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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)

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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:

- 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 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 means any inspection post as defined in post'

 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 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).

⁽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 quarantine'

means the quarantine of equidae which

- (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 quarantine'

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

- 1. 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 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 reentry 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 slaughterhouse 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.

▼ M2

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), Article 17(1)(d) and Article 19(2)(a) shall apply from 14 December 2019.

▼B

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

▼<u>M2</u>

LIST OF THIRD COUNTRIES AND PARTS OF THE TERRITORY OF THIRD COUNTRIES (¹) FROM WHICH THE ENTRY INTO THE UNION OF CONSIGNMENTS OF EQUIDAE AND OF SEMEN, OVA AND EMBRYOS OF EQUIDAE IS AUTHORISED

		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	RH	RH	RH	ES	RE +		SEMEN		O/E	Equi- dae	
		country							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	X	X	X	X	X	X	X	X	X	X	
BA	Bosnia and Herze- govina	BA-0	Whole country	В	X	X	X	_	_	_	_	_	_	X	
ВВ	Barbados	BB-0	Whole country	D	X	X	X	_	_	_	_	_	_	X	
ВН	Bahrain	BH-0	Whole country	Е	X	X	X	_	_	_	_	_	_	X	
BM	Bermuda	BM-0	Whole country	D	X	X	X	_	_	_	_	_	_	X	
ВО	Bolivia	BO-0	Whole country	D	X	X	X	_	_	_	_	_	_	X	
BR	Brazil	BR-0	Whole country	_	_	_	_		_	_	_	_	_	_	
		BR-1	The states of:												
			Paraná and Rio de Janeiro	D	X	X	X	_	_	_	_	_	_	X	
BY	Belarus	BY-0	Whole country	В	X	X	X	X	X	_	_	_	_	X	
CA	Canada	CA-0	Whole country	С	X	X	X	X	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 country	Description of the part of the territory of the third country	SG	RH	RH	RH	ES	RE + EBP		SEMEN	ı	O/E	Equi- dae	
		country							EBP	RH	RE	EBP			
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16
СН	Switzerland (2)	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	_	X	_	_	_	_	_	_	_	_	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	Equi- dae	
		country							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)	E	Х	_	X	_	_	_	_		_	X	
FK	Falkland Islands	FK-0	Whole country	A	X	X	X	_	X	_	_	_	_	X	
GL	Greenland	GL-0	Whole country	A	X	X	X	X	X	_	_	_	_	X	
НК	Hong Kong	HK-0	Whole country	G	X	X	X	_	_	_	_		_	X	
IL	Israel (3)	IL-0	Whole country	Е	X	X	X	X	X	X	X	_	_	X	
IS	Iceland (4)	IS-0	Whole country	A	X	X	X	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	Equi- dae	
		country							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	
ЈО	Jordan	JO-0	Whole country	Е	X	X	X	_	_	_	_	_	_	X	
JP	Japan	JP-0	Whole country	G	X	X	X	_	_	_	_	_	_	X	
KG	Kyrgyzstan	KG-0	Whole country	_	_	_	_	_		_		ĺ	_	_	
		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	_	_	_	_		_			_	_	
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	_	_	ĺ	_	X	
MK	North Macedonia	MK-0	Whole country	В	X	X	X	X	X	_	_	_	_	X	
МО	Macao	MO-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	Equi- dae	
		country							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
		MX-2	The whole country except the States of Chiapas, Oaxaca, Tabasco, Campeche, Yucatan, Quintana Roo, Veracruz and Tamaulipas	С	X	X	Х	_	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	_	_	_	X	
RS	Serbia (5)	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	Equi- dae	
		country							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, Nijninovgorod, 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	X	X	X	X	_	_	_	_	X	
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	RH	RH	RH	ES	RE +		SEMEN		O/E	Equi- dae	
		country							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	X	X	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	X	X	X	_	_	_	_	_	_	X	
UA	Ukraine	UA-0	Whole country	В	X	X	X	X	X	X	X	X	_	X	
US	United States of America	US-0	Whole country	С	Х	X	X	X	X	X	Х	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⁾ Where official regionalisation applies in accordance with Article 13(2)(a) of Directive 2009/156/EC.

⁽²⁾ Without prejudice to specific certification and control requirements provided for in Decision 2002/309/EC, Euratom of the Council, and of the Commission (OJ L 114, 30.4.2002, p. 1).

⁽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) Without prejudice to specific certification requirements provided for in Article 17 of the Agreement on the European Economic Area (OJ No L 1, 3.1.1994, p. 3).

⁽⁵⁾ As defined in Article 135 of the Stabilisation and Association Agreement between the European Communities and their Member States of the one part, and the Republic of Serbia, of the other part (OJ L 278, 18.10.2013, p.16).

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.
ЕВР	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

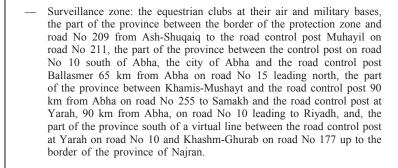
Sanitary Groups

Sanitary Group	Specific animal health guarantees required for entry of equidae into the Union
A	equine infectious anaemia, equine viral arteritis
В	equine infectious anaemia, equine viral arteritis, glanders, dourine
C	equine infectious anaemia, equine viral arteritis, Eastern and Western equine encephalomyelitis, vesicular stomatitis
D	equine infectious anaemia, equine viral arteritis, glanders, dourine, Eastern and Western equine encephalomyelitis, Venezuelan equine encephalomyelitis, vesicular stomatitis
E	equine infectious anaemia, equine viral arteritis, glanders, dourine, African horse sickness
F	equine infectious anaemia, dourine, African horse sickness
G	equine infectious anaemia, equine viral arteritis, glanders, dourine, Japanese encephalitis

▼<u>B</u>

CN	China	CN-1	The specific equine disease-free zone in the Guangdong Province with the following delimitation:				
			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 International 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 zo	one 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.			

BOX	2		
EG	Egypt	EG-1	The Equine Disease Free Zone (EDFZ) of about 0,1 km ² 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:
			Riyadh Airport King Abdulaziz Race Track (Janadrijah)
			2. King Adduraziz Race Hack (Janadiljan)
		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
			 1. Province of Jizan — Protection zone: the whole province, except the part north of the road
			 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



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 stations: 1. Kenilworth Quarantine Station Delimitation of the Metropolitan area of Cape-Town (ZA-1): Northern boundary: Blaauwberg Road (M14); Koeberg Road (M14), Plattekloof Road (M14), N7 Eastern boundary: 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

▼<u>M2</u>

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

COUNTRY: Veterinary certificate to EU Certificate reference No 1.2.a. Consignor 1.2. Name Address Central competent authority Tel. I.4. Local competent authority Part I: Details of dispatched consignment 1.6. Consignee Name Address Postcode I.7. Country I.8. Region of Code I.9. Country of ISO Region of Cod of origin code destination code destination I.11. Place of origin I.12. Place of destination Name Name Approval number Address Address Postcode I.13. Place of loading I.14. Date of departure I.15. Means of transport I.16. Entry BIP in EU Ship Railway Aeroplane wagon Road vehicle Other I.17. No(s) of CITES Identification Documentary references I.18. Description of animal I.19. Commodity code (HS code) 01 01 I.20. Quantity 1.21. I.22. Number of packages 1.24. I.23. Seal/Container No I.25. Animal certified for: Registered horse 1.26. I.27. For import or admission into EU I.28. Identification of the animal Species (Scientific name) Identification system Identification number Age Sex Equus caballus

COUNTRY

			II.a.	Certificate reference number	II.b. Local reference number			
ļ	(3							
II.	Attestatio	n of anima	al health and w	elfare				
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): - was examined today ⁽¹⁾ and found free of clinical signs of diseases and of obvious signs of infestation; - is not intended for slaughter under a national programme of infectious or contagious disease - 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 representations.								
	 is a registered horse as defined in Article 2(c) of Commission Implementing Regulation (EU) 2018/65 was examined today⁽¹⁾ and found free of clinical signs of diseases and of obvious signs of ectop infestation; 							
	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 c	ountry or part of	the territory of third country and ho	lding of dispatch			
II.1.1.	The animal is dispatched from							
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;							
II.1.3.	the animal is dispatched from a country or part of the territory 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; 							
	b)	in which Venezuelan equine encephalomyelitis has not occurred during the period of 2 years prior to the date of dispatch;						
	c)	in which	ich dourine has not occurred during the period of 6 months prior to the date of dispatch;					
	d)	38 35 35 35		t occurred during the period of 6 mo	성이 가는 이 경우 전문에 가는 이 사람들은 아이가 아이들은 아이라면 아이들을 만나 하다니다.			
(3)either	[e)	in which vesicular stomatitis has not occurred during the period of 6 months prior to the date of dispatch;]						
⁽³⁾ Or	[e)	in which vesicular stomatitis has occurred during the period of 6 months prior to the date dispatch, and a blood sample taken from the animal on(insert date), within period of 21 days prior to the date of dispatch, was tested with negative result for antibody to the vesicular stomatitis virus						
		(3)either	[in a virus neut	ralisation test at a serum dilution of	1 in 32;]]			
		⁽³⁾ or		accordance with the relevant Chapte or Terrestrial Animals of the OIE;]]	er of the Manual of Diagnostic Tests			
II.1.4. the animal does not come from a holding and to the best of my knowledge for the time periods referred to a 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:								
((4)[II.1.4.1. in the case of equidae suspected of having contracted dourine,							
		⁽³⁾ either		inning on the date of the last actual aving contracted dourine or infected				
		(3)and/or	[in the case of	a stallion, until the animal is castrate	ed;]			
1		(3)and/or						

COUNTRY

			II.a. Certificate reference number II.b. Local reference number					
(⁽⁴⁾ [II.1.4.2.	in the cas	se of glanders,					
		(3)either [6 months beginning on the day on which the equidae suffering from the disease of 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;]						
		(3)and/or	[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,						
			[6 months beginning on the day on which the equidae suffering from the disease have been slaughtered;]					
		(3)and/or	(3) and/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;]					
		(3)and/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.	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;						
	II.1.4.5.	in the case of vesicular stomatitis.						
		(3)either	[6 months following the last case;]					
		⁽³⁾ and/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.1.4.6.	in the case of rabies, 30 days following the last case and the date of completion of the cleansin and disinfection of the premises;						
	II.1.4.7.		se of anthrax, 15 days following the last case and the date of completion of the cleansing fection of the premises;					
II.1.5.		t of my knowledge, during the period of 15 days prior to the date of dispatch the animal has n intact with equidae infected or suspected of an infectious or contagious disease.						
II.2.	Attestation	of residen	nce and pre-export isolation					
⁽³⁾ either	[II.2.1.	During a period of at least 40 days prior to the date of dispatch, the animal has been resident on holdings under veterinary supervision situated in the country or part of the territory of the country of dispatch which is assigned to Sanitary Group A, B, C, D, E or G, and						
	(3)either	[in a Member State of the Union;]						
	(3)and/or	[in a country or part of the territory of a country with Code:						
		⁽³⁾ either	[assigned to the same Sanitary Group ⁽²⁾ as the country or part of the territory the country of dispatch;]]					
		(3)and/or	[assigned to Sanitary Group A, B or C;]]					
		(3)and/or	[China ⁽⁵⁾⁽⁶⁾ , Hong Kong, Japan, Korea, Macao, Malaysia (Peninsula), Singapori Thailand or the United Arab Emirates;]]]					

COUNTRY

			II.a.	Certificate reference number	II.b.	Local reference number				
⁽³⁾⁽⁷⁾ Or	[II.2.1.	During a period of at least 60 days prior to the date of dispatch, the animal has been resident on holdings under veterinary supervision situated in the country or part of the territory of the 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⁾⁽⁷⁾ either	[11.2.2.	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	from vector insects into the country or p with point II.2.1 from	for a p art of t a Mer	in the country or part of the territory of period of at least 40 days prior to the the territory of the country of dispatc mber State of the Union or a country ary Group A, B, C, D, E or G;]]	e date h, if it w	of dispatch, or since entra as imported in accordance				
	⁽³⁾ or	40 days prior to the country of dispatch, Union or a country of C, D, E or G, and the	date of if it was or part ne cou	ed premises under official veterinary of dispatch, or since entry into the co- tas imported in accordance with point of the territory of a country which is ntry or part of the territory of the co- f African horse sickness;]]	untry o II.2.1 fi assign	or part of the territory of the rom a Member State of the ed to Sanitary Group A, E				
⁽³⁾⁽⁷⁾ or	[11.2.2.		the animal is dispatched from a country or part of the territory of country which is assigned to Sanitary Group F and was kept:							
	⁽³⁾ either	[in the approved vector-protected quarantine station of								
⁽³⁾ or	name of o	uarantine station) dur	ing the	ed vector-proof quarantine station e period of at least 14 days prior to as proven absence of vectors insid	the date	e of dispatch and constan				
II.3.	Attestation	n of vaccination and h	ealth t	ests						
(3)either [II.3.1. The animal was not vaccinated against African horse sickness in the continuous there is no information suggesting previous vaccination;]				ne country of dispatch and						
⁽³⁾ or	[II.3.1.			d against African horse sickness, an or to the date of dispatch;]]	d this v	accination was carried out				
	⁽³⁾ or	[more than 60 days	and le	ess than 12 months prior to the date erred to in point II.1.3.(a), from when						
⁽³⁾⁽⁷⁾ or	[II.3.1.	The animal is dispatched from a country or part of the territory of a country which is assigned Sanitary Group F and was vaccinated against African horse sickness on(insert danot more than 24 months and at least 40 days prior to the date of entry in the vector-protect quarantine by administration of a registered vaccine according to manufacturer's instruction which is protective against the circulating serotypes of the African horse sickness virus;]								
	II.3.2.	the animal was not of 60 days prior to the		ated against Venezuelan equine er e of dispatch from	cephal	omyelitis during the period				
	⁽³⁾ either			ts of the territory are free of Venezus prior to the date of dispatch;]	elan eq	uine encephalomyelitis fo				
	⁽³⁾⁽⁷⁾ or	Venezuelan equine	encep uine e	country which is assigned to Sanita halomyelitis for a period of at least 2 encephalomyelitis occurs in the rem	years p	prior to the date of dispatch				

COUNTRY

		II.a.	Certificate reference number	II.b. Local reference number			
	⁽³⁾ either	course and rev 60 days and n vector-protecte and during tha daily, remained same holding v	against Venezuelan equine encephal accinated according to manufacturer o more than 12 months prior to the digital quarantine for a period of at least 2 to period remained clinically healthy, digital within the normal physiological ranythich showed a rise in body temperatorize virus isolation for Venezuelan equin	's récommendations not less than date of dispatch, and was kept in I days prior to the date of dispatch, and its body temperature, taken ge, and any equine animal on the ture, taken daily, was subjected to			
	⁽³⁾ <i>or</i>	vector-protecter remained clinic normal physiolo a rise in body to for Venezuelar dispatched wencephalomye days after the	ated against Venezuelan equine end quarantine for a period of at leas ally healthy, and its body temperaturogical range, and any equine animal comperature, taken daily, was subjected as subjected to a diagnostic litis with negative result conducted on date of entry into of the vector-provector insects until dispatch;]	t 21 days, and during that period e, taken daily, remained within the on the same holding which showed de to a blood test for virus isolation ative results, and the animal to be test for Venezuelan equine n a sample taken not less than 14			
	⁽³⁾ or	encephalomye taken on two of (insert to the date of of transcription-ye encephalomye within 48 hours vector attacks combined use	d to a haemagglutination inhibition inhibition litis carried out by the same laborate occasions with an interval of 21 day date), the second of which was take dispatch, without an increase in antibiolymerase chain reaction) test for the litis virus genome, carried out with reprior to dispatch, on (insert different the moment of the RT-PCR same of approved insect repellents and not the stable and the means in whice	ory on the same day on samples or on (insert date) and on on during a period of 10 days prior body titre, and a RT-PCR (reverse e detection of Venezuelan equine legative result on a sample taken late), and has been protected from only pling until loading for dispatch, by insecticides on the animal and			
(3)[II.3.3.	the anima	al is an uncastra	ted male equine animal older than 18	30 days, and			
⁽³⁾ either	[is dispat	tched from a co and has not bee	untry in which equine viral arteritis in officially reported during the perio	(EVA) is a compulsorily notifiable			
⁽³⁾ or		te of dispatch, by	mple taken on(insert dat virus neutralisation test for EVA with				
⁽³⁾ Or	of 21 day	s prior to the da	an aliquot of its entire semen taken on (insert date), within a period to the date of dispatch, by virus isolation test, polymerase chain reaction (PCR) or EVA with negative result;]]				
⁽³⁾ or	[was vaccinated against EVA on						
	⁽³⁾ either		per 2018, on the day a blood sample s neutralisation test for EVA with negative sample.				

COUNTRY

				II.a.	Certificate reference number	II.b.	Local reference number
		⁽³⁾ or	official which w	eterinar as testec	er 2018, during a period of isology supervision, commencing on during that isolation period in a tale serum dilution of 1 in 4;]]]	the day a	blood sample was taker
		⁽³⁾ or	supervis EVA car same da	ion, duri ried out by by the	30 to 270 days, during a periong which the animal was subjection with negative result at a serum same laboratory with stable or days apart;]]]	ected to a dilution of	virus neutralisation test fo 1 in 4, or carried out on the
		⁽³⁾ or	result at	a serum after com	was subjected to a virus neu dilution of 1 in 4, carried out on mencing a period of uninterrupt ation;]]]	a blood sa	mple taken not earlier than
		⁽³⁾ or	test for E on the s	EVA carr ame day	0 to 250 days, after the animal vided out with negative result at a py by the same laboratory with states 14 days apart;]]]	serum dilu	tion of 1 in 4, or carried ou
(³⁾ or	EVA car blood sa to the da	rried out wample of th	ith nega at anima atch, wa	solation test, polymerase chair tive result on an aliquot of its eal taken on(insert as tested in a virus neutralisation in 4;]]	ntire seme	n collected after the date a a period of 6 months price
	⁽³⁾ or		eviously te ted against		sitive for antibodies against th	e equine a	arteritis virus or has bee
		a)	consecution prior to an serological	ve days, nd until a al tests f	f 6 months prior to the date of to at least two mares which we at least 28 days after test mat or EVA with negative results a at the time of test mating and	ere kept in ing and wh t a serum	isolation during the 7 day lich were subjected to two dilution of 1 in 4 on blood
		b)	within 21	days pric	a virus neutralisation test for EV or to the date of dispatch on sitive result at a serum dilution o	(inse	rt date),
			(3) or	[with neg	gative result at a serum dilution	of 1 in 4;]]]	A
	⁽³⁾ Or	legislation that the specified participa	on animal is d in that ating in suc	tempora legal ac	ng for EVA or vaccination agai. (insert reference to the application of the union for the union for the union for the union for the union is keptiand that any breeding activity or any residence in the Union;]]	cable Union participati separated	n legal act) on the ground on in the equestrian even d from other equidae no
⁽³⁾⁽⁷⁾ either	[11.3.4.	anaemia	a, where it	t was co	om Iceland, which is certified a intinuously resident since birth ad Iceland from other countries;	, and did r	
⁽³⁾ or	[II.3.4.	the anin Coggins on	nal was su test) or to (ins [a period	ubjected an ELIS ert date, d of 90 d	with negative result to an aga A for equine infectious anaemi), this being within ays prior to the date of dispatch th is assigned to Sanitary Grou	ar gel imm a carried or n from a co	ut on a blood sample take untry or part of the territor
		(3)or					

COUNTRY

			II.a.	Certificate reference number	II.b. Local reference number		
⁽³⁾ [II.3.5.	Sanitary reported complen on a blo	Group B of during a nent fixation	or E, or from period on test for taken o	om a country or part of the territory om Brazil, China or Thailand, or froi of 3 years prior to the date of dir glanders carried out with negative n	m a country in which glanders was ispatch, and was subjected to a result at a serum dilution of 1 in 5		
⁽³⁾ [II.3.6.	from a configuration from a co	ountry or p in China or orior to the out with n	oart of the or Thailan e date of negative . (insert of breeding	ed male or a female equine anima e territory of a country which is assi d, or from a country in which dourir dispatch, and was subjected to a co- result at a serum dilution of 1 in date), within a period of 30 days prion g during the period of at least 30 days	gned to Sanitary Group B, D, E or ne was reported during a period of emplement fixation test for dourine 5 on a blood sample taken on or to the date of dispatch, and has		
⁽³⁾⁽⁷⁾ [II.3.7.		Group C ([Western country	or D, and n and Eas or part o	om a country or part of the territory stern equine encephalomyelitis have f the territory of the country of disp date of dispatch;]]	e not been officially reported in the		
	⁽³⁾ or	[the animal was vaccinated with a complete primary course and revaccinated according to manufacturer's instructions within a 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	vector-p	rotected n tests fo	kept for a period of at least 21 days quarantine and during this period wo for Western and Eastern equine end on the same day	as subjected to haemagglutination		
		⁽³⁾ either		ample of blood taken ons prior to the date of dispatch, with			
		⁽³⁾ or	days o	mples of blood taken on two occasi in(insert date) and or if of which was taken within a period th, without increase in antibody titre	n (insert date), the od of 10 days prior to the date of		
⁽³⁾ [II.3.8.	Sanitary	Group G,	or from	om a country or part of the territory a country in which Japanese encep of at least 2 years prior to the date	halitis has been officially reported		
	⁽³⁾ either	that hold	ding whe	olding situated in the centre of an ar- re there has been no case of Japa o the date of dispatch;]]			
	⁽³⁾ or	the date	[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				
		⁽³⁾ either	enceph of bloc which without	aemagglutination inhibition or virus nalitis carried out by the same labora of taken on two occasions with an (insert date) and on	atory on the same day on samples in interval of at least 14 days on (insert date), the second of ays prior to the date of dispatch, in antibody titre between the two		

COUNTRY

					1
			II.a.	Certificate reference number	II.b. Local reference number
		⁽³⁾ or	Japanes sample t	-M capture ELISA test for the se encephalitis virus with negative aken not earlier than 7 days after(insert date), and remained p	ve result, carried out on a bloo the date the isolation commence
	⁽³⁾ Or	revaccin	ated acco	gainst Japanese encephalitis wit rding to manufacturer's recommer not more than 12 months prior to	ndations during a period of not les
⁽³⁾⁽⁷⁾ either [II.3.9.	Sanitary	Group E,	and was si	n a country or part of the territory ubjected to a serological test for A /156/EC, which was carried out by	frican horse sickness as describe
	⁽³⁾ either	on		taken on two occasions with an ir (insert date) and on	(insert date), the second
		(3)either	[with neg	gative results in each case.]]]	
		(3)or	[with a p	ositive result in the first sample, a	and
			⁽³⁾ either		ubsequently tested with negation test as described in Annex IV
			⁽³⁾ or	increase in antibody titre in	ed without more than a two-fon a virus neutralisation test a apter 2.5.1. of the OIE Terrestriand Vaccines.]]]]
	⁽³⁾ or	prior to t	the date o	le taken on	rt of the territory of the country
⁽³⁾⁽⁷⁾ or [II.3.9.		al is dispa Group F,		n a country or part of the territory	of a country which is assigned
	⁽³⁾ either	IV to Dire day on b days, on sample	ective 2009 blood samp not taker ne, the se	a serological test for African hors 9/156/EC, which was carried out b ples taken on two occasions with (insert date) and on less than 7 days after introcoond sample taken within a periodocond.	y the same laboratory on the sam an interval of between 21 and 3 (insert date), the fir duction into the vector-protecte
		(3)either	[with neg	gative results in each case.]]]	
		⁽³⁾ or	[with a p	ositive result in the first sample, a	and
			⁽³⁾ either		quently tested with negative resu described in Annex IV to Directiv
			⁽³⁾ or		d without more than a two-forus neutralisation test as describe

COUNTRY

				II.a.	Certificate reference	number	II.b.	Local reference number	er
		⁽³⁾ or	sickness result in than 28	as des each ca days af	cribed in Annex IV to Dase on a blood sample	Directive 2009/ taken on action into the	156/E0	ion test for African ho C, carried out with negat (insert date) not le -protected quarantine a	tive less
		⁽³⁾ or	in Annex	x IV to taken or tion into	Directive 2009/156/EC	C, carried out ert date) not le	with n	orse sickness as descrit legative result on a blo n 14 days after the date ore than 72 hours bef	ood e of
II.4.	Attestation	of the trai	nsport cor	ditions					
⁽³⁾⁽⁷⁾ either	[II.4.1.	Sanitary the Union into cont	. It is dispatched from a country or part of the territory of a country which is assigned to ry Group A, B, C, D, E or G and arrangements have been made to transport it directly to ion, without passing through a market, marshalling or assembly centre and without coming ontact with other equidae not complying with at least the same health requirements as used in this health certificate.]						
⁽³⁾⁽⁷⁾ or	[II.4.1.	Sanitary protected	Group F d quarantin certificate [to the a for the a	and arrane station either for irport un aircraft to ed in the	angements have been on without coming into our or imports or for tempo nder vector-protected or o be cleansed and dis the third country of disp	n made to tran contact with of crary admission conditions and sinfected in ad-	nsport ther eq n into t d arran vance	ountry which is assigned it directly from the vect uidae not accompanied he Union gements have been ma with a disinfectant officia against vector insects j	tor- d by ade ially
		⁽³⁾ or	condition schedule or part of in stalls	ns and a ed direct f the ten which w ed in the	arrangements have be tly to a port in the Unior ritory of a country not a vere cleansed and disi	een made to to n without callin approved for th infected in adv	transpo ng into ne entry vance v	ntry under vector-protect of it on a vessel which a port situated in a cour into the Union of equid with a disinfectant officinst vector insects just p	h is ntry dae, ially
	II.4.2.	complyin	g with at le	east the		ents as describ		ct with other equidae his health certificate dur	
	II.4.3.	disinfecte	ed before y are so	loading	with a disinfectant office	cially recognise	ed in th	e loaded were cleaned a ne third country of dispa er cannot escape dur	atch
II.5.	Attestation	of animal	welfare						
								ansported on the intend ctively at all stages of	
Notes:									
Part I:									
Box I.8.:					art of the territory of the ting Regulation (EU) 20		ispatch	as appearing in colum	ın 3
Box I.15.:							nber (a	ircraft) or name (ship) a	and rder

▼ M2

COUNTRY

Temporary admission - Registered horse

II.a. Certificate reference number II.b. Local reference number 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 etc.) 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:

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

- (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)Delete statement if the attestation in point II.1.3. applies to the entire country of dispatch.
- (5) Part of the territory of country authorised for temporary admission as appearing in columns 3 and 6 respectively of Annex I to Commission Implementing Regulation (EU) 2018/659.
- (6) Only authorised if country of dispatch is assigned to Sanitary Group G.
- (7) 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:

- 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; (b)
- 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 (e) 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.

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

				ne owner or representative rary admission of a registe				
Ide	ntification of	the ani	mal ⁽¹⁾					
Species (Scientific Identification system Identification number Age Sex name)								
Equ	ıus caballus	5						
I, th	e undersign the horse	ed own	er ⁽²⁾ or representative of the	owner ⁽²⁾ of the registered he	orse described above, he	ereby declare, that:		
	(2)either			(insert name of country or prior to the date of dispatc		a country of dispatch)		
	⁽²⁾ or			rt name of country or part of east 40 days prior to the date		of dispatch) during the		
		(a)		rom (insert ritory of country of dispatch)		where horse entered		
		(b)		rom (insert ritory of country of dispatch)		where horse entered		
		(c)		rom (insert ritory of country of dispatch)		where horse entered		
-			of 15 days prior to the date agious diseases transmissil	e of dispatch the horse has a ble to equidae;	not been in contact with	animals suffering from		
-	the transp stages of			vay that health and well-being	g of the horse can be pro	tected effectively at all		
-				isolation as applicable in ac territory of the country of dis		of the accompanying		
-			the transport as applicable the territory of the country of	in accordance with point II.4 of dispatch are fulfilled;	1. of the accompanying h	ealth certificate for the		
•	during its premises:	residen	ce inside the Union for a pe	eriod of less than 90 days the	e horse will be accommo	dated on the following		
	(b) from (c) from	1 1	(date) to (date) ir (date) to (date) ir	n(place of holing place	ding) in (Mem ding) in (Mem	nber State) nber State)		
-	I am awar outlined in	e that i this de	n the event that the horse claration, it must be accomp	moves from one Member Stoanied by a health certificate st be notified to the Member	tate of the Union to anot issued by an official vete	ther Member State, as		
-				Union on I place of border post of exit		the border post of		
Nar	me and addr	ess of t	he owner ⁽²⁾ or representativ	e ⁽²⁾ :				
Dat	e:		(dd/mm/yyyy)					
				(Signature)				
(1)	document a system (such If a passport validated it. Age: Date	as defir ch as ea ort acco	ned in Article 2(b) of Commar tag, tattoo, brand, transp	an individual identifier which nission Implementing Regula onder) and the anatomic pla umber should be stated an	ation (EU) 2018/659. Sp ce used on the animal.	ecify the identification		
(2)	Delete as a	ppropri	ate.					

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

COU	INTRY:	Veterinary certificate to EU
	I.1. Consignor Name	I.2. Certificate reference No I.2.a.
	Address	I.3. Central competent authority
	Tel.	I.4. Local competent authority
Part I : Details of dispatched consignment	I.5. Consignee Name Address Postcode Tel.	I.6. Person responsible for the load in EU Name Address Postcode Tel.
s of disp	I.7. Country of ISO code I.8. Region Code origin of origin	I.9. Country of ISO code I.10. Region of Code destination
tails	I.11. Place of origin	1.12.
Part I: De	Name Approval number Address	
	I.13. Place of loading	I.14. Date of departure
15	I.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 animals	I.19. Commodity code (HS code) 01 01
		I.20. Quantity
	1.21.	1 I.22. Number of packages
	I.23. Seal/Container No	1.24.
	I.25. Animals certified for:	
	Registered equidae breeding and pro	oduction
	I.26. For transit through EU to third country X	1.27.
	Third country ISO code	
	I.28. Identification of the animal	
	Species (Scientific name) Identification system	n Identification number Age Sex

	COUNTR	RY					Transit - Equidae
				II.a.	Certificate reference number	II.b.	Local reference number
	11.	Attesta	tion of animal	healt	h and welfare		
Part II: Certification	I, the un	was exa infestati is not in meets th	amined today ⁽¹⁾ ion; tended for slau he requirements	and f	, hereby certify, that the equine animal found free of clinical signs of diseases under a national programme of infectionsted in points II.1. to II.5. of this certificate declaration, signed by the owner of the company of the content of the company of the compa	and of our coreste;	obvious signs of ectoparasite ntagious disease eradication;
6. T	II.1.		ion on third cou	intry (or part of the territory of third country a	nd holdir	ng of dispatch
	II.1.1. The animal is dispatched from						
	II.1.2.	dourine types in	(Trypanosoma	equi	the following diseases are compulsorii iperdum), glanders (Burkholderia mala equine encephalomyelitis), equine infe	ei), equi	ne encephalomyelitis (of all
	II.1.3.	the anin	nal is dispatche	d from	m a country or part of the territory of a c	ountry	
		a)	2009/156/E or epidemic date of disp	C an ologic oatch	dered free from African horse sicks of in which there has been no clinical, stal evidence of African horse sickness d and in which there have been no vaccinaths prior to the date of dispatch;	serologic uring the	al (in unvaccinated equidae) period of 2 years prior to the
		b)			elan equine encephalomyelitis has not of dispatch;	occurred	I during the period of 2 years
		c)	in which do	urine	has not occurred during the period of 6	months	prior to the date of dispatch;
		d)	in which gla	ander	s has not occurred during the period of	6 months	prior to the date of dispatch;
	(3)either	[e)	in which ve of dispatch		ar stomatitis has not occurred during the	e period	of 6 months prior to the date
	⁽³⁾ or	[e)	dispatch, a period of 2	nd a l	ar stomatitis has occurred during the p blood sample taken from the animal or s prior to the date of dispatch, was test matitis virus	٠	(insert date), within a
			(3)either [i	n a vi	irus neutralisation test at a serum dilutio	on of 1 in	32;]]

(3)or

[in an ELISA in accordance with the relevant Chapter of the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals of the OIE;]]

COUNT	нү			Transit - Equida
			II.a. Certificate reference number	II.b. Local reference number
II.1.4.	to in poin	ts II.1.4.1.	come from a holding, and to the best of my know to II.1.4.7. was not in contact with animals fror the reasons referred to in points II.1.4.1. to II.1.4	m holdings, which were subject to
	⁽⁴⁾ [II.1.4.1.	in the cas	se of equidae suspected of having contracted do	urine,
		⁽³⁾ either	[6 months beginning on the date of the last actual suspected of having contracted douring equiperdum;]	
		(3)and/or	[in the case of a stallion, until the animal is cast	rated;]
		(3)and/or	[30 days following the date of completion of th premises after all animals of susceptible specie	
	(4)[II.1.4.2.	in the cas	se of glanders,	
	•	⁽³⁾ either	[6 months beginning on the day on which the ed subjected with positive result to a test for the d Burkholderia mallei or antibodies to that pathog	etection of the causative pathoge
		(3)and/or	[30 days following the date of completion of th premises after all animals of susceptible specie	
	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 chave been slaughtered;]	equidae suffering from the diseas
		(3)and/or	[6 months beginning on the day on which th causing West Nile Fever, Eastern equine encencephalomyelitis have died, been removed fro	ephalomyelitis or Western equin
		(3)and/or	[30 days following the date of completion of th premises after all animals of susceptible specie	
	II.1.4.4.	been slau an agar	se of equine infectious anaemia, until the date on ughtered, the remaining animals on the holding gel immunodiffusion test (AGID or Coggins te on two occasions 3 months apart;	have shown a negative reaction i
	II.1.4.5.		se of vesicular stomatitis,	
		(3)either	[6 months following the last case;]	
		(3)and/or	[30 days following the date of completion of th premises after all animals of susceptible specie	
	II.1.4.6.		se of rabies, 30 days following the last case a and disinfection of the premises;	and the date of completion of the
	II.1.4.7.		se of anthrax, 15 days following the last case and disinfection of the premises;	and the date of completion of the
II.1.5.			wledge, during the period of 15 days prior to the equidae infected or suspected of an infectious or	
II.2.	Attestation	n of residen	ce and pre-export isolation	
⁽³⁾ eithei		on holdin country o	period of at least 40 days prior to the date of disp gs under veterinary supervision situated in a c f dispatch which is assigned to Sanitary Group A	country or part of the territory of
	(3)either		nber State of the Union;]]	
	⁽³⁾ and/or	for tempo into the c strict as t	ntry or part of the territory of country with Code: . orary admission into the Union of registered hors ountry or part of the territory of the country of di- horse required in accordance with the Union legisted horses from this country or part of the territory is is:	es, and from which it was importe spatch under conditions at least a slation for the temporary admission
		(3)either	[assigned to the same Sanitary Group(2)	as the country or part of the territo

(3)either

[assigned to the same Sanitary Group $\dots\dots^{(2)}$ as the country or part of the territory of the country of dispatch;]]]

COUNTRY Transit - Equidae

			II.a.	C	ertificate refer	ence number		II.b.	Local reference number
		(3)and/or (3)and/or	[ass	igne	d to Sanitary		r G and the		mal is a registered horse a Regulation (EU) 2018/659;]
⁽³⁾⁽⁵⁾ or	[II.2.1.	on holding country of days prior	period of gs und f dispa	of at der v atch e da	least 60 days eterinary sup which is assig te of dispatch	prior to the da ervision situate ned to Sanita from a Mem	ate of dispa ed in a co ary Group I ber State	tch, the	the animal has been resider or part of the territory of was imported during the 6 be Union before entering the with point II.2.2.;]
⁽³⁾⁽⁵⁾ either	[11.2.2.	the animal				untry or part o	of the territor	ory of	a country which is assigne
	⁽³⁾ either	protected since entry in accorda	from v y into t ance w	vecto the c vith p	r insects for a ountry or part point II.2.1 fror	period of at le of the territory	east 40 day of the cou State of the	ys prid intry o Unio	y of the country of dispatc or to the date of dispatch, of dispatch, if it was importe n or a country or part of the c, D, E or G;]]
	⁽³⁾ or	least 40 da of the cou State of th Group A, I	ays pri intry of ie Unio B, C, I d by th [the	f disponsor on or O, E ne Ol anii	the date of dispatch, if it was a country or p or G, and the IE as officially mal is a regi	spatch, or since s imported in a art of the territo country or par free of African	e entry into accordance ory of a count of the ter horse sich as defined	the control the with untry writory kness	supervision for a period of a country or part of the territor point II.2.1 from a Membe which is assigned to Sanitar of the country of dispatch i and Article 2(c) of Commissio
		⁽³⁾ or							ntry in which African horserior to the date of dispatch;
⁽³⁾⁽⁵⁾ or	[11.2.2.		l is dis	patc					a country which is assigne
	⁽³⁾ either	of quarant date) to two hours veterinary insecticide isolation fr	prior to super e effect rom eq	o sur rvisio ctive quida	during the 40 (insert date nset until two h on, following to against Culic te not being pi	days prior to to days prior to to do confined to do confined to do confined days prior to	the date of of the vector- irise and ex of insect of the removement to the l	dispat prote ercise repelle al fro Union	ch (insert name ch from
	⁽³⁾ or	[permaner (insert nar dispatch a	ntly come of and co	nfine quar onsta	ed in the appro cantine station ant monitoring	oved vector-produced) during the pe	oof quarar eriod of at protection	ntine s least	station of
II.3.	Attestation	of vaccinat	tion an	nd he	ealth tests				
(3)either	[II.3.1.				•	ainst African h previous vacc		ess in	the country of dispatch an
⁽³⁾ or	[II.3.1.					·		, and	this vaccination was carried
	(3)either					date of dispato			
	⁽³⁾ or								admission into the part of the is dispatched;]]
⁽³⁾⁽⁵⁾ or	[II.3.1.	to Sanitary date) not a protected	y Grou more t quara	p F a than ntine	and was vaccing 24 months are by administr	nated against And at least 40 of at least 40 of a regi	African hors days prior istered vac	se sicle to the cine	a country which is assigne kness on(inset a date of entry in the vector according to manufacturer of the African horse sickness

	II.a.	Certificate reference number	II.b. Local reference number
	II.a.	Certificate reference number	II.b. Local reference number
II.3.2.		not vaccinated against Venezuelan eques prior to the date of dispatch from	uine encephalomyelitis during the
⁽³⁾ either		ich all parts of the territory are free of Ver t least 2 years prior to the date of dispatch	
⁽³⁾⁽⁵⁾ Or	Venezuelan equi	ritory of a country which is assigned to Sa uine encephalomyelitis for a period of at enezuelan equine encephalomyelitis occ ountry of dispatch, and	least 2 years prior to the date of
	prima not le and w the da tempe equin daily,	accinated against Venezuelan equine en ry course and revaccinated according to ss than 60 days and no more than 12 m ras kept in vector-protected quarantine for ate of dispatch, and during that period rema- terature, taken daily, remained within the ne e animal on the same holding which showed was subjected to a blood test for virus shalomyelitis with negative results;]]	manufacturer's recommendations on the prior to the date of dispatch, a period of at least 21 days prior to dinied clinically healthy, and its body ormal physiological range, and anyed a rise in body temperature, taken
	vecto remai the no show virus the a equin less t	t vaccinated against Venezuelan equine r-protected quarantine for a period of at le ned clinically healthy, and its body temper ormal physiological range, and any equine ed a rise in body temperature, taken daily isolation for Venezuelan equine encephal nimal to be dispatched was subjected to be encephalomyelitis with negative result han 14 days after the date of entry intend protected from vector insects until dispatched protected from vector insects until dispatched.	ast 21 days, and during that period rature, taken daily, remained within a animal on the same holding which, was subjected to a blood test for omyelitis with negative results, and a diagnostic test for Venezuelan conducted on a sample taken not be vector-protected quarantine and
	encep taken on prior ' PCR Venez result and r samp and ir	subjected to a haemagglutination inhib shalomyelitis carried out by the same labo on two occasions with an interval of 21 (insert date), the second of which we to the date of dispatch, without an increa (reverse transcription-polymerase chain zuelan equine encephalomyelitis virus g on a sample taken within 48 hours prior that been protected from vector attacks ling until loading for dispatch, by combine is ecticides on the horse and disinsectization it is transported;]]	ratory on the same day on samples days on (insert date) and as taken during a period of 10 days se in the antibody titre, and a RT-reaction) test for the detection denome, carried out with negative o dispatch, on (insert date), from the moment of the RT-PCR d use of approved insect repellents
⁸⁾⁽⁵⁾ either [II.3.3.	anaemia, where	patched from Iceland, which is certified as it was continuously resident since birth have entered Iceland from other countries;	and did not come into contact with
³⁾ or [II.3.3.	the animal was Coggins test) o	subjected with negative result to an agar r to an ELISA for equine infectious anael (insert date), this being within	gel immunodiffusion test (AGID or

(3)or

[a period of 30 days prior to the date of dispatch from a country or part of the territory of a country which is assigned to Sanitary Group D, E or F;]]

COUNTRY Transit - Equidae

		T		
		II.a. C	Certificate reference number	II.b. Local reference number
⁽³⁾ [II.3.4.	to Sanital was repo complem in 5 on a	ry Group B rted during ent fixation	ched from a country or part of the terri or E, or from Brazil, China or Thailand, g a period of 3 years prior to the date on test for glanders carried out with neg ple taken on	or from a country in which glanders of dispatch, and was subjected to a ative result at a serum dilution of 1
⁽³⁾⁽⁵⁾ [II.3.5.			ched from a country or part of the terri	tory of a country which is assigned
	⁽³⁾ either	the count	and Eastern equine encephalomyelitis ry or part of the territory of the countr ears prior to the date of dispatch;]]	
	⁽³⁾ or	according 30 days and East	nal was vaccinated with a complete g to manufacturer's instructions within prior to the date of dispatch with ina ern equine encephalomyelitis, the (insert date);]]	a period of 6 months and at least activated vaccine against Western
	⁽³⁾ or	a vector	al was kept for a period of at least 21 of -protected quarantine and during lutination inhibition tests for V omyelitis carried out by the same labor	this period was subjected to Vestern and Eastern equine
		⁽³⁾ either	[on a sample of blood taken on of 10 days prior to the date of dispatc	
		⁽³⁾ Or	[on samples of blood taken on two or 21 days on (insert date) the second of which was taken within of dispatch, without increase in antibo	and on (insert date), a period of 10 days prior to the date
⁽³⁾ [II.3.6.	to Sanita	ry Group (ched from a country or part of the terri G, or from a country in which Japane during a period of at least 2 years pr	se encephalitis has been officially
	⁽³⁾ either	around th	om a holding situated in the centre o at holding where there has been no ca of 21 days prior to the date of dispatch	se of Japanese encephalitis during
	⁽³⁾ or	to the da	in a vector-protected quarantine during te of dispatch, and during that period within the normal physiological range	the body temperature, taken daily,
		⁽³⁾ either	[to a haemagglutination inhibition or vi encephalitis carried out by the same samples of blood taken on two occas days on	e laboratory on the same day on sions with an interval of at least 14 l on(insert date), the eriod of 10 days prior to the date of d increase in antibody titre between
		⁽³⁾ or	[to a Ig-M capture ELISA test for the Japanese encephalitis virus with neg sample taken not earlier than 7 commenced on(insert date vector insects until dispatch;]]]	ative result, carried out on a blood days after the date the isolation

II.a. Certificate reference number II.b. Local reference number II.b. Local reference number II.b. Local reference number II.b. Local reference number revaccinated according to manufacturer's recommendations during a period of less than 21 days and not more than 12 months prior to the date of dispatch;] II.a.	COUNTRY						Transit - Equida
revaccinated according to manufacturer's recommendations during a period of reless than 21 days and not more than 12 months prior to the date of dispatch;]] the animal is dispatched from a country or part of the territory of a country which is assign to Sanitary Group E and was subjected to a serological test for African horse sickness described in Annex IV to Directive 2009/156/EC, which was carried out by the same laborate on the same day **Beither** [Interest date) and on			II.a.	Certificate	reference number	II.b.	Local reference number
to Sanitary Group E and was subjected to a serological test for African horse sickness described in Annex IV to Directive 2009/156/EC, which was carried out by the same laborate on the same day on the same day "elither" [on blood samples taken on two occasions with an interval of between 21 and days, on		⁽³⁾ or	revaccir	ated acco	rding to manufacturer's recor	nmend	lations during a period of no
(a) (a) (b) (a) (a) (a) (a) (a) (a) (a) (a) (a) (a	⁽³⁾⁽⁵⁾ either [II.3.7.	to Sanita describe on the s	ary Group ed in Anne ame day [on bloo days, or	E and wa (IV to Direct d samples	s subjected to a serological ctive 2009/156/EC, which was taken on two occasions with (insert date) and or	test for s carrie n an in n	or African horse sickness a ed out by the same laborator terval of between 21 and 3 (insert date), the
(3)either [the second sample was subsequently tested with negative result in an agent identification test as described in Annex to Directive 2009/156/EC.]]]] (3)or [the two samples were tested without more than a two-forcease in antibody titre in a virus neutralisation test described in point 2.4 of Chapter 2.5.1. of the OIE Terrests Manual for Diagnostic Tests and Vaccines.]]]] (3)or [on a blood sample taken on			(3)either	[with neg	gative results in each case.]]]		
result in an agent identification test as described in Annex to Directive 2009/156/EC.]]]] (3) or [the two samples were tested without more than a two-force increase in antibody titre in a virus neutralisation test described in point 2.4 of Chapter 2.5.1. of the OIE Terrest Manual for Diagnostic Tests and Vaccines.]]]] (3) or [on a blood sample taken on			(3) <i>or</i>	[with a p	ositive result in the first samp	ole, and	t
increase in antibody titre in a virus neutralisation test described in point 2.4 of Chapter 2.5.1. of the OIE Terrest Manual for Diagnostic Tests and Vaccines.]]] (3) or [on a blood sample taken on				⁽³⁾ either	result in an agent identific	cation t	
prior to the date of dispatch, and the country or part of the territory of the country dispatch is recognised by the OIE as officially free of African horse sickness and "implementing Regulation (EU) 2018/659;]]] (3) or [the country of dispatch is not adjacent to a country in which African hor sickness has occurred during the period of 2 years prior to the date dispatch;]]] (3)(5) or [II.3.7. the animal is dispatched from a country or part of the territory of a country which is assign to Sanitary Group F, and [was subjected to a serological test for African horse sickness as described Annex IV to Directive 2009/156/EC, which was carried out by the same laborate on the same day on blood samples taken on two occasions with an inter of between 21 and 30 days, on				⁽³⁾ or	increase in antibody titre described in point 2.4 of 0	e in a Chapte	virus neutralisation test a r 2.5.1. of the OIE Terrestri
[II.3.7. Ithe animal is dispatched from a country or part of the territory of a country which is assign to Sanitary Group F, and (3)either [was subjected to a serological test for African horse sickness as described Annex IV to Directive 2009/156/EC, which was carried out by the same laborate on the same day on blood samples taken on two occasions with an intern of between 21 and 30 days, on		⁽³⁾ or	prior to t	he date of	dispatch, and the country or	part of	the territory of the country
sickness has occurred during the period of 2 years prior to the date dispatch;]]] the animal is dispatched from a country or part of the territory of a country which is assign to Sanitary Group F, and (**]**bether** [was subjected to a serological test for African horse sickness as described Annex IV to Directive 2009/156/EC, which was carried out by the same laborate on the same day on blood samples taken on two occasions with an inter of between 21 and 30 days, on			⁽³⁾ either				in Article 2(c) of Commission
to Sanitary Group F, and (3)either [was subjected to a serological test for African horse sickness as described Annex IV to Directive 2009/156/EC, which was carried out by the same laborate on the same day on blood samples taken on two occasions with an inter of between 21 and 30 days, on			⁽³⁾ or	sickness	has occurred during the pe		
(3) or [with a positive 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 Directive 2009/156/EC.]]]] (3) or [the two samples were tested without more than a two-fincrease in antibody titre in a virus neutralisation test described in point 2.4 of Chapter 2.5.1. of the OIE Terresti	⁽³⁾⁽⁵⁾ or [II.3.7.	to Sanita	ary Group [was su Annex I' on the of betwee on introduc	F, and bjected to V to Direct same day een 21	a serological test for Africa ive 2009/156/EC, which was on blood samples taken cand 30 days, on . (insert date), the first sample vector-protected quarantine	carried two	se sickness as described dout by the same laborato occasions with an interv (insert date) ar taken less than 7 days aftr
(3)either [the second sample was subsequently tested with negative result in an agent identification test as described in Annex IV Directive 2009/156/EC.]]]] (3)or [the two samples were tested without more than a two-fincrease in antibody titre in a virus neutralisation test described in point 2.4 of Chapter 2.5.1. of the OIE Terresti			(3)either	[with neg	gative results in each case.]]]		
result in an agent identification test as described in Annex IV Directive 2009/156/EC.]]]] (3)or [the two samples were tested without more than a two-found increase in antibody titre in a virus neutralisation test described in point 2.4 of Chapter 2.5.1. of the OIE Terresti			(3) <i>or</i>	[with a p	ositive result in the first samp	ole, and	d
increase in antibody titre in a virus neutralisation test described in point 2.4 of Chapter 2.5.1. of the OIE Terresti				⁽³⁾ either	result in an agent identificat		
				⁽³⁾ or	increase in antibody titre described in point 2.4 of C	in a hapter	virus neutralisation test a 2.5.1. of the OIE Terrestri

COUNTRY Transit - Equidae

			II.a. Certificate reference number II.b. Local reference number
		⁽³⁾ or	[was subjected to a serological and an agent identification test for African hors sickness as described in Annex IV to Directive 2009/156/EC, carried out with negative result in each case on a blood sample taken on
		⁽³⁾ or	[was subjected to an agent identification test for African horse sickness a described in Annex IV to Directive 2009/156/EC, carried out with negative result o a blood sample taken on
II.4.	Attestation	on of the tra	ansport conditions
⁽³⁾⁽⁵⁾ eithe	er[II.4.1.	to Sanita to the U coming	mal is dispatched from a country or part of the territory of a country which is assigned ary Group A, B, C, D, E or G and arrangements have been made to transport it directly inion, without passing through a market, marshalling or assembly centre and without into contact with other equidae not complying with at least the same health nents as described in this health certificate.]
⁽³⁾⁽⁵⁾ or [II.4.1.		to Sanita protecte by a hea	mal is dispatched from a country or part of the territory of a country which is assigne ary Group F and arrangements have been made to transport it directly from the vector ad quarantine station without coming into contact with other equidae not accompanie alth certificate either for imports or for temporary admission into the Union or for trans the Union [to the airport under vector-protected conditions and arrangements have bee made for the aircraft to be cleansed and disinfected in advance with a disinfectar officially recognised in the third country of dispatch, and sprayed against vector insects just prior to take off.]]
		⁽³⁾ Or	[to a sea port in that country or part of the territory of the country under vector protected conditions and arrangements have been made to transport it on a vesse which is scheduled directly to a port in the Union without calling into a port situate in a country or part of the territory of a country not approved for the entry into th Union of equidae, in stalls which were cleansed and disinfected in advance with disinfectant officially recognised in the third country of dispatch and sprayed against vector insects just prior to departure.]]
	II.4.2.	complyi	ments have been made and verified to prevent any contact with other equidae noing with at least the same health requirements as described in this health certificate the period from certification until dispatch to the Union.
	II.4.3.	and disi	asport vehicles or containers in which the animal is going to be loaded were cleaner infected before loading with a disinfectant officially recognised in the third country on and they are so constructed that faeces, urine, litter or fodder cannot escape during tation.
	11.4.4	the Unio	ine animal is proceeding to
II.5.	Attestatio	on of anima	l welfare
			ed in Box I.28. was examined today ⁽¹⁾ and found fit to be transported on the intended ements were made to protect its health and well-being effectively at all stages of the

COUNTRY Transit - Equidae

		II.a.	Certificate reference number	II.b.	Local reference number
Notes:					
Part I:					
Box I.6.:	Person responsible	for the	e load in Union.		
Box I.8.:			untry or part of the territory of the country on Implementing Regulation (EU) 2018/6		oatch as appearing in column
Box I.15.	and information is t	to be p	vay wagons or container and lorries), fli rovided. In case of unloading and reloa entry into the Union.		
Box 1.23.	: The container numb	ber and	the seal number (if applicable) should	be inclu	ded.
Box 1.28.			Equus caballus, Equus asinus, Equus uus zebra, Equus grevyi, or indicate an		
	the identification do	ocumer the ide	animal must bear an individual identifient as defined in Article 2(b) of Commisentification system (such as ear tag, the animal.	sion Im	plementing Regulation (EU)
	If a passport according authority which valid		s the animal, its number should be stat.	ted and	the name of the competent
	Age: Date of birth (dd/mm/	/yyyy).		
	Sex ($M = male, F =$	female	e, C = castrated).		
(2) (3) (4) (5) (5)	lay before loading of the entry into the Union the date of authorisation the country referred to into the country or gainst the error code of the country or no columns 3 and 5 responded as appropriate. Delete as appropriate. Delete statement if the Statements that relate	ne animal of these of the sen for train point of the sen for train point of the sen for th	on the day of loading or in the case of a nal for dispatch to the Member State of c se animals shall not be allowed when the ansit through the Union from the respect II.1.1., or during a period where restrict equidae from this country or this part of the territory of the country of dispatch, ally of Annex I to Commission Implement tion in point II.1.3. applies to the entire of and exclusively to a Sanitary Group or part of its territory, is assigned, may be is maintained.	destinati e anima etive cou ive mea he territo and the s ing Regi country of	on in the Union. als were loaded either prior to untry or part of the territory of sures have been adopted by ory of the country of dispatch. Sanitary Group as appearing ulation (EU) 2018/659. of dispatch. from the Sanitary Group to
This hea	Ith certificate shall:				
th	ne Member State of de nd undergo the veterin	estination			
	e made out to a single			~.	
(d) c	consist of a single she ndivisible by inserting p	et of p	olour different to the colour of the printing paper or all sheets of paper required a umbers and total number of pages, and f the page and those pages are stapled	re part each p	age shall bear the certificate
Official v	eterinarian				
	me (in capital letters):			O	ualification and title:
	27 5) 5)				
Da	ite:			Sig	gnature:

Stamp:

				e owner or representative or ough the Union of an equin		
Ide	ntification of	the anin	nal ⁽¹⁾			
	ecies (Scient ne)	ific	Identification system	Identification number	Age	Sex

I, th	ne undersign the anima	l		owner ⁽²⁾ of the animal describ		
	either			(insert name of country or s prior to the date of dispatch;		country or dispatch)
	⁽²⁾ or			name of country or part of the ast 40 days prior to the date of		dispatch) during the
		(a)		om (insert rittory of country of dispatch)	ame of country from wh	nere animal entered
		(b)		om (insert rittory of country of dispatch)	ame of country from wh	ere animal entered
		(c)		om (insert rittory of country of dispatch);]	ame of country from wh	nere animal entered
-			of 15 days prior to the date	of dispatch the animal has no le to equidae;	ot been in contact with an	imals suffering from
•				solation as applicable in accidentitions of the country of disp		f the accompanying
-	the conditi	ons for		n accordance with point II.4.		Ith certificate for the
ū		ortation	will be effected in such a w	ay that health and well-being	of the animal can be pro	otected effectively at
ė.				on on place of border post of exit);	(insert date) at	the border post of
Na	me and addr	ess of th	ne owner ⁽²⁾ or representative	9(2)-		
Dat	te:		(dd/mm/yyyy)			
				(Signature)		
(1)	quagga, Ed Identification document a	quus zel on syste as defin	bra, Equus grevyi, or indicat m: The animal must bear a ed in Article 2(b) of Comm	Equus asinus, Equus africar te any cross between those. In individual identifier which p ission Implementing Regulat Inder) and the anatomic place	permits to link the animal ion (EU) 2018/659. Spec	to the identification
	If a passport validated it.		mpanies the animal, its nu	mber should be stated and	the name of the compe	tent authority which
			dd/mm/yyyy). female, C = castrated).			
(2)	Delete as a					

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 not more than 30 days

NTRY	Y:			Veterinary certificate to E
1.1		Consignor Name	1.2	I.2. Certificate reference No I.2.a.
		Address	1.3	I.3. Central competent authority
		Tel.	1.4	I.4. Local competent authority
1.5		Consignee Name	1.6	1.6.
		Address		
		Postcode Tel.		
1.5		Country of ISO code I.8. Region of origin Code	9 1.9	I.9. Country of ISO code I.10. Region of Code destination
11	1	Place of origin	11	I.12. Place of destination
	•	•	'	Name
		Name Approval number Address		Name Address
				Postcode
1.1	3.	Place of loading	1.1	I.14. Date of departure
1.1	5.	Means of transport	I.1	I.16. Entry BIP in EU
		Aeroplane Ship Railway wagon I Road vehicle Other Identification Documentary references		I.17. No(s) of CITES
1.1	8.	Description of animal		I.19. Commodity code (HS code)
				I.20. Quantity
1.2	1.			I.22. Number of packages
1.2	3.	Seal/Container No		1.24.
1.2	5.	Animal certified for:		
		Registered horse		
1.2	6.			I.27. For import or admission into EU
1.2	8.	Identification of the animal		
	(5		ntifica	ication number Age Sex

COUNTRY

Re-entry after temporary export of not more than 30 days Registered horse

				Trogiotorou moros						
			II.a. Certificate reference number	II.b. Local reference number						
	II.	Attestation of animal	health and welfare							
	I, the unders	igned official veterinaria	n, hereby certify, that the animal described in Box I.28	3.:						
	_	is a registered horse as	s defined in Article 2(c) of Commission Implementing	Regulation (EU) 2018/659;						
uo	_	was examined today infestation;	was examined today (1) and found free of clinical signs of diseases and of obvious signs of ectoparasite infestation;							
tificati	_	is not intended for slau	ghter under a national programme of infectious or cor	ntagious disease eradication;						
Part II: Certification	_	meets the requirement	s attested in points II.1. to II.3. of this certificate;							
Part	_	is accompanied by the	written declaration, signed by the owner of the horse	or the representative of the owner.						
	II.1.	Attestation on third cou	untry or part of the territory of third country and holding	g of dispatch						
	II.1.1.	The animal is dispatched from								
	II.1.2.	in the country of dispatch the following diseases are compulsorily notifiable: African horse sickness, doc (Trypanosoma equiperdum), glanders (Burkholderia mallei), equine encephalomyelitis (of all types inclu Venezuelan equine encephalomyelitis), equine infectious anaemia, vesicular stomatitis, rabies and anthrax;								
	II.1.3.	the animal is dispatche	d from a country or part of the territory of a country:							
		which there African ho	nsidered free from African horse sickness in accordate has been no clinical, serological (in unvaccinated educes sickness during the period of 2 years prior to the no vaccinations against the disease during the period of the p	quidae) or epidemiological evidence of e date of dispatch and in which there						
		b) in which Vothe date of	enezuelan equine encephalomyelitis has not occurre dispatch;	d during the period of 2 years prior to						
		c) in which do	ourine has not occurred during the period of 6 months	prior to the date of dispatch;						
		d) in which gla	anders 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	ome from a holding, and to the best of my knowled .7. was not in contact with animals from holdings, who to in points II.1.4.1. to II.1.4.7. and which last for:							
		(3) [II.1.4.1. in the case	of equidae suspected of having contracted dourine,							
			6 months beginning on the date of the last actua suspected of having contracted dourine or infected wi							
		(⁴) and/or	in the case of a stallion, until the animal is castrated;]							
			30 days following the date of completion of the clear after all animals of susceptible species have been sla							

COUNTRY

Re-entry after temporary export of not more than 30 days Registered horse

			II.a. Certificate reference number	II.b. Local reference number
	(³) [II.1.4.2.	in the case	of glanders,	
			[6 months beginning on the day on which the eq subjected with positive results to a test for the of Burkholderia mallei or antibodies to that pathogen, we	detection of the causative pathogen
			[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	of equine encephalomyelitis of any type,	
			[6 months beginning on the day on which the equidae slaughtered;]	suffering from the disease have been
		, ,	6 months beginning on the day on which the equidae Nile Fever, Eastern equine encephalomyelitis or We died, been removed from the holding or fully recovere	estern equine encephalomyelitis have
			[30 days following the date of completion of the clear after all animals of susceptible species have been slat	
	II.1.4.4.	slaughtere gel immur	e of equine infectious anaemia, until the date on whi d, the remaining equine animals on the holding have nodiffusion test (AGID or Coggins test) carried c ons 3 months apart;	shown a negative reaction in an agar
	II.1.4.5.	in the case	of vesicular stomatitis,	
		(4) either	[6 months following the last case;]	
			[30 days following the date of completion of the clear after all animals of susceptible species have been slat	
	II.1.4.6.		of rabies, 30 days following the last case and the day of the premises;	ate of completion of the cleansing and
	II.1.4.7.		of anthrax, 15 days following the last case and the day of the premises;	ate of completion of the cleansing and
II.1.5.			rledge, during the period of 15 days prior to the date of fected or suspected of an infectious or contagious dis	
II.2.	Attestation	n of residenc	e and pre-export isolation	
II.2.1.	The anima	al was import	ed on (insert date)	
	(4) either	[directly fro	m the EU Member State(ii	nsert name of EU Member State);]
	(⁴) or		untry or part of the territory of a countrytilitions at least as strict as those set out in this certifica	
II.2.2.	part of the veterinary	territory of a supervision	the Union less than 30 days ago, and since exit from a country (¹) other than those of the same Sanitary G accommodated in separated stables without cominuring racing, competition or the cultural event.	Group, and resident on holdings under
II.3.	Attestation	n of animal w	elfare	
			in Box I.28. was examined today ⁽¹⁾ and found fit to be e made to protect its health and well-being effectively	

COUNTRY

Stamp:

Re-entry after temporary export of not more than 30 days

COUNTRY		ке-епту аптет тетпро	orary export of not more than 30 days Registered horse
		II.a. Certificate reference number	II.b. Local reference number
Notes:			
Part I:			
Box I.8.:		ne country or part of the territory of the country as tring Regulation (EU) 2018/659.	appearing in column 3 of Annex I to
Box I.15.:		railway wagons or container and lorries), flight rovided. In case of unloading and reloading, the considerion.	
Box I.23.:	The container number	and the seal number (if applicable) should be include	ed.
Box I.28.:	identification documen the identification syste	The animal must bear an individual identifier wh t as defined in Article 2(b) of Commission Implement m (such as ear tag, tattoo, brand, transponder) and t ccompanying passport must be stated and the na	ting Regulation (EU) 2018/659. Specify the anatomic place used on the animal.
	Age: Date of birth (dd/	mm/yyyy).	
	Sex (M = male, F = fer	nale, C = castrated).	
Part II:			
	ertificate must be issued ember State of destination	on the day of loading or on the last working day befor on in the Union.	ore loading of the animal for dispatch to
to the referre	date of authorisation for ed to in point II.1.1., or	export of this registered horse shall not be allowed wore-entry into the Union from the respective country of during a period where restrictive measures have be country or this part of the territory of the country of di	or the part of the territory of the country een adopted by the Union against the
		of the territory of the country and the Sanitary Grounission Implementing Regulation (EU) 2018/659.	oup as appearing in columns 3 and 5
(3) Delete	e statement if the attesta	tion in point II.1.3. applies to the entire country of disp	patch.
(4) Delete	e as appropriate.		
This health	certificate shall:		
State		guage understood by the certifying officer and one of the second by the certifying officer and one of the second by the certifying officer and one of the second by the se	
(b) be ma	ade out to a single consig	nee;	
(c) be sig	gned and stamped in a co	olour different to the colour of the printing;	
insert		paper or all sheets of paper required are part of a otal number of pages, and each page shall bear the re stapled and stamped.	
Official vete	rinarian		
Name	e (in capital letters):		Qualification and title:
Date:			Signature:

			for the re-entry after t	owner or representative of the o emporary export of a registered empetition and cultural events		
Ider	itification o	f the animal (1)				
Sp	ecies (Scie	ntific name)	Identification system	Identification number	Age	Sex
	Equus c	aballus				
I, th	e undersig	ned owner (²) or	representative of the owne	r (2) of the registered horse descri	bed above, hereby dec	lare, that:
_	the horse					
	(²) either		ily exported from the Union prior to this declaration;]	to the country of dispatch on		(insert date) less
	(²) or		ountry of dispatch on There horse entered country	(insert date) fro of dispatch);]	om	(insert name of
_			ays prior to the date of disparansmissible to equidae;	atch the horse has not been in cor	ntact with animals suffe	ring from infectious
_	the transpof the jou		effected in such a way that	health and well-being of the horse	e can be protected effe	ctively at all stages
_				n as applicable in accordance w country of dispatch are fulfilled.	ith point II.2. of the acc	companying health
Nan	ne and add	lress of the own	er (²) or representative (²):			
Date	ə:		(dd/mm/yyyy)			
(1)	Article 2(to transponder If a passponder Age: Date Sex (M = r	o) of Commission er) and the anaton	n Implementing Regulation (E nic place used on the animal. ne animal, its number should be yyy).	dentifier which permits to link the anir (U) 2018/659. Specify the identifical e stated and the name of the competen	tion system (such as ea	r tag, tattoo, brand,

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

▼<u>M2</u>

Chapter 1

Model health certificate and model declaration applicable to re-entry into the Union of registered horses for competition after temporary export for a period of not more than 90 days to participate in equestrian events organised under the auspices of the Fédération Equestre Internationale (FEI)

(Test event in preparation of the Olympic Games, Olympic Games, Paralympics, World Equestrian Games/World Championship, Asian Equestrian Games, American Equestrian Games (including the PanAmerican Games, South American Games, Central American and Caribbean Games), Endurance World Cup in United Arab Emirates, LG Global Champions Tour)

cou	NTRY:	Veterina	ary certificate to EU
	I.1. Consignor Name	I.2. Certificate reference No	I.2.a.
	Address Tel.	I.3. Central competent authority	
nent		I.4. Local competent authority	
Part I: Details of dispatched consignment	I.5. Consignee Name Address Postcode Tel.	1.6.	
ails of disp	I.7. Country ISO code I.8. Region Code of origin	I.9. Country of ISO code I destination	.10. Region Code of destination
: Det	I.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	
	I.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	1.19.	Commodity code (HS code) 01 01
			I.20. Quantity
	I.21.		I.22. Number of packages
	I.23. Seal/Container No		1.24.
	I.25. Animal certified for:		
	Registered horse		
	1.26.	1.27. For import or admission	n into EU
	I.28. Identification of the animal	1	
	Species (Scientific name) Identification system	Identification number A	ge Sex

Equus caballus

COUNTRY

Re-entry after temporary export of not more than 90 days Specific competitions - Registered horse

				II.a.	Certificate reference number	II.b.	Local reference number
Part II: Certification	II. I, the ui	is a register was exam infestation is not inter meets the	fficial veteri ered horse a ined today ; nded for sla requiremen	as defined in A 1) and found ughter under ts attested in	welfare y certify, that the animal described in Article 2(c) of Commission Implement free of clinical signs of diseases ar a national programme of infectious or points II.1. to II.3. of this certificate; laration, signed by the owner of the	ting Regu nd of obv	pus disease eradication;
	II.1. II.1.1.	The anima	al is dispate part of the	hed from e territory of	of the territory of third country and ho (insert name of country or µ a country which on the date of iss d to Sanitary Group(2);	part of th	e territory of a country), a
	II.1.2.	in the cour (<i>Trypanos</i> Venezuela	ntry of dispa oma equipe in equine e	atch the follow erdum), gland ncephalomyel	ving diseases are compulsorily notifia ers (<i>Burkholderia mallei</i>), equine enc iitis), equine infectious anaemia, vesio	ephalomy cular ston	yelitis (of all types including
	II.1.3.	the animal	which is c and in wh evidence in which t	onsidered fre ich there has of African hor	untry or part of the territory of a count e from African horse sickness in acco been no clinical, serological (in unvac ses sickness during the period of 2 ye sen no vaccinations against the disea atch;	ordance v ccinated e ars prior	equidae) or epidemiological to the date of dispatch and
		b) c)	in which V to the date	enezuelan ed e of dispatch;	quine encephalomyelitis has not occur	BMB 86	5
	II.1.4.	d) the animal points II.1.	in which g does not c 4.1. to II.1.	landers has r ome from a he 4.7. was not	not occurred during the period of 6 mo- lolding, and to the best of my knowled in contact with animals from holding in points II.1.4.1. to II.1.4.7. and which	onths prio ge for the s, which	or to the date of dispatch; e time periods referred to in
		⁽³⁾ [II.1.4.1.		[6 months be	suspected of having contracted douring eginning on the date of the last actual having contracted dourine or infected	l or possi	
			(4)and/or	[in the case of	of a stallion, until the animal is castrat	ed;]	
					owing the date of completion of the er all animals of susceptible species h		
		(3)[II.1.4.2.	in the cas	e of glanders,			10.400
			⁽⁴⁾ either	subjected with	eginning on the day on which the eq th positive results to a test for the d mallei or antibodies to that pathogen	letection	of the causative pathogen
					owing the date of completion of the er all animals of susceptible species h		
		II.1.4.3.	in the cas	e of equine e	ncephalomyelitis of any type,		77290 gsc wa 507 176
			⁽⁴⁾ either	[6 months be been slaught	ginning on the day on which the equi ered;]	dae suffe	ring from the disease have
			(4)and/or	West Nile	eginning on the day on which the equ Fever, Eastern equine encept velitis have died, been removed from	nalomyelit	tis or Western equine
			(4)and/or		owing the date of completion of the er all animals of susceptible species h		

COUNTRY

Re-entry after temporary export of not more than 90 days Specific competitions - Registered horse

			II.a.	Certificate reference number	II.b.	Local reference numbe		
	II.1.4.4.	been slaugh in an agar g	ntered, the re	fectious anaemia, until the date on emaining equine animals on the holo fusion test (AGID or Coggins test) ca ths apart;	ding have	shown a negative reaction		
	II.1.4.5.		of vesicular s months follo	stomatitis, owing the last case;]				
				owing the date of completion of the rall animals of susceptible species I				
	II.1.4.6.		of rabies, 30 ction of the p	days following the last case and the remises;	date of	completion of the cleansing		
	II.1.4.7.		of anthrax, 1stion of the p	5 days following the last case and the remises;	e date of	completion of the cleansin		
II.1.5.				g the period of 15 days prior to the d or suspected of an infectious or co				
11.2.	Attestation	n of residence	and pre-ex	port isolation				
II.2.1.		nal was impo		ne country or part of the territor t date)	ry of the	e country of dispatch of		
	(4)either	[directly from	n the EU Me	mber State(ii	nsert nan	ne of EU Member State);]		
	⁽⁴⁾ or			of the territory of a country t as strict as those set out in this cer		(insert name of country		
11.2.2.	the anima	l exited from t	he Union					
	⁽⁴⁾ either	territory of a territory of a accommoda status exce participating	a country oth the country ated in sepa pt during co g in the LG G	and since exit from the Union wa er than those of the same Sanitary of dispatch it was resident on hole rated stables without coming into competition, and it has taken part in a slobal Champions Tour	Group. I dings und ontact with or was st	n the country or part of the der veterinary supervision th equidae of lower healt		
		(4)either [ir	n the Metrop	olitan area of Mexico City, Mexico;]]				
		(4)and/or [ir	n Miami, Uni	tes States of America;]				
		(4) <i>or</i> [ir	n Shanghai,	China;]]				
	⁽⁴⁾ Or	[less than 60 days ago, and since exit from the Union was never in a country, or part of territory of a country other than those of the same Sanitary Group. In the country or part of territory of the country of dispatch it was resident on holdings under veterinary supervisi accommodated in separated stables without coming into contact with equidae of lower heat status except during competition and it has taken part in or was stabled together with horse participating in (4)either [the Asian Games in						
		(4)or [tl	he American	Games ⁽⁵⁾ in		(insert place).ll		
		10000		e World Cup in United Arab Emirate		V		
	(4)							
	⁽⁴⁾ Or	territory of a	country ⁽¹⁾ o	, and since exit from the Union wa ther than those of the same Sanitan of dispatch it was resident on hol	Group.	In the country or part of the der veterinary supervision		
		status exce participating	pt during co in	rated stables without coming into competition and it has taken part in contact to the contact to	or was st	abled together with horse		
		status exce participating	pt during co in		or was st	abled together with horse		

COUNTRY

Re-entry after temporary export of not more than 90 days Specific competitions - Registered horse

		II.a.	Certificate reference number	II.b.	Local reference number
	(4)or [the	Paralymp	pics in		(insert place).]]
	(4) <i>or</i> [the	e World Ed	uestrian Games/World Championsh	nips in	(insert place).]]
II.3.	Attestation of animal welf	are			
			was examined today(1) and found finade to protect its health and well-li		
Notes:					
Part I:					
Box I.8.			or part of the territory of the country egulation (EU) 2018/659.	as appea	ring in column 3 of Annex I
Box I.1		rovided. In	gons or container and lorries), flight case of unloading and reloading, the Union.		
Box I.23	3.: The container number	and the s	eal number (if applicable) should be	included.	
Box I.28	identification documer Specify the identificat	nt as define ion systen The numb	al must bear an individual identifier w ed in Article 2(b) of Commission Impl n (such as ear tag, tattoo, brand, tr er of the accompanying passport n ated it.	ementing ansponde	Regulation (EU) 2018/659. er) and the anatomic place
	Age: Date of birth (dd/	/mm/yyyy)	2		
	Sex ($M = male, F = fe$	male, C =	castrated).		
Part II:					
(1)	The certificate must be issudispatch to the Member Sta		day of loading or on the last working	day befo	ere loading of the animal for
	The re-entry after temporar either prior to the date of au territory of the country refer	ry export outhorisation in	of this registered horse shall not be a in for re-entry into the Union from the point II.1.1., or during a period wh ry of equidae from this country or this	respective ere restric	ve country or the part of the ctive measures have been
(2)	Code of the country or part		ritory of the country, and the Sanita mission Implementing Regulation (E		
(3)	Delete statement if the atte		point II.1.3. applies to the entire cour		
(4) (5)	Delete as appropriate. Including the PanAmerican	Games, S	South American Games, Central Ame	erican and	d Caribbean Games.
This he	alth certificate shall:				
No.		n and of th	nderstood by the certifying officer and the Member State where the registered:		
(b)	be made out to a single con	nsignee;			
(c) (d)	consist of a single sheet of by inserting page numbers	paper or a s and tota	ferent to the colour of the printing; ill sheets of paper required are part of I number of pages, and each page ose pages are stapled and stamped.	shall be	
Official	veterinarian				
N	ame (in capital letters):			Qualific	ation and title:
D	ate:			Signatu	re:
0	tamo				

		for the		e owner or representative prary export of a registered		ion
Ide	ntification of the	e animal ⁽¹⁾				
	ecies (Scientific me)	e lo	lentification system	Identification number	Age	Sex
	uus caballus	33				
I, ti	the horse (2) either (2) or	[was temp than 60 d [entered t where ho	porarily exported from ays ⁽²⁾ or 90 days ⁽²⁾ pric the country of dispatch arse entered country o	on (insert date) from	lispatch on	(insert date) less
	(2)either (2)or (2)or (2)or (2)or (2)or (2)or (2)or (2)or	[the Asiar [the Amer [the Endu [the Test [the Olym [the Paral [the World	n Games in	Inited Arab Emirates;] Games in World Championships inr in ea of Mexico City, Mexico;]	(insert place);](insert place);](insert place);](insert place);]	
	infectious or the condition health certific	contagious as for reside cate for the ation will be	diseases transmissib ence and pre-export country or part of the	of dispatch the horse has n le to equidae; isolation as applicable in acc territory of the country of dis ay that health and well-being	cordance with point I patch are fulfilled;	I.2. of the accompanying
Na	me and addres	s of the ow	ner ⁽²⁾ or representative	9 ⁽²⁾ :		••
Da	te:	((dd/mm/yyyy)			
				(Signature)		
(1)	document as system (such If a passport validated it.	defined in as ear tag, accompan	Article 2(b) of Comm tattoo, brand, transpo ies the animal, its nu	an individual identifier which ission Implementing Regula onder) and the anatomic plac umber should be stated and	tion (EU) 2018/659. e used on the anima	Specify the identification I.
	Age: Date of Sex ($M = mal$		m/yyyy). ile, C = castrated).			
(2)	Delete as app	ropriate.		erican Games, Central Amer	ican and Caribbean	Games.

Veterinary certificate to EU

▼<u>M1</u>

COUNTRY:

Chapter 2

Model health certificate and model declaration applicable to re-entry into the Union of registered horses for racing after temporary export for a period of not more 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)

	l.1.	Consignor Name		.2. Certificate reference No I.2.a.	
		Address	1.3	.3. Central competent authority	
ŧ		Tel.	1.4	.4. Local competent authority	
nsignme	1.5.	Consignee Name Address	1.6	.6.	
Part I : Details of dispatched consignment		Postcode Tel.			
Is of disp	1.7.	Country of origin ISO code I.8. Region of Code origin	1.9	.9. Country of ISO code I.10. Region of Codestination destination	ode
Detai	1.11.	. Place of origin	L.	.12. Place of destination	
Part I: I		Name Approval number Address		Name Address	
				Postcode	
	I.13.	. Place of loading	l.	.14. Date of departure	
	I.15.	. Means of transport	1.	.16. Entry BIP in EU	
		Aeroplane Ship Railway wagon	ı		
		Road vehicle Other Identification Documentary references	1.1	.17. No(s) of CITES	
	I.18.	. Description of animal		I.19. Commodity code (HS code) 01 01	
				I.20. Quantity	
	1.21.			I.22. Number of pac	kages
	1.23.	. Seal/Container No		I.24.	
	1.25.	. Animal certified for:		,	
		Registered horse			
	1.26.			1.27. For import or admission into EU □	
	1.28.	. Identification of the animal		'	
		Species Identification system Ide (Scientific name) Equus caballus	entifica	cation number Age Sex	

COUNTRY

Re-entry after temporary export of not more than 90 days Specific races — Registered horse

			II.a. Certificate reference number	II.b. Local reference number							
	II.	Attestation of anim	al health and welfare								
	I, the unders	signed official veterinar	ian, hereby certify, that the animal described in E	3ox I.28.:							
	_	 is a registered horse as defined in Article 2(c) of Commission Implementing Regulation (EU) 2018/659; 									
Part II: Certification	_	 was examined today (¹) and found free of clinical signs of diseases and of obvious signs of ectopara infestation; 									
Certif	_	 is not intended for slaughter under a national programme of infectious or contagious disease eradication; 									
Part II:	_	meets the requireme	nts attested in points II.1. to II.3. of this certificate	9;							
	_	is accompanied by th	e written declaration, signed by the owner of the	horse or the representative of the owner.							
	II.1.	Attestation on countr	y or part of the territory of the country and holdin	g of dispatch							
	II.1.1.	II.1.1. the animal is dispatched from									
	II.1.2.	(Trypanosoma equip	spatch the following diseases are compulsorily berdum), glanders (<i>Burkholderia mallei</i>), equine infectious anaemia, vo	e encephalomyelitis (of all types including							
	II.1.3.	the animal is dispatc	ned from a country or part of the territory of a cou	untry:							
		which the African h	considered free from African horse sickness in accordance with Directive 2009/156/EC and it re has been no clinical, serological (in unvaccinated equidae) or epidemiological evidence corse sickness during the period of 2 years prior to the date of dispatch and in which them in no vaccinations against the disease during the period of 12 months prior to the date of								
			Venezuelan equine encephalomyelitis has not confiding dispatch;	occurred during the period of 2 years prior to							
		c) in which	dourine has not occurred during the period of 6 r	months prior to the date of dispatch;							
		d) in which	glanders has not occurred during the period of 6	months prior to the date of dispatch;							
	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 holding to in points II.1.4.1. to II.1.4.7. and which last	ngs, which were subject to prohibition orders							
		(³) [II.1.4.1. in the ca	se of equidae suspected of having contracted do	urine,							
		(⁴) either	[6 months beginning on the date of the last suspected of having contracted dourine or inference of the contracted dourine or inference of the contracted dourine or inference of the contracted douring the co								
		(⁴) and/o	[in the case of a stallion, until the animal is cas	trated;]							
		(⁴) and/o	[30 days following the date of completion of the after all animals of susceptible species have be								

COUNTRY

Re-entry after temporary export of not more than 90 days Specific races — Registered horse

		II.a. Certificate reference number	II.b. Local reference number						
(³) [II.1.4.2. in the case of glanders,									
	(⁴) either	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;]							
	(⁴) and/or	[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 cas	e of equine encephalomyelitis of any type,							
	(⁴) either	[6 months beginning on the day on which the e slaughtered;]	quidae suffering from the disease have been						
	(⁴) and/or	[6 months beginning on the day on which the only Nile Fever, Eastern equine encephalomyelitis died, been removed from the holding or fully results.]	or Western equine encephalomyelitis have						
	(⁴) and/or	[30 days following the date of completion of th after all animals of susceptible species have be							
II.1.4.4.	slaughtere gel immu	n the case of equine infectious anaemia, until the date on which, the infected animals having bee laughtered, the remaining equine animals on the holding have shown a negative reaction in an age el immunodiffusion test (AGID or Coggins test) carried out on blood samples collected ow occasions 3 months apart;							
II.1.4.5.	in the cas	e of vesicular stomatitis,							
	(4) either	(4) either [6 months following the last case;]							
	(⁴) and/or	[30 days following the date of completion of th after all animals of susceptible species have be							
II.1.4.6.		e of rabies, 30 days following the last case and n of the premises;	the date of completion of the cleansing and						
II.1.4.7.		e of anthrax, 15 days following the last case and n of the premises;	d the date of completion of the cleansing and						
		wledge, during the period of 15 days prior to the infected or suspected of an infectious or contagi							
II.2. Attestation	n of residend	ce and pre-export isolation							
II.2.1. The anima (insert dat		ted into the country or part of the territory of the	country of dispatch on						
(⁴) either	[directly fr the partici	om the EU Member State pation in	(insert name of EU Member State) for						
	(⁴) either	[The Japan Cup;]							
	(4) or	[The Melbourne Cup;]							
	(4) or	[The Dubai Racing World-Cup;]							
	(4) or	[The Hong Kong International Races;]							

▼M1

COUNTRY

Re-entry after temporary export of not more than 90 days Specific races — Registered horse ther II.b. Local reference number

(4) or [from Australia (4), Canada (4), the United States of America (4), Hong Kong (4), Japan (4), Singapore (4), United Arab Emirates (4) or Qatar (4) for the participation in International Group/Grade 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 (4) or representative of the owner (4) 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 respectively of Annex I to Commission Implementing Regulation (EU) 2018/659.
- (3) Delete statement if the attestation in point II.1.3. applies to the entire country of dispatch.
- (4) Delete as appropriate.

COUNTRY

Re-entry after temporary export of not more than 90 days Specific races — Registered horse

		II.a. Certificate reference number	II.b. Local reference number							
This	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 registered horse will enter Union territory and undergo the veterinary border checks;										
(b)	be made out to a single consi	gnee;								
(c)	be signed and stamped in a co	plour different to the colour of the printing;								
(d)		paper or all sheets of paper required are par otal number of pages, and each page shall bear are stapled and stamped.								
Offici	al 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								
Ider	tification o	f the animal (1)							
Spe	ecies (Scie	ntific name)	Identification system	Identification number	Age	Sex			
	_								
	Equus ca	apailus							
I, the	e undersigr	ned owner (²) o	representative of the owner	(2) of the registered horse descri	bed above, hereby decl	are, that:			
_	the horse								
	(²) either		ily exported from the Union a ays prior to this declaration;]	to the country of dispatch on		(insert date)			
	(²) or		ountry of dispatch on where horse entered country	(insert date) from of dispatch);]		(insert name of			
_	the horse	has been temp	orarily exported from the Un	ion to take part in					
	(²) either	[The Japan Cu	ıp;]						
	(²) or	[The Melbourn	e Cup;]						
	(2) or	[The Dubai Ra	cing World-Cup;]						
	(2) or	[The Hong Ko	ng International Races;]						
	(²) or		Group/Grade meetings in Agapore (²), United Arab Emir	ustralia (2), Canada (2), the Unitates (2) or Qatar (2);]	ed States of America (²), Hong Kong (²),			
_			ays prior to the date of disparansmissible to equidae;	tch the horse has not been in cor	ntact with animals suffer	ing from infectious			
_				n as applicable in accordance wi country of dispatch are fulfilled;	ith point II.2. of the acc	companying health			
_	the transpof the jour		effected in such a way that	health and well-being of the horse	e can be protected effec	ctively at all stages			
Nan	ne and add	ress of the own	er (²) or representative (²): .						
Date	e:		(dd/mm/yyyy)						
(1)	Article 2(b transponde If a passpo Age: Date Sex (M = n	o) of Commission er) and the anator	n Implementing Regulation (E nic place used on the animal. ne animal, its number should be /yy).	dentifier which permits to link the anir U) 2018/659. Specify the identificat stated and the name of the competen	tion system (such as ear	tag, tattoo, brand,			
` ′									

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

COU	NTRY:	Veterinary certificate to EU
	I.1. Consignor Name	I.2. Certificate reference No I.2.a.
	Address	I.3. Central competent authority
	Tel.	I.4. Local competent authority
Part I : Details of dispatched consignment	I.5. Consignee Name Address Postcode Tel.	1.6.
tails of dispa	I.7. Country ISO code I.8. Region Code of origin	I.9. Country of ISO code I.10. Region of Code destination
Part I: Det	I.11. Place of origin Name Approval number Address	I.12. Place of destination Name Address Postcode
	I.13. Place of loading	I.14. Date of departure
	I.15. Means of transport Aeroplane	I.16. Entry BIP in EU
	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
	1.21.	I.22. Number of packages
	I.23. Seal/Container No	1.24.
	I.25. Animal certified for: Registered horse registered equine anim	nal breeding and production
	1.26.	I.27. For import or admission into EU
	I.28. Identification of the animal	
	Species (Scientific name) Identification sys	stem Identification number Age Sex

COUNTRY

Import - Registered horse, registered equine animal or equine animal for breeding and production

					- C** C X - C X				
				II.a.	Certificate reference number	II.b.	Local reference number		
	II.	Attestatio	n of anim	al health and	d welfare				
Part II: Certification	I, the undersigned official veterinarian, hereby certify, that the animal described in Box I.28.: - (1)either [is a registered equine animal, other than horse, as defined in Article 2(c) of Directive 2009/156/E (1)or [is a registered horse as defined in Article 2(c) of Commission Implementing Regulation (EU) 2018/(1)or [is an equine animal for breeding and production as defined in Article 2(e) of Directive 2009/156/E comes from a country or part of the territory of a country which is authorised for imports into the Union of category of equidae specified in the first indent above; - was examined today ⁽²⁾ and found free of clinical signs of diseases and of obvious signs of ectoparasite infests is not intended for slaughter under a national programme of infectious 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 II.1.1. The animal is dispatched from(insert name of country or part of the territory of a country a country or part of the territory of a country, which on the date of issuing this certificate has the Code: and is assigned to Sanitary Group ⁽⁹⁾ :								
	II.1.2.	(<i>Trypanos</i> Venezuela	oma equip an equine e	erdum), glan encephalomy	wing diseases are compulsorily ders (<i>Burkholderia mallei</i>), equi elitis), equine infectious anaemi	ne encephal a, vesicular :	omyelitis (of all types including		
	II.1.3. the animal is dispatched from a country or part of the territory of country a) which is considered free from African horse sickness in accordance with Directive 2009 and in which there has been no clinical, serological (in unvaccinated equidae) or epidem evidence of African horse sickness during the period of 2 years prior to the date of disp. in which there have been no vaccinations against the disease during the period of 12 prior to the date of dispatch:						ed equidae) or epidemiological rior to the date of dispatch and		
		b)		Venezuelan e te of dispatch	equine encephalomyelitis has no n;	t occurred d	uring the period of 2 years prior		
		c) d)	in which	glanders has	not occurred during the period on not occurred during the period	of 6 months	prior to the date of dispatch;		
	(*)either [e) in which vesicular stomatitis has not occurred during the period of 6 months prior to the date dispatch;]								
	(1)or [e) in which vesicular stomatitis has occurred during the period of 6 months prior to the date of dispatch, and a blood sample taken from the animal on(insert date), within a period of 21 days prior to the date of dispatch, was tested with negative result for antibody to the vesicular stomatitis virus								
			(1)either	[in an ELISA	eutralisation test at a serum dilu A in accordance with the relevan es for Terrestrial Animals of the	t Chapter of	an varia established transfer of the		

COUNTRY

Import - Registered horse, registered equine animal or equine animal for breeding and production

				II.a.	Certificate r	reference number	II.b.	Local reference number
					in contact w	ith animals from he	oldings, which	the time periods referred to in the were subject to prohibition or:
	(4)[II.1.4.1.	in the cas	se of ed	quidae	suspected of	having contracted	dourine,	
		(1)either						ssible contact with an animal Trypanosoma equiperdum;]
		(1)and/or	[in the	case	of a stallion,	until the animal is c	astrated;]	
		(1)and/or				late of completion s of susceptible spe		nsing and disinfection of the een slaughtered;]]
	(4)[II.1.4.2.	in the cas	se of gl	anders	S.			
(1)either				norths beginning on the day on which the equidae suffering from the disease or cted with positive results to a test for the detection of the causative pathogen holderia mallei or antibodies to that pathogen, were killed and destroyed;]				
		(1)and/or						nsing and disinfection of the een killed and destroyed;]]
	II.1.4.3.	in the cas	se of ed	quine e	encephalomy	elitis of any type,		
		⁽¹⁾ either			eginning on tl tered;]	he day on which the	e equidae su	ffering from the disease have
		(1)and/or	West	Nile	Fever, Ea	astern equine e	ncephalomy	nfected with the virus causing elitis or Western equine Iding or fully recovered;]
		(1)and/or				late of completion s of susceptible spe		nsing and disinfection of the een slaughtered;]
	II.1.4.4.	been slau in an aga	ughtere r gel im	d, the muno	remaining ec	uine animals on th	e holding ha	, the infected animals having ve shown a negative reaction ut on blood samples collected
	II.1.4.5.	in the cas	se of ve	esicula	r stomatitis,			
		(1)either	[6 mo	nths fo	llowing the la	st case;]		
		(1)and/or				late of completion s of susceptible spe		nsing and disinfection of the een slaughtered;]
	II.1.4.6.				30 days follow premises;	ving the last case a	nd the date o	of completion of the cleansing
	II.1.4.7.				15 days follow premises;	wing the last case a	and the date	of completion of the cleansing
II.1.5.						of 15 days prior to ted of an infectious		f dispatch the animal has not us disease.

COUNTRY

Import - Registered horse, registered equine animal or equine animal for breeding and production

			1	I.a.	Certificate reference num	nber	II.b.	Local reference number
II.2. Attestation of residence and pre-export isolation								
⁽¹⁾ either	[II.2.1.	During a period of at least the 90 days prior to the date of dispatch, or since birth if the animal is less than 90 days old, or since entry if the animal was imported directly from the Union during a period of 90 days prior to the date of dispatch, the animal has been resident on holdings under veterinary supervision situated in a country or part of the territory of a country which is:						
	(1)(5)either				Group A, and during the part from equidae not of eq			30 days prior to the date of atus;]]
}	⁽¹⁾⁽⁵⁾ or	date of dis	spatch,	it wa		on under	veterinar	of at least 30 days prior to the y supervision without coming
1	⁽¹⁾⁽⁵⁾ or	place of o	rigin in E	Box I	.11., protected from vector	insects		isolation centre described as
		(1)either	[during	the p	period of at least 40 days pr	rior to the	date of	dispatch;]]]
								of dispatch from a country of of African horse sickness and
			⁽¹⁾ either		e animal is a registered he plementing Regulation (EU			n Article 2(c) of Commission
			⁽¹⁾ or	sic				ountry in which African horse 2 years prior to the date of
(1)(5) or	[II.2.1.	assigned dispatch, veterinary dispatch, prior to the free of Afr	The animal is dispatched from a country of which at least a part of the territory of the country is assigned to Sanitary Group F, and during the period of at least 90 days prior to the date of dispatch, or since birth if the animal is less than 90 days old, it was resident on holdings under veterinary supervision and was kept during the period of at least 60 days prior to the date of dispatch, or since 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 horse sickness in accordance with the Union legislation and underwent the pre-export isolation [in the approved vector-protected quarantine station of					
	⁽¹⁾ either	quarantine from premises provided i combination						
	⁽¹⁾ or	(insert nai dispatch a	s required for temporary admission or imports into the Union.]] permanently confined in the approved vector-proof quarantine station of					
II.3.	Attestation	of vaccina	tion and	l hea	Ith tests			
(1)either	[II.3.1.				accinated against African I suggesting previous vaccir		ckness in	the country of dispatch and
⁽¹⁾ or	[II.3.1. (1)either (1)or	The anima [more than [more than	al was v n 12 mo n 60 da	accir onths ys ar	nated against African horse prior to the date of dispatch	sickness h;]] ior to the	date of	s vaccination was carried out: admission into the country or where it is dispatched;]]

COUNTRY

			II.a.	Certificate reference number	II.b.	Local reference number					
⁽¹⁾⁽⁵⁾ or	[II.3.1.	Sanitary Group not more than quarantine by	The animal is dispatched from a country or part of the territory of a country which is assigned a 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									
	(1)either			Il parts of the territory are free o		n equine encephalomyelitis fo					
	⁽¹⁾⁽⁵⁾ Or	Venezuelan ed	uine e in equ	of a country which is assigned ncephalomyelitis for a period of ine encephalomyelitis occurs in and	at least 2 ye	ars prior to the date of dispato					
		cour 60 d vecte and daily sam	se and ays ar or-prot during , rema e holdi od tes	ted against Venezuelan equine I revaccinated according to mar Ind no more than 12 months price ected quarantine for a period of that period remained clinically ined within the normal physiolong which showed a rise in body at for virus isolation for Venezue	ufacturer's r or to the date at least 21 da healthy, an gical range, temperature	ecommendations not less that e of dispatch, and was kept ays prior to the date of dispatch di its body temperature, take and any equine animal on the e, taken daily, was subjected					
		vectorema norm a rist for V disp; with of e	or-proteined of all physe in both enezuatched negation or a control of the contro	ccinated against Venezuelan e ected quarantine for a period of dinically healthy, and its body te siological range, and any equine dy temperature, taken daily, was elan equine encephalomyelitis was subjected to a diagnostic te we result conducted on a sample to the vector protected quarar il dispatch;]]	of at least 2 mperature, to animal on the subjected to with negative at for Venezue taken not least	1 days, and during that pericaken daily, remained within the same holding which showe oo a blood test for virus isolatice results, and the animal to buelan equine encephalomyelitess than 14 days after the data					
		ence take to th trans ence withi vect	phalor on to (insee date cription phalor on 48 hor attacoined	ected to a haemagglutination myelitis carried out by the same wo occasions with an interval of sert date), the second of which we dispatch, without an increase in-polymerase chain reaction) to myelitis virus genome, carried cours prior to dispatch, on	e laboratory of 21 days of vas taken du in antibody est for the de out with nega (insert date PCR samplinents and inserts	on the same day on sample on (insert date) and ouring the period of 10 days prictitre, and a RT-PCR (reverse etection of Venezuelan equinative result on a sample take), and has been protected frong until loading for dispatch, besecticides on the animal an					
((1)[II.3.3.	the animal is a	ı unca	strated male equine animal olde	er than 180 d	days, and					
	(1)either			a country in which equine viral been officially reported during							
	⁽¹⁾ or	[was tested on		d sample taken on(h, by virus neutralisation test for							

COUNTRY

				II.a.	(Certificate	referenc	e number	II.b.		Local reference number
(1	or	21 days p	orior to	the da	ate		h, by vir	us isolation			sert date), within a period of ase chain reaction (PCR) or
(1)or	and re-va	ccinate by the [befor	ed at re comp e 1 Oc	eg pet ctc	ular interva ent author ober 2018,	als accor ity, and t on the c	ding to the he initial va lay a blood	manufact accination sample v	turer's was was t	ficial veterinary supervision, s instructions, with a vaccine carried out aken that was subsequently esult at a serum dilution of 1
			in 4;]]]								
		⁽¹⁾ Or	officia was t	l veter ested	rina	ary supervi uring that	ision, cor isolation	nmencing	on the day a virus	y a blo	ot more than 15 days under good sample was taken which ralisation test for EVA with
		⁽¹⁾ Or	super EVA o same	vision, arried day b	d o	luring which	th the argative readorator	nimal was sult at a se	subjected rum diluti	to a ion of	ion under official veterinary virus neutralisation test for 1 in 4, or carried out on the titres on two blood samples
		⁽¹⁾ or	at a se	erum o	dilu ner	ution of 1 in	4, carrie	ed out on a	blood sar	mple	t for EVA with negative result taken not earlier than 7 days which lasted until 21 days
		⁽¹⁾ Or	test fo	r EVA	A c	arried out	with neg same la	ative result aboratory v	at a seru	ım dil	cted to a virus neutralisation lution of 1 in 4 or carried out declining titres on two blood
(1	or (EVA carr blood sar	ied out nple of te of di	with in that a spatch	ne ani h,	gative resumal taken was tested	ult on an	aliquot of (ins	its entire ert date),	seme withi	(PCR) or real-time PCR for en collected after the date a n a period of 6 months prior EVA with positive result at a
(1	or or	[has prev					r antibo	dies again	st the eq	quine	arteritis virus or has been
		, c F S	consecution to serolog	utive of and utical te	day uni	ys, to at le til at least s for EVA	ast two in 28 days with neg	mares which after test gative resu	th were k mating a lits at a s	ept ir ind w erum	h, was test mated, on two isolation during the 7 days thich were subjected to two idilution of 1 in 4 on blood 3 days after the test mating,
								tion test fo dispatch or			out on a blood sample taken ert date),
		(1	either)	[wit	th	positive res	sult at a	serum dilut	tion of at I	least	1 in 4;]]]
		(1	¹)or	[wit	th	negative re	esult at a	serum dilu	ition of 1	in 4;]]]]
(1)either	[II.3.4.	anaemia	, wher	e it w	vas	continuo	usly resi		birth and		free from equine infectious not come into contact with
⁽¹⁾ or	[11.3.4.	Coggins	test) o	r to ar	n E	ELISA for e	quine in	ectious an	aemia cai	rried	nunodiffusion test (AGID or out on a blood sample taken for to the date of dispatch;]

		II.a.	Certificate reference r	number	II.b.	Local reference number
⁽¹⁾ [II.3.5.	Sanitary Gro reported dur complement	up B, D ng a pe fixation t sample	or E, or from China or eriod of 3 years prior t est for glanders carried taken on	Thailand, to the da I out with	or from a co ate of dispate negative res	country which is assigned to buntry in which glanders was ch, and was subjected to a sult at a serum dilution of 1 in thin a period of 30 days prior
⁽¹⁾ [II.3.6.	from a count or F, or from of 2 years p dourine carri	y or par China or ior to the ed out wi (ir d for bre	t of the territory of a co Thailand, or from a cou e date of dispatch, and ith negative result at a s asert date), within a peri	untry who untry in wid was su serum dilition of 30	ich is assigne hich dourine bjected to a ution of 1 in 5 days prior to	er than 270 days dispatched ed to Sanitary Group B, D, E was reported during a period complement fixation test for 5 on a blood sample taken on the date of dispatch, and has brior to and after the date the
⁽¹⁾⁽⁵⁾ [II.3.7.	Sanitary Gro (1)either [We cou	up C or I stern an ntry or p	D, and d Eastern equine encep	halomye e country	litis have not	country which is assigned to been officially reported in the during a period of at least 2
	to n the	anufacti date of	urer's instructions withir dispatch with inactivat	n a period ed vacci	d of 6 months ne against V	e and revaccinated according and at least 30 days prior to Vestern and Eastern equine (insert date);]]
	vec inhi	or prote	cted quarantine, and casts for Western and Ea	during thi	is period sub	or to the date of dispatch in a ojected to haemagglutination alomyelitis carried out by the
	⁽¹⁾ eii		n a sample of blood tale days prior to the date			sert date), within a period of tive result;]]]
	⁽¹⁾ Or	da se	ys on (ins	sert date en within	and on a period of	with an interval of at least 21 (insert date), the 10 days prior to the date of
⁽¹⁾ [II.3.8.	Sanitary Grou	G, or f		Japanes		country which is assigned to s has been officially reported
	that	holding		no case o	f Japanese e	at least 30 km radius around encephalitis during a period of
	the	date of		that per	riod the bod	od of at least 21 days prior to ly temperature, taken daily, subjected
	⁽¹⁾ eji	en of wh	blood taken on two of the community of t	y the same ccasions and on a period r-fold ind	ne laboratory with an inte of 10 days p crease in ant	atralisation test for Japanese on the same day on samples erval of at least 14 days on (insert date), the second of orior to the date of dispatch, tibody titre between the two nsects until dispatch;]]]

			8	II.a. Certif	icate reference number	II.b.	Local reference number					
			⁽¹⁾ or	Japanese sample ta	e encephalitis virus with aken not earlier than 7 da (insert date), and rem	negative r	tection of antibodies agains esult, carried out on a blood date the isolation commence acted from vector insects unti-					
		⁽¹⁾ or	revacc	inated accord		commendat	complete primary course and ions during a period of not les date of dispatch;]]					
⁽¹⁾⁽⁵⁾ either	[11.3.9.	Sanitary describe on the s	y Group ed in An same day [on blo on	E, and wannex IV to Dir	s subjected to a serolo ective 2009/156/EC, which taken on two occasions w	gical test for the state of the	a country which is assigned to radical Arrican horse sickness a sed out by the same laborator and of between 21 and 30 days (insert date), the second of date of dispatch					
			(1)eithei	with neg	ative results in each case	e;]]]						
			(1)or	[with posi	tive result in the first sam	ple, and						
				⁽¹⁾ either		ntification te	equently tested with negativest as described in Annex IV to					
				⁽¹⁾ or	increase in antibody	titre in a of Chapte	without more than a two-fo virus neutralisation test a r 2.5.1. of the OIE Terrestri Vaccines;]]]]					
		⁽¹⁾ or	to the o	date of dispat		art of the ter	within a period of 21 days pri ritory of the country of dispate se sickness and					
			⁽¹⁾ eithei		nal is a registered horse nting Regulation (EU) 20		in Article 2(c) of Commission					
			⁽¹⁾ or		has occurred during th		country in which African hors 2 years prior to the date					
(1)(5) or	[II.3.9.		nal is dispatched from a country or part of the territory of a country which is assign Group F and [was subjected to a serological test for African horse sickness as described in A IV to Directive 2009/156/EC, which was carried out by the same laboratory on the day on blood samples taken on two occasions with an interval of between 21 a days, on									
			(1)eithei	with neg	ative results in each case	e;]]]						
			⁽¹⁾ or	[with posi	tive result in the first sam	ple, and						
				(1)either [the second sample was subsequently tested with negative in an agent identification test as described in Annex IV to Direction 1.5]								
					2009/156/EC;]]]]							

COUNTRY

				II.a.	Certificate refere	nce number	II.b.	Local reference number				
		⁽¹⁾ or	sickne result than 2	ess as in eac 28 day	described in Anne h case on a bloo	ex IV to Direction d sample takes of introduction	ve 2009/156/ n on into the vect	cation test for African horse EC, carried out with negative(insert date) not less tor-protected quarantine and				
		⁽¹⁾ or	in An	nex IV le take uction	to Directive 200 n on	9/156/EC, car (insert date	rried out with e) not less th	horse sickness as described n negative result on a blood nan 14 days after the date of t more than 72 hours before				
II.4.	Attestation	of the tran	sport o	conditio	ons							
⁽¹⁾ either [(II.4.1.	Sanitary (a market,	The animal is dispatched from a country or part of the territory of a country which is assigned to Sanitary Group A, B, C, D, E or G and is transported directly to the Union, without passing through a market, marshalling or assembly centre and without coming into contact with other equidae not complying with at least the same health requirements as described in this health certificate.]									
⁽¹⁾⁽⁵⁾ or [[II.4.1.	Sanitary of coming in	Group Into control con	F and intact with admission admissio	is transported directly of the strange of the strange of the transfer of the transfer of the strange of the str	ctly from the vot accompanient of accompanient	rector-protect ed by a healt tions and arra ed in advance	country which is assigned to led quarantine station without h certificate either for imports angements have been made e with a disinfectant officially gainst vector insects just prior				
		⁽¹⁾ or	condit sched or par in stal recogn	tions a luled d t of the lls whice	nd arrangements irectly to a port in territory of a cour ch were cleansed in the third country	have been methe Union with htry not approve and disinfected	nade to trans lout calling in led for the en led in advance	country under vector-protected sport it on a vessel which is to a port situated in a country itry into the Union of equidae, e with a disinfectant officially gainst vector insects just prior				
1	II.4.2.	complying	g with a	at least		equirements a	is described in	ntact with other equidae not n this health certificate during				
	II.4.3.	disinfecte	ed befor	re load	ling with a disinfe	ctant officially	recognised in	be loaded were cleaned and the third country of dispatch dder cannot escape during				
II.5.	Attestation	of animal	welfare	9								
j								transported on the intended ffectively at all stages of the				
Notes:												
Part I:												
Box I.8.:					or the part of the te egulation (EU) 20		ountry as app	pearing in column 3 of Annex				
Box I.15.:	Registrat	ion numbe on is to be	r (railw	ay wa	gons or container	and lorries), f		(aircraft) or name (ship) and or must inform the BIP of entry				

▼ M2

COUNTRY

Import - Registered horse, registered equine animal or equine animal for breeding and production

Local reference number

II.b.

Certificate reference number

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).

II.a.

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.

- 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.
- Delete statement if the attestation in point II.1.3. applies to the entire country of dispatch.
- 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:

- 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:
- 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

Name (in capital letters): Qualification and title:

Date: Signature:

Stamp

			ne owner or representative nto the Union of an equine		
Ide	ntification of the anima	al ⁽¹⁾			
Spe	ecies (Scientific me)	Identification system	Identification number	Age	Sex

I, th	the animal (2)either [has rer	mained in the country or	e owner ⁽²⁾ of the animal descr part of the territory of the co	ountry of dispatch during	a period of at least 90
			e territory of the country of d of dispatch from a Member :		red residence period of
-	during the period of infectious or contag	15 days prior to the date ious diseases transmissil	of dispatch the animal has ole to equidae:	not been in contact with	animals suffering from
-	the conditions for re	esidence and pre-export	isolation as applicable in activities territory of the country of dis		. of the accompanying
-	the conditions for th		in accordance with point II.4	() ■ () () () () () () () () () () () () ()	ealth certificate for the
-		vill be effected in such a	way that health and well-beir	ng of the animal can be	protected effectively at
Nai	me and address of the	owner ⁽²⁾ or representativ	e ⁽²⁾ :		
Dat	te:	(dd/mm/yyyy)			
			(Signature)		
(1)	quagga, Equus zebr Identification system document as defined system (such as ear	a, Equus grevyi, or indica : The animal must bear d in Article 2(b) of Comn tag, tattoo, brand, transp	Equus asinus, Equus africate any cross between those an individual identifier which nission Implementing Regulationder) and the anatomic plant	n permits to link the anination (EU) 2018/659. Space used on the animal.	nal to the identification pecify the identification
	validated it.	Å	umber should be stated an	d the name of the com	petent authority which
	Age: Date of birth (de				
(2)	- (이) [1] [1] [1] [1] [1] [1] [1] [1] [1] [1]	emale, C = castrated).			
(2)	Delete as appropriate	b.			

Section B

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

COL	INTRY:	Veterinary certificate to EU
	I.1. Consignor Name	I.2. Certificate reference No I.2.a.
	Address	I.3. Central competent authority
	Tel.	I.4. Local competent authority
Part I: Details of dispatched consignment	I.5. Consignee Name Address Postcode Tel.	1.6.
tails of disp	I.7. Country of ISO code I.8. Region Code origin of origin	I.9. Country of destination ISO code I.10. Region of destination Code
Part I: Det	I.11. Place of origin Name Approval number Address	I.12. Place of destination Name Address Postcode
	I.13. Place of loading	I.14. Date of departure
	I.15. Means of transport Aeroplane Ship Railway wagon	I.16. Entry BIP in EU
	Road vehicle Other Identification Documentary references	I.17. No(s) of CITES
	I.18. Description of animals	I.19. Commodity code (HS code) 01 01
		I.20. Quantity
	1.21.	I.22. Number of packages
	I.23. Seal/Container No	1.24.
	I.25. Animals certified for: Slaughter	
	1.26.	I.27. For import or admission into EU
	I.28. Identification of the animals	1
	Species (Scientific name) Identification system	Identification number Age Sex

COUNTRY

				9200	2000 1100	CONTRACTOR MARKET CO.		500000	N. CONSCIONATION STORY CONTROL				
				II.a.	Certifi	cate referen	ce number	II.b.	Local reference number				
	II.	Attestation	on of animal	health, an	imal we	Ifare and po	ublic health						
Part II: Certification	- are	e equidae for ere examina estation;	or slaughter a ed today ⁽¹⁾ ar	s defined in and found fr	n Article ee of cl	2(d) of Directinical signs		6/EC; and of ob	ovious signs of ectoparasite				
<u>:</u>			_					or contag	gious disease eradication;				
Part	- are	 meet the requirements attested in points II.1. to II.5. of this certificate; are accompanied by the written declaration, signed by the owner of the animals or the representative of the owner. 											
	II.1.	Attestation	n on third cou	ıntry or par	t of the t	territory of th	ird country a	nd holdin	g of dispatch				
	II.1.1.	country),		part of the t	erritory o	of a country,	which on the		or part of the territory of a suing this certificate has the				
	II.1.2.	dourine (Trypanosoma	equiperd	inders (Burk	kholderia ma	llei), equir	ole: African horse sickness ne encephalomyelitis (of al naemia, vesicular stomatitis					
	II.1.3.	the anima	als are dispate	ched from a	country	or part of th	ne territory of	country					
		a)	2009/156/E or epidemidate of disp	EC and in vological evidual exict of the contract of the contr	which the dence of n which t	ere has bee f African hor	n no clinical, se sickness c een no vacci	serologica during the	accordance with Directive al (in unvaccinated equidae period of 2 years prior to the gainst the disease during the				
		b)	in which Ve prior to the			encephalomy	elitis has not	occurred	during the period of 2 years				
		c)	in which do	ourine has r	not occu	rred during t	he period of	6 months	prior to the date of dispatch;				
	7000X 88405	d)							prior to the date of dispatch				
	(3)either	[e)	in which ve of dispatch		matitis h	as not occu	rred during th	e period o	of 6 months prior to the date				
	⁽³⁾ Or	[e)	dispatch, a	nd a blood riod of 21	sample days pri	taken from or to the da	each of the a	inimals on	6 months prior to the date on(insert date) ted with negative results fo				
			(3)either [in a virus n	eutralisa	ation test at a	a serum diluti	on of 1 in	32;]]				
							n the relevantial Animals of		of the Manual of Diagnostic				
	II.1.4.	in points I	II.1.4.1. to II.	1.4.7. have	not bee	n in contact	with animals	from hold	the time periods referred to dings, which were subject to which last for:				
	(⁴⁾ [II.1.4.1.	in the case	of equidae	suspec	ted of havin	g contracted	dourine,					
									ssible contact with an anima Trypanosoma equiperdum				
			(3)and/or [in the case	of a sta	llion, until th	e animal is ca	astrated;]					
									nsing and disinfection of the been slaughtered;]]				

COUNTRY

				II.a	Certific	ate reference number	II.b.	Local reference number					
		(4)[II.1.4.2.	in the case	of glande	rs,								
			(3)either [6	6 months l subjected	beginning with position	ve results to a test for the	ne detection	uffering from the disease or n of the causative pathogen e killed and destroyed;]					
				(3) and/or [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	encephal	omyelitis of any type,							
				6 months nave been			he equidae	suffering from the disease					
			V	Nest Nile	Fever,	Eastern equine end	cephalomye	ected with the virus causing slitis or Western equine olding or fully recovered;]					
						ne date of completion of mals of susceptible spe		sing and disinfection of the been slaughtered;]					
		II.1.4.4.	been slaug reaction in	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;									
		II.1.4.5.	in the case	of vesicul	ar stomati	tis,							
			(3)either [6	6 months	following t	he last case;]							
						ne date of completion of mals of susceptible spe		sing and disinfection of the been slaughtered;]					
		II.1.4.6.				s following the last car e premises;	se and the	date of completion of the					
		II.1.4.7.				ys following the last ca e premises;	se and the	date of completion of the					
	II.1.5.					eriod of 15 days prior to r suspected of an infect		dispatch the animals have tagious disease.					
	11.2.	Attestation	of residence	e and pre-	export isc	lation							
	II.2.1.	period of 9	0 days prior nder veterina	to the dat	e of dispa	tch, or since birth if the	animals ar	untry of dispatch during the e less than 90 days old, on y or part of the territory of a					
		⁽³⁾ either				and during the period rom equidae not of equ		30 days prior to the date of th status;]					
		⁽³⁾ or	date of disp	patch they	were ke		n under vet	at least 30 days prior to the erinary supervision without					
		⁽³⁾ or						prior to the date of dispatch I.11., protected from vector					
-													

COUNTRY

			1	I.a.	Certificate reference number	II.b.	Local reference number
II.3.	Attestation	n of vaccina	ation and	hea	lth tests		
(3)either	[11.3.1.				vaccinated against African hor ation suggesting previous vaccir		s in the country of dispatc
⁽³⁾ or	[II.3.1.				cinated against African horse sic ths prior to dispatch;]	kness, and	this vaccination was carrie
	II.3.2.		als were r or to dispa		vaccinated against Venezuelan e from	equine enc	ephalomyelitis during the 6
	(3)either				parts of the territory are free of 2 years prior to the date of disparent		an equine encephalomyelit
	⁽³⁾⁽⁵⁾ or	Venezue dispatch	lan equin and Ven	e e ezu	of a country which is assigned to ncephalomyelitis for a period o elan equine encephalomyelitis of dispatch, and	f at least	2 years prior to the date
		⁽³⁾ either	primary of less that were kep date of of tempera equine a daily, we	count of in disp ture anim as	nated against Venezuelan equi rse and revaccinated according to 0 days and not more than 12 mo o vector-protected quarantine for atch, and during that period rem to taken daily, remained within the lal on the same holding which she subjected to a blood test for a light provided in the same holding which she subjected to a blood test for a	o manuface on the prior a period o ained clini e normal powed a rise	turer's recommendations note the date of dispatch, are fat least 21 days prior to the cally healthy, and their boothysiological range, and are in body temperature, takes
		⁽³⁾ or	in vector dispatch tempera equine a daily. we encepha subjecte negative entry into	ture anim as a alom ed to e res	accinated against Venezuelan ecotected quarantine for a period and during that period remained taken daily. remained within the properties of the same holding which should be the properties with negative results, are a diagnostic test for Venezuelt conducted on a sample taken he vector-protected quarantine dispatch;]]	of at least ed clinical per normal properties of a rise principal to the animulation of less to the state of	21 days prior to the date by healthy, and their boo physiological range, and are ein body temperature, take tion for Venezuelan equirnals to be dispatched we line encephalomyelitis withan 14 days after the date
⁽³⁾⁽⁵⁾ eithei	r [II.3.3.	infectious	s anaemia	a, wi	atched from Iceland, which is onere they have been continuouslae which have entered Iceland in	y resident	since birth and did not com
⁽³⁾ or	[11.3.3.	the anim an ELISA	als were s A for equin taken on	subj	ected to an agar gel immunodiff fectious anaemia carried out wit(insert date), this being	usion test h negative	(AGID or Coggins test) or tresult in each case on block
C	³⁾ [II.3.4.	to Sanita period of test for g blood sa	ary Group 3 years p landers ca mples tak	B, orion arrie cen	ched from a country or part of the D or E, or from a country in what to the date of dispatch, and we do out with negative result in each on	nich glandere subject ch case at	ers was reported during the red to a complement fixation a serum dilution of 1 in 5 co

COUNTRY

			l.a. Certificate reference number II.b. Local reference numb
⁽³⁾ [II.3.5.	from a co E or from of dispat negative	ountry or a countr ch, and result in	castrated males or female equine animals older than 270 days dispatch part of the territory of a country which is assigned to Sanitary Group B, D in which dourine was reported during the period of 2 years prior to the devere subjected to a complement fixation test for dourine carried out we each case at a serum dilution of 1 in 5 on blood samples tak (insert date), this being within the period of 21 days prior to the date
⁽³⁾⁽⁵⁾ [II.3.6.		ry Group [Wester the cou	patched from a country or part of the territory of a country which is assign C or D, and and Eastern equine encephalomyelitis have not been officially reported try or part of the territory of the country of dispatch during the period of to the date of dispatch;]]
	⁽³⁾ or	according 30 days Eastern	nals were vaccinated with a complete primary course and revaccinat g to manufacturer's instructions within the period of 6 months and at leprior to the date of dispatch with inactivated vaccine against Western a equine encephalomyelitis, the last vaccination was applied (insert date);]]
	⁽³⁾ or	this per	nals were kept for at least 21 days protected from vector insects and duri od subjected to haemagglutination inhibition tests for Western and Easte encephalomyelitis on
		⁽³⁾ either	[a sample of blood taken from each of the animals in the consignmen
		⁽³⁾ Or	[samples of blood taken from each of the animals in the consignment two occasions with an interval of at least 21 days on
⁽³⁾ [II.3.7.	to Sanita	ry Group	patched from a country or part of the territory of a country which is assign G, or from a country in which Japanese encephalitis has been official during the past 2 years, and the animals
	⁽³⁾ either	those h	om holdings situated in the centre of an area of at least 30 km radius arou oldings where there has been no case of Japanese encephalitis during f at least 21 days prior to the date of dispatch;]]
	⁽³⁾ or	the date	pt in a vector–protected quarantine during a period of at least 21 days prior of dispatch, and during that period the body temperature of each of the anima ily, remained within the normal physiological range, and were subjected
		⁽³⁾ either	[to a haemagglutination inhibition or virus neutralisation test for Japane encephalitis carried out by the same laboratory on the same day samples of blood taken on two occasions with an interval of at least days on

							_				
				II.a. Certific	ate reference number	II.b. Local reference numbe	r				
			⁽³⁾ or	Japanese sample ta commence	encephalitis virus with n ken not earlier than 7	r the detection of antibodies again negative result, carried out on a blood days after the date the isolation, and remained protected from vectors.	od on				
		⁽³⁾ or	revacci	nated accordi	ing to manufacturer's rec	itis with a complete primary course ar commendations during a period of n ths prior to the date of dispatch;]]					
(3)(5)	[11.3.8.	Sanitary (Group E, IV to D	is are dispatched from a country or part of the territory of a country which is assigned to iroup E, and were subjected to a serological test for African horse sickness as described IV to Directive 2009/156/EC, which was carried out by the same laboratory on the							
		⁽³⁾ either	Deither [on blood samples taken from each of the animals in the consignment on occasions with an interval of between 21 and 30 days, on								
			(3)either	[with negat	tive result in each case;]]]]					
			(3) <i>or</i>	[with positi	ve results in the first sam	nple, and					
				⁽³⁾ either		rere subsequently tested with negativen agent identification test as described 2009/156/EC;]]]]					
				⁽³⁾ or	tested without more that in a virus neutralisation	ach animal of the consignment we an a two-fold increase in antibody tit on test as described in point 2.4 OIE Terrestrial Manual for Diagnost	re of				
		⁽³⁾ or	in the country to the dispatch not adjacent	consignment of date of dispatch is recognise acent to a cou	in (insert d tch, and the country or ed by the OIE as officially	sample taken from each of the anima late), within the period of 10 days pri- part of the territory of the country y free of African horse sickness and orse sickness has occurred during the]	or of is				
II.4.	Attestation	of the trai	sport co	onditions							
	[II.4.1.	slaughter	Arrangements were made and verified to ensure that the animals are transported directly to a slaughterhouse on the territory of the Union, without passing through a market, marshalling or assembly centre referred to in Article 7(1) of Directive 2009/156/EC, and without coming into								
⁽³⁾ or	[II.4.1.	slaughter marshalli the same	tact with other equidae not authorised for the entry into the Union.] Ingements were made and verified to ensure that before the animals are transported to a Ighterhouse on the territory of the Union they pass only through a single approved market, Ishalling or assembly centre referred to in Article 7(1) of Directive 2009/156/EC situated in Same Member State, from where they are transferred directly to the slaughterhouse lout coming into contact with other equidae not authorised for the entry into the Union.]								
	II.4.2.	complyin	g with a	t least the sai		any contact with other equidae n as described in this health certifica Union.					

Import - Equidae for slaughter

II.a.	Certificate reference number	II.b.	Local reference number
1			

II.4.3. The transport vehicles or containers in which the animals are 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 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 of public health

The animals described in Box I.28. have not received any stilbene or thyrostatic substances nor any oestrogenic, androgenic, gestagenic or beta-agonist substances for purposes other than therapeutic or zootechnical treatment as defined in Article 1(2)(b) and(c) of Directive 96/22/EC.

The guarantees covering live equidae provided by the residue plan submitted and approved in accordance with Article 29 of Directive 96/23/EC are fulfilled.

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.: Species: Select amongst: "Equus caballus", "Equus asinus" or "Equus caballus x Equus asinus".

 Identification system: Each of the animals must bear an individual identifier which permits to link the animal to the identification document. 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/yyyy).

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

Part II:

The certificate must be issued on the day of loading of the animals for dispatch to the Member State of destination in the Union.

The import of these equine animals for slaughter shall not be allowed when the animals were loaded either prior to the date of authorisation for imports of live equidae for slaughter into the Union from the respective country or part of the territory of a country mentioned under point II.1.1., or during a period where restrictive measures have been adopted by the Union against the entryof 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 respectively of Annex I to Commission Implementing Regulation (EU) 2018/659.
- (3) Delete as appropriate.
- (4) Delete statement if the attestation in point II.1.3. applies to the entire country of dispatch.
- (5) 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.

COUNTRY

		II.a.	Certificate reference number	II.b.	Local reference number
Thi (a)	the Member State of destination undergo the veterinary bord	ation and er check	understood by the certifying offic d of the Member State where the s;		
(c) (d)	be signed and stamped in a consist of a single sheet o indivisible by inserting page	colour of paper numbe	different to the colour of the printi or all sheets of paper required rs and total number of pages, a page and those pages are stapled	are part	age shall bear the certificate
Off	ficial veterinarian				
	Name (in capital letters):			Qua	alification and title:
	Date:			Sig	nature:
	Stamp:				
1					

			e owner or representative of consignments of live equals to the equals of the equals o		
Identification of	the anim	nals ⁽¹⁾			
Species (Scien	tific	Identification system	Identification number	Age	Sex
		***************************************		***************************************	***********
I, the undersign	ed owne	r ⁽²⁾ or representative of the	owner ⁽²⁾ of the animals descr	ibed above, hereby declar	re, that:
- the anima		emained in the country or p	eart of the territory of the coun	atry of dispatch for at least	90 days prior to the
		f 15 days prior to the date o	of dispatch the animals have rate to equidae;	not been in contact with an	imals suffering from
			solation as applicable in acc		the accompanying
- the condit	ions for t		in accordance with point II.4.		Ith certificate for the
	ortation	will be effected in such a w	ray that health and well-being	g of the animal can be pro	tected effectively a
the anima					
(2)either		y from the premises of dispequidae not of the same hea	patch to the slaughterhouse of alth status;]	of destination without comi	ng into contact with
⁽²⁾ or	marsha		he slaughterhouse of destinat eferred to in Article 7(1) of Di the same health status;]		
Name and add	ress of the	e owner ⁽²⁾ or representative	₅ (2):		
Date:		(dd/mm/yyyy)			
			(Signature)		
Identification document	on system as define	m: The animal must bear a ed in Article 2(b) of Comm	uus asinus, or indicate any cr in individual identifier which r ission Implementing Regulat inder) and the anatomic place	permits to link the animal ion (EU) 2018/659. Spec	
			er should be stated and the na		ority which validated
	of birth (c	dd/mm/yyyy).			
Sex (M = n	nale, F =	female, C = castrated).			
(2) Doloto oo	nnronria	to			

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 (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 watermark.
- 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:								Veterinary certific	ate to El		
	l.1.	Name				1.2.	Certificate reference	e No	I.2.a.			
		Address				1.3.	I.3. Central competent authority					
ent		Tel.				1.4.	Local competent au	ıthority				
Part I : Details of dispatched consignment	1.5.	Consignee Name Address				1.6.	Person responsible Name Address	for the load	l in EU			
atched o		Postal code Tel.					Postal code Tel.					
s of disp	1.7.	Country of origin	ISO code	I.8. Region of origin	f Code	1.9.	Country of destination	ISO code	I.10. Region of destination	Code		
: Detail	I.11. Place of origin Semen centre				1.12	Place of destination Semen centre ☐	1	Holding	1			
Part	Name Approval number Address Postal code I.13. Place of loading						Name Approval number Address					
							Postal code					
						1.14	I.14. Date of departure					
	I.15.	. Means of tra	ansport			I.16	I.16. Entry BIP in EU					
		Aeroplane D			y wagon 🗖							
		Road vehicl Identification Documentar		_		1.17	1.17.					
	I.18.	Description	of commodity	у			I.	19. Commo	odity code (HS code) 05 11 99 85			
									I.20. Quantity			
	I.21.								I.22. Number of pack	ages		
	1.23	Seal/Contai	ner No						1.24.			
	1.25	. Commoditie Artificial rep		r: □								
	I.26. For transit through EU to third country					I.27. For import or admission into EU						
		Third countr	ry I	SO code								
	1.28	Identification	n of the comr	nodities								
	s	Species (Scier	ntific name)	Don	or identity		Date of collec	ction	Quantity			
	1											

Equine semen – Section A

	II. Health information	1	II.a. Certificate reference No	II.b.						
	I, the undersigned, offic	cial veterinarian, of th	ne exporting country (2)	hereby						
			(name of exp	orting country)						
	certify that:									
E.	export to	the Union is approv	(3), in which the semen described above yed and supervised by the competent auth Annex D to Directive 92/65/EEC (4);	was collected, processed and stored for nority in accordance with the conditions of						
Part II: Certification	date the		ng 30 days prior to the date of first collection was dispatched or until the 30 days sto							
Part II:	II.2.1.	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:								
	 not considered to be infected with African horse sickness in accordance with Article 5(2)(a) and (b) of Directive 2009/156/EC, 									
	 free from Venezuelan equine encephalomyelitis for a period of at least 2 years, 									
	free from glanders and dourine for a period of at least 6 months;									
	II.2.2. fulfilled the conditions for a holding laid down in Article 4(5) of Directive 2009/156/EC and in particular									
	(¹) either		ng a case of a disease mentioned below necease located in the holding were slaughter							
		t	from any type of equine encephalomye beginning on the day on which the e slaughtered,							
		1	from equine infectious anaemia (EIA) for negative result in an agar gel immunodiffus on samples taken after the infected anin 3 months apart from each of the remaining	ion test (AGID or Coggins test) carried out nals were slaughtered on two occasions						
			from vesicular stomatitis (VS) for a period case,	of at least 6 months from the last recorded						
		— f	from rabies for a period of at least one mon	th from the last recorded case,						
		— f	from anthrax for a period of at least 15 days	s from the last recorded case,]						
	(¹) or	disease and th enceph the cas	ng a case of a disease mentioned below all e located in the holding have been slaughte e holding was free for a period of at nalomyelitis, equine infectious anaemia, ve se of anthrax, beginning on the day on whi infection of the premises was satisfactorily	ered or killed and the premises disinfected, least 30 days from any type of equine sicular stomatitis and rabies or 15 days in ch following the destruction of the animals						
	II.2.3.	contained only ed metritis,	quidae which were free of clinical signs of e	equine viral arteritis and contagious equine						
	II.3. Prior to e	ntering the semen co	ollection centre the donor stallions and any	other equidae located in the centre:						

COUNTRY Equine semen – Section A

II. Health	n information		II.a. Certificate reference No	II.b.
	II.3.1.	a Member State regionalisation in	y resident for a period of 3 months (or since e of the Union during the 3 months period) in t accordance with Article 13 of Directive 2009 ntry which was during that period:	he exporting country or, in the case o
			red to be infected with African horse sicknessirective 2009/156/EC,	ess in accordance with Article 5(2)(a
		— free from Ve	enezuelan equine encephalomyelitis for a perio	od of at least 2 years,
		— free from gla	anders and dourine for a period of at least 6 m	onths;
(¹) either	[II.3.2.		ne country of export which was on the day of is (VS) for a period of at least 6 months,]	of admission into the centre free fron
(¹) or	[II.3.2.	result at a serum with the relevant	o 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 d sample taken (⁶) within 14 days prior to ente	ut with a negative result in accordance and Vaccines for Terrestrial Animals o
	II.3.3.	originated from h point II.2.2;	oldings which on the day of admission onto	the centre fulfilled the requirements o
II.4.	The seme	n described above	was collected from donor stallions which:	
	II.4.1.		clinical sign of an infectious or contagious dis centre and on the day the semen was collecte	
	II.4.2.		eriod of at least 30 days prior to the date of s shown any clinical sign of equine viral arteri	
	II.4.3.	collection and be	r natural mating during a period of at least 3 ween the dates of the first sample referred to of the collection period;	
	II.4.4.	Manual of Diagno which is recognis	llowing tests, which meet at least the require stic Tests and Vaccines for Terrestrial Animal ed by the competent authority and has the testivalent to that provided for in Article 12 of	s of the OIE, carried out in a laborator its referred to hereinafter included in it
	(test) or	uine infectious anaemia (EIA), an agar-gel im an enzyme-linked immunosorbent assay (ELI tive result;]	
		II.4.4.2. for equ	ine viral arteritis (EVA),	
	((1) either [II.4.4.2	2.1. a serum neutralisation test with a negatin four;]	ative result at a serum dilution of on
	((1) and/or [II.4.4.2	a virus isolation test, polymerase chain r negative result on an aliquot of the entire	
		three s	ntagious equine metritis (CEM), an ager pecimens (swabs) taken from the donor stallic s than 7 days at least from the penile sheath	on on two occasions with an interval o

COUNTRY Equine semen – Section /

cour	NTRY				Equine semen – Section A
H.	Health information	ı	11	I.a. Certificate reference No	II.b.
			(local treatransport r	les were in no case taken earlier than 7 days (systettment) after antimicrobial treatment of the donor st medium with activated charcoal, such as Amies medi where they were subjected with a negative result to a	tallion and were placed in ium, before dispatch to the
		(¹) either	[II.4.4.3.1.	the isolation of <i>Taylorella equigenitalis</i> after cultival conditions for a period of at least 7 days, set up with specimens from the donor animal, or 48 hours who cool during transport;]	nin 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 specimen	
	II.4.5.	programm		the results specified in point II.4.4 in each case to respectively in points 1.6(a), (b) and (c) of Chapter s:	
		(⁹) [II.4.5.1.	at least 30 the semen	stallion was continuously resident on the semen colled days prior to the date of the first collection and during a described above, and no equidae on the semen colleto direct contact with equidae of lower health status the	g the period of collection of lection 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 fres ss than 14 days following the date of the commencem 30 days prior to the first semen collection.]	season or prior to the first sh, chilled or frozen semen
		(°) [II.4.5.2.	30 days pr semen des centre vete	r stallion was resident on the semen collection cent 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-	period of collection of the der the responsibility of the s, and/or other equidae on
			stallion at the first co semen and	described in point II.4.4 were carried out on sample least once a year at the beginning of the breeding se illection of semen intended for imports into the Unior d not less than 14 days following the date of the commit least 30 days prior to the first semen collection,	eason or prior to the date of n of fresh, chilled or frozen
		and	chilled or	period of collection of the semen intended for impo frozen semen the donor stallion was subjected , as follows:	
			(a)	for equine infectious anaemia, one of the tests des 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 sample 30 days prior to the date of the collection of the semi-	
			(¹) or	[in point II.4.4.2.2 was carried out on an aliquot of donor stallion taken (6) not more than 6 months collection of the semen described above and a bloodonor stallion during the 6 months period reacted serum neutralisation test for equine viral arteritis a than one in four;]	s prior to the date of the d sample taken (6) from the with a positive result in a

COUNTRY

Equine semen – Section A

II. Health	n informatior		п.	a. Certificate re	oforonco No			II h			
II. Healti	1 Information	1	111.4	a. Certificate re	elerence No			II.b.			
			` '	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 occasion	ons;]						
			(1) or	[on a single oc	casion and s	subjected to a	PCR or real-	time PCR.]]			
				onor stallion does not meet the conditions set out in points 1.6(a) and (b) of Chapter II ex D to Directive 92/65/EEC and the semen is collected for imports into the Union of semen.							
				described in po om the donor							
			the donor s from the da semen colle	escribed in point tallion during thate of the collection centre, f the semen des	ne storage po ction of the not less tha	eriod of the s semen and b an 14 days a	emen of a mefore the se	inimum perioo men is remov	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.]											
				[the non-shedd was confirmed negative resul stallion taken (stallion has re- four in a serum	by virus ison t on sample b) twice a yeacted with a	plation test, Po es of an alique ear at an inter positive resu	CR or real-timuot of the erwal of at leas It at a serum	ne PCR carrie ntire semen of t 4 months ar dilution of at	ed out with a of the donor nd the donor		
	II.4.6.	underwent dates:	the testing	provided for i	n points II.3	.2 (¹) and II.4	1.5 on sample	es taken on t	the following		
of		Start	t date (⁶)			Date of samplin	g for health tes	its (6)			
Identification of semen	Test	Donor	Semen	VS (¹)	EIA		A II. 1.2.		EM .4.3.		
Identii	proc	residence	collectio		II.4.4.1.	Blood sample	Semen sample	1. sample	2. sample		

COUNTRY Equine semen – Section A

II. Health	information	1	II.a. Certificate reference No	II.b.						
(1) either	[II.5.	No antibiotics were	added to the semen;]							
(¹) or	[11.5.	The following antib	iotic or combination of antibiotics was added to produce of less than (10):	a concentration in the final						
				;]						
II.6.	The seme	en described above w	as:							
	II.6.1.	 collected, processed, stored and transported under conditions which comply with the requirements of Chapters II(I)(1) and III(I) of Annex D to Directive 92/65/EEC; 								
	II.6.2.	sent to the place of loading in a sealed container in accordance with point 1.4 of Chapter III(I) Annex D to Directive 92/65/EEC and bearing the number indicated in Box I.23.								
Notes										
Part I:										
Box I.11.: The place of origin shall correspond to the semen collection centre of the semen origin.										
Box I.22.:	The number of packages shall correspond to the number of containers.									
Box I.23.:	1.23.: The identification of container and seal number shall be indicated.									
Box I.28.:	1.28.: 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:										
Guidance fo	r the compl	etion of the table in p	oint II.4.6.							
Abbreviation	ns:									
VS	Vesic	ular stomatitis (VS) te	esting if required in accordance with point II.3.2							
EIA-1	Equin	e infectious anaemia	(EIA) testing first occasion							
EIA-2	EIA te	esting second occasion	on							
EVA-B ²	1 Equin	e viral arteritis (EVA)	testing on blood sample first occasion							
EVA-B2	2 EVA t	esting on blood samp	ole second occasion							
EVA-S1	I EVA t	esting on semen sam	nple first occasion							
EVA-S2	2 EVA t	esting on semen sam	pple 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	1 CEM	testing second occas	ion first sample							
CEM-2	2 CEM	testing second occas	ion second sample taken 7 days after CEM-21							
Instructions	:									
For eac	ch semen i	dentified in column	A in correspondence with Box I.28, the test program	me (points II.4.5.1, II.4.5.2						

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.

COUNTRY

Equine semen - Section A

Qualification and title:

Signature:

	• • • • • • • • • • • • • • • • • • • •							_	quine semei	i – occion
II.	Health in	formation	ı	II.a. (II.a. Certificate reference No II.b.					
	required in the boxes. The dates shall be expected to the content of the content	in points I s marked v s when sa entered in	amples were to amples were to the lower line example belo	2 and II.4.5.3, 'A-B1 or EVA- aken for repea of columns 5	shall be en S1 and CEM at laboratory	tered in the /I-11 and CE / testing as r	upper line of M-12 in the erequired in ac	columns 5 to example below ccordance with	9 of the tabl /. n point II.4.5.2	e, this being
	Start date Date of sampling for						ing for health te	ests		
	Identification of semen	Test programme	Donor	Semen	VS II.3.2.	EIA		VA .4.2.	CEM II.4.4.3.	
		prog	residence	collection		11.4.4.1.	Blood sample	Semen sample	1. sample	2. sample
	А	T_	0	Б	Ve	EIA-1	EVA-B1	EVA-S1	CEM-11	CEM-12
	A	В	С	D	VS	EIA-2	EVA-B2	EVA-S2	CEM-21	CEM-22
(²) (³) (⁴)	Regulat column Only ap Commis Council the Cor Commu	ion (EU) 2 4 of that A oproved s ssion web Directive nmunity c nity rules	e semen are a 2018/657 prov Annex from a c semen collecti site: http://ec.e 92/65/EEC of of animals, se referred to in a	ided that the solution centres libertopa.eu/food 13 July 1992 men, ova and Annex A(I) to I	semen was of the categoral sted in accord/animal/ser laying down a month of the categoral sembryos robrective 90.	collected in topy of equida cordance with the men_ova/equidation animal head not subject to 1/425/EEC (C	the part of the ae indicated in the Article 17 uine/index_er alth requirement of animal head of L 268, 14.9	e territory of the columns 11, (3)(b) of Direct. ents governing alth requirements, 1992, p. 54).	e third countries the third countries of the ective 92/65/light trade in and ents laid down	y detailed in nat Annex. EEC on the imports into n in specific
(⁵)			e 2009/156/E0 third countries					ditions gover	ning the mo	vement and
(⁶)	Insert d	ate in tabl	e in point II.4.6	6 (follow Guida	ance in Part	II of the Not	es).			
(⁷)	perform	ed to ens	No 882/2004 sure the verifi 004, p. 1).							
(8)	donor e equine	quidae wh infectious	nunodiffusion to nich have cont anaemia and nd during the p	inuously resid no equidae a	ed in Icelan and their se	d since birth men, ova ar	, provided that	at Iceland has	remained off	icially free o

Cross out the programmes that do not apply to the consignment.

The signature and the stamp must be in a different colour to that of the printing.

(10) Insert names and concentrations.

Name (in capital letters):

Official veterinarian

Date:

Stamp:

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	1.2.	Certificate reference No	I.2.a.		
		Address	1.3.	Central competent authority			
		Tel.	I.4. Local competent authority				
signment	1.5.	Consignee Name Address	I.6.	Person responsible for the load i Name Address	n EU		
ched con		Postal code Tel.		Postal code Tel.			
Part I : Details of dispatched consignment	1.7.	Country of ISO code I.8. Region of origin Code	I.9.	Country of ISO code destination	.10. Region of Code destination		
t I : Details	l.11.	Place of origin Semen centre □	I.12.	Place of destination Semen centre □	Holding		
Parl		Name Approval number Address		Name Approva Address	ıl number		
		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 ☐ Railway wagon ☐ Road vehicle ☐ Other ☐	I.17.				
		Identification Documentary references	1.17.				
	I.18.	Description of commodity	I.19. Commodity code (HS code) 05 11 99 85				
					I.20. Quantity		
	I.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					
	s	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 unders	signed, offici	ial veterinar	rian, of the	e exporting country (²)(name of exporting co					
					(name of exponding co	ountry)				
	certify that :									
uo	II.1.	export to	the Europe	ean Union	(3), in which the semen described above was collecte is approved and supervised by the competent author d Chapter I(II)(1) of Annex D to Directive 92/65/EEC,					
Part II: Certification	II.2.				30 days prior to the date of first collection of the semen semen elapsed, the semen collection centre:	en described above until the				
Part II: O		II.2.1.	was situated in the exporting country or, in the case of regionalisation according to Article 13 Directive 2009/156/EC (8), in that part of the territory of the exporting country which was:							
			 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 Venezuelan equine encephalomyelitis for 2 years, 									
	— free from glanders 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	her [II.2.2.1. following a case of a disease mentioned below not all the animals of species susceptible the disease located on the holding were slaughtered or killed and the holding has befree:							
					om any type of equine encephalomyelitis for at least ay on which the equidae suffering from the disease are					
				re ta	om equine infectious anaemia for at least the period result in an agar gel immunodiffusion test (Coggins to ken after the infected animals were slaughtered on two meach of the remaining animals,	est) carried out on samples				
				— fro	om vesicular stomatitis for at least 6 months from the la	st recorded case,				
				— fro	om rabies for at least one month from the last recorded	case,				
				— fro	om anthrax for at least 15 days from the last recorded c	ase,]				
		(1) or [II.2.2.1. following a case of a disease mentioned below all the animals of species susceptible to the disease located on the holding have been slaughtered or killed and the premise disinfected, the holding has been free for at least 30 days from any type of equivalence encephalomyelitis, equine infectious anaemia, vesicular stomatitis and rabies or 15 days the case of anthrax, beginning on the day on which following the destruction of the animal the disinfection of the premises was satisfactorily completed;]								
		II.2.3.	contained metritis,	d only equ	ildae which were free of clinical signs of equine viral an	eritis and contagious equine				
	II.3.	Prior to en	ntering the s	semen col	llection centre the donor stallions and any other equidae	e located in the centre:				

COUNTRY

Equine semen – Section B

II. H	ealth informatior	1		II.a. Certificate reference No	II.b.
	II.3.1.	State of the regionalis	he Europe ation acco	esident for 3 months (or since entry if they were dir an Union during the 3 months period) in the expor ording to Article 13 of Directive 2009/156/EC (*), in hich was during that period	rting country or, in the case o
				d to be infected with African horse sickness in active 2009/156/EC (8),	ccordance with Article 5(2)(a
		— free	from Vene	zuelan equine encephalomyelitis for at least 2 years	3,
		— free	from gland	ders and dourine for at least 6 months;	
(¹) eithe	er [II.3.2.			country of export which was on the day of adm (VS) for at least 6 months,]	nission into the centre free o
(¹) or	[II.3.2.			a virus neutralisation test for vesicular stomatitis (llution of 1 in 12 on a blood sample taken (⁴) within	
	II.3.3.	originated point II.2.2		dings which on the day of admission onto the cent	re fulfilled the requirements o
II.4.	The seme	en described	above wa	is collected from donor stallions, which:	
	II.4.1.			r clinical sign of an infectious or contagious disease le day the semen was collected;	at the time of admission onto
	II.4.2.			30 days prior to the date of semen collection on hol cal sign of equine viral arteritis or contagious equine	
	II.4.3.	and betwe	een the da	for natural mating during at least 30 days prior to the tes of the first sample referred to in points II.4.5.1, tition period;	
	II.4.4.	the Manu samples	ıal of Diag taken in a	e following tests, which meet at least the requirement gnostic Tests and Vaccines for Terrestrial Animal accordance with one of the programmes specified competent authority:	ls of the OIE, carried out or
	(¹) (⁵) either	[II.4.4.1.	an agar-o	gel immuno-diffusion test (Coggins test) for equine result;]	infectious anaemia (EIA) with
	(¹) (⁵) or	[11.4.4.1.	an ELISA	for equine infectious anaemia (EIA) with negative re	esult;]
and	(¹) either	[II.4.4.2.		neutralisation test for equine viral arteritis (EVA) wf one in four;]	<i>i</i> ith negative result at a serum
	(¹) or	[11.4.4.2.		colation test for equine viral arteritis (EVA) carried the entire semen of the donor stallion;]	out with negative result on ar

COUNTRY Equine semen – Section B

II. Health information			II.a. Certificate reference No	II.b.
and	6 8	occasion equigenit and from	t identification test for contagious equine metritis (0 s on samples collected with an interval of 7 days alis after a cultivation of 7 to 14 days from pre-ejaculato genital swabs taken at least from the penile sheath, tive result in each case;	by isolation of <i>Taylorella</i> ry fluid or a semen sample
			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
	3	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 collection direct contact with equidae of lower health status than the	period of collection of the on centre came during that
	f	irst seme	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.	
	ti a	he date above, b continuou	or stallion was resident on the semen collection centre for the first collection and during the period of collectic ut has left the centre under the responsibility of the us period of less than 14 days, or other equidae on the stact with equidae of lower health status.	on of the semen described centre veterinarian for a
	C S	date of the semen d	described in point II.4.4 have been carried out on sar the first semen collection of the breeding season or colle escribed above was collected and at least 14 days between the of the residence period of at least 30 days,	ction period in the year the
and	S		described in point II.4.4.1 for equine infectious anaemia f blood taken (4) not more than 90 days before the se	
and			he tests described in point II.4.4.2 for equine viral arteritiaken $(^4)$ not more than 30 days before the semen describ	
	s s	aliquot of semen d eacted p	solation test for equine viral arteritis was carried out the entire semen of the donor stallion taken (4) not more escribed above was collected and a blood sample tate in a serum neutralisation test for equine viral art none 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	
	C	date of th	described in point II.4.4 have been carried out on sar the first semen collection of the breeding season or collected, escribed above was collected,	
and			described in point II.4.4 have been carried out on s 0 days after the collection of the semen described above	

COUNTRY

Equine semen – Section B

II. Health	n information		II.a. C	Certificate r	eference No			II.b.	
	II.4.6.	have under following da	gone the testi tes:	ng provide	ed for in poi	ints II.3.2 (1)	and II.4.5 (on samples to	aken on the
of		Start o	date (4)		[Date of samplin	g for health te	sts (4)	
Identification of semen	Test	Donor	Semen	VS (¹)	EIA	EVA II. 4.4.2.		CEM II.4.4.3.	
Identii	broç	residence	collection	II.3.Ž	II.4.4.1.	Blood sample	Semen sample	1. sample	2. sample
(¹) either	[II.5.	No antibiotio	s were added	to the sem	en;]				
(¹) or	[II.5.		g antibiotic or e en of not less tl		n of antibiotion	cs was added	I to produce	a concentratio	n in the fina
									::
II.6.	The seme	n described a	bove was:						
	II.6.1.		rocessed, store				which compl	y with the req	uirements o
	II.6.2.		place of loadii Directive 92/65						apter III(I) o
Notes									
Part I:									
Box I.11.:	The place	of origin shall	correspond to	the semen	collection ce	entre of the se	emen origin.		
Box I.22.:	The numb	er of package	s shall corresp	ond to the	number of co	ontainers.			
Box I.23.:	The identit	fication of con	tainer and seal	l number sl	nall be indica	ted.			
Box I.28.:	The donor	identity shall	correspond to	the official	identification	of the anima	l.		
	The date of	of collection sh	nall be indicate	d in the foll	owing format	t: dd/mm/yyyy	<i>1</i> .		

▼C2

COUNTRY Equine semen – Section B

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

Part II:

Guidance for the completion of the table in point II.4.6.

Abbreviations:

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 testing second occasion second sample taken 7 days after CEM-21

Instructions:

CEM-22

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	n)	Start	date		Date of sampling for health tests						
tification	Semmen Donor Sem-residence collection		Semen	VS	EIA	EVA II.4.4.2.		CEM II.4.4.3.			
Identi	pro	residence	collection	II.3.2.	II.4.4.1.	Blood sample	Semen sample	1. sample	2. sample		
Α	В	С	D	VS	EIA-1	EVA-B1	EVA-S1	CEM-11	CEM-12		
A	ь			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 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.

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.			
Offic	cial veterinarian				
	Name (in capital letters):		Qualification 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

COUN	TRY:									Vet	erinary certifica	te to EU
	l.1.	Consignor Name					1.2.	Certificate reference	ce No	1.2.8	a	
		Address			1.3.	Central competent	authority					
		Tel.					1.4.	Local competent a	uthority			
Part I : Details of dispatched consignment	1.5.	Consignee Name Address					I.6.	Person responsible Name Address	e for the load	in EU		
tched co		Postal code Tel.						Postal code Tel.				
of dispat	1.7.	Country of origin	ISO code		Region of origin	Code	1.9.	Country of destination	ISO code		egion of estination	Code
l : Details	l.11.	Place of orig					I.12.	Place of destination		ding 🗖		
Part	Name Approval number Address				Name Address	Approval	numbe	r				
		Postal code						Postal code				
	I.13.	Place of load	ding				I.14.	Date of departure				
	I.15.	Means of tra	ansport				I.16.	Entry BIP in EU				
		Aeroplane C Road vehicle Identification	e 🗆		Railway wago er □	n 🗆	I.17.					
			y references									
	I.18.	Description	of commodity	1					I.19. Commo		de (HS code) I 1 99 85	
										1.20. (Quantity	
	I.21.									1.22. 1	Number of packa	ages
	1.23.	Seal/Contain	ner No							1.24.		
	1.25.	Commoditie Artificial rep										
	1.26.	For transit the	_	third c	-			I.27. For import or	admission into	EU		
		Identification		nodities	B Donor ide	entity		Date of colle	ection		Quantity	

COUNTRY Equine semen – Section C

	II. Health information		II.a. Certificate reference No	II.b.			
	I, the unders	signed, official veterinarian, of the ex	xporting country (²)(name of exporting co	-			
	certify that:						
uo	II.1.	The semen collection centre in who to the European Union:	nich the semen described above was collected, pr	ocessed and stored for export			
ertificati	II.1.1.	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.		case of regionalisation according to Article 13 of y of export which was on the day the semen w				
		 African horse sickness, in ac 	ccordance with EU legislation,				
		 Venezuelan equine encepha 	alomyelitis for 2 years,				
		 glanders and dourine for 6 n 	nonths;				
	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 s prohibition lasted for:	usceptible to the disease located in the holding v	vere slaughtered or killed, the			
		 6 months, beginning on the day on which the equidae suffering from the disease are slaughtered, in the case of equine encephalomyelitis, 					
		 a period required to carry out with negative result two Coggins tests 3 months apart in the animals remaining after the infected animals have been slaughtered, in the case of infectious equine anaemia, 					
		— 6 months, in the case of ves	icular stomatitis,				
		 one month from the last reco 	orded case, in the case of rabies,				
		 15 days from the last record 	ed case, in the case of anthrax.				
	II.1.3.2.	the premises disinfected, the prol	ptible to the disease located in the holding have be hibition lasted for 30 days, or 15 days in the case stion of the animals the disinfection of the premises	of anthrax, beginning on the			
	II.1.4.		nmencing 30 days prior to semen collection and free of clinical signs of equine viral arteritis and co				
	II.2.	Prior to entering the semen collect	tion 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 g that period free of:				
		 African horse sickness, in ac 	ccordance with EU legislation,				
		Venezuelan equine encepha	alomyelitis 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	[11.2.2.	originated from the ten	ritory of the country of export which was on the datitis for 6 months,]	ly of admission into the centre
(¹) or	[11.2.2.		irus neutralisation test for vesicular stomatitis (4), this being within 14 days prior to entering the 12;]	
II.2.3.	originated	from holdings which on	the day of admission onto the centre fulfilled the re	equirements of point II.1.3;
II.3.	The seme	n described above was o	collected from donor stallions, which:	
II.3.1.	on the day	y the semen was collecte	d have not shown clinical signs of an infectious or	contagious disease,
II.3.2.	during at I	east 30 days prior to coll	ection of the semen have not been used for natura	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:	ecognised by the competent
II.3.6.1.	an agar-ge	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 at	: 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 gen n the urethral fossa with negative result in each ca	ital swabs taken at least from
II.3.7.	have beer	n subjected to one of the	following test programmes (5):	
II.3.7.1.	collection,	and during the collection	asly resident on the collection centre for at least on period, and no equidae on the collection cent r 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 educe of lower health status than the donor stallions.	luidae on the collection centre
			ave been carried out on samples taken one first semen collection and at least at the beginni	
		equired in point II.3.6.1 vn 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(4);]	was last carried out not more than 30 days before	e the semen was collected on

Equine semen - Section C

II.	Health	information	II.a. Certificate reference No	II.b.
(¹) 01	r		opositive stallion for equine viral arteritis was co	
II.3.7	7.3.		nave been carried out during the 30 days man- after the collection of the semen on samples take	
II.4.			collected, processed, stored and transported and III of Annex D to Directive 92/65/EEC.	under conditions which comply
Note	es			
Part	l:			
Box	l.11.:	The place of origin shall correspon	d to the semen collection centre of the semen or	igin.
Box	1.22.:	The number of packages shall con	respond to the number of containers.	
Box	1.23.:	The identification of container and	seal number shall be indicated.	
Box	1.28.:	The donor identity shall correspond	d to the official identification of the animal.	
		The date of collection shall be indi-	cate in the following format: dd/mm/yyyy.	
Part	II:			
(¹)	Delete	as necessary.		
(2)	Regula	ation (EU) 2018/659 provided the s	from a third country listed in column 2 of Annex semen was collected in the part of the territory on of the category of equidae indicated in column	of the third country detailed in
(3)	equida equine	e which have continuously resided	ns test) or the ELISA for equine infectious anald in Iceland since birth, provided that Iceland are and their semen, ova and embryos have becomen was collected.	has remained officially free of
(4)	Insert	date.		
(5)	Cross	out the programmes that do not app	bly to the consignment.	
(6)	OJ L 1	92, 23.7.2010, p. 1.		
_	The sig	gnature and the stamp must be in a	different colour to that of the printing.	
Offic	ial veter	inarian		
	Name	(in capital letters):		Qualification and title:
	Date:			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

COUN	TRY:		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			
Part I: Details of dispatched consignment	1.5.	Consignee Name Address	I.6. Person responsible for the load in EU Name Address			
tched co		Postal code Tel.	Postal code Tel.			
s of dispa	1.7.	Country of ISO code I.8. Region of origin Code	I.9. Country of destination ISO code I.10. Region of destination Code			
: I : Detail	l.11.	Place of origin Semen centre □	I.12. Place of destination Semen centre ☐ Holding ☐			
Part		Name Approval number Address	Name Approval number Address			
		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 Railway wagon Road vehicle Other Identification	I.17. No(s) of CITES			
		Documentary references				
	I.18.	Description of commodity	I.19. Commodity code (HS code) 05 11 99 85			
			I.20. Quantity			
	I.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 Third country ISO code	I.27. For import or admission into EU			
	1.28.	Identification of the commodities				
	s	pecies (Scientific name) Donor identity	Date of collection Quantity			

Equine semen - Section D

	II. Health	information	II.a. Certificate reference No	II.b.					
	I, the unders	signed official veterinarian of the	exporting country (²)						
(name of exporting country)									
	certify that:								
Ē	II.1.	The centre (3) described in Box I.11 at which the semen to be exported to the Union was stored:							
tificatio	(1) either	[II.1.1. meets the conditions laid down in Chapter I(I)(1) and is operated and supervised in accordance with the conditions laid down in Chapter I(II)(1) of Annex D to Directive 92/65/EEC (4);]							
Part II: Certification	(¹) or	[II.1.1. meets the conditions laid down in Chapter I(I)(2) and is operated and supervised in accordance with the conditions laid down in Chapter I(II)(2) of Annex D to Directive 92/65/EEC;]							
- Pa	II.2.	The semen to be exported to the	ne Union:						
	II.2.1.		and stored for a minimum period of 30 days immeditre (5) operated and supervised in accordance with C, which is						
	(¹) either	[located in the exporting countr	y;]						
	(¹) or								
	II.2.2.	was moved to the centre descr	ibed 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.	was stored under conditions wh	nich satisfy the terms of Annex D to Directive 92/65/6	EEC;					
	II.2.4.		n a sealed container in accordance with point 1.4 ng the number indicated in Box I.23.	of Chapter III(I) of Annex D to					
	Notes								
	Part I:								
	Box I.11.:	The place of origin shall corres	pond to the semen storage centre of semen dispatch	n.					
	Box I.17.:	Box I.17.: The serial number of the individual official document(s) or health certificate(s) that accompanied the semi described above from the approved semen collection centre of its origin to the described above semen storage centre shall be indicated. The original(s) of this/these document(s) or certificate(s) or the officially endorse copy/copies of thereof must be attached to this certificate.							

▼<u>C2</u>

COUNTRY Equine semen – Section D

II.	Health	ealth information II.a. Certificate reference No II.b.						
Вох	1.22.: The number of packages shall correspond to the number of containers.							
Вох	1.23.:	.23.: The identification of container and seal number shall be indicated.						
Box	1.28.:	28.: The donor identity shall correspond to the official identification of the animal.						
		The date of collection shall be indic	cated in the following format: dd/mm/yyyy.					
Part	II:							
(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 column 11, 12 or 13 of that Annex.							
(3)		pproved semen collection or storage mmission website:	ge centres listed in accordance with Article 1	7(3)(b) of Directive 92/65/EEC on				
	http://e	ec.europa.eu/food/animal/semen_ov	a/equine/index_en.htm					
(4)	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).							
(5)		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							
(⁶)	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.							
Offic	ial veter	inarian						
	Name (in capital letters): Qualification and title:							
	Date: Signature:							
	Stamp	:						

PART 2

Model health certificate for imports of ova and embryos

Section A

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

COUN	TRY:					Veterinary certifica	te to EU	
	l.1.	Name		Certificate referen		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	1.6.	Person responsibl Name Address	e for the load	in EU		
atched co		Postal code Tel.		Postal code Tel.				
s of dispa	1.7.	Country of ISO code I.8. Region of Code origin congin	1.9.	Country of destination	ISO code	I.10. Region of destination	Code	
l : Detail	l.11.	Place of origin Embryo team □	I.12.	Place of destination	on Embryo team			
Part		Name Approval number Address		Name Address	Approval num	nber		
		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 ☐ Railway wagon ☐						
		Road vehicle Other I Identification Documentary references	l.17.					
	I.18.	Description of commodity	I.19. Commodity code (HS code) 05 11 99 85					
						I.20. Quantity		
	l.21.					I.22. Number of packa	iges	
	1.23.	Seal/Container No	1.24.					
	I.25. Commodities certified for: Artificial reproduction I.26. For transit through EU to third country Third country ISO code							
				I.27. For import or admission into EU				
	1.28.	Identification of the commodities						
	s	Species (Scientific Category D name)	onor i	dentity D	ate of collectio	on Quantity		

▶" Equine ova/embryos - Section A ◄

	II. Health	information		II.a. Certificate reference No	II.b.					
	I, the under	signed, offic	ial veterinarian, of the e	exporting country (²)(nai	ne of exporting country)	ру				
	certify that:			,						
ion	II.1. The ova (¹)/embryos (¹) described above:									
Part II: Certification	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.	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.	.4. were collected at a place separated from other parts of the premises or holding which is in good repair and was cleaned and disinfected prior to the collection;								
	II.1.5.	prohibition	n or quarantine measur quipment and materials	d packed in laboratory facilities which res as set out in Box II.1.6, in a section used in contact with donor animals and	which is separated from the secti	on for				
	II.1.6.	come from	n donor mares which:							
		II.1.6.1.	a Member State of the regionalisation in according to the control of the control	sident for a period of 3 months (or since the Union during the 3 months period) in cordance with Article 13 of Directive 2009 which was during that period	the exporting country or, in the ca	ase of				
				to be infected with African horse sick tive 2009/156/EC,	ness in accordance with Article 5	5(2)(a)				
			 free from Venez 	zuelan equine encephalomyelitis for a per	od of at least 2 years,					
			 free from glande 	ers and dourine for a period of at least 6 r	nonths;					
	(1) either	[II.1.6.2.	collection free from vesicular stor	matitis						
	result at a service with the relevant			virus neutralisation test for vesicular sto ution of 1 in 32 or a VS ELISA carried of apter of the Manual of Diagnostic Tests ample taken on	ut with a negative result in accordand Vaccines for Terrestrial Anim	dance				
	(1) either	[II.1.6.3.	veterinary supervisio	e past 30 days prior to the date of the c n which fulfilled from the day of the colle n the conditions for a holding laid down	ction of the ova (1)/embryos (1) un	ntil the				
	(¹) or	[II.1.6.3.	collection were kept collection of the ova	ova (¹)/embryos (¹), during a period of t in holdings under veterinary supervisi a (¹)/embryos (¹) until the end of the pe the conditions for a holding laid down in	on which fulfilled, from the day a priod of 30 days mandatory store	of the age at				

▶" Equine ova/embryos - Section A •

II.	Health information	ı		II.a. Certificate reference No II.b.
	(¹) either	[II.1.6.3.1.	suscepti	ng a case of a disease mentioned below not all the animals of species tible to that disease located in the holding were slaughtered or killed and the has been free:
			begi	m any type of equine encephalomyelitis for a period of at least 6 months, ginning on the day on which the equidae suffering from the disease are ughtered,
			resu sam	m equine infectious anaemia for at least the period required to obtain a negative sult in an agar gel immunodiffusion test (AGID or Coggins tests) carried out on mples taken after the infected animals were slaughtered on two occasions nonths apart from each of the remaining equidae,
			— from	m vesicular stomatitis for a period of at least 6 months from the last recorded se,
			— from	m rabies for a period of at least one month from the last recorded case,
			— from	m anthrax for a period of at least 15 days from the last recorded case,]
	(¹) or	[II.1.6.3.1.	that dis- disinfect encepha period of following	ng a case of a disease mentioned below all the animals of species susceptible to sease located in the holding were slaughtered or killed and the premises sted, the holding was free for a period of at least 30 days from any type of equine lalomyelitis, equine infectious anaemia, vesicular stomatitis and rabies or a of at least 15 days in the case of anthrax, beginning on the day on which go the destruction of the animals the disinfection of the premises was storily completed;]
	II.1.6.4.		e of the e	past 30 days prior to the collection the ova (¹)/embryos (¹) were kept in holdings equidae has shown clinical signs of contagious equine metritis for a period of at
	II.1.6.5.	of the ova (1)/embryos	ural breeding during a period of at least 30 days prior to the date of the collection os (1) and between the date of the first samples referred to in points II.1.6.6.1 date of the collection of the ova (1)/embryos (1);
	II.1.6.6.	Manual of Di- which is reco	agnostic T gnised by	tests, which meet at least the requirements of the relevant Chapters of the Tests and Vaccines for Terrestrial Animals of the OIE, carried out in a laboratory by the competent authority and has the tests referred to hereinafter included in its ent to that provided for in Article 12 of Regulation (EC) No 882/2004(7), as
		(⁸) [II.1.6.6.1.	Coggins carried of being n referred	uine infectious anaemia (EIA), an agar-gel immuno-diffusion test (AGID or s test) or an enzyme-linked immunosorbent assay (ELISA) with a negative result out on a blood sample taken on
		II.1.6.6.2.	negative point II.	tagious equine metritis (CEM), an agent identification test carried out with a e result on at least two specimens (swabs) taken during the period referred to in .1.6.5 from at least the mucosal surfaces of the clitoral fossa and the clitoral s of the donor mare
		(1) either	[II.1.6.6.	on two occasions with an interval of not less than 7 days on

▶⁽ⁱ⁾ Equine ova/embryos - Section A •

II. Health	information	Ĭ		II.a.	Certificate reference No		II.b.
		(1) and/or	[II.1.6.6	2.2.	on one occasion ongenome of <i>Taylorella equigenitalis</i> by or real-time PCR , carried out within from the donor animal,]	a po	lymerase chain reaction (PCR)
			earlier t	han i	s referred to in points II.1.6.6.2.1 and 7 days (systemic treatment) or 21 days the donor stallion and were placed thas Amies medium, before dispatch to	(loc	al treatment) after antimicrobial ansport medium with activated
	II.1.6.7.				e and as far as I could ascertain, were agious disease during the period of		
	II.1.6.8.	on the day o contagious di		ectio	n of the ova (¹)/embryos (¹) did not sh	ow c	clinical signs of an infectious or
II.1.7.					e date on which the embryo collection nt authority of the exporting country;	(1)/p	roduction (1) team described in
II.1.8.	collection		(1), and	wer	proved conditions for a period of at le e transported under conditions which 2/65/EEC;		
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 (³) 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) (¹¹¹);						
(¹²) [II.3.					the embryos described above comply ne requirements set up in points II.1.1 to		
Notes							
Part I:							
Box I.11.:	ova/embr	yos were collec	cted/prod	uced	to the embryo collection team or eml, processed, stored and approved in commission website:		
	http://ec.e	europa.eu/food/a	animal/se	men _.	_ova/equine/index_en.htm		
Box I.22.:	The numb	per of packages	shall cor	resp	and to the number of containers.		
Box I.23.:	The identi	ification of conta	ainer and	seal	number shall be indicated.		
Box I.28.:		gory: specify sipulated embry		o de	erived embryos, in vivo derived ov	a, in	o vitro produced embryos or
	The dono	r identity shall o	orrespon	d to t	he official identification of the animal.		
	The date	of collection sha	all be indi	cate	in the following format: dd/mm/yyyy.		

Signature:

▼<u>B</u>

II.	Health information	II.a. Certificate reference No	II.b.		
ll.	Health Information	II.a. Certificate reference No	11.0.		
Part	II:				
(¹)	Delete as appropriate.				
(²)	Only third countries or parts of the terr Implementing Regulation (EU) 2018/659, and production are also authorised and as	respectively from which imports of regis	tered equidae and equidae for breeding		
(³)	Only approved embryo collection teams Directive 92/65/EEC on the Commission v		in accordance with Article 17(3)(b) of		
	http://ec.europa.eu/food/animal/semen_ov	va/equine/index_en.htm			
(4)	Council Directive 92/65/EEC of 13 July 19 the Community of animals, semen, ova Community rules referred to in Annex A(I)	and embryos not subject to animal he	alth requirements laid down in specific		
(⁵)	Council Directive 2009/156/EC of 30 Nimportation from third countries of equidae		nditions governing the movement and		
(⁶)	Insert date. (follow Guidance in Part II of t	he Notes).			
(⁷)	Regulation (EC) No 882/2004 of the E performed to ensure the verification of (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 or 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 on the Commission websites:	listed in accordance with Article 11(4) or	r Article 17(3)(b) of Directive 92/65/EEC		
	https://ec.europa.eu/food/animals/live_ani http://ec.europa.eu/food/animal/semen_ov				
(¹⁰)	Imports of equine semen are authorised from third countries listed in column 2 of Annex I to Commission Implementing Regulation (EU) 2018/659 provided that the semen was collected in the part of the territory of the third country detailed in column 4 from a donor stallion of the category of equidae positively indicated in column 11, 12 or 13 of Annex I thereto.				
(11)	Does not apply to ova.				
(¹²)	Delete if none of the embryos in the consignment was produced by in vitro fertilisation of ova.				
_	The signature and the stamp must be in a different colour to that of the printing.				
Offic	ial veterinarian				

Date:

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	TRY:			Veterinary certificate to EU			
	l.1.	Consignor Name	1.2.	Certificate reference No I.2.a.			
	11111111			Central competent authority			
		Tel.	1.4.	Local competent authority			
ment	I.5. Consignee I. Name			Person responsible for the load in EU Name			
nsign		Address		Address			
tched 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 ISO code I.10. Region of Code destination			
etails	l.11.	Place of origin	l.12.	. Place of destination			
G: :		Embryo team □		Holding ☐ Embryo team ☐			
Par	Name Approval number Address			Name Approval number Address			
		Postal code		Postal code			
	I.13.	Place of loading	l.14.	. Date of departure			
	I.15.	Means of transport	I.16. Entry BIP in EU				
		Aeroplane ☐ Ship ☐ Railway wagon ☐					
		Road vehicle Other Identification	I.17.				
	1.40	Documentary references		140 0			
	1.18.	Description of commodity		1.19. Commodity code (HS code) 05 11 99 85			
				I.20. Quantity			
	I.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					
	S	species (Scientific Category Dename)	onor i	identity Date of collection Quantity			

▶⁽ⁱ⁾ Equine ova/embryos - Section B **◄**

	II. Health	information			II.a. Certificate reference No		II.b.				
	I, the unders	signed, officia	al veterinar	ian, of the exp	porting country (²)(name of e			by			
					(hame of e	exporting	country)				
	certify that:										
	II.1.	The ova (1)/embryos	(1) described a	above:						
on		II.1.2.	supervise	rere collected (¹)/produced (¹) by the team (³) described in Box I.11, which has been apprupervised in accordance with Chapter I(III) of Annex D to Directive 92/65/EEC and is suspection by an official veterinarian at least once every calendar year;							
Part II: Certification		II.1.3.		collected (1)/produced (1), processed and stored in accordance with the requirent III(II) of Annex D to Directive 92/65/EEC;							
Part II: 0		II.1.4.			ce separated from other parts of the premision fected prior to the collection;	ses or ho	olding which is in good	t repair			
		II.1.5.	to prohibi section for	tion or quarant or storing equi	ssed and packed in laboratory facilities what tine measures as set out in Box II.1.6, in a ipment and materials used in contact will are handled;	a section	which is separated fr	om the			
		II.1.6.	come from	m donor mares	s which:						
			II.1.6.1.	Member Stat or, in the cas	rously resident for 3 months (or since entry te of the European Union during the 3 mo se of regionalisation according to Article 13 rritory of the exporting country which was	onths per 3 of Direc	iod) in the exporting of tive 2009/156/EC (8),	country			
					onsidered to be infected with African to 5(2)(a) and (b) of Directive 2009/156/EC,	horse sid	ckness in accordanc	e with			
				— free fro	om Venezuelan equine encephalomyelitis	for at leas	st 2 years,				
				— free fro	om glanders and dourine for at least 6 mor	nths;					
		(¹) either	[II.1.6.2.		om a country of export which was on the at least 6 months;]	ne day of	collection free of ve	sicular			
		(¹) or	[II.1.6.2.		by a virus neutralisation test for vesicular 						
		(¹) either	[II.1.6.3.	supervision v	ast 30 days prior to collection have been which fulfilled from the day of collection of the conditions for a holding laid down in ular:]	f ova (1)/	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 which fulfilled from the day of collection of va (¹)/embryos (¹), the period of 30 da apsed, the conditions for a holding laid and in particular:]	f ova (¹)/e ays mand	embryos (¹) until, in the latory storage at ap	ne case proved			

▶⁽ⁱ⁾ Equine ova/embryos - Section B ∢

II. Health inform	mation		II.a. Certificate reference No	II.b.
	(¹) either	• DOCUMENT OF STREET	ollowing a case of a disease mentioned below susceptible to the disease located on the holding he holding has been free:	BB (1일 : 10일 : 14일 : 15일 :
			 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 immunodiffus out on samples taken after the infected two occasions 3 months apart from each of 	ion test (Coggins tests) carried animals were slaughtered on
			 from vesicular stomatitis for at least 6 month 	ns from the last recorded case,
			- from rabies for at least one month from the	ast recorded case,
			 from anthrax for at least 15 days from the la 	st recorded case,]
	(¹) or		collowing a case of a disease mentioned belowing a case of a disease located in the holdifulled and the premises disinfected, the holdir to days from any type of equine encephalomyelizesicular stomatitis and rabies or 15 days in 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 tis, 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 of	en used for natural breeding during at least ova or embryos and between the date of th 6 and II.1.6.7 and the date of the collection of over	e first samples referred to in
	II.1.6.6.	test) or an oncollection of	subjected with negative result to an agar-gel in ELISA for equine infectious anaemia carried (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 of negative res the first colle sinuses on t and on an	subjected to an agent identification test for of Taylorella equigenitalis after a cultivation of Taylorella equigenitalis after a cultivation of Taylorella equipenitalis after a cultivation of Taylorella equipenitalism and the pection of ova or embryos from mucosal surfaces were consecutives oestrus periods on	7 to 14 days carried out with ast 30 days prior to the date of of the clitoral fossa and clitoral (4) and on(4).
	II.1.6.8.		f my knowledge and as far as I could ascertain, ering from an infectious or contagious disease of e collection;	
	II.1.6.9.	have on the or contagiou	day of collection of ova (¹)/embryos (¹) not show s disease;	n clinical signs of an infectious
II.1.			uced (1) after the date on which the embryo co as approved by the competent authority of the e	

Health information

▶" Equine ova/embryos - Section B •

II.b.

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 Checker (IIII) of Appendix De Dispersive 20/55/5500
	in Chapter III(II) of Annex D to Directive 92/65/EEC;

II.a. Certificate reference No

- II.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);
- II.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

Part I:

Box I.11.: The place of origin shall correspond to the embryo collection team or embryo production team by which the 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.
- (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 Council Directive 92/65/EEC on the Commission website:

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

- (4) Insert date.
- (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.

▼B

COUNTRY ▶⁽¹⁾ Equine ova/embryos - Section B ◀ Health information II.a. Certificate reference No II.b. 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: (⁶) 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 different colour to that of the printing. Official veterinarian Name (in capital letters): Qualification and title: Signature: Date: Stamp:

►(1) <u>M1</u>

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.

- (i) The original of the health certificate shall accompany the consignment until it reaches the border inspection post of entry into the Union.
- 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.

⁽¹⁾ OJ L 13, 16.1.1997, p. 28.

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		
I, the undersigned, master of ship (name), declare that the animals referred to in the attached health certificate No		
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)

	Se	erial Number:
	Re	eference No of Air Cargo Transfer Manifest:(1)
Country where transhipment takes p	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	
I, the undersigned, official veterina took place under my supervision a		e above mentioned airport (²)/port (²) declare that the transhipment ying conditions:
(a) the equidae were during the tra	anshipment protected from attac	cks by insect vectors of diseases transmissible to equidae;
(b) the equidae did not come into contact with equidae of a different health status;		
(c) the crates, containers or jet-stalls and the surrounding airspace in the transport compartment were sprayed with an appropriate insect repellent in combination with an insecticide immediately after the closing of the doors of the aircraft (²)/vessel (²).		
The consignment has been transhipped in full and apparent good order and conditions except as noted in the "Remarks" column.		
Done at		. on
		Stamp
(signature of the official veter		
(name in capital le	etters and title)	
(¹) Keep empty if transhipment from v (²) Delete as appropriate	essel to vessel	