae and their semen, ova and embryos have been introduced into Iceland from outside prior to and during the period the semen was collected.
- Cross out the programmes that do not apply to the consignment. (⁹)
- Insert names and concentrations.

_	The signature and the stamp must be in a different colour to that of the printing.									
Offici	Official veterinarian									
	Name (in capital letters):	Qualification and title:								
	Date:	Signature:								
	Stamp:									

▼<u>C2</u>

Section B

MODEL 2 – Model health certificate for imports of consignments of stocks of semen of equidae collected, processed and stored in accordance with Directive 92/65/EEC after 31 August 2010 and before 1 October 2014 and dispatched after 31 August 2010 from an approved semen collection centre of origin of the semen

COUN	TRY:					Veterinary certificate to EU		
	l.1.	Consignor Name		Certificate reference No		I.2.a.		
		Address	I.3. Central competent authority					
		Tel.	I.4. Local competent authority					
Jent	1.5.	Consignee	1.6.	Person responsible for the lo	oad i	in EU		
guu		Name		Name				
isi		Address		Address				
ched cc		Postal code Tel.		Postal code Tel.				
Part I : Details of dispatched consignment	1.7.	Country of ISO code I.8. Region of code origin Code	1.9.	Country of destination	ie	I.10. Region of Code destination		
ails	111	Place of origin	112	Place of destination	_			
l : Det	1.11.	Semen centre	1. 12.	Semen centre	ŀ	Holding 🗖		
Part		Name Approval number Address		Name App Address	rova	al number		
		Postal code		Postal code				
	I.13.	Place of loading	1.14.	Date of departure				
	I.15.	Means of transport	I.16.	Entry BIP in EU				
		Aeroplane ☐ Ship ☐ Railway wagon ☐						
		Road vehicle Other	1.17.					
		Identification						
	140	Documentary references		140.0		dit d - // 10 d -)		
	1.18.	Description of commodity		1.19. Com	moc	dity code (HS code) 05 11 99 85		
						I.20. Quantity		
						n.20. Quantity		
	1.21.					I.22. Number of packages		
	1.23.	Seal/Container No				1.24.		
	1.25.	Commodities certified for: Artificial reproduction						
	1.26.	For transit through EU to third country		I.27. For import or admission	into	EU 🔲		
		Third country ISO code						
	1.28.	Identification of the commodities						
		pecies (Scientific name) Donor identity		Date of collection		Quantity		

Equine semen – Section B

	II. Health information			II.a. Certificate reference No	II.b.					
	I, the under	signed, offic	ial veterina	rian, of the	e exporting country (²)(name of exporting co					
					(name or exporting con	тиу)				
	certify that	:								
u	II.1. The semen collection centre (³), in which the semen described above was collected, processed and stored export to the European Union is approved and supervised by the competent authority in accordance with conditions of Chapter I(I)(1) and Chapter I(II)(1) of Annex D to Directive 92/65/EEC,									
Part II: Certification	II.2.				g 30 days prior to the date of first collection of the seme ten semen elapsed, the semen collection centre:	n described above until the				
Part II: 0		II.2.1.			he exporting country or, in the case of regionalisation S/EC ($^{\rm o}$), in that part of the territory of the exporting country					
			 not considered to be infected with African horse sickness in accordance with Article 5(2)(a) and (b) of Directive 2009/156/EC (⁸), 							
			— free	from Ven	nezuelan equine encephalomyelitis for 2 years,					
free from glanders and dourine for 6 months;					nders and dourine for 6 months;					
	II.2.2. fulfilled the conditions for a holding laid down in Article 4(5) of Directive 2009/156/EC (8) and particular:									
		(¹) either	[II.2.2.1.		g a case of a disease mentioned below not all the anima ease located on the holding were slaughtered or killed					
					om any type of equine encephalomyelitis for at least 6 ay on which the equidae suffering from the disease are s					
				re ta	om equine infectious anaemia for at least the period re esult in an agar gel immunodiffusion test (Coggins tes aken after the infected animals were slaughtered on two om each of the remaining animals,	st) carried out on samples				
				— fro	om vesicular stomatitis for at least 6 months from the las	t recorded case,				
				— fr	om rabies for at least one month from the last recorded o	ase,				
				— fr	om anthrax for at least 15 days from the last recorded ca	se,]				
		(¹) or	[II.2.2.1.	disease disinfect encepha the case	g a case of a disease mentioned below all the animals o located on the holding have been slaughtered or ted, the holding has been free for at least 30 days alomyelitis, equine infectious anaemia, vesicular stomatice of anthrax, beginning on the day on which following the infection of the premises was satisfactorily completed;	killed and the premises from any type of equine tis and rabies or 15 days in				
		II.2.3.	contained metritis,	d only equ	uidae which were free of clinical signs of equine viral arte	eritis and contagious equine				
	II.3.	Prior to er	ntering the s	semen col	llection centre the donor stallions and any other equidae	located in the centre:				

Equine semen – Section B

II. He	alth information		II.a. Certificate reference No	II.b.
	II.3.1.	State of the Euro regionalisation a	y resident for 3 months (or since entry if they were pean Union during the 3 months period) in the excording to Article 13 of Directive 2009/156/EC (8) which was during that period	xporting country or, in the case of
			red to be infected with African horse sickness in irective 2009/156/EC (8),	n accordance with Article 5(2)(a)
		— free from Ve	enezuelan equine encephalomyelitis for at least 2 years	ears,
		— free from gl	anders and dourine for at least 6 months;	
(¹) eithe	r [II.3.2.		he country of export which was on the day of a is (VS) for at least 6 months,]	admission into the centre free of
(¹) or	[II.3.2.		o a virus neutralisation test for vesicular stomatii dilution of 1 in 12 on a blood sample taken (⁴) w	
	II.3.3.	originated from h point II.2.2;	oldings which on the day of admission onto the c	centre fulfilled the requirements of
II.4.	The semer	n described above	was collected from donor stallions, which:	
	II.4.1.		any clinical sign of an infectious or contagious dise the day the semen was collected;	ease at the time of admission onto
	II.4.2.		or 30 days prior to the date of semen collection on inical sign of equine viral arteritis or contagious equ	
	II.4.3.		ed for natural mating during at least 30 days prior t dates of the first sample referred to in points II.4.5 lection period;	
	II.4.4.	the Manual of E samples taken in	the following tests, which meet at least the requiring in accordance with one of the programmes specific competent authority:	imals of the OIE, carried out on
	(¹) (⁵) either		r-gel immuno-diffusion test (Coggins test) for equeriersult;]	nine infectious anaemia (EIA) with
	(¹) (⁵) or	[II.4.4.1. an ELI	SA for equine infectious anaemia (EIA) with negativ	ve result;]
and	(¹) either		m neutralisation test for equine viral arteritis (EVA of one in four;]	with negative result at a serum
	(¹) or		isolation test for equine viral arteritis (EVA) carri of the entire semen of the donor stallion;]	ed out with negative result on an

COUNTRY Equine semen – Section B

H.	Health information			II.a. Certificate reference No	II.b.
and		II.4.4.3.	occasion equigenia and from	t identification test for contagious equine metritis (0 s on samples collected with an interval of 7 days talis after a cultivation of 7 to 14 days from pre-ejaculato genital swabs taken at least from the penile sheath, ative result in each case;	by isolation of <i>Taylorella</i> ry fluid or a semen sample
	II.4.5.	have bee	n subject nes (⁶) det	ed with the results specified in II.4.4. in each case to ailed in points II.4.5.1, II.4.5.2 and II.4.5.3 as follows:	o at least one of the test
		II.4.5.1.	30 days semen d	or stallion was continuously resident on the semen co prior to the date of the first collection and during the escribed above, and no equidae on the semen collectic direct contact with equidae of lower health status than the	period of collection of the on centre came during that
			first sem	s described in point II.4.4 have been carried out on sar en collection and at least 14 days following the date of e period of at least 30 days.	
		II.4.5.2.	the date above, b continuo	or stallion was resident on the semen collection centre f of the first collection and during the period of collectio but has left the centre under the responsibility of the sus period of less than 14 days, or other equidae on the intact with equidae of lower health status.	n of the semen described centre veterinarian for a
			date of the	s described in point II.4.4 have been carried out on sar ne first semen collection of the breeding season or colle- lescribed above was collected and at least 14 days cement of the residence period of at least 30 days,	ction period in the year the
	and			described in point II.4.4.1 for equine infectious anaemia of blood taken (4) not more than 90 days before the set;	
	and	(¹) either		ne tests described in point II.4.4.2 for equine viral arteriti aken (4) not more than 30 days before the semen describ	
		(¹) or	aliquot or semen or reacted p	isolation test for equine viral arteritis was carried out of the entire semen of the donor stallion taken (4) not more lescribed above was collected and a blood sample ta positive in a serum neutralisation test for equine viral art n one in four,]	e than 6 months before the ken on the same date (4)
	and			described in point II.4.4.3 for contagious equine metrit taken (4) , not more than 60 days before the semen desc	
		II.4.5.3.	date of the	s described in point II.4.4 have been carried out on sar ne first semen collection of the breeding season or collection escribed above was collected,	
	and			described in point II.4.4 have been carried out on so 0 days after the collection of the semen described above	

Equine semen – Section B

II. Health	information		II.a.	Certificate r	eference No			II.b.		
	II.4.6.	have under following da		ting provide	ed for in po	ints II.3.2 (¹)	and II.4.5	on samples ta	aken on the	
of	0	Start	date (4)		ı	ests (4)				
Identification of semen	Test	Donor	Semen	VS (1)	EIA		EVA II. 4.4.2.		CEM II.4.4.3.	
Identif	T prog	residence	collection	II.3.2	II.4.4.1.	Blood sample	Semen sample	1. sample	2. sample	
(1) either	[II.5.	No antibiotion	s were added	I to the sem	en;]					
(1) or	[II.5.		g antibiotic or en of not less		n of antibioti	cs was added	I to produce	a concentratio	n in the final	
									;]	
II.6.	The seme	n described a	bove was:							
	II.6.1.		rocessed, stor l)(1) and III(I)				which comp	ly with the req	uirements of	
	II.6.2.		place of load Directive 92/6					oint 1.4 of Cha 23.	apter III(I) of	
Notes										
Part I:										
Box I.11.:	The place	of origin shall	correspond to	o the semer	collection co	entre of the s	emen origin.			
Box I.22.:	The numb	er of package	s shall corres	pond to the	number of co	ontainers.				
Box I.23.:	The identi	fication of con	tainer and sea	al number sl	nall be indica	ited.				
Box I.28.:	The donor	ridentity shall	correspond to	the official	identification	of the anima	l.			
	The date	of collection sh	nall be indicate	ed in the foll	owing forma	t: dd/mm/yyy	<i>/</i> .			

Equine semen - Section B

II. Health information	II.a. Certificate reference No	II.b.
Part II:		
Guidance for the completion of the table in po	pint II.4.6.	

Αb

b	breviations:	
	VS	Vesicular stomatitis (VS) testing if required in accordance with point II.3.2
	EIA-1	Equine infectious anaemia (EIA) testing first occasion
	EIA-2	EIA testing second occasion
	EVA-B1	Equine viral arteritis (EVA) testing on blood sample first occasion
	EVA-B2	EVA testing on blood sample second occasion
	EVA-S1	EVA testing on semen sample first occasion
	EVA-S2	EVA testing on semen sample second occasion
	CEM-11	Contagious equine metritis (CEM) testing first occasion first sample
	CEM-12	CEM testing first occasion second sample taken 7 days after CEM-11
	CEM-21	CEM testing second occasion first sample
	CEM-22	CEM testing second occasion second sample taken 7 days after CEM-21
٠.	tructions:	

Instructions:

For each semen identified in column A in correspondence with Box I.28, the test programme (II.4.5.1., II.4.5.2. and/or II.4.5.3.) must be specified in column B, and columns C and D must be completed with the dates required.

The dates when samples were taken for laboratory testing prior to the first collection of the semen described above as required in II.4.5.1., II.4.5.2. and II.4.5.3., are entered in the upper line of columns 5 to 9 of the table, this being the boxes marked with EIA-1, EVA-B1 or EVA-S1 and CEM-11 and CEM-12 in the example below.

The dates when samples were taken for repeat laboratory testing as required in accordance with II.4.5.2. or II.4.5.3. are entered in the lower line of columns 5 to 9 in table, this being the boxes EIA-2, EVA-B2 or EVA-S2 and CEM-21 and CEM-22 in the example below.

of	0	Stan	t date	Date of sampling for health tests						
Identification	Test	Donor	Semen collection	VS II.3.2.	EIA II.4.4.1.	EVA II.4.4.2.		CEM II.4.4.3.		
Identi	buo	residence				Blood sample	Semen sample	1. sample	2. sample	
	В		C D \	\ <u>'</u> C	EIA-1	EVA-B1	EVA-S1	CEM-11	CEM-12	
A				VS	EIA-2	EVA-B2	EVA-S2	CEM-21	CEM-22	

^{(&}lt;sup>1</sup>) Delete as necessary.

Imports of equine semen are authorised from a third country listed in column 2 of Annex I to Commission Implementing Regulation (EU) 2018/657 provided the semen was collected in the part of the territory of the third country detailed in column 4 of that Annex from a donor stallion of the category of equidae indicated in columns 11, 12 or 13 of that Annex.

▼<u>C2</u>

COUNTRY Equine semen – Section B

II.	Health information	II.a. Certificate reference No	II.b.					
(3)	Only approved semen collection centres listed in accordance with Article 17(3)(b) of Council Directive 92/65/EEC on the Commission website: http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm							
(4)	Insert date in table in point II.4.6 (follow	Guidance in Part II of the Notes)						
(5)	The agar gel immunodiffusion test (Coggins test) or the ELISA for equine infectious anaemia are not required for donor equidae which have continuously resided in Iceland since birth, provided that Iceland has remained officially free of equine infectious anaemia and no equidae and their semen, ova and embryos have been introduced into Iceland from outside prior to and during the period the semen was collected.							
(⁶)	Cross out the programmes that do not	apply to the consignment.						
(7)	Insert names and concentrations.							
(8)	OJ L 192, 23.7.2010, p. 1.							
_	The signature and the stamp must be in	n a different colour to that of the printing.						
Offi	cial veterinarian							
	Name (in capital letters):	Qu	ualification and title:					
	Date: Signature:							
	Stamp:							

Section C

MODEL 3 – Model health certificate for imports of consignments of stocks of semen of equidae collected, processed and stored in accordance with Directive 92/65/EEC before 1 September 2010 and dispatched after 31 August 2010 from an approved semen collection centre of origin of the semen

UN.	ΓRY:									Veterinary	certifica	te to EU
	l.1.	Consignor Name						Certificate referen		I.2.a.		
		Address					I.3. Central competent authority					
		Tel.					1.4.	Local competent a	authority			
Part I : Details of dispatched consignment	1.5.	Consignee Name Address					1.6.	Person responsib Name Address	le for the load	in EU		
tched co		Postal code Tel.						Postal code Tel.				
s of dispat	1.7.	Country of origin	ISO code		Region of origin	Code	1.9.	Country of destination	ISO code	I.10. Region o destination		Code
: I : Detail	I.11. Place of origin Semen centre □				l.12.	Place of destination		ding 🗖				
Parl	Name Approval number Address				Name Address	Approval	number					
		Postal code						Postal code				
	I.13. Place of loading			I.14.	Date of departure							
	I.15.	15. Means of transport Aeroplane □ Ship □ Railway wagon □		on \square	I.16.	Entry BIP in EU						
		Road vehicl Identification	e 🗆	Oth	er 🗖	on ப	1.17.					
	I.18.	Description	of commodity	/					I.19. Commo	dity code (HS o		
										I.20. Quantity	<i>'</i>	
	I.21.									I.22. Number	of packa	ages
	1.23.	Seal/Contai	ner No							1.24.		
	1.25.	Commoditie	es certified for	: 								
	1.26.	For transit th	hrough EU to	third c	country			I.27. For import or	admission int	o EU 🔲		
		Third countr	ry I	SO cod	de							
		Identification	n of the comm	noditie	s Donor id	entity	I	Date of coll	ection	Qu	antity	

Equine semen - Section C

	II. Health information		II.a. Certificate reference No	II.b.						
	I, the unders	signed, official veterinarian, of the ex	porting country (²)	-						
	(name of exporting country)									
	certify that:									
uo	II.1.	I.1. The semen collection centre in which the semen described above was collected, processed and stored for export to the European Union:								
ertificati	II.1.1.	is approved and supervised by the competent authority according to the conditions of Chapter I, Annex D to Directive 92/65/EEC,								
Part II: Certification	II.1.2.	II.1.2. is situated in the territory or in the case of regionalisation according to Article 13 of Directive 2009/156/EC (⁶) in a part of the territory of the country of export which was on the day the semen was collected until the date of dispatch free of:								
		 African horse sickness, in ac 	cordance with EU legislation,							
		 Venezuelan equine encepha 	lomyelitis for 2 years,							
		 glanders and dourine for 6 m 	onths;							
	II.1.3.		g 30 days prior to the date of collection of the sem or animal health reasons which laid down one of th							
	II.1.3.1.	if not all the animals of species suprohibition lasted for:	usceptible to the disease located in the holding w	vere slaughtered or killed, the						
		 6 months, beginning on the case of equine encephalomy 	day on which the equidae suffering from the diselitis,	sease are slaughtered, in the						
			out with negative result two Coggins tests 3 animals have been slaughtered, in the case of infe							
		 6 months, in the case of vesi 	cular stomatitis,							
		 one month from the last reco 	rded case, in the case of rabies,							
		 15 days from the last recorded 	ed case, in the case of anthrax.							
	II.1.3.2.	the premises disinfected, the proh	otible to the disease located in the holding have be ibition lasted for 30 days, or 15 days in the case tion of the animals the disinfection of the premises	of anthrax, beginning on the						
	II.1.4.		mencing 30 days prior to semen collection and free of clinical signs of equine viral arteritis and co							
	II.2.	Prior to entering the semen collecti	ion centre the donor stallions and any other equid	ae located in the centre:						
	II.2.1.		nonths (or since entry if they were directly importe in the territory or in the case of regionalisation in that period free of:							
		 African horse sickness, in ac 	cordance with EU legislation,							
		 Venezuelan equine encepha 	lomyelitis for 2 years,							
		 glanders for 6 months, 								
		dourine for 6 months;								

Equine semen - Section C

II. Health information			II.a. Certificate reference No	II.b.				
(¹) either	[II.2.2. originated from the territory of the country of export which was on the day of admission into the free of vesicular stomatitis for 6 months,]							
(¹) or	[II.2.2.	[II.2.2. were tested by a virus neutralisation test for vesicular stomatitis in a blood sample taken on						
II.2.3.	originated	from holdings which on	the day of admission onto the centre fulfilled the r	equirements of point II.1.3;				
II.3.	The seme	en described above was o	collected from donor stallions, which:					
II.3.1.	on the da	y the semen was collecte	ed have not shown clinical signs of an infectious or	r contagious disease,				
II.3.2.	during at	least 30 days prior to coll	ection of the semen have not been used for natur	al service,				
II.3.3.		e last 30 days prior to d linical signs of equine vira	collection of the semen have been kept on hold al arteritis,	ings where no equine animal				
II.3.4.		e last 60 days prior to d linical signs of contagious	collection of the semen have been kept on hold sequine metritis,	ings where no equine animal				
II.3.5.			as far as I could ascertain have not been in conta e the 15 days immediately preceding the collection					
II.3.6.			nimal health tests carried out in a laboratory r t programme as specified in point II.3.7:	recognised by the competent				
II.3.6.1.	an agar-g	el immuno-diffusion test	(Coggins test) for equine infectious anaemia with	negative result (³);				
(1) either	[11.3.6.2.	a serum neutralisation	test for equine viral arteritis with negative result a	t a serum dilution of 1 in 4;]				
(¹) or	[11.3.6.2.	a virus isolation test fo semen;]	r equine viral arteritis carried out with negative re	sult on an aliquot of the entire				
II.3.6.3.	Taylorella	equigenitalis from pre-e	ritis carried out on two occasions with an inter jaculatory fluid or a semen sample and from ger n the urethral fossa with negative result in each ca	ital swabs taken at least from				
II.3.7.	have bee	n subjected to one of the	following test programmes (5):					
II.3.7.1.	collection	, and during the collection	isly resident on the collection centre for at leas on period, and no equidae on the collection cent or health status than the donor stallions.					
		4 days after the comme	ave been carried out on samples taken on encement of the above residence period and at					
II.3.7.2.			ously resident on the collection centre or other ed lae of lower health status than the donor stallions.	quidae on the collection centre				
			ave been carried out on samples taken one first semen collection and at least at the beginn					
		required in point II.3.6.1 vn n was collected on	was last carried out on a sample of blood taken n(⁴);	ot more than 120 days before				
(1) either		required in point II.3.6.2	was last carried out not more than 30 days befor	e the semen was collected on				

Equine semen - Section C

II.	Health i	nformation	II.a. Certificate referer	nce No	II.b.					
(¹) or		[The non-shedder state of the sero which was carried out not more that								
II.3.7	.3.	The tests required in point II.3.6 h semen and not less than 14 days a on								
II.4.		The semen described above was with the requirements of Chapter II			nder conditions which comply					
Notes										
Part I	l:									
Box I	.11.:	The place of origin shall correspon	to the semen collectio	n centre of the semen orig	gin.					
Box I	.22.:	The number of packages shall corr	spond to the number of	of containers.						
Box I	.23.:	The identification of container and	eal number shall be ind	dicated.						
Box I	.28.:	The donor identity shall correspond	to the official identifica	tion of the animal.						
		The date of collection shall be indic	ate in the following forn	nat: dd/mm/yyyy.						
Part	II:									
(¹)	Delete	as necessary.								
(2)	Regula	rts of equine semen are authorised from a third country listed in column 2 of Annex I to Commission Implementing lation (EU) 2018/659 provided the semen was collected in the part of the territory of the third country detailed in nn 4 of that Annex from a donor stallion of the category of equidae indicated in columns 11, 12 or 13 of that Annex.								
(3)	equida equine	the agar gel immunodiffusion test (Coggins test) or the ELISA for equine infectious anaemia are not required for donor quidae which have continuously resided in Iceland since birth, provided that Iceland has remained officially free of quine infectious anaemia and no equidae and their semen, ova and embryos have been introduced into Iceland from utside prior to and during the period the semen was collected.								
(⁴)	Insert date.									
(⁵)	Cross out the programmes that do not apply to the consignment.									
(⁶)	OJ L 192, 23.7.2010, p. 1.									
_	The sig	he signature and the stamp must be in a different colour to that of the printing.								
Official veterinarian										
	Name	in capital letters):		(Qualification and title:					
	Date:			S	Signature:					
	Stamp:									

Section D

MODEL 4 - Model health certificate for imports of consignments of semen of equidae collected, processed and stored in accordance with Directive 92/65/EEC after 30 September 2014 and of consignments of stocks of semen of animals of the equine species collected, processed and stored in accordance with Directive 92/65/EEC after 31 August 2010 and before 1 October 2014 or before 1 September 2010 and dispatched after 31 August 2010 from an approved semen storage centre

COUNTRY: Veterinary certificate to											
	l.1.	Consignor Name Address		I.2.	Certificate referen	ce No	I.2.a.				
				I.3. Central competent authority							
nsignment		Tel.		1.4.	I.4. Local competent authority						
	1.5.	Consignee Name Address		I.6.	Person responsibl Name Address	e for the load	in EU				
atched co		Postal code Tel.			Postal code Tel.						
Part I : Details of dispatched consignment	1.7.	Country of ISO code origin	I.8. Region of origin	Code	1.9.	Country of destination	ISO code	I.10. Region of destination	Code		
	l.11.	Place of origin Semen centre			l.12.	Place of destination		olding \square			
Parl		Name Approval number Address			Name Address	Appro	val number				
		Postal code				Postal code					
	I.13.	. Place of loading		I.14.	Date of departure						
	I.15.	.15. Means of transport Aeroplane □ Ship □ Railway wagon □ Road vehicle □ Other □		I.16.	Entry BIP in EU						
				l.17.	No(s) of CITES						
		Identification Documentary references									
	I.18.	18. Description of commodity				I.19. Commodity code (HS code) 05 11 99 85					
								I.20. Quantity			
	I.21.	21.						I.22. Number of packa	iges		
	I.23.	3. Seal/Container No						1.24.			
	1.25.	Commodities certified for:									
		Artificial reproduction									
	1.26.	For transit through EU to third country			I.27. For import or admission into EU						
		Third country IS	O code								
	1.28.	Identification of the comm	odities								
	s	pecies (Scientific name)	Donor iden	ntity		Date of coll	ection	Quantity			

COUNTRY

Equine semen - Section D

	COUNTRY			Equine semen – Section D						
	II. Health	information	II.a. Certificate reference No	II.b.						
	I, the unders	igned official veterinarian of the exp	porting country (2)	hereby						
	(name of exporting country)									
certify that:										
	II.1.	The centre (3) described in Boy I 1	11 at which the semen to be exported to the Union	was stored:						
=	11.1.	The centre () described in Box i.	That which the sement to be experted to the officin	was stored.						
tificatio	(1) either		laid down in Chapter I(I)(1) and is operated and s wn in Chapter I(II)(1) of Annex D to Directive 92/65/							
Part II: Certification	(¹) or		laid down in Chapter I(I)(2) and is operated and s wn in Chapter I(II)(2) of Annex D to Directive 92/65/							
Pa	II.2.	The semen to be exported to the	Union:							
	II.2.1.		nd stored for a minimum period of 30 days immedia $(^5)$ operated and supervised in accordance with which is							
	(¹) either	[located in the exporting country;]								
	(¹) or	[located in								
	II.2.2.	was moved to the centre described in Box I.11 under conditions at least as strict as described in:								
	(1) either	[Model 1 in Section A of Part 1 of	Annex III to Regulation (EU) 2018/657 (⁶);]							
	(1) or	[Model 2 in Section B of Part 1 of	Annex III to Regulation (EU) 2018/657 (⁶);]							
	(1) or	[Model 3 in Section C of Part 1 of	Annex III to Regulation (EU) 2018/657 (6);]							
	(1) or	[Model 1 in Section A of Part 2 of	Annex II to Decision 2010/471/EU (⁶);]							
	(1) or	[Model 2 in Section B of Part 2 of	Annex II to Decision 2010/471/EU (6);]							
	(1) or	[Model 3 in Section C of Part 2 of	Annex II to Decision 2010/471/EU (6);]							
	(1) or	[Commission Decision 96/539/EC	(⁶);]							
	II.2.3.		h satisfy the terms of Annex D to Directive 92/65/El							
	II.2.4.	sent to the place of loading in a Directive 92/65/EEC and bearing	sealed container in accordance with point 1.4 o the number indicated in Box I.23.	f Chapter III(I) of Annex D to						
	Notes									
	Part I:									
	Box I.11.:	The place of origin shall correspond	nd to the semen storage centre of semen dispatch.							
	Box I.17.:	described above from the approv	dual official document(s) or health certificate(s) to red semen collection centre of its origin to the desoriginal(s) of this/these document(s) or certificate tached to this certificate.	scribed above semen storage						

Signature:

▼<u>C2</u>

Date:

Stamp:

COUNTRY Equine semen - Section D П. Health information II.a. Certificate reference No II.b Box I.22.: The number of packages shall correspond to the number of containers. Box I.23.: The identification of container and seal number shall be indicated. Box I.28.: The donor identity shall correspond to the official identification of the animal. The date of collection shall be indicated in the following format: dd/mm/yyyy. Part II: (¹) Delete as necessary Imports of equine semen are authorised from a third country listed in column 2 of Annex I to Commission Implementing Regulation (EU) 2018/657 provided that the semen was collected in the part of the territory of the third country detailed in column 4 of that Annex from a donor stallion of the category of equidae indicated in column 11, 12 or 13 of that Annex. $(^{3})$ Only approved semen collection or storage centres listed in accordance with Article 17(3)(b) of Directive 92/65/EEC on the Commission website: http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm Council Directive 92/65/EEC of 13 July 1992 laying down animal health requirements governing trade in and imports into the Community of animals, semen, ova and embryos not subject to animal health requirements laid down in specific Community rules referred to in Annex A(I) to Directive 90/425/EEC (OJ L 268, 14.9.1992, p. 54). Only approved semen collection centres listed in accordance with Article 11(4) and Article 17(3)(b) of Directive 92/65/EEC on the Commission websites: https://ec.europa.eu/food/animals/live_animals/approved-establishments_en; http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm The original(s) of the document(s) or the health certificate(s) or the officially endorsed copy/copies thereof that accompanied the semen described above from the approved semen collection centre of the semen origin to the centre of the semen dispatch described in Box I.11 must be attached to this certificate. The signature and the stamp must be in a different colour to that of the printing Official veterinarian Name (in capital letters): Qualification and title:

PART 2

Model health certificate for imports of ova and embryos

Section A

MODEL 1- Model health certificate for imports of consignments of ova and embryos of equidae collected or produced in accordance with Directive 92/65/EEC after 30 September 2014 and dispatched by an approved embryo collection or production team of origin of the ova or embryos

IIKY:					veterinary certifica	ate to Eu	
l.1.	Consignor Name	1.2.	Certificate reference	ce No	I.2.a.		
	Address	1.3.	Central competent	authority			
	Tel.	1.4.	Local competent a	uthority			
1.5.	Consignee Name Address	1.6.	Person responsible Name Address	e for the load	in EU		
	Postal code Tel.		Postal code Tel.				
1.7.	Country of ISO code I.8. Region of Code origin configure	1.9.	Country of destination	ISO code	I.10. Region of destination	Code	
l.11.	Place of origin Embryo team □	1.12	Place of destination	n Embryo team			
	Name Approval number Address		Name Address	Approval num	nber		
	Postal code	Postal code					
I.13.	B. Place of loading		Date of departure				
I.15.	Means of transport	1.16	. Entry BIP in EU				
	Aeroplane Ship Railway wagon Road vehicle Other Identification Documentary references	1.17					
I.18.	Description of commodity			I.19. Commod	dity code (HS code) 05 11 99 85		
					I.20. Quantity		
I.21.					I.22. Number of pack	ages	
1.23.	Seal/Container No				1.24.		
1.25.	I.25. Commodities certified for: Artificial reproduction						
1.26.	For transit through EU to third country Third country ISO code		I.27. For import or a	admission into	o EU 🔲		
1.28.	Identification of the commodities						
s	pecies (Scientific Category D name)	onor	identity Da	ate of collection	on Quantity	/	

	II. Health information			II.a. Certificate reference No	II.b.					
	I, the undersigned, official veterinarian, of the exporting country (²)									
	certify that:									
tion	II.1.	1. The ova (¹)/embryos (¹) described above:								
Part II: Certification	II.1.2.	II.1.2. were collected (¹)/produced (¹) by the team (³) described in Box I.11, which has been approved and supervised in accordance with Chapter I(III) of Annex D to Directive 92/65/EEC (⁴) and is subject to inspection by an official veterinarian at least once every calendar year;								
Part	II.1.3.	were collected (¹)/produced (¹), processed and stored in accordance with the requirements of Chapter III(II) of Annex D to Directive 92/65/EEC;								
	II.1.4.		ected at a place separa nd disinfected prior to th	ated from other parts of the premises or ho he collection;	olding which is in good repair and was					
	II.1.5. were examined, processed and packed in laboratory facilities which are not situated in a zone subjeting prohibition or quarantine measures as set out in Box II.1.6, in a section which is separated from the section storing equipment and materials used in contact with donor animals and from the area where the donor an are handled;									
	II.1.6.	come from	n donor mares which:							
		II.1.6.1. were continuously resident for a period of 3 months (or since entry if they were directly import a Member State of the Union during the 3 months period) in the exporting country or, in the regionalisation in accordance with Article 13 of Directive 2009/156/EC(5), in that part of the ter the exporting country which was during that period								
				to be infected with African horse sicknetive 2009/156/EC,	ess in accordance with Article 5(2)(a)					
			free from Venez	zuelan equine encephalomyelitis for a perio	d of at least 2 years,					
			 free from glande 	ers and dourine for a period of at least 6 mo	onths;					
	(1) either	[II.1.6.2.	originated from a cou (VS) for a period of at	untry of export which was on the day of c t least 6 months;]	ollection free from vesicular stomatitis					
	(¹) or [II.1.6.2. were subjected to a virus neutralisation test for vesicular stomatitis (VS) carried out with a negative result in according with the relevant Chapter of the Manual of Diagnostic Tests and Vaccines for Terrestrial Aniithe OIE on a blood sample taken on									
	(1) either	[II.1.6.3.	[II.1.6.3. during a period of the past 30 days prior to the date of the collection were located in holdings ur veterinary supervision which fulfilled from the day of the collection of the ova (¹)/embryos (¹) until date of their dispatch the conditions for a holding laid down in Article 4(5) of Directive 2009/156/ and in particular:]							
	(¹) or	[II.1.6.3. in the case of frozen ova (¹)/embryos (¹), during a period of the past 30 days prior to the date of the collection were kept in holdings under veterinary supervision which fulfilled, from the day of the collection of the ova (¹)/embryos (¹) until the end of the period of 30 days mandatory storage at approved premises, the conditions for a holding laid down in Article 4(5) of Directive 2009/156/EC, and in particular:]								

			· · · · · · · · · · · · · · · · · · ·
II. Health information	on	II.a. Certificate reference No	II.b.
(¹) either	[II.1.6.3.1.	following a case of a disease mentioned belo susceptible to that disease located in the holding holding has been free:	
		 from any type of equine encephalomyelitis to beginning on the day on which the equidate slaughtered, 	
		 from equine infectious anaemia for at least the result in an agar gel immunodiffusion test (AG samples taken after the infected animals w 3 months apart from each of the remaining equ 	GID or Coggins tests) carried out or ere slaughtered on two occasions
		 from vesicular stomatitis for a period of at lead case, 	ast 6 months from the last recorded
		— from rabies for a period of at least one month fr	om the last recorded case,
		 from anthrax for a period of at least 15 days fro 	m the last recorded case,]
(¹) or	[II.1.6.3.1.	following a case of a disease mentioned below all that disease located in the holding were slaugi disinfected, the holding was free for a period of at le encephalomyelitis, equine infectious anaemia, we period of at least 15 days in the case of anthra following the destruction of the animals the satisfactorily completed;]	htered or killed and the premises east 30 days from any type of equine esicular stomatitis and rabies or a ax, beginning on the day on which
II.1.6.4.		od of the past 30 days prior to the collection the ova e of the equidae has shown clinical signs of contagion;	
II.1.6.5.	of the ova (d for natural breeding during a period of at least 30 d)/embryos (¹) and between the date of the first sar and the date of the collection of the ova (¹)/embryos	mples referred to in points II.1.6.6.
II.1.6.6.	Manual of Di which is reco	one the tests, which meet at least the requirement agnostic Tests and Vaccines for Terrestrial Animals of gnised by the competent authority and has the tests equivalent to that provided for in Article 12 of R	of the OIE, carried out in a laboratory referred to hereinafter included in its
	(⁸) [II.1.6.6.1.	for equine infectious anaemia (EIA), an agar-g Coggins test) or an enzyme-linked immunosorbent carried out on a blood sample taken onbeing not less than 14 days following the date referred to in point II.1.6.5, and the test was last car	assay (ELISA) with a negative resul
	II.1.6.6.2.	for contagious equine metritis (CEM), an agent i negative result on at least two specimens (swabs) i point II.1.6.5 from at least the mucosal surfaces sinuses of the donor mare	taken during the period referred to ir
	(¹) either	[II.1.6.6.2.1. on two occasions with an inte on	

II. Health	information		II.a.	Certificate reference	e No	II.b.
				on one occasion or	1 (⁶)	, in the case of detection of
					carried out within 48 ho	olymerase chain reaction (PCR) purs after taking the specimens
		earlier treatm	than ent of	7 days (systemic tre the donor stallion	atment) or 21 days (loc	6.6.2.2 were in no case taken all treatment) after antimicrobial ansport medium with activated laboratory.
	II.1.6.7.					n contact with equidae suffering ays immediately preceding the
	II.1.6.8.	on the day of the co contagious disease;	llectio	n of the ova (¹)/emb	oryos (1) did not show o	clinical signs of an infectious or
II.1.7.		ected (¹)/produced (¹) a vas approved by the co				production (1) team described in
II.1.8.	collection		d we	re transported und		30 days immediately after their atisfy the terms laid down in
II.2.	The embryos described above were conceived by artificial insemination(1)/as a result of <i>in vitro</i> fertilisation (1) using semen meeting the requirements of Directive 92/65/EEC and coming from semen collection centres approved in accordance with Article 11(2) or 17(3)(b) of Directive 92/65/EEC (9) and located respectively in a Member State of the Union or in a third country or parts of the territory of a third country listed in columns 2 and 4 of Annex I to Commission Implementing Regulation (EU) 2018/659 from which the import of equine semen collected from registered horses, registered equidae or equidae for breeding and production is authorised in accordance with Article 4 of Commission Implementing Regulation (EU) 2018/659 and indicated in columns 11, 12 and 13 of Annex I thereto. (10) (11);					
(¹²) [II.3.					ped above comply with up in points II.1.1 to II.1	the requirements of Annex D to .8 of this certificate.]
Notes						
Part I:						
Box I.11.:	ova/embry		duced	, processed, stored	and approved in acco	production team by which the ordance with Article 17(3)(b) of
	http://ec.e	uropa.eu/food/animal/s	emen	_ova/equine/index_e	en.htm	
Box I.22.:	The numb	er of packages shall co	rresp	ond to the number o	containers.	
Box I.23.:	The identi	fication of container an	d seal	number shall be ind	icated.	
Box I.28.:		gory: specify if <i>in v</i> ipulated embryos.	ivo d	erived embryos, <i>in</i>	vivo derived ova, in	n vitro produced embryos or
	The donor	r identity shall correspo	nd to	the official identificat	ion of the animal.	
	The date of	of collection shall be in	dicate	in the following form	at: dd/mm/yyyy.	

COUNTRY	Equine ova/embryos	
II. Health information	II.a. Certificate reference No	II.b.

Part	II:					
(¹)	Delete as appropriate.					
(2)	Only third countries or parts of the territory of third countries listed in columns 2 and 4 of Annex I to Commission Implementing Regulation (EU) 2018/659, respectively from which imports of registered equidae and equidae for breeding and production are also authorised and as indicated in column 14 of Annex I thereto.					
(3)	Only approved embryo collection teams and embryo production teams listed in accordance with Article 17(3)(b) of Directive 92/65/EEC on the Commission website:					
	http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm					
(4)	Council Directive 92/65/EEC of 13 July 1992 laying down animal health requirements gover the Community of animals, semen, ova and embryos not subject to animal health requirements governmently rules referred to in Annex A(I) to Directive 90/425/EEC (OJ L 268, 14.9.1992,	uirements laid down in specific				
(⁵)	Council Directive 2009/156/EC of 30 November 2009 on animal health conditions importation from third countries of equidae (OJ L 192, 23.7.2010, p. 1).	governing the movement and				
(⁶)	Insert date. (follow Guidance in Part II of the Notes).					
(⁷)	Regulation (EC) No 882/2004 of the European Parliament and of the Council of 29 April 2004 on official controls performed to ensure the verification of compliance with feed and food law, animal health and animal welfare rules (OJ L 165, 30.4.2004, p. 1).					
(8)	The agar gel immunodiffusion test (AGID or Coggins test) or the ELISA for equine infectious anaemia are not required for donor equidae which have continuously resided in Iceland since birth, provided that Iceland has remained officially free of equine infectious anaemia and no equidae and their semen, ova and embryos have been introduced into Iceland from outside prior to and during the period the ova or embryos were collected and the semen was used for fertilisation.					
(⁹)	Only approved semen collection centres listed in accordance with Article 11(4) or Article on the Commission websites:	17(3)(b) of Directive 92/65/EEC				
	https://ec.europa.eu/food/animals/live_animals/approved-establishments_en; http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm					
(10)	Imports of equine semen are authorised from third countries listed in column 2 of Annex Regulation (EU) 2018/659 provided that the semen was collected in the part of the territor column 4 from a donor stallion of the category of equidae positively indicated in column 11	ry of the third country detailed in				
(11)	Does not apply to ova.					
(12)	Delete if none of the embryos in the consignment was produced by in vitro fertilisation of o	va.				
_	The signature and the stamp must be in a different colour to that of the printing.					
Offic	al veterinarian					
	Name (in capital letters):	Qualification and title:				
	Date:	Signature:				
	Stamp:					

Section B

MODEL 2 – Model health certificate for imports of consignments of stocks of ova and embryos of equidae collected, processed and stored in accordance with Directive 92/65/EEC after 31 August 2010 and before 1 October 2014 and dispatched after 31 August 2010 by an approved embryo collection or production team of origin of the ova or embryos

COUN	UNTRY: Veterinary certificate to EU								
	I.1. Consignor Name					Certificate referen	ce No	I.2.a.	
		Address			1.3.	Central competen	t authority		
		Tel.	1.4.	Local competent a	authority				
Part I : Details of dispatched consignment	1.5.	Consignee Name Address				Person responsibl Name Address	le for the load	in EU	
ched co		Postal code Tel.				Postal code Tel.			
of dispat	1.7.	Country of ISO code origin	I.8. Region of origin	Code	1.9.	Country of destination	ISO code	I.10. Region of destination	Code
tails	1.11.	Place of origin	1		1.12.	Place of destination	on		
 D		Embryo team □				Holding \square	Embryo team	n 🗆	
Part		Name Approval number Address				Name Address	Approval nun	mber	
		Postal code			Postal code				
	I.13.	Place of loading			I.14. Date of departure				
	I.15.	Means of transport			I.16. Entry BIP in EU				
		Aeroplane Ship C Road vehicle Ot Identification Documentary references	her 🗆	n 🗖	1.17.				
	I.18.	Description of commodit	у		I.19. Commodity code (HS code) 05 11 99 85				
						·		I.20. Quantity	
	I.21.							I.22. Number of page	ckages
	1.23.	Seal/Container No						1.24.	
	I.25. Commodities certified for: Artificial reproduction								
	I.26. For transit through EU to third country				I.27. For import or admission into EU				
	Third country ISO code								
	1.28.	Identification of the com	nodities						
	S	Species (Scientific name)	Category	D	onor	identity D	ate of collection	on Quant	iity

	II. Health information			II.a. Certificate reference No	II.b.					
	I, the unders	signed, offici	al veterina	rian, of the exp	orting country (²)	-				
	(name of exporting country)									
	certify that:									
	II.1.	The ova ()/embryos	(¹) described a	bove:					
uo		II.1.2.	supervise	were collected (¹)/produced (¹) by the team (³) described in Box I.11, which has been approved supervised in accordance with Chapter I(III) of Annex D to Directive 92/65/EEC and is subjection by an official veterinarian at least once every calendar year;						
Part II: Certification	II.1.3. were collected (¹)/produced (¹), processed and stored in accordance with the re Chapter III(II) of Annex D to Directive 92/65/EEC;					ce with the requirements of				
Part II: (II.1.4.			e separated from other parts of the premises or h sinfected prior to the collection;	nolding which is in good repair				
		II.1.5.	to prohib	ition or quaran	sed and packed in laboratory facilities which are tine measures as set out in Box II.1.6, in a sectio pment and materials used in contact with donc s are handled;	n which is separated from the				
		II.1.6.	come fro	m donor mares	s which:					
			II.1.6.1.	Member Stat or, in the cas	ously resident for 3 months (or since entry if they e of the European Union during the 3 months pe e of regionalisation according to Article 13 of Directions of the exporting country which was during t	eriod) in the exporting country ective 2009/156/EC (8), in that				
					nsidered to be infected with African horse s 5(2)(a) and (b) of Directive 2009/156/EC,	sickness in accordance with				
				— free fro	m Venezuelan equine encephalomyelitis for at le	ast 2 years,				
				— free fro	m glanders and dourine for at least 6 months;					
		(1) either	[II.1.6.2.		om a country of export which was on the day of at least 6 months;]	of collection free of vesicular				
		(¹) or	[II.1.6.2.		by a virus neutralisation test for vesicular stoma					
		(¹) either	[II.1.6.3.	supervision v	ast 30 days prior to collection have been located which fulfilled from the day of collection of ova (1 the conditions for a holding laid down in Article 4 lar:])/embryos (¹) until the date of				
		(¹) or	[II.1.6.3.	supervision v of frozen ov premises ela	ast 30 days prior to collection have been located which fulfilled from the day of collection of ova (1) a (1)/embryos (1), the period of 30 days man apsed, the conditions for a holding laid down and in particular:]	/embryos (1) until, in the case				

II. Health information			II.a. Certificate reference No	II.b.
	(¹) either	[II.1.6.3.1.	following a case of a disease mentioned below susceptible to the disease located on the holding the holding has been free:	
			 from any type of equine encephalomye beginning on the day on which the equidae slaughtered, 	
			 from equine infectious anaemia for at least negative result in an agar gel immunodiffusi out on samples taken after the infected two occasions 3 months apart from each of 	on test (Coggins tests) carried animals were slaughtered on
			 from vesicular stomatitis for at least 6 month 	s from the last recorded case,
			— from rabies for at least one month from the l	ast recorded case,
			— from anthrax for at least 15 days from the la	st recorded case,]
	(¹) or	[II.1.6.3.1.	following a case of a disease mentioned below susceptible to the disease located in the holdin killed and the premises disinfected, the holding 30 days from any type of equine encephalomyelity vesicular stomatitis and rabies or 15 days in the the day on which following the destruction of the premises was satisfactorily completed;]	ng have been slaughtered or g has been free for at least its, equine infectious anaemia, case of anthrax, beginning on
	II.1.6.4.		ast 30 days prior to collection have been kept in m clinical signs of contagious equine metritis for	
	II.1.6.5.	collection	een used for natural breeding during at least ova or embryos and between the date of the 6 and II.1.6.7 and the date of the collection of ova	e first samples referred to in
	II.1.6.6.	test) or an on collection of	subjected with negative result to an agar-gel in ELISA for equine infectious anaemia carried community. (4), being during the past 30 days ova or embryos and the test was last carried out	out on a blood sample taken prior to the date of the first on a sample of blood taken on
	II.1.6.7.	isolation or negative re the first col sinuses on and on an	subjected to an agent identification test for control and agent after a cultivation of 7 ults in each case on samples taken during the pasterion of ova or embryos from mucosal surfaces of two consecutives oestrus periods on additional culture specimen taken during one of cervix on agents.	to 14 days carried out with ast 30 days prior to the date of of the clitoral fossa and clitoral (4), and on
	II.1.6.8.	equidae su	of my knowledge and as far as I could ascertain, rering from an infectious or contagious disease d e collection;	
	II.1.6.9.	have on the	day of collection of ova (1)/embryos (1) not shows s disease;	n clinical signs of an infectious
II.1.7.			luced (1) after the date on which the embryo colvas approved by the competent authority of the expression of the express	

IJЬ

COUNTRY Equine ova/embryos II.a. Certificate reference No

II.1.8.	were processed and stored under approved conditions for at least 30 days immediately after their collection (¹)/production (¹), and were transported under conditions which satisfy the terms laid down in Chapter III(II) of Annex D to Directive 92/65/EEC;
---------	---

- 11.2. The embryos described above were conceived by artificial insemination (1)/as a result of in vitro fertilisation (1) using semen meeting the requirements of Directive 92/65/EEC and coming from semen collection centres approved in accordance with Article 11(2) or 17(3)(b) of Directive 92/65/EEC and located respectively in a Member State of the European Union or in a third country or parts of the territory of third country listed in columns 2 and 4 of Annex I to Commission Implementing Regulation (EU) 2018/659 from which the import of equine semen collected from registered horses, registered equidae or equidae for breeding and production is authorised in accordance with Article 4 of Commission Implementing Regulation (EU) 2018/659 and indicated in columns 11, 12 and 13 of Annex I thereto. (6) (7);
- 11.3. The ova used for in vitro production of the embryos described above comply with the requirements of Annex D to Directive 92/65/EEC and in particular the requirements set up in points II.1.1 to II.1.8 of this certificate (1).

Notes

Health information

Part I:

The place of origin shall correspond to the embryo collection team or embryo production team by which the Box I.11.: ova/embryos were collected/produced, processed, stored and approved in accordance with Article 17(3)(b) of Council Directive 92/65/EEC and listed on the Commission website:

http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm

- Box I.22.: The number of packages shall correspond to the number of containers
- Box I.23.: The identification of container and seal number shall be indicated
- Box I.28.: The category: specify if in vivo derived embryos, in vivo derived ova, in vitro produced embryos or micromanipulated embryos.

The donor identity shall correspond to the official identification of the animal.

The date of collection shall be indicate in the following format: dd/mm/yyyy.

Part II:

- (¹) Delete as appropriate.
- Only third countries or parts of the territory of third countries listed in columns 2 and 4 of Annex I to Commission Implementing Regulation (EU) 2018/659 respectively from which imports of registered equidae and equidae for breeding and production are also authorised and as indicated in column 14 of Annex I thereto.
- $(^3)$ Only approved embryo collection teams and embryo production teams listed in accordance with Article 17(3)(b) of Council Directive 92/65/EEC on the Commission website:

http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm

- $(^{4})$ Insert date.
- The agar gel immunodiffusion test (Coggins test) or the ELISA for equine infectious anaemia are not required for donor equidae which have continuously resided in Iceland since birth, provided that Iceland has remained officially free of equine infectious anaemia and no equidae and their semen, ova and embryos have been introduced into Iceland from outside prior to and during the period the semen was collected.

COUN	COUNTRY Equine ova/embry							
II.	Health information	II.a. Certificate reference No	II.b.					
(⁶)	Only approved semen collection centres listed in accordance with Article 11(4) or Article 17(3)(b) of Council Directive 92/65/EEC on the Commission websites:							
	https://ec.europa.eu/food/animals/live_animals/approved-establishments_en; http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm							
(7)	Does not apply to ova.							
(8)	OJ L 192, 23.7.2010, p. 1.							
_	The signature and the stamp must be in a d	ifferent colour to that of the printing.						
Offic	cial veterinarian							
	Name (in capital letters): Qualification and title:							
	Date: Signature:							
	Stamp:							

PART 3

Explanatory notes for the certification

(a) The health certificates shall be issued by the competent authority of the exporting country, based on the model set out in Part 1 or 2 of Annex III, according to the layout of the model that corresponds to the commodity concerned.

They shall contain, in the numbered order that appears in the model, the attestations that are required for any country and, as the case may be, those supplementary guarantees that are required for the exporting country or part of the territory of the country.

- (b) A separate and unique health certificate shall be issued for each consignment of semen, oocytes or embryos that are exported from a single territory appearing in columns 2 and 4 of Annex I which are consigned to the same destination and transported in the same railway wagon, lorry, aircraft or ship.
- (c) The original of the health certificate shall consist of a single sheet of paper, or, where more text is required, it shall be in such a form that all sheets of paper required are part of an integrated whole and indivisible.
- (d) Where the model health certificate states that certain statements shall be kept as appropriate, statements which are not relevant, may be crossed out and initialled and stamped by the official veterinarian, or completely deleted from the health certificate.
- (e) The health certificate shall be drawn up in at least one of the official languages of the EU Member State of the border inspection post of entry of the consignment into the Union and of the Member State of destination. However, those Member States may authorise the health certificate to be drawn up in the official language of another Member State, and accompanied, if necessary, by an official translation.
- (f) If for the reasons of identification of the items of the consignment (schedule in Box I.28 of the model health certificate), additional sheets of paper are attached to the health certificate, those sheets of paper shall also be considered as forming part of the original of the health certificate by application of the signature and stamp of the official veterinarian, on each of the pages.

- (g) When the health certificate, including additional schedules referred to in (f), comprises more than one page, each page shall be numbered -(page number) of (total number of pages)-, at the end of the page and shall bear the certificate reference number that has been designated by the competent authority on the top of the pages.
- (h) The original of the health certificate shall be completed and signed by an official veterinarian within 24 hours prior to loading of the consignment for exportation to the Union. The competent authorities of the exporting country shall ensure that the principles of certification equivalent to those laid down in Council Directive 96/93/EC (¹) are followed.

The colour of the signature and the stamp of the official veterinarian shall be different to that of the printing on the health certificate. This requirement also applies to stamps other than those embossed or watermarks.

- The original of the health certificate shall accompany the consignment until it reaches the border inspection post of entry into the Union.
- (j) The certificate reference number referred to in Box I.2 and Box II.a of the model health certificate shall be provided by the competent authority of the exporting country.

ANNEX IV

CATEGORIES OF MALE EQUIDAE TO WHICH THE CONDITIONS FOR EQUINE VIRAL ARTERITIS APPLY IN ACCORDANCE WITH ARTICLE 15(b)(ii) OF DIRECTIVE 2009/156/EC

- The requirement relating to equine viral arteritis laid down in Article 15(b)(ii)
 of Directive 2009/156/EC shall apply to uncastrated male equidae with the
 exception of:
 - (a) equidae vaccinated against equine viral arteritis under official supervision with a vaccine approved by the competent authority in accordance with one of the following protocols:
 - (i) the equidae shall be vaccinated during isolation of at least 28 days after they had been tested either in a serum neutralisation test for equine viral arteritis carried out with negative result at a serum dilution of 1 in 4 on a sample of blood taken not earlier than 7 days of commencing isolation, or in a virus isolation test carried out with negative result on an aliquot of the entire semen collected not earlier than 7 days of commencing isolation, and were kept separated from other equidae for 21 days following vaccination;
 - (ii) the equidae shall be vaccinated at the age of 180 to 270 days, after having been subjected to a virus neutralisation test for equine viral arteritis carried out with negative result at a serum dilution of 1 in 4, or carried out with stable or declining titres on two blood samples taken at least 14 days apart. The equidae shall be separated from other equidae until 21 days after vaccination.
 - (b) equidae less than 180 days old;
 - (c) equidae for slaughter sent directly to a slaughterhouse.
- The test shall be carried out and certified, and the result and vaccination certified, under official veterinary supervision. Vaccination shall be repeated at regular intervals according to manufacturer instructions.

Batch numbers of the approved vaccine, the details of the vaccination and revaccination and the results of serological or agent-identification tests shall be documented, where available in the identification document (passport), and made available for certification purposes.

3. Test mating as described in point 4(a) of Article 12.9.2. of the Terrestrial Animal Health Code of the World Organisation for Animal Health (OIE) is considered equivalent to the virus isolation test referred to in point 1(a)(i) to prove absence of the equine arteritis virus in semen.

ANNEX V

MODEL DECLARATIONS

PART 1

Declaration by the captain of the aircraft

(To be completed and attached to the health certificate when transport to the Union frontier includes, even for part of the journey, transportation by air.)

Declaration by the captain of the aircraft		
I, the undersigned, captain of the aircraft (name), declare that the crate or container and the area around the crate or container containing the animals referred to in the attached health certificate No		
Done at	on	
(Airport of departure)	(Date of departure)	
	(signature of captain)	
(stamp)		
	(name in capital letters and title)	

PART 2

Declaration by the captain of the vessel

(To be completed and attached to the health certificate when transport to the Union frontier includes, even for part of the journey, transportation by ship.)

Declaration by the master of the ship	
certificate No from and that the ship did not call at any place outside), declare that the animals referred to in the attached health
Done at	on
(Port of arrival)	(Date of arrival)
	(signature of master)
(stamp)	
	(name in capital letters and title)

PART 3

Model Transhipment Manifest

(To be completed and attached to the health certificate when transport to the Union frontier includes transhipment from one aircraft to another aircraft or from one vessel to another vessel in a country not listed in Annex I to Commission Implementing Regulation (EU) 2018/659)

		Serial Number:	
		Reference No of Air Cargo Transfer Manifest:	(1)
Country where transhipment take	es place:		
Airport (2)/Port (2) of arrival:			
Date of arrival:			
Date of transhipment:			
Transferring Carrier:			
Receiving Carrier:			
Description of consignment:			
Serial No of Health Certificate	Remarks		
	rinarian (²)/customs officer (²) at on and in compliance with the fol	the above mentioned airport (²)/port (²) declare that the tranship lowing conditions:	ment
(a) the equidae were during the	e transhipment protected from a	ttacks by insect vectors of diseases transmissible to equidae;	
(b) the equidae did not come ir	nto contact with equidae of a diff	erent health status;	
		space in the transport compartment were sprayed with an appropely after the closing of the doors of the aircraft $(^2)$ /vessel $(^2)$.	oriate
The consignment has been trai	nshipped in full and apparent go	od order and conditions except as noted in the "Remarks" column	۱.
Done at		on	
		Stamp	
	eterinarian or customs officer)		
(name in capit	al letters and title)		
(¹) Keep empty if transhipment fro (²) Delete as appropriate	m vessel to vessel		