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)

# COMMISSION IMPLEMENTING REGULATION (EU) 2018/659

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

(Text with EEA relevance)

### THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to Council Directive 91/496/EEC of 15 July 1991 laying down the principles governing the organisation of veterinary checks on animals entering the Community from third countries and amending Directives 89/662/EEC, 90/425/EEC and 90/675/EEC<sup>(1)</sup>, and in particular Article 3(2) and Article 9(1)(c) thereof,

Having regard to 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<sup>(2)</sup>, and in particular Article 17(3) thereof,

Having regard to Council Directive 2009/156/EC of 30 November 2009 on animal health conditions governing the movement and importation from third countries of equidae<sup>(3)</sup>, and in particular Article 2(i), Article 12(1), (4) and (5), Article 13(2), Articles 15, 16, 17 and 19 thereof,

# Whereas:

- (1) Directive 2009/156/EC lays down the animal health requirements governing imports into the Union of equidae. It provides that only equidae that come from a third country or part of a third country on a list of third countries drawn up in accordance with that Directive, and accompanied by a health certificate corresponding to a model also drawn up in accordance with that Directive, may be imported into the Union. The health certificate must attest that the equidae comply with the health conditions set out in accordance with that Directive in the corresponding health certificate.
- (2) The list of third countries from which Member States authorise imports of live equidae and semen, ova and embryos of the equine species and the regionalisation of certain of those third countries should be established on the basis of the animal health status of those third countries and be based on the list of third countries and parts of the territory of third countries set out in Commission Decision 2004/211/EC<sup>(4)</sup>.
- (3) In accordance with Article 12(2)(a) of Directive 2009/156/EC, the animal health requirements laid down in this Regulation should be based on a risk assessment. The principle of grouping countries in health zones, as provided for in Article 12(4)

of Directive 2009/156/EC, according to common risks has proven to be effective. However, as the word 'zones' suggests certain contiguity and certain risks of the same kind may prevail in distant areas, countries should be assigned to specific 'sanitary groups'.

- (4) Directive 92/65/EEC lays down the animal health requirements governing imports into the Union of semen, ova and embryos of the equine species. It provides that only commodities that come from a third country or part of a third country on a list of third countries drawn up in accordance with that Directive, and accompanied by a health certificate corresponding to a model also drawn up in accordance with that Directive, may be imported into the Union. The health certificate must attest that the commodities come from approved collection and storage centres or collection and production teams offering guarantees at least equivalent to those established in Annex D(I) to that Directive.
- (5) Directive 92/65/EEC, as amended by Council Directive 2008/73/EC<sup>(5)</sup>, introduced a simplified procedure for the listing of semen collection and storage centres and embryo collection and production teams in third countries, approved for imports of the commodities into the Union. The lists are available at the Commission website<sup>(6)</sup>.
- (6) Annex D to Directive 92/65/EEC sets out certain requirements for semen, ova and embryos of equidae and provides for rules for the approval, supervision and operation of semen collection and storage centres and embryo collection and production teams and detailed conditions for the health status of the donor animals. Accordingly, it is necessary to establish model health certificates for imports into the Union of semen, ova and embryos of equidae.
- (7) In addition, provision should be made for imports into the Union of existing stocks of commodities that comply with the provisions of Directive 92/65/EEC established prior to the entry into force of the amendments introduced by Commission Regulation (EU) No 176/2010<sup>(7)</sup>. Accordingly, it is necessary to set out separate model health certificates for imports of consignments of semen, ova and embryos of equidae collected or produced, processed and stored in accordance with Annex D to Directive 92/65/EEC prior to 1 September 2010.
- (8) The long lasting stocking capabilities for such commodities make it impossible at present to fix a date for the exhaustion of the existing stocks. Therefore, it is not possible to fix a date for the termination of the use of those model health certificates for the existing stocks.
- (9) In order to ensure full traceability of the commodities, model health certificates should be set out in this Regulation for imports into the Union of semen of equidae collected in approved semen collection centres and dispatched from an approved semen storage centre, whether or not the latter constitutes part of a semen collection centre approved under a different approval number.
- (10) In addition, it is appropriate that consignments of the commodities imported into the Union from Switzerland are accompanied by the health certificates drawn up in accordance with the models used for trade within the Union in semen, ova and embryos

of animals of the equine species and set out in Commission Decision 2010/470/EU<sup>(8)</sup>, with the adaptations set out in points 8 and 9 of Chapter IX(B) of Appendix 2 of Annex 11 to the Agreement between the European Community and the Swiss Confederation on trade in Agricultural Products, as approved by Decision 2002/309/EC, Euratom of the Council and of the Commission<sup>(9)</sup>.

- (11) Semen, ova and embryos of animals of the equine species consigned from Canada to the Union may be accompanied by health certificates laid down in accordance with the Agreement between the European Community and the Government of Canada on sanitary measures to protect public and animal health in respect of trade in live animals and animal products<sup>(10)</sup>, as approved by Council Decision 1999/201/EC<sup>(11)</sup>.
- (12) Semen, ova and embryos of animals of the equine species consigned from New Zealand to the Union may be accompanied by health certificates laid down in accordance with the Agreement between the European Community and New Zealand on sanitary measures applicable to trade in live animals and animal products<sup>(12)</sup>, as approved by Council Decision 97/132/EC<sup>(13)</sup>.
- (13) With the view to simplifying Union legislation, it is appropriate to group together in a single Regulation the animal health and certification requirements applicable for the entry into the Union of consignments of equidae and of semen, ova and embryos of equidae, including the list of third countries and parts of the territory of third countries from which Member States are to authorise the introduction into the Union of such consignments.
- (14) In order to preserve the certified health status of equidae during their movement from the exporting third country to the Union, it is necessary to lay down animal health requirements concerning the transport of equidae.
- (15) Provisions should be made for the quality of health testing and the recording of vaccinations. Provisions should also be made for the confirmation of test results by the European Union reference laboratory for equine diseases other than African horse sickness, designated in accordance with Commission Regulation (EC) No 180/2008<sup>(14)</sup> where risk based sampling of equidae, in accordance with Commission Decision 97/794/EC<sup>(15)</sup>, produced results different to those certified by the dispatching third country.
- (16) The tests used for the diagnosis of equine viral arteritis and the categories of male equidae to which the test requirements for equine viral arteritis apply should be defined based on the recommendations of the Scientific Veterinary Committee<sup>(16)</sup>, which are contained in Commission Decision 95/329/EC<sup>(17)</sup>, and the latest recommendations in Chapter 12.9. of the Terrestrial Animal Health Code of the World Organisation for Animal Health (OIE), 2016 Edition<sup>(18)</sup>.
- (17) A specific model health certificate should be laid down for transit through the Union of live equidae from one third country, or part of the territory of a third country, to another third country or to another part of the territory of the same third country.
- (18) For the controls necessary to ensure a uniform implementation by Member States of the provisions on temporary admission of registered horses, the re-entry of registered

horses after temporary export, the transit of equidae and the conversion of temporary admission of registered horses into permanent entry, it is necessary to lay down specific and additional provisions on the use of the integrated computerised veterinary system 'TRACES' provided for in Commission Decisions 2003/24/EC<sup>(19)</sup> and 2004/292/EC<sup>(20)</sup> from the veterinary border inspection post of entry, approved in accordance with Commission Decision 2009/821/EC<sup>(21)</sup> till the exit point from the Union.

- (19) In the interests of consistency and simplification of Union legislation, the format of the model health certificates for entry into the Union of equidae and of semen, ova and embryos of the equidae should be based on the standard models for veterinary certificates set out in Annex I to Commission Decision 2007/240/EC<sup>(22)</sup>.
- (20) Provisions should be made to determine, in accordance with Article 19(c) of Directive 2009/156/EEC, the conditions for converting temporary entry into permanent entry, including provisions on the necessary information in TRACES and the exchange of the Common Veterinary Entry Document (CVED) set out in Annex I to Commission Regulation (EC) No 282/2004<sup>(23)</sup>.
- (21) Specific animal health conditions should be laid down for the re-entry of registered horses after temporary export to third countries in order to participate in races, competitions and cultural events, and the corresponding model certificates should be set out in an Annex to this Regulation.
- (22) Commission Decision 93/444/EEC<sup>(24)</sup> defines the 'exit point' and requires, inter alia, that animals destined for export to a third country are, on their way to the exit point, to be accompanied by a health certificate applicable at least to trade in animals for slaughter of the species concerned. It also requires the competent authority at the place of dispatch to notify the exit point of the intended movement. It is necessary to clarify that in order to ensure traceability the 'exit point' should be a border inspection post and that the health certificate referred to in Article 2(1) of Decision 93/444/EEC should be the health certificate set out in Annex III to Directive 2009/156/EC also in the case of registered horses intended for temporary export.
- (23) For reasons of legal certainty, Commission Decisions 92/260/EEC<sup>(25)</sup>, 93/195/EEC<sup>(26)</sup>, 93/196/EEC<sup>(27)</sup>, 93/197/EEC<sup>(28)</sup>, 94/699/EC<sup>(29)</sup>, 95/329/EC, 2003/13/EC<sup>(30)</sup>, 2004/177/ EC<sup>(31)</sup>, 2004/211/EC, 2010/57/EU<sup>(32)</sup> and 2010/471/EU<sup>(33)</sup> should be repealed.
- (24) In order to enable economic operators to adapt to the new rules laid down in this Regulation, it is appropriate to provide for a transitional period during which Member States are to authorise the entry into the Union of equidae and semen, ova and embryos of equidae which comply with the conditions set out in the model health certificates applicable before the date of application of this Regulation.
- (25) The measures provided for in this Regulation are in accordance with the opinion of the Standing Committee on Plants, Animals, Food and Feed,

#### HAS ADOPTED THIS REGULATION:

#### 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 wit a distinct health status with respect to on or more specific diseases and subject to appropriate surveillance, disease control and biosecurity measures;
(b)	'identification document'	means any document which may be used to prove the identity of an equine animal and which includes at least the following information:  (i) a narrative describing the animal and recording its marks depicted in a completed outline diagram;  (ii) a reference to specific marks, characteristics or identifiers which establish an unambiguous link between the animal and the document;  (iii) the information set out in points 1, 2, 3 and 6 to 10 of Part A and in

a 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).

b 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).

c 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).

d https://ec.europa.eu/food/sites/food/files/animals/docs/ad\_control-measures\_bt\_guidance\_vpe\_7068\_2012.pdf

		points 12 to 18 in Part B of Section 1 of Annex I to Commission Implementing Regulation (EU) 2015/262 <sup>a</sup> ;
(c)	'registered horse'	means an animal of the species <i>Equus caballus</i> registered as defined in Council Directive 90/427/EEC <sup>b</sup> , 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 <sup>c</sup> ;
(e)	'type of entry'	means respectively the temporary admission, the re-entry after temporary export, imports and transit;
(f)	'temporary admission'	means the status of a registered horse originating in a third country and moved into the Union territory for a period of less than 90 days;
(g)	'temporary export'	means the movement of a registered horse out of the Union for a period of less than 90 days;
(h)	're-entry'	means the movement of a registered horse from a third country into the Union after temporary export from the Union;
(i)	'imports'	means the movement of a consignment of equidae or their semen, ova or embryos into the Union for an undetermined period;
(j)	'transit'	means the movement of a consignment of equidae across Union territory by road,
a	Commission Implementing Regulation (EU) 2015/262 of Directives 90/427/EEC and 2009/156/EC as regards the n Regulation) (OJ L 59, 3.3.2015, p. 1).	17 February 2015 laying down rules pursuant to Council
b	Council Directive 90/427/EEC of 26 June 1990 on the zo Community trade in equidae (OJ L 224, 18.8.1990, p. 55)	
с	Council Directive 97/78/EC of 18 December 1997 laying checks on products entering the Community from third co	down the principles governing the organisation of veterinary puntries (OJ L 24, 30.1.1998, p. 9).

https://ec.europa.eu/food/sites/food/files/animals/docs/ad\_control-measures\_bt\_guidance\_vpe\_7068\_2012.pdf

		rail or waterway transport from one third country to another or from one part of the territory of a third country to another part of the territory of the same third country;
(k)	'border inspection post'	means any inspection post as defined in Article 2(2)(f) of Directive 91/496/EEC and Article 2(2)(g) of Directive 97/78/EC and approved for the commodity concerned in accordance with Decision 2009/821/EC;
(1)	'category of equidae'	means respectively registered equidae, equidae for slaughter and equidae for breeding and production as defined in Article 2 of Directive 2009/156/EC, and registered horses;
(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;
(0)	'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,

a 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).

**b** 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).

c 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).

d https://ec.europa.eu/food/sites/food/files/animals/docs/ad\_control-measures\_bt\_guidance\_vpe\_7068\_2012.pdf

		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 body-coverage;
(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 <sup>d</sup> ,  — 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.

- a 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).
- b 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).
- c 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).
- $\textbf{d} \qquad \text{https://ec.europa.eu/food/sites/food/files/animals/docs/ad\_control-measures\_bt\_guidance\_vpe\_7068\_2012.pdf}$

#### 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.

#### Article 4

# 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.

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.

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
- 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.

#### Article 10

# 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

- 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).
- 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

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 (34).

- 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<sup>(35)</sup>;
  - 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

- 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).
- 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.
- 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

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);
- 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).
- 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.
- 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.
- 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.
- 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.

### Article 17

# Operator responsibilities for temporarily admitted registered horses

- 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:
  - the registered horse shall leave the Union through a border inspection post indicated in the health certificate referred to in Article 3(1)(a) not later than 89 days following the date of entry into the Union indicated on the corresponding CVED.
- 2 The operator referred to in paragraph 1 shall remain responsible for the movement of the registered horse during its temporary admission in the Union, and in particular shall inform:
  - a the competent authority referred to in points (i) and (iii) of Article 16(1)(b) regarding any changes to be made to the movements stated in the declaration accompanying the health certificate referred to in Article 3(1)(a);
  - b the border inspection post of exit regarding the date when the temporarily admitted registered horse is to depart from the Union;
  - 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.

#### Article 18

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

- 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).
- 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).

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).

#### Article 19

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

- 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).
- 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.
- 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;
  - the health certificate for temporary export signed by the official veterinarian in the Member State of origin, or an authorised copy thereof, is presented on request of the border inspection post of re-entry into the Union.
- 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.
- 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.

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

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

Done at Brussels, 12 April 2018.

For the Commission

The President

Jean-Claude JUNCKER

### ANNEX I

# LIST OF THIRD COUNTRIES<sup>0</sup>AND PARTS OF THE TERRITORY OF THIRD COUNTRIES<sup>0</sup>FOR THE ENTRY INTO THE UNION OF CONSIGNMENTS OF EQUIDAE AND OF SEMEN, OVA AND EMBRYOS OF EQUIDAE

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

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

c 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.

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

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

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

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- **d** The former Yugoslav Republic of Macedonia the definitive nomenclature for this country will be agreed following current negotiations at UN level.
- e Without prejudice to specific certification requirements provided for in Article 17 of the Agreement on the European Economic Area (OJ L 1, 3.1.1994, p. 3).
- **f** Excluding Kosovo as defined by United Nations Security Council Resolution 1244 of 10 June 1999.

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

c 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.

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

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

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

MK	fYRO	)MK-	0Who		X	X	X	X	X					X	
МО	Maca	фО-	0Who coun		X	X	X	_	-		_	_	-	X	
MY	Mala	yMY-	0Who		_	_	_	_			_	-			
		MY-	1Penii	ıSüla	X	X	X	_	<u> </u>		_	_		X	
MU	Maur	i <b>MU</b> S-	0Who coun		_	_	X		-		_	_	-	X	
MX	Mexi	dolX-	0Who		_	_		_	_	_	_	_	_		
		MX-	lMetr area of Mex City		an	X									Only if certified in accordance with Chapter 1 of Section B of Part 2 of Annex II
NO	Norw	<b>ayo-</b> 1	Who coun	l <b>&amp;</b> try	X	X	X	X	X	X	X	X	X	X	
NZ	New Zeala	NZ-0	Who coun		X	X	X	X	X					X	
OM	Oma	nOM-	0Who		X	X	X	_	_		_	_	_	X	
PE	Peru	PE-0	Who		_	_			_	_	_	_	-		
		PE-1	Region of Lima		X	X	X	_			_	_		X	

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

- **b** Where official regionalisation applies in accordance with Article 13(2)(a) of Directive 2009/156/EC.
- c 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.
- **d** The former Yugoslav Republic of Macedonia the definitive nomenclature for this country will be agreed following current negotiations at UN level.
- e Without prejudice to specific certification requirements provided for in Article 17 of the Agreement on the European Economic Area (OJ L 1, 3.1.1994, p. 3).
- f Excluding Kosovo as defined by United Nations Security Council Resolution 1244 of 10 June 1999.

	1									,		1		
PM	St Pierro & Miqu	e	)Who		_	_	X	_	X				X	
PY	Parag	g <b>uPak</b> y-0	Who		X	X	X	X	X				X	
QA	Qatai	QA-0	)Who		X	X	X	_	_		_	 	X	
RS	Serbi	aRS-0	Who		X	X	X	X	X				X	
RU	Russ	i <b>R</b> U-0	Who		_	_	_	_	_	_	_	 _		
		RU-1	Arkh Volog Murr Lenir Novg Psko Briar Vlad Ivano Tver, Kalu Kostr Mosk Orjol Riasa Smol Tula, Jaros	ningrad angels gda, nansk ngrad, sorod, v, isk, imir, ovo, ga, roma, ensk, lavl, novgo v, orod, nesh, k,	sk,	X	X	X	X				X	

- a Without prejudice to specific certification requirements provided for in Decision 2002/309/EC, Euratom of the Council, and of the Commission.
- $\label{eq:barrier} \textbf{b} \qquad \text{Where official regionalisation applies in accordance with Article 13(2)(a) of Directive 2009/156/EC.}$
- e 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.
- **d** The former Yugoslav Republic of Macedonia the definitive nomenclature for this country will be agreed following current negotiations at UN level.
- e Without prejudice to specific certification requirements provided for in Article 17 of the Agreement on the European Economic Area (OJ L 1, 3.1.1994, p. 3).
- f Excluding Kosovo as defined by United Nations Security Council Resolution 1244 of 10 June 1999.

		Tambov, Astrahan, Volgograd, Penza, Saratov, Uljanovsk, Rostov, Orenburg, Perm and Kurgan											
	RU-2	Regions of Stavropol and Krasnodar	X	X	X	X	X					X	
		Repultics of Karelia, Marij-El, Mordovia, Chuvachia, Kalmykia, Tatarstan, Dagestan, Kabardino-Balkaria, Severnaya Osetia, Ingushetia and Karachaeve Cherkesia		X	X	X	X					X	
SA	Saudi SA-0 Arabia	country	_					_	_	_	_		
	SA-1	Whole country, except SA-2	X	X	X			X	_	_		X	

- a Without prejudice to specific certification requirements provided for in Decision 2002/309/EC, Euratom of the Council, and of the Commission.
- **b** Where official regionalisation applies in accordance with Article 13(2)(a) of Directive 2009/156/EC.
- c 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.
- **d** The former Yugoslav Republic of Macedonia the definitive nomenclature for this country will be agreed following current negotiations at UN level.
- **e** Without prejudice to specific certification requirements provided for in Article 17 of the Agreement on the European Economic Area (OJ L 1, 3.1.1994, p. 3).
- f Excluding Kosovo as defined by United Nations Security Council Resolution 1244 of 10 June 1999.

	SA-2	Protection and		_									
		surveilland	e										
		zones											
		in											
		the											
		provinces											
		of Jizan											
		Asir											
		and											
		Najran											
		as											
		described											
		in											
		BOX											
		3											
SG	Singar Sore	WholeG	X	X	X						_	X	
50		country	11	11	11							11	
TII	The Heat		37	37	37							V	
TH	Thailah H-0		X	X	X	_			_	_	_	X	
		country											
TN	TunisiaN-0		X	X	X	X	X					X	
		country											
TR	Turke\R-0	Whole-	_		_								
		country											
	TP 1	Provides											
	1 IX-1	of											
		Ankara,											
		Edirne,											
		Istanbul,											
		Izmir,											
		Kirklareli											
		and											
		Tekirdag											
UA	UkraineA-(	)Whol&B	X	X	X	X	X	X	X	X		X	
		country											
US	UnitedUS-0		X	X	X	X	X	X	X	X	X	X	
US	States	country	A	Λ	Λ	Λ	Λ	Λ	Λ	Λ	Λ	Λ	
	States	Country y											

- a Without prejudice to specific certification requirements provided for in Decision 2002/309/EC, Euratom of the Council, and of the Commission.
- **b** Where official regionalisation applies in accordance with Article 13(2)(a) of Directive 2009/156/EC.
- e 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.
- **d** The former Yugoslav Republic of Macedonia the definitive nomenclature for this country will be agreed following current negotiations at UN level.
- e Without prejudice to specific certification requirements provided for in Article 17 of the Agreement on the European Economic Area (OJ L 1, 3.1.1994, p. 3).
- **f** Excluding Kosovo as defined by United Nations Security Council Resolution 1244 of 10 June 1999.

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	of Ame	rica													
UY	Urug	ulalyY-(	)Who	1	X	X	X	X	X	X	X	X		X	
ZA	Soutl Afric	nZA-0 a	Who			_	_	_			_	_	_		
		ZA-1	Metrarea of Cape Town (see BOX 4 for detai	1	a <del>n</del>	_	_	_	_	_	_	_			Commission Decision 2008/698/ EC

- a Without prejudice to specific certification requirements provided for in Decision 2002/309/EC, Euratom of the Council, and of the Commission.
- **b** Where official regionalisation applies in accordance with Article 13(2)(a) of Directive 2009/156/EC.
- c 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.
- **d** The former Yugoslav Republic of Macedonia the definitive nomenclature for this country will be agreed following current negotiations at UN level.
- e Without prejudice to specific certification requirements provided for in Article 17 of the Agreement on the European Economic Area (OJ L 1, 3.1.1994, p. 3).
- f Excluding Kosovo as defined by United Nations Security Council Resolution 1244 of 10 June 1999.

#### **LEGEND TO ANNEX I:**

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

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

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A		equine infectious a arteritis	equine infectious anaemia, equine viral arteritis				
В			equine infectious anaemia, equine viral arteritis, glanders, dourine				
C		arteritis, Eastern ar	equine infectious anaemia, equine viral arteritis, Eastern and Western equine encephalomyelitis, vesicular stomatitis				
D		arteritis, glanders, and Western equine Venezuelan equine	equine infectious anaemia, equine viral arteritis, glanders, dourine, Eastern and Western equine encephalomyelitis, Venezuelan equine encephalomyelitis, vesicular stomatitis				
E			naemia, equine viral dourine, African horse				
F		equine infectious a horse sickness	naemia, dourine, African				
G			equine infectious anaemia, equine viral arteritis, glanders, dourine, Japanese encephalitis				
BOX 1							
CN	China	CN-1	The specific equine disease-free zone in the Guangdong Province with the following delimitation:				
			Core: equestrian zone 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 administrative divisions in Conghua City surrounding the core zone				

	of 2 00 Protection are zone bounda the follocontigue administration surroun	ds ries of owing ous strative ns
	_ _ _	District of Conghua City, Huadu District of Guangzhou City, Zengcheng City, administrative divisions in
l l	— — Biose <del>cu</del> rity nighway passage	Qingcheng District of Qingyuan City, Fogang County, Xinfeng County, Longmen County from the equestrian site
		in the core zone to Guangzhou Baiyun International Airport through to the

<b>Q</b>
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
mgm way

BOX 2

			with the equine exclusion zone on both sides of that highway of at least one km width; Pre-: the quarantine entry facilities in the quaraptitection 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 zone in the Metropolitan area of Shanghai:
			Westernuangpu River boundary its estuary in the North to the bifurcation of the Dazhi River, Southfrom the boundarfurcation of the Huangpu River to the estuary of the Dazhi River in the East, Northernst line. and Eastern boundaries

EG	Egypt	EG-1	The Equine Disease Free Zone (EDFZ) of about 0,1 km² size, established around the Egyptian Armed Forces Veterinary Hospital at El-Nasr Road, across Al Ahly Club, on the Eastern outskirts of Cairo, (localised at 30°04′19.6″N 31°21′16.5″E) and the passage of 10 km on the El-Nasr Road and the Airport Road to Cairo International Airport.  (a) Delineation of the boundaries of the EDFZ: From the crossing of El-Nasr Road with El-Shaheed Ibrahim El-Shaikh Road (at 30°04′13.6″N 31°21′04.3″E) along the El-Shaheed Ibrahim El-Shaikh Road for about 500 m to the North until the first junction with the Passage Inside
			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
Road for 300 m to
Road for
Road for 300 m to South-
Road for 300 m to South- West until
Road for 300 m to South- West until opposite of
Road for 300 m to South- West until opposite of
Road for 300 m to South- West until opposite of the junction
Road for 300 m to South- West until opposite of the junction of El-Nasr
Road for 300 m to South- West until opposite of the junction of El-Nasr
Road for 300 m to South- West until opposite of the junction of El-Nasr Road with
Road for 300 m to South- West until opposite of the junction of El-Nasr Road with Hassan
Road for 300 m to South- West until opposite of the junction of El-Nasr Road with
Road for 300 m to South- West until opposite of the junction of El-Nasr Road with Hassan Ma'moon
Road for 300 m to South- West until opposite of the junction of El-Nasr Road with Hassan Ma'moon Road,
Road for 300 m to South- West until opposite of the junction of El-Nasr Road with Hassan Ma'moon Road, turning
Road for 300 m to South- West until opposite of the junction of El-Nasr Road with Hassan Ma'moon Road, turning
Road for 300 m to South- West until opposite of the junction of El-Nasr Road with Hassan Ma'moon Road, turning right and
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
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
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
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
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
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,
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
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
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
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
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
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
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
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,
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
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
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
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

the Passage

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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 preexport 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.
BOX 3			
SA	Saudi Arabia	SA-1	Approved Quarantine stations:  1. Riyadh Airport  2. King Abdulaziz Race Track (Janadrijah)
		SA-2	Delimitation of the protection and surveillance zones established in accordance with points (a) and (b) of the second paragraph of Article 5(2) of Directive 2009/156/ EC:  1. Province of Jizan  — Protection zone: the whole province, except the part north of the road control post at Ash-

2.	Shuqaiq at road No 5 and north of road No 10; — Surveillance zone: the part of the province north of the road control post at Ash- Shuqaiq at road No 5, controlled by the road control post at Al Qahmah, and north of road No 10. Province of Asir — Protection zone: the part
----	--

the
province
delineated
by
road
No
10,
between
Ad
Darb,
Abha
and
Khamis-
Mushayt
to
the
north,
except
the
equestrian
clubs
at
their
air
and
military
bases,
and
the
part
of
the .
province
delineated
to
the
north
by
road
No
15
leading
from
Khamis-
Mushayt
through
Jarash,
Al
Utfah
and
Dhahran
Al
Janoub

of Najran, and, the part of the province delineated to the north by the road leading from Al Utfah through Al Fayd to Badr Al Janoub (Province of Najran); — Surveillance zone: the equestrian clubs at their air and military	
clubs at their air and	
the part of the province between the	

border
of
the
protection
zone
and
road
No
209
from
Ash-
Shuqaiq
to
the
road
control
post
Muhayil
on
road
No 211,
the
part
of
the
province
between
the
control
post
on
road
No 10
south
of
Abha,
the
city
of
Abha
and
the
road
control
post
Ballasmer
65 km
from
Abha
on
road
No
15

leading
north,
the
part
of
the
province
between
Detween
Khamis
Mushay
and
the
road
control
post
90
km
from
Abha
on
road
No 255
to
Samakh
and
the
road
control
post
at
Yarah,
90
km
from
Abha,
on
road
No 10
leading
to
Riyadh,
and,
the
part
of
the
province
south
of
a
virtual
line
between

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	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 177 leading south to the part of the part of the province south of road No 15
	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, on road No 177, from the border of the province of Najran until the road control post at Khashm-Ghurab, on the border of the province of Najran until the road control post at Khashm-Ghurab,
--	--	--	--	---

BOX 4			km from Najran, and west of road No 175 leading to Sharourah.
ZA	South Africa	ZA-1	Approved Quarantine stations:
			1. Kenilworth Quarantine Station Delimitation of the Metropolitan area of Cape-Town (ZA-1):  Northeanwberg bouraddy (M14); Easkeonberg bouraddy (M14), Plattekloof Road (M14), N7 Highway, N1 Highway and M5 Highway, Southteny bouraddy Prince George's Drive, Wetton Road, Riverstone 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;
		Westonstline
	1	oou <b>inda</b> ry
		Camps
		Bay
		to
		Blaauwberg
		Road.

# ANNEX II

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

# PART 1

# Temporary admission and transit

# Section A

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

OUN	TRY:		Veterinary certificate to E	
	l.1.	Consignor Name	1.2.	Certificate reference No I.2.a.
		Address	1.3.	Central competent authority
ent		Tel.	1.4.	Local competent authority
шÉ	1.5.	Consignee	1.6.	
nsić		Name Address		
9				
atche		Postcode Tel.		
Part I : Details of dispatched consignment	1.7.	Country of ISO code I.8. Region of Code origin crigin	1.9.	Country of ISO code I.10. Region of Code destination destination
etails	l.11.	11. Place of origin		. Place of destination
art I: De		Name Approval number Address		Name Address
ъ.				Postcode
	I.13.	Place of loading	1.14.	. Date of departure
	I.15.	Means of transport	1.16.	. Entry BIP in EU
		Aeroplane ☐ Ship ☐ Railway wagon ☐		
		Road vehicle Other Identification	l.17.	. No(s) of CITES
		Documentary references		T
	I.18.	Description of animal		I.19. Commodity code (HS code) 01 01
				I.20. Quantity
	I.21.			I.22. Number of packages
	1.23.	Seal/Container No		1.24.
	1.25.	Animal certified for:		
		Registered horse		
	1.26.			I.27. For import or admission into EU
	1.28.	Identification of the animal		1
		pecies (Scientific Identification system Identi name) Equus caballus	fication	ion number Age Sex

EUROPEAN UNION

		II.a. Certificate reference number	II.b. Local reference number								
II.	Attestation of a	nimal health and welfare									
I, the under	the undersigned official veterinarian, hereby certify, that the animal described in Box I.28:										
_	<ul> <li>is a registered horse as defined in Article 2(c) of Commission Implementing Regulation (EU) 2018/659;</li> </ul>										
_	was examined today (1) and found free of clinical signs of diseases and of obvious signs of ectoparasite										
	infestation;										
_	is not intended f	or slaughter under a national programme of infe	ectious or contagious disease eradication;								
_	meets the requir	rements attested in points II.1 to II.5 of this certi	ificate;								
_	is accompanied	by the written declaration, signed by the owner	of the animal or the representative of the owner.								
II.1.	Attestation on th	ird country or part of the territory of third countr	ry and holding of dispatch								
II.1.1.	a country), a co		(insert name of country or part of the territory of on the date of issuing this certificate has the Code (²);								
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:										
	in w of A have	hich there has been no clinical, serological (in u frican horse sickness during the period of 2 yea	ess in accordance with Directive 2009/156/EC and unvaccinated equidae) or epidemiological evidence ars prior to the date of dispatch and in which there uring the period of 12 months prior to the date of								
		hich Venezuelan equine encephalomyelitis has date of dispatch;	s not occurred during the period of 2 years prior t								
	c) in w	hich dourine has not occurred during the period	d of 6 months prior to the date of dispatch;								
	d) in w	hich glanders has not occurred during the perio	od of 6 months prior to the date of dispatch;								
(3) either		rhich vesicular stomatitis has not occurred duatch;]	uring the period of 6 months prior to the date of								
(³) or	and 21 c	a blood sample taken from the animal on	the period of 6 months prior to the date of dispatch(insert date), within a period of with negative result for antibody to the vesicular								
	(³) e	ither [in a virus neutralisation test at a serum of	dilution of 1 in 32;]]								
	(³) o	r [in an ELISA in accordance with the rel and Vaccines for Terrestrial Animals of the	levant Chapter of the Manual of Diagnostic Test he OIE;]]								
II.1.4.		s not come from a holding and to the best of II.1.4.7 was not in contact with animals from	my knowledge for the time periods referred to i								

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# **EUROPEAN UNION**

EUROPEAN U	INION		Registered equidae, equidae for breedi	ing an	d production ed	quidae for slaughter	
			II.a. Certificate reference number	II.b.	Local reference	number	
	II.1.4.1.	in the cas	se of equidae suspected of having contracted douri	ine,			
		( <sup>3</sup> ) either	[6 months beginning on the date of the last a suspected of having contracted dourine or infected				
		(³) or	[in the case of a stallion, until the animal is castra	ited;]			
		( <sup>3</sup> ) or	[30 days following the date of completion of the after all animals of susceptible species have been		•	tion of the premises	
	II.1.4.2.	in the cas	se of glanders,				
		(³) either	[6 months beginning on the day on which the subjected with positive results to a test for the Burkholderia mallei or antibodies to that pathoger	he de	tection of the	causative pathogen	
		( <sup>3</sup> ) or	[30 days following the date of completion of the after all animals of susceptible species have been				
	II.1.4.3.	in the cas	e case of equine encephalomyelitis of any type,				
		( <sup>3</sup> ) either	[6 months beginning on the day on which the equal slaughtered;]	ıidae s	suffering from the	disease have been	
		(³) or	[6 months beginning on the day on which the equal Nile Fever, Eastern equine encephalomyelitis or died, been removed from the holding or fully reco	r Wes	tern equine enc		
		( <sup>3</sup> ) or	[30 days following the date of completion of the after all animals of susceptible species have been			tion of the premises	
	II.1.4.4.	slaughter gel immi	se of equine infectious anaemia, until the date on red, the remaining equine animals on the holding h unodiffusion test (AGID or Coggins test) carrie sions 3 months apart;	nave s	hown a negative	reaction in an agar	
	II.1.4.5.	in the cas	se of vesicular stomatitis,				
		(³) either	[6 months following the last case;]				
		( <sup>3</sup> ) or	[30 days following the date of completion of the after all animals of susceptible species have been			tion of the premises	
	II.1.4.6.		se of rabies, 30 days following the last case and thon of the premises;	ne date	e of completion of	of the cleansing and	
	II.1.4.7.		se of anthrax, 15 days following the last case and to on of the premises;	he dat	e of completion	of the cleansing and	
II.1.5.			wledge, during the period of 15 days prior to the d infected or suspected of an infectious or contagiou			mal has not been in	
II.2.	Attestation	of residen	ce and pre-export isolation				
( <sup>3</sup> ) either	[II.2.1.	holdings	period of at least 40 days prior to the date of di under veterinary supervision situated in the count which is assigned to Sanitary Group A, B, C, D, E of	try or p	part of the territo		

**EUROPEAN UNION** 

		Registered equidae, equidae for breeding	g and production equidae for slaughter
		II.a. Certificate reference number	II.b. Local reference number
(3)	either [in a Mem	ber State of the Union;]]	
(3)	for tempo country or required i	try or part of the territory of a country with Code: rary admission into the Union of registered horses, r part of the territory of the country of dispatch und n accordance with the Union legislation for the ten country or part of the territory of the country directly to	and from which it was imported into the ler conditions at least as strict as those mporary admission of registered horses
	(³) either	[assigned to the same Sanitary Groupterritory of the country of dispatch;]]]	(2) as the country or part of the
	(³) and/or	[assigned to Sanitary Group A, B or C;]]]	
	(³) andlor	[China ( <sup>5</sup> ), Hong Kong, Japan, Korea, Macao, Mala the United Arab Emirates;]]]	ysia (Peninsula), Singapore, Thailand or
( <sup>3</sup> ) ( <sup>4</sup> ) or [II.2	holdings u dispatch v of dispatc	period of at least 60 days prior to the date of dispunder veterinary supervision situated in the country which is assigned to Sanitary Group F, or was import h from a Member State of the Union before entering station in accordance with point II.2.2;]	or part of the territory of the country of rted during the 60 days prior to the date
(3) (4) either [II.2		ll is dispatched from a country or part of the terri Group E and	itory of a country which is assigned to
(3)	from vector country of point II.2.	kept in isolation in the country or part of the territor insects for a period of at least 40 days prior to the r part of the territory of the country of dispatch, I from a Member State of the Union or a country or to Sanitary Group A, B, C, D, E or G;]]	e date of dispatch, or since entry into the if it was imported in accordance with
(3)	40 days p country of Union or a E or G, an officially f	kept in designated premises under official vetering rior to the date of dispatch, or since entry into the fidispatch, if it was imported in accordance with paracountry or part of the territory of a country which is not the country or part of the territory of the country ree of African horse sickness and is not adjacently as occurred during the period of 2 years prior to the	e country or part of the territory of the coint II.2.1 from a Member State of the sassigned to Sanitary Group A, B, C, D, of dispatch is recognised by the OIE as at to a country in which African horse
( <sup>3</sup> ) ( <sup>4</sup> ) or [II.2		I is dispatched from a country or part of the territory nd was kept:	of country which is assigned to Sanitary
(3)	station) from vector-pro exercise repellents stables, a	during at least the last 40 days particles of the last from two hours prior to so was provided under official veterinary supervision in combination with an insecticide effective against and in strict isolation from equidae not being prepare equired for the temporary admission or imports into the	prior to the date of dispatch (insert date), confined to the sunset until two hours after sunrise and on, following the application of insect Culicoides prior to the removal from the d for export under conditions at least as
(3)	( <i>insert na</i> dispatch a	ntly confined in the approved vector-proof quarantine me of quarantine station) during the period of a ind constant monitoring of the vector protection ha stected part of the quarantine station.]]	at least 14 days prior to the date of
II.3. Att	estation of vaccina	tion and health tests	
(3) either [II.3		al was not vaccinated against African horse sicknes ation suggesting previous vaccination;]	s in the country of dispatch and there is

II.a. Certificate reference number

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[II.3.1.

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(3) or

# Registered equidae, equidae for breeding and production equidae for slaughter

The animal was vaccinated against African horse sickness, and this vaccination was carried out:

II.b. Local reference number

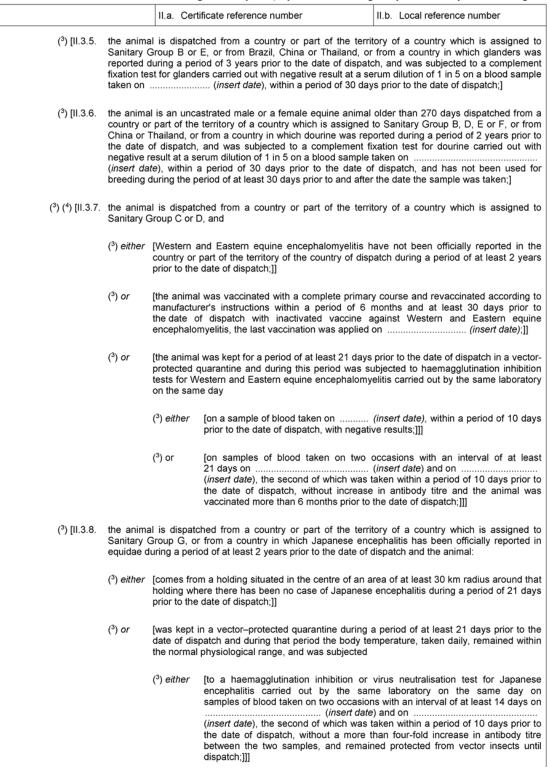
	(²) either	[more than 12 months prior to the date of dispatch;]]			
	(²) or	[more than 60 days and less than 12 months prior to the date of admission into the part of the territory of the country referred to in point II.1.3.(a), from where it is dispatched;]]			
(³) (⁴) or	[II.3.1.	The animal is dispatched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

# EUROPEAN UNION

			II.a. Certificate reference number	II.b. Local reference number	
	(³) or		d on a blood sample taken on (insert date, patch, by virus neutralisation test for EVA with negati		
	( <sup>3</sup> ) or	of 21 days	d on an aliquot of its entire semen taken ons s prior to the date of dispatch, by virus isolation tes PCR for EVA with negative result;]]		
	(³) or	and re-va	nated against EVA on (insert date) ccinated at regular intervals according to the manuby the competent authority, and the initial vaccination	ufacturer's instructions, with a vaccine	
			[before 31 December 2017, on the day a blood sar tested in a virus neutralisation test for EVA with of 1 in 4;]]]		
			[before 31 December 2017, during a period of isol official veterinary supervision, commencing on the was tested during that isolation period in a virus ne result at a serum dilution of 1 in 4;]]]	day a blood sample was taken which	
		, ,	[at the age of 180 to 270 days, during a period supervision, during which the animal was subjected carried out with negative result at a serum dilution oby the same laboratory with stable or declining titre 10 days apart;]]]	d to a virus neutralisation test for EVA f 1 in 4, or carried out on the same day	
		.,	[after the animal was subjected to a virus neutralisa a serum dilution of 1 in 4, carried out on a blood sar commencing a period of uninterrupted isolation vaccination;]]]	mple taken not earlier than 7 days after	
		, ,	[at the age of 180 to 250 days, after the animal was for EVA carried out with negative result at a serum same day by the same laboratory with stable or decl at least 14 days apart;]]]	dilution of 1 in 4, or carried out on the	
	( <sup>3</sup> ) or	[was subjected to a virus isolation test, polymerase chain reaction (PCR) or real-time PCR from carried out with negative result on an aliquot of its entire semen collected after the date a sample of that animal taken on (insert date), within a period of 6 months prior to the of dispatch, was tested in a virus neutralisation test for EVA with positive result at a serum dilutat least 1 in 4;]]			
	( <sup>3</sup> ) or	legislation animal is t legal act a that any l	irements for testing for EVA or vaccination again(insert reference to the applicable emporarily admitted into the Union for participation ind that the animal is kept separated from other equipreeding activity, including the collection of seme in the Union;]]	Union legal act) on the ground that the n the equestrian event specified in that dae not participating in such event and	
( <sup>3</sup> ) ( <sup>4</sup> ) either	[II.3.4.	anaemia,	I is dispatched from Iceland, which is certified as where it was continuously resident since birth, and a e entered Iceland from other countries;]		
( <sup>3</sup> ) or	[II.3.4.	Coggins to	al was subjected with negative result to an aga est) or to an ELISA for equine infectious anaemia c (insert date), this being within		
		(³) either	[a period of 90 days prior to the date of dispatch;]]		
		(3) or	[a period of 30 days prior to the date of dispatch fro	m a country or part of the territory of a	

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#### **EUROPEAN UNION**



# **EUROPEAN UNION**

	1,00	gistered eq	uluae, equiuae i	or breeding and	production eq	juidae for slaughte
	II.a. Certit	ficate refere	nce number	II.b. Local refer	ence number	
	( <sup>3</sup> ) or	encephali not earlier	tis virus with neg	gative result, car the date the iso	ried out on a b lation commence	s against Japanese blood sample taken ed on til dispatch;]]]
	( <sup>3</sup> ) or	and reva	ccinated accordi not less than 21	ng to manufact	urer's recomme	elete primary course endations during a ths prior to the date
Sanitary	nal is dispatched from a country or part of the territory of a country which is assigned of Group E, and was subjected to a serological test for African horse sickness as described to Directive 2009/156/EC, which was carried out by the same laboratory on the same day		ess as described in			
(³) either	[on blood s	samples tak	en on two occas	ons with an inte	rval of between	21 and 30 days, on
			(insert date) an aken within a per			(insert date), the f dispatch:
	(³) either	[with nega	ative results in ea	ch case.]]]		
	(3) or	[with a po	sitive result in the	first sample, and	t	
		(³) either		fication test as		th negative result in nex IV to Directive
		( <sup>3</sup> ) or	in antibody titr point 2.4 of C	e in a virus n	eutralisation tes f the OIE Terr	a two-fold increase st as described in restrial Manual for
( <sup>3</sup> ) or	prior to the dispatch is adjacent to	e date of one of the control of the country	dispatch, and the d by the OIE as	country or par officially free of	t of the territory African horse s	period of 21 days y of the country of sickness and is not luring the period of
	al is dispato Group F, an		a country or part	of the territory	of a country wi	hich is assigned to
(³) either	Directive 2 day on blo on not taken	2009/156/E od samples (in less than	C, which was c taken on two oc sert date) and on	arried out by the casions with an incommendation into the case of	ne same labora nterval of betwe (insert date) e vector-protect	ribed in Annex IV to atory on the same en 21 and 30 days, the first sample ted quarantine, the atch,
	(³) either	[with nega	ative results in ea	ch case.]]]		
	(3) or	[with a po	sitive result in the	first sample, and	t	
		(³) either		fication test as		th negative result in nex IV to Directive
		( <sup>3</sup> ) or	in antibody titr point 2.4 of C	e in a virus n	eutralisation tes f the OIE Terr	a two-fold increase st as described in restrial Manual for

# EUROPEAN UNION

Box I.23.:

#### Registered equidae, equidae for breeding and production equidae for slaughter

EUROPEAN UNION			Registered equidae, equidae for breeding and production equidae for slaughter					
			II.a. Certificate reference number	II.b. Local reference number				
		( <sup>3</sup> ) or	[was subjected to a serological and an agent identific described in Annex IV to Directive 2009/156/EC, c case on a blood sample taken on	arried out with negative result in each (insert date) not less than				
		( <sup>3</sup> ) or	[was subjected to an agent identification test for A Annex IV to Directive 2009/156/EC, carried out witaken on	th negative result on a blood sample less than 14 days after the date of				
II.4.	Attestatio	n of the tran	sport conditions					
( <sup>3</sup> ) ( <sup>4</sup> ) either	[II.4.1.	Sanitary Union, w	nal is dispatched from a country or part of the territ Group A, B, C, D, E or G and arrangements have be ithout passing through a market, marshalling or ass ith other equidae of a different health status.]	een made to transport it directly to the				
(³) (⁴) or	[II.4.1.	Sanitary quarantin	nal is dispatched from a country or part of the territ Group F and arrangements have been made to transpers estation without coming into contact with other estation into the	port it directly from the vector-protected quidae not accompanied by a health				
		(²) either	[to the airport under vector-protected conditions and the aircraft being cleansed and disinfected in a recognised in the third country of dispatch, and spratake off.]]	advance with a disinfectant officially				
		( <sup>3</sup> ) or	[to a sea port in that country or part of the territory conditions and arrangements have been made to scheduled directly to a port in the Union without cal part of the territory of a country not approved for to stalls which were cleansed and disinfected in recognised in the third country of dispatch and spradeparture.]]	to transport it on a vessel which is ling into a port situated in a country or he entry into the Union of equidae, in advance with a disinfectant officially				
	II.4.2.	with at le	nents have been made and verified to prevent any co last the same health requirements as described in t fication until dispatch to the Union.					
	II.4.3.	disinfecte	sport vehicles or containers in which the animal is and before loading with a disinfectant officially recognises constructed that faeces, urine, litter or fodder cannot	sed in the third country of dispatch and				
II.5.	Attestatio	n of animal	welfare					
			d in Box I.28 was examined today (1) and found fit to ere made to protect its health and well-being effectively					
Notes:								
Part I:								
Box I.8.:			the country or part of the territory of the country of on Implementing Regulation (EU) 2018/659.	dispatch as appearing in column 3 of				
Box I.15.:	informatio		(railway wagons or container and lorries), flight rovided. In case of unloading and reloading, the consi Union.					

The container number and the seal number (if applicable) should be included.

#### **EUROPEAN UNION**

# Registered equidae, equidae for breeding and production equidae for slaughter

II.a. Certificate reference number

II.b. Local reference number

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

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:

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

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

- (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) 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.
- (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.

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 consignee;
- (c) accompany the registered horse in the original throughout its temporary admission in the Union;
- (d) be signed and stamped in a colour different to the colour of the printing;
- (e) 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:					

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	Declaration by the owner or representative of the owner for the temporary admission of a registered horse											
Iden	tification o	f the	animal (1)									
Spe	Species (Scientific name) Identification system Identification number Age Sex											
	Equus caballus											
I, the undersigned owner (2) or representative of the owner (2) of the registered horse described above, hereby declare, that:												
_	the horse											
	(²) either			in( <i>inse</i> ys prior to the date		country or part of the	e territory of	a country	y of dispatch	n) during a p	period of	
	(²) or					country or part of prior to the date of d		ry of a c	country of d	ispatch) du	iring the	
		(a)				n country of dispatch)		name o	of country	from wher	e horse	
		(b)				n country of dispatch)		name o	of country	from wher	e horse	
		(c)				n country of dispatch)		name o	of country	from wher	e horse	
_				ays prior to the date		the horse has not b	een in cont	act with a	nimals suffe	ering from ir	nfectious	
_	•	oorta	tion will be	·		alth and well-being o	of the horse	can be p	rotected effe	ectively at a	ll stages	
_	the condi	tions	for reside			s applicable in acco		h point II	.2 of the ac	companyin	g health	
_	the condi	tions	for the tra		e in accorda	nce with point II.4 o	,	panying	health certifi	cate for the	country	
_		resi	,	, ,		less than 90 days	the horse	will be a	ccommodate	ed on the t	following	
	(a) from		(d	ate) to	(date) i	n	. (place of h	oldina) in	١	(Membe	r State)	
				,	, ,	n				,	,	
						n						
			•	*	, ,	n				•	,	
_	I am aware that in the event that the horse moves from one Member State of the Union to another Member State, as outlined in this declaration, it must be accompanied by a health certificate issued by an official veterinarian of the Member State of dispatch and that this movement must be notified to the Member State of destination;											
_	the horse is scheduled to leave the Union on											
Name and address of the owner (²) or representative (²):												
Date	e:			(dd/mm/yyyy)	)							
(1)	Article 2(b) of Commission Implementing Regulation (EU) 2018/659. Specify the identification system (such as ear tag, tattoo, brand, transponder) and the anatomic place used on the animal.  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).											
(2)	Sex (M = male, F = female, C = castrated).  (2) Delete as appropriate.											

# Section B

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

COUN	TRY:									V	eterinary	certifica	te to EU		
	l.1.	Consignor Name					1.2.	Certificate referen	ice No	1.3	2.a.				
		Address					I.3. Central competent authority								
_		Tel.					1.4.	Local competent a	authority						
Part I : Details of dispatched consignment	1.5.	Consignee Name Address					1.6.	Person responsib Name Address	le for the load	in El	J				
atched		Postcode Tel.					Postcode Tel.								
s of disp	1.7.	Country of origin	ISO code	1.8.	Region of origin	Code	1.9.	Country of destination	ISO code	I.10.	Region of destinatio	n	Code		
Detail	1.11.	Place of orig	l ain				I.12.								
Part I:		Name Approval number Address													
	I.13.	. Place of loading						Date of departure							
	I.15.	Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification Documentary references					I.16.	Entry BIP in EU							
							l.17.	No(s) of CITES							
	I.18.	Description	of animals						I.19. Commo	dity o	ode (HS c	ode)			
								,		1.20	. Quantity				
	I.21.									1.22	. Number	of packa	ges		
	1.23.	Seal/Contain	ner No							1.24					
	1.25.	25. Animals certified for:  Registered equidae □ breeding and						uction $\square$		sla	ughter				
	1.26.	For transit th	nrough EU to	third	country	Χ		1.27.							
	Third country ISO code														
	1.28.	Identification	n of the anima	als											
	S	pecies (Scie name)	ntific Id	entifi	Species (Scientific Identification system Identification number Age Sex name)										

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			II.a. Certificate reference number	II.b. Local reference number								
	II. Attestation of animal health and welfare											
	I, the under	I, the undersigned official veterinarian, hereby certify, that the equine animal described in Box I.28:										
	<ul> <li>was examined today (¹) and found free of clinical signs of diseases and of obvious signs of ectoparasite infestation;</li> </ul>											
cation	_	<ul> <li>is not intended for slaughter under a national programme of infectious or contagious disease eradication;</li> </ul>										
Part II: Certification	_	<ul> <li>meets the requirements attested in points II.1 to II.5 of this certificate;</li> </ul>										
art II:	_	<ul> <li>is accompanied by the written declaration, signed by the owner of the animal or the representative of the owner.</li> </ul>										
_	II.1.	II.1. Attestation on third country or part of the territory of third country and holding of dispatch										
	II.1.1.	I.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.	3. the animal is dispatched from a country or part of the territory of a country										
		in which th of African	onsidered free from African horse sickness in accord lere has been no clinical, serological (in unvaccinated horse sickness during the period of 2 years prior to the no vaccinations against the disease during the per	equidae) or epidemiological evidence ne date of dispatch and in which there								
		b) in which Venezuelan equine encephalomyelitis has not occurred during the period of 2 years prior the date of dispatch;										
		c) in which dourine has not occurred during the period of 6 months prior to the date of dispatch;										
		d) in which gl	anders has not occurred during the period of 6 month	s prior to the date of dispatch;								
	(3) either	[e) in which with dispatch;]	resicular stomatitis has not occurred during the per	riod of 6 months prior to the date of								
	(³) or	(3) or [e) in which vesicular stomatitis has occurred during the period of 6 months prior to the date of dis and a blood sample taken from the animal on										
		(³) either	[in a virus neutralisation test at a serum dilution of 1 in	n 32;]]								
			[in an ELISA in accordance with the relevant Chapt and Vaccines for Terrestrial Animals of the OIE;]]	ter of the Manual of Diagnostic Tests								
	II.1.4.	points II.1.4.1 to II.1.4	come from a holding, and to the best of my knowled .7 was not in contact with animals from holdings, who do in points II.1.4.1 to II.1.4.7 and which last for:									

EUROPEAN U	NION		Registered equidae, equidae for breeding and production equidae for slaughter					
			II.a. Certificate reference number II.b. Local reference number					
	II.1.4.1.	I.1.4.1. in the case of equidae suspected of having contracted dourine,						
		(3) either [6 months beginning on the date of the last actual or possible contact with an anim suspected of having contracted dourine or infected with Trypanosoma equiperdum;]						
		(3) or	[in the case of a stallion, until the animal is castrated;]					
		(³) or	[30 days following the date of completion of the cleansing and disinfection of the premises after all animals of susceptible species have been slaughtered;]					
	II.1.4.2.	in the cas	ase of glanders,					
		( <sup>3</sup> ) either	6 months beginning on the day on which the equidae suffering from the disease or subjected with positive result to a test for the detection of the causative pathogen surkholderia mallei or antibodies to that pathogen, were killed and destroyed;]					
		( <sup>3</sup> ) or	30 days following the date of completion of the cleansing and disinfection of the premises fter all animals of susceptible species have been killed and destroyed;]					
	II.1.4.3.	in the cas	case of equine encephalomyelitis of any type,					
		( <sup>3</sup> ) either	either [6 months beginning on the day on which the equidae suffering from the disease have been slaughtered;]					
		(³) or	[6 months beginning on the day on which the equidae infected with the virus causing Wes Nile Fever, Eastern equine encephalomyelitis or Western equine encephalomyelitis have died, been removed from the holding or fully recovered;]					
		( <sup>3</sup> ) 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.	slaughter immunodi	the case of equine infectious anaemia, until the date on which, the infected animals having been ughtered, the remaining animals on the holding have shown a negative reaction in an agar gel munodiffusion test (AGID or Coggins test) carried out on blood samples collected on two occasions nonths apart;					
	II.1.4.5.	in the cas	e of vesicular stomatitis,					
		(³) either	[6 months following the last case;]					
		( <sup>3</sup> ) 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.		se of rabies, 30 days following the last case and the date of completion of the cleansing and on of the premises;					
	II.1.4.7.		ee of anthrax, 15 days following the last case and the date of completion of the cleansing and on of the premises;					
II.1.5.			wledge, during the period of 15 days prior to the date of dispatch the animal has not been in infected or suspected of an infectious or contagious disease.					
II.2.	Attestation	n of residen	ce and pre-export isolation					
(3) either	[II.2.1.	holdings	period of at least 40 days prior to the date of dispatch, the animal has been resident on under veterinary supervision situated in a country or part of the territory of a country of which is assigned to Sanitary Group A, B, C, D, E or G and					

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# **EUROPEAN UNION**

			II.a. Certificate reference number	II.b. Local reference number
	(3) either	[in a Memb	er State of the Union;]]	
	( <sup>3</sup> ) and/or	for tempora country or required in	y or part of the territory of country with Code: ary admission into the Union of registered horse part of the territory of the country of dispatch u accordance with the Union legislation for the to puntry or part of the territory of the country directle	s, and from which it was imported into the nder conditions at least as strict as those emporary admission of registered horses
			assigned to the same Sanitary Grouphe territory of the country of dispatch;]]]	(²) as the country or part of
		(3) and/or [	assigned to Sanitary Group A, B or C;]]]	
			assigned to Sanitary Group D, E or G and the Article 2(c) of Commission Implementing Regulat	
( <sup>3</sup> ) ( <sup>4</sup> ) or	[II.2.1.	holdings un dispatch who of dispatch	eriod of at least 60 days prior to the date of d nder veterinary supervision situated in a count nich is assigned to Sanitary Group F, or was imp from a Member State of the Union before ent station in accordance with point II.2.2;]	ry or part of the territory of a country of corted during the 60 days prior to the date
( <sup>3</sup> ) ( <sup>4</sup> ) either	[11.2.2.	the animal Sanitary Gr	is dispatched from a country or part of the teroup E and	rritory of a country which is assigned to
	(3) either	from vector country or point II.2.1	kept in isolation in the country or part of the ter insects for a period of at least 40 days prior to part of the territory of the country of dispatc from a Member State of the Union or a country Sanitary Group A, B, C, D, E or G;]]	he date of dispatch, or since entry into the n, if it was imported in accordance with
	( <sup>3</sup> ) or	40 days pr country of Union or a E or G, and officially fre	kept in designated premises under official vete ior to the date of dispatch, or since entry into dispatch, if it was imported in accordance with country or part of the territory of a country which d the country or part of the territory of the country ee of African horse sickness and is not adjact as occurred during the period of 2 years prior to t	the country or part of the territory of the point II.2.1 from a Member State of the is assigned to Sanitary Group A, B, C, D, y of dispatch is recognised by the OIE as ent to a country in which African horse
( <sup>3</sup> ) ( <sup>4</sup> ) or	[11.2.2.		is dispatched from a country or part of the te coup F and was kept	rritory of a country which is assigned to
	( <sup>3</sup> ) either	to sunset u following the Culicoides prepared for	soved vector-protected quarantine station of	spatch from
	(³) or	of quaranti monitoring	ly confined in the approved vector-proof quarant ne station) during the period of at least 14 days of the vector protection has proven absence of ine station.]]	prior to the date of dispatch and constant
II.3.	Attestation	of vaccination	on and health tests	
( <sup>3</sup> ) either	[II.3.1.		was not vaccinated against African horse sickn ion suggesting previous vaccination;]	ess in the country of dispatch and there is

# **EUROPEAN UNION**

EUROPEAN U	NION		Registered equidae, equidae for breeding	and production equidae for slaughter				
			II.a. Certificate reference number	II.b. Local reference number				
(3) or	[II.3.1.	The anim	al was vaccinated against African horse sickness, and	this vaccination was carried out				
	(³) either	[more tha	in 12 months prior to the date of dispatch;]]					
	(³) or		in 60 days and less than 12 months prior to the date o untry referred to in point II.1.3.(a), from where it is disp					
( <sup>3</sup> ) ( <sup>4</sup> ) or	ory of a country which is assigned to kness on							
	II.3.2.		al was not vaccinated against Venezuelan equine e prior to the date of dispatch from	encephalomyelitis during the period of				
	(³) either		y of which all parts of the territory are free of Vene at least 2 years prior to the date of dispatch;]	zuelan equine encephalomyelitis for a				
	( <sup>3</sup> ) ( <sup>4</sup> ) or	Venezuel	f the territory of a country which is assigned to Sanitary Group C or D which is free lan equine encephalomyelitis for a period of at least 2 years prior to the date of dispatch lan equine encephalomyelitis occurs in the remaining parts of the territory of the country and					
		( <sup>3</sup> ) either	[is vaccinated against Venezuelan equine encept course and revaccinated according to manufactur 60 days and no more than 12 months prior to the day protected quarantine for a period of at least 21 daying that period remained clinically healthy, an remained within the normal physiological range, a holding which showed a rise in body temperature, talfor virus isolation for Venezuelan equine encephalom	rer's recommendations not less than ate of dispatch, and was kept in vectorays prior to the date of dispatch, and dits body temperature, taken daily, and any equine animal on the same ken daily, was subjected to a blood test				
		( <sup>3</sup> ) or	[is not vaccinated against Venezuelan equine ence protected quarantine for a period of at least 21 d. clinically healthy, and its body temperature, take physiological range, and any equine animal on the body temperature, taken daily, was subjected to Venezuelan equine encephalomyelitis with nega dispatched was subjected to a diagnostic test for with negative result conducted on a sample taken rentry into vector-protected quarantine and remained dispatch;]]	ays, and during that period remained en daily, remained within the normal same holding which showed a rise in a blood test for virus isolation for tive results, and the animal to be Venezuelan equine encephalomyelitis not less than 14 days after the date of				
		( <sup>3</sup> ) or	[was subjected to a haemagglutination inhib encephalomyelitis carried out by the same laborator on two occasions with an interval of 21 days on	ry on the same day on samples taken				
( <sup>3</sup> ) ( <sup>4</sup> ) either	[11.3.3.	anaemia,	al is dispatched from Iceland, which is certified as where it was continuously resident since birth and o we entered Iceland from other countries;]					

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# EUROPEAN UNION

EUROPEA	AN UNION	Registered equidae, equidae for breeding and production equidae f						
			II.a. Ce	rtificate reference number	II.b. Local reference number			
(3) or	[II.3.3.	the animal was subjected with negative result to an agar gel immunodiffusion test (AG Coggins test) or to an ELISA for equine infectious anaemia carried out on a blood sample take (insert date), this being within						
		(³) either	[a period o	of 90 days prior to the date of dispatch;]]				
		(³) or		of 30 days prior to the date of dispatch from nich is assigned to Sanitary Group D, E or F				
	(³) [II.3.4.	(3) [II.3.4. the animal is dispatched from a country or part of the territory of a country which is assigned Sanitary Group B or E, or from Brazil, China or Thailand, or from a country in which glander reported during a period of 3 years prior to the date of dispatch, and was subjected to a comp fixation test for glanders carried out with negative result at a serum dilution of 1 in 5 on a sample taken on						
	(3) (4) [II.3.5. the animal is dispatched from a country or part of the territory of a country which Sanitary Group C or D, and  (3) either [Western and Eastern equine encephalomyelitis have not been officially							
		(³) either	country or	and Eastern equine encephalomyelitis ha part of the territory of the country of dispa e date of dispatch;]]				
		( <sup>3</sup> ) or	manufactu date of	Il was vaccinated with a complete primary rer's instructions within a period of 6 mor dispatch with inactivated vaccine aga myelitis, the last vaccination was applied or	nths and at least 30 days prior to the inst Western and Eastern equine			
		( <sup>3</sup> ) or	protected	Il was kept for a period of at least 21 days p quarantine and during this period was sub /estern and Eastern equine encephalomye ne day	ejected to haemagglutination inhibition			
			(³) either	[on a sample of blood taken onperiod of 10 days prior to the date of disp				
			( <sup>3</sup> ) or	[on samples of blood taken on two of 21 days on	e) and on			
	( <sup>3</sup> ) [II.3.6.	ory of a country which is assigned to phalitis has been officially reported in spatch, and the animal						
		(³) either	holding wh	om a holding situated in the centre of an are here there has been no case of Japanese e e date of dispatch;]]				
		(³) or	date of dis	in a vector-protected quarantine during a patch, and during that period the body tem I physiological range, and was subjected				
			( <sup>3</sup> ) either	[to a haemagglutination inhibition Japanese encephalitis carried out by the samples of blood taken on two occasions	same laboratory on the same day on with an interval of at least 14 days on (insert date), the eriod of 10 days prior to the date of ncrease in antibody titre between the			
l								

# **EUROPEAN UNION**

			II.a. Cer	tificate refe	rence number	II.b. Local reference number			
			( <sup>3</sup> ) or	Japanese sample ta	encephalitis virus with nega ken not earlier than 7 days after	he detection of antibodies against tive result, carried out on a blood the date the isolation commenced on d remained protected from vector			
		( <sup>3</sup> ) or	revaccinate	ed accordin		ith a complete primary course and lations during a period of not less than e of dispatch;]]			
( <sup>3</sup> ) ( <sup>4</sup> ) either	[II.3.7.	Sanitary (	Group E and	is dispatched from a country or part of the territory of a country which is assigned to oup E and was subjected to a serological test for African horse sickness as described in Directive 2009/156/EC, which was carried out by the same laboratory on the same day					
		( <sup>3</sup> ) either	-	on blood samples taken on two occasions with an interval of between 21 and 30 days, on					
			(³) either	either [with negative results in each case.]]]					
			(³) or	[with a pos	sitive result in the first sample, a	and			
				( <sup>3</sup> ) either		equently tested with negative result in s described in Annex IV to Directive			
				( <sup>3</sup> ) or	in antibody titre in a virus	without more than a two-fold increase neutralisation test as described in of the OIE Terrestrial Manual for s.]]]]			
		( <sup>3</sup> ) or	prior to the dispatch is adjacent to	e date of or recognise a country or	dispatch, and the country or p d by the OIE as officially free	nsert date), within a period of 21 days part of the territory of the country of of African horse sickness and is not ry in which African horse sickness has of dispatch.]]			
(³) (⁴) or	[11.3.7.		al is dispato Group F, and		a country or part of the territo	ry of a country which is assigned to			
		( <sup>3</sup> ) either	[was subjected to a serological test for African horse sickness as described in Annex IV to Directive 2009/156/EC, which was carried out by the same laboratory on the same day on blood samples taken on two occasions with an interval of between 21 and 30 days, on						
			(3) either	[with nega	ative results in each case.]]]				
			(³) or	[with a pos	sitive result in the first sample, a	and			
				(3) either		equently tested with negative result in s described in Annex IV to Directive			
				( <sup>3</sup> ) or	in antibody titre in a virus	without more than a two-fold increase neutralisation test as described in of the OIE Terrestrial Manual for s.]]]]			

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# **EUROPEAN UNION** Registered equidae, equidae for breeding and production equidae for slaughter II.a. Certificate reference number II.b. Local reference number (3) or [was subjected to a serological and an agent identification test for African horse 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 ..... (insert date) not less than 28 days after the date of introduction into the vector-protected quarantine and within a period of 10 days prior to the date of dispatch.]] (3) or [was subjected to an agent identification test for African horse sickness as described in Annex IV to Directive 2009/156/EC, carried out with negative result on a blood sample taken on ...... (insert date) not less than 14 days after the date of introduction into the vector-protected quarantine and not more than 72 hours before dispatch.]] 11.4. Attestation of the transport conditions (3) (4) either [II.4.1. 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 arrangements have been made to transport it directly to the Union, without passing through a market, marshalling or assembly centre and without coming into contact with other equidae of a different health status.] (3) (4) or The animal is dispatched from a country or part of the territory of a country which is assigned to [1].4.1. Sanitary Group F and arrangements have been made to transport it directly from the vector-protected quarantine station without coming into contact with other equidae not accompanied by a health certificate either for imports or for temporary admission into the Union (3) either [to the airport under vector-protected conditions and the aircraft being cleansed and disinfected in advance with a disinfectant officially recognised in the third country of dispatch, and sprayed against vector insects just prior to take off.]] [to a sea port in that country or part of the territory of the country under vector-protected (3) or conditions and arrangements have been made to transport it on a vessel which is scheduled directly to a port in the Union without calling into a port situated in a country or part of the territory of a country not approved for the entry into the Union of equidae, in stalls which were cleansed and disinfected in advance with a disinfectant officially recognised in the third country of dispatch and sprayed against vector insects just prior to departure.11 11.4.2. Arrangements have been made and verified to prevent any contact with other equidae not complying with at least the same health requirements as described in this health certificate during the period from certification until dispatch to the Union. The transport vehicles or containers in which the animal is going to be loaded were cleaned and 11.4.3. 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. 11.4.4. The equine animal is proceeding to ...... (insert country of destination outside the Union). Arrangements have been made and the necessary animal health conditions certified to ensure that the animal transits the Union without delay. 11.5. 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.6.: Person responsible for the load in Union.

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#### **EUROPEAN UNION**

### Registered equidae, equidae for breeding and production equidae for slaughter

II.a. Certificate reference number

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

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

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

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

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

#### Part II:

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

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

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

This health certificate shall:

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

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

					owner or representative of the ogh the Union of an equine anim		
Iden	tification o	f the	animal (1)				
Spe	Species (Scientific name) Identification system Identification number Age Sex						
I, the	e undersigi	ned c	wner (²) o	r representative of the owne	r (²) of the animal described above	ve, hereby declare, that:	
_	the anima	al					
	(²) either			I in(insert name	of country or part of the territory ch;]	of a country of dispatch) d	uring a period of
	(²) or				e of country or part of the terrials prior to the date of dispatch:	tory of a country of dispa	atch) during the
		(a)	on entered	(insert date) country or part of the territor	from(inse y of country of dispatch)	ert name of country fron	n where animal
		(b)	on entered	(insert date) country or part of the territor	from(inse y of country of dispatch)	ert name of country fron	n where animal
		(c)		(insert date) country or part of the territor	from(inse y of country of dispatch);]	ert name of country fron	n where animal
_				days prior to the date of diseases transmissible to ed	dispatch the animal has not be juidae;	en in contact with animal	s suffering from
_					n as applicable in accordance v country of dispatch are fulfilled;	with point II.2 of the accor	mpanying health
_				nsport as applicable in according the country of dispatch are f	ordance with point II.4 of the accountilled;	ompanying health certificate	e for the country
_	the transpostages of			e effected in such a way th	nat health and well-being of the	animal can be protected	effectively at all
_				to leave the Union on of border post of exit);	(insert date) at t	the border post of	
Nan	ne and add	ress	of the own	ner (2) or representative (2):			
Date	<b>9</b> :			(dd/mm/yyyy)			
(¹)					Equus africanus, Equus hemionus, Ec	quus kiang, Equus quagga, Ec	guus 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 passpo Age: Date	ort acc	companies t h (dd/mm/y	he animal, its number should be yyy).	e stated and the name of the compete	nt authority which validated it.	
(2)	Sex (M = male, F = female, C = castrated).  Delete as appropriate.						

ANNEX I
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# PART 2

# Re-entry after temporary export

# Section A

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

COUN	NTRY: Veterinary certificate to EU						
	l.1.	Consignor Name	1.2.	Certificate reference No	I.2.a.		
		Address	1.3.	Central competent authority			
ŧ		Tel.	1.4.	Local competent authority			
Part I : Details of dispatched consignment	1.5.	Consignee Name Address	1.6.				
atched		Postcode Tel.					
Is of disp	1.7.	Country of ISO code I.8. Region of Code origin Code	1.9.	Country of ISO code destination	.10. Region of Code destination		
: Detai	l.11.	Place of origin	I.12.	Place of destination			
Partl		Name Approval number Address		Name Address			
				Postcode			
	I.13.	Place of loading	1.14.	Date of departure			
	I.15.	Means of transport	I.16.	Entry BIP in EU			
		Aeroplane Ship Railway wagon Road vehicle Other Identification  Documentary references		No(s) of CITES			
	I.18.	Description of animal		I.19. Commo	dity code (HS code) 01 01		
					I.20. Quantity 1		
	I.21.				I.22. Number of packages		
	1.23.	Seal/Container No			1.24.		
	1.25.	Animal certified for:					
		Registered horse					
	1.26.			I.27. For import or admission int	o EU 🔲		
	1.28.	Identification of the animal					
	Species (Scientific Identification system Identification number Age Sex name)  Equus caballus						

### **EUROPEAN UNION**

				· · · · · · · · · · · · · · · · · · ·			
			II.a. Certificate reference number	II.b. Local reference number			
	II.	Attestation of anima	al health and welfare				
	I, the undersigned official veterinarian, hereby certify, that the animal described in Box I.28:						
	_	is a registered horse	Regulation (EU) 2018/659;				
u <sub>o</sub>	_	was examined today infestation;	y (1) and found free of clinical signs of diseases a	and of obvious signs of ectoparasite			
Part II: Certification	_	is not intended for sla	ughter under a national programme of infectious or co	entagious disease eradication;			
E. Ce	_	meets the requirement	nts attested in points II.1 to II.3 of this certificate;				
Part	_	is accompanied by th	e written declaration, signed by the owner of the horse	or the representative of the owner.			
	II.1.	Attestation on third co	ountry or part of the territory of third country and holdin	g of dispatch			
	II.1.1.	or part of the territory	hed from(insert name of country or part of a country which on the date of issuing this certificat y Group(²);				
	II.1.2.	(Trypanosoma equip	spatch the following diseases are compulsorily notificerdum), glanders ( <i>Burkholderia mallei</i> ), equine encophalomyelitis), equine infectious anaemia, vesicula	ephalomyelitis (of all types including			
	II.1.3.	the animal is dispatch	ned from a country or part of the territory of a country:				
		which the African h	considered free from African horse sickness in accorda are has been no clinical, serological (in unvaccinated e orse sickness during the period of 2 years prior to the an no vaccinations against the disease during the pe	quidae) or epidemiological evidence on the date of dispatch and in which there			
			Venezuelan equine encephalomyelitis has not occurre of dispatch;	ed during the period of 2 years prior t			
		c) in which o	dourine has not occurred during the period of 6 months	s prior to the date of dispatch;			
		d) in which (	glanders has not occurred during the period of 6 month	ns prior to the date of dispatch;			
	II.1.4.	points II.1.4.1 to II.1.4	come from a holding, and to the best of my knowler 4.7 was not in contact with animals from holdings, whice to in points II.1.4.1 to II.1.4.7 and which last for:				
		II.1.4.1. in the cas	se of equidae suspected of having contracted dourine,				
		(³) either	[6 months beginning on the date of the last actual suspected of having contracted dourine or infected w				
		(³) or	[in the case of a stallion, until the animal is castrated;	]			
		(³) or	[30 days following the date of completion of the clea after all animals of susceptible species have been sla				
		(°) or					

EUROPEAN UNION	Registered equidae, equidae for breeding	and production equidae for slaughter
	II.a. Certificate reference number	II.b. Local reference number
II.1.4.2. in the	case of glanders,	
(³) eith	ther [6 months beginning on the day on which the end subjected with positive results to a test for the Burkholderia mallei or antibodies to that pathogen, w	detection of the causative pathogen
(³) or	[30 days following the date of completion of the clear after all animals of susceptible species have been kill	
II.1.4.3. in the	case of equine encephalomyelitis of any type,	
(³) eiti	ther [6 months beginning on the day on which the equidate slaughtered;]	e suffering from the disease have been
(³) or	[6 months beginning on the day on which the equida Nile Fever, Eastern equine encephalomyelitis or W died, been removed from the holding or fully recover	estern equine encephalomyelitis have
(³) or	[30 days following the date of completion of the clear after all animals of susceptible species have been sla	
slaugl gel ir	e case of equine infectious anaemia, until the date on whitered, the remaining equine animals on the holding have mmunodiffusion test (AGID or Coggins test) carried ccasions 3 months apart;	e shown a negative reaction in an agar
II.1.4.5. in the	case of vesicular stomatitis,	
(³) eiti	ther [6 months following the last case;]	
(³) or	[30 days following the date of completion of the clear after all animals of susceptible species have been sla	
	e case of rabies, 30 days following the last case and the dection of the premises;	date of completion of the cleansing and
	case of anthrax, 15 days following the last case and the dection of the premises;	date of completion of the cleansing and
	knowledge, during the period of 15 days prior to the date dae infected or suspected of an infectious or contagious di	
II.2. Attestation of resid	idence and pre-export isolation	
II.2.1. The animal was in	mported on (insert date)	
(3) either [direct	tly from the EU Member State (	insert name of EU Member State);]
	a country or part of the territory of a countryr conditions at least as strict as those set out in this certific	
part of the territor veterinary superv	from the Union less than 30 days ago, and since exit from ry of a country (1) other than those of the same Sanitary vision, accommodated in separated stables without comicept during racing, competition or the cultural event.	Group, and resident on holdings under
II.3. Attestation of anim	mal welfare	
	ribed in Box I.28 was examined today (¹) and found fit to I is were made to protect its health and well-being effectively	

Qualification and title:

Signature:

Status: This is the original version (as it was originally adopted).

### **EUROPEAN UNION**

Official veterinarian

Date:

Stamp:

Name (in capital letters):

### Registered equidae, equidae for breeding and production equidae for slaughter

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

Status: This is the original version (as it was originally adopted).

	Declaration by the owner or representative of the owner for the re-entry after temporary export of a registered horse for racing, competition and cultural events							
Iden	tification of	the animal (1)						
Spe	ecies (Scie	ntific name)	Identification system	Identification number	Age	Sex		
	Equus ca	aballus						
I, the	e undersigr	ned owner (2) or	representative of the owne	er (2) of the registered horse descri	bed above, hereby decla	are, 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 onrhere horse entered country	of dispatch);]	om	(insert name of		
_			ays prior to the date of disparansmissible to equidae;	atch the horse has not been in cor	ntact with animals sufferi	ing from infectious		
_	the transp of the jour		effected in such a way that	health and well-being of the horse	e can be protected effec	tively at all stages		
_				on as applicable in accordance we country of dispatch are fulfilled.	ith point II.2 of the acc	ompanying health		
Nam	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	of Commission or) and the anatom rt accompanies th of birth (dd/mm/yy nale, F = female, (	<ul> <li>Implementing Regulation (E nic place used on the animal.</li> <li>ne animal, its number should be ryy).</li> </ul>	identifier which permits to link the anir (U) 2018/659. Specify the identificat e stated and the name of the competen	ion system (such as ear	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

# Chapter 1

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

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

COUNT	JNTRY:										Veterinary cert	ificate to EU
	l.1.	Consignor Name					1.2.	Certificate refe	erence N	lo	I.2.a.	
		Address					I.3. Central competent authority					
_		Tel.			1.4.	Local compete	ent autho	ority				
neu	I.5. Consignee Name Address			1.6.								
consiç												
atched		Postcode Tel.										
Part I: Details of dispatched consignment	1.7.	Country of ISC origin	) code		Region of origin	Code	1.9.	Country of destination	ISO co	de I	I.10. Region of destination	Code
: Detai	l.11.	Place of origin					I.12.	Place of destir	nation			
Part		Name Approval number Address					Name Address					
								Postcode				
	I.13.	I.13. Place of loading					I.14. Date of departure					
	I.15.	Means of transpo	ort				I.16. Entry BIP in EU					
		Aeroplane 🗆	Ship		Railway wag	gon 🗖						
		Road vehicle  Identification Documentary refe		er 🗖			I.17.	No(s) of CITES	S			
	I.18.	Description of an	imal							I.19. (	Commodity code	(HS code)
											I.20. Quantity	
	I.21.										I.22. Number of	packages
	I.23. Seal/Container No										1.24.	
	1.25.	Animal certified for	or:									
		Registered horse										
	1.26.							I.27. For impor	rt or adm	nission	into EU	
	1.28.	Identification of th	he animal									
		Species Scientific name)	Iden	tificat	ion system	Identifica	tion n	umber	Age	•	S	эх

### **EUROPEAN UNION**

			II.a. Certificate reference number	II.b. Local reference number						
	II.	Attestation of anima	I health and welfare							
	I, the undersigned official veterinarian, hereby certify, that the animal described in Box I.28:									
	_	is a registered horse	as defined in Article 2(c) of Commission Implementing	Regulation (EU) 2018/659;						
	_	was examined today infestation;	r (1) and found free of clinical signs of diseases a	nd of obvious signs of ectoparasite						
ation	_	is not intended for sla	ughter under a national programme of infectious or cor	ntagious disease eradication;						
Part II: Certification	_	meets the requiremen	ats attested in points II.1 to II.3 of this certificate;							
art II: 0	is accompanied by the written declaration, signed by the owner of the horse, or the representative of the over									
ď	II.1.	Attestation on third co	ountry or part of the territory of third country and holding	g of dispatch						
	II.1.1.	The animal is dispatched from (insert name of country or part of the territory of a country), a country or part of the territory of a country which on the date of issuing this certificate has the Code:								
	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 dispatch	ed from a country or part of the territory of a country:							
		which the African h	onsidered free from African horse sickness in accordar re has been no clinical, serological (in unvaccinated ec orse sickness during the period of 2 years prior to the n no vaccinations against the disease during the per	quidae) or epidemiological evidence of e date of dispatch and in which there						
			/enezuelan equine encephalomyelitis has not occurred fidispatch;	d during the period of 2 years prior to						
		c) in which o	lourine has not occurred during the period of 6 months	prior to the date of dispatch;						
		d) in which (	landers has not occurred during the period of 6 months	s prior to the date of dispatch;						
	II.1.4.	points II.1.4.1 to II.1.4	come from a holding, and to the best of my knowled .7 was not in contact with animals from holdings, which o in points II.1.4.1 to II.1.4.7 and which last for:							
		II.1.4.1. in the cas	e of equidae suspected of having contracted dourine,							
		(³) either	[6 months beginning on the date of the last actual suspected of having contracted dourine or infected with							
		(³) or	[in the case of a stallion, until the animal is castrated;]							
		(³) or	[30 days following the date of completion of the clear after all animals of susceptible species have been slar							

### **EUROPEAN UNION**

			II.a. Certificate reference number	II.b. Local reference number
	II.1.4.2.	in the cas	e of glanders,	
		(³) either	[6 months beginning on the day on which the subjected with positive results to a test for the Burkholderia mallei or antibodies to that pathogen	ne detection of the causative pathogen
		(³) or	[30 days following the date of completion of the cafter all animals of susceptible species have been	
	II.1.4.3.	in the cas	e of equine encephalomyelitis of any type,	
		(³) either	[6 months beginning on the day on which the equi slaughtered;]	dae suffering from the disease have been
		( <sup>3</sup> ) or	[6 months beginning on the day on which the equ Nile Fever, Eastern equine encephalomyelitis or died, been removed from the holding or fully recov	Western equine encephalomyelitis have
		( <sup>3</sup> ) or	[30 days following the date of completion of the cafter all animals of susceptible species have been	
	II.1.4.4.	slaughter gel immu	se of equine infectious anaemia, until the date on ed, the remaining equine animals on the holding handiffusion test (AGID or Coggins test) carriedions 3 months apart;	ave shown a negative reaction in an agar
	II.1.4.5.	in the cas	e of vesicular stomatitis,	
		(³) either	[6 months following the last case;]	
		( <sup>3</sup> ) or	[30 days following the date of completion of the cafter all animals of susceptible species have been	
	II.1.4.6.		e of rabies, 30 days following the last case and then of the premises;	e date of completion of the cleansing and
	II.1.4.7.		e of anthrax, 15 days following the last case and then of the premises;	ne date of completion of the cleansing and
II.1.5.			wledge, during the period of 15 days prior to the da infected or suspected of an infectious or contagious	
II.2.	Attestation	n of residen	ce and pre-export isolation	
II.2.1.	The anima		rted into the country or part of the territory of the co	untry of dispatch on
	(³) either	[directly fr	om the EU Member State	(insert name of EU Member State);]
	(³) or		ountry or part of the territory of a country under conditions at least as strict as those set out in	

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### **EUROPEAN UNION**

(?) either [less than 30 days ago, and since evit from the Union was never in a country, or part of the territory or part of the strategy and resident on holdings under veterina supervision, accommodated in separated stables without coming into contact with equidace of low health status except during competition and has taken part in or was stabled together with hors participating in the LG Global Champions Tour  (?) either [in the Metropolitan area of Mexico City, Mexico:]]  (?) or [in Shanghai, China;]]  (?) or [in Shanghai, China;]]  (?) or [less than 60 days ago, and since exit from the Union was never in a country, or part of the territory a country (?) other than those of the same Sanitary Group, and resident on holdings under veterina supervision, accommodated in separated stables without coming into contact with equidace of health status except during competition and has taken part in or was stabled together with hors participating in  (?) or [the Asian Games in	EUROPEAN UNI	ON		Registered equidae, equidae for breeding	and production equidae for slaughte
a country (¹) other than those of the same Sanitary Group, and resident on holdings under veterins supervision, accommodated in separated stables without coming into contact with equidae of low health status except during competition and has taken part in or was stabled together with hors participating in the LG Global Champions Tour  (²) either [in the Metropolitan area of Mexico City, Mexico;]]  (²) or [in Shanghai, China;]]  (²) or [in Shanghai, China;]]  (²) or [less than 60 days ago, and since evit from the Union was never in a country, or part of the territory a country (¹) other than those of the same Sanitary Group, and resident on holdings under vetreting supervision, accommodated in separated stables without coming into contact with equidae of low health status except during competition and has taken part in or was stabled together with hore participating in  (²) either [the Asian Games in				II.a. Certificate reference number	II.b. Local reference number
(*) and/or [in Miami, Unites States of America;]  (*) or [in Shanghai, China;]]  (*) or [less than 60 days ago, and since exit from the Union was never in a country, or part of the territory a country (*) other than those of the same Sanitary Group, and resident on holdings under vertices supervision, accommodated in separated stables without coming into contact with equidae of low health status except during competition and has taken part in or was stabled together with hors participating in  (*) either [the Asian Games in	(	( <sup>3</sup> ) either	a country supervision health sta	(¹) other than those of the same Sanitary Group, and on, accommodated in separated stables without com- tus except during competition and has taken part in	resident on holdings under veterinary ing into contact with equidae of lower
(*) or [in Shanghai, China;]]  (*) or [less than 60 days ago, and since exit from the Union was never in a country, or part of the territory a country (*) other than those of the same Sanitary Group, and resident on holdings under veterins supervision, accommodated in separated stables without coming into contact with equidae of low health status except during competition and has taken part in or was stabled together with hors participating in  (*) either [the Asian Games in			(³) either	[in the Metropolitan area of Mexico City, Mexico;]]	
(*) or [less than 60 days ago, and since exit from the Union was never in a country, or part of the territory a country (*) other than those of the same Sanitary Group, and resident on holdings under veterins supervision, accommodated in separated stables without coming into contact with equidae of low health status except during competition and has taken part in or was stabled together with hors participating in  (*) either [the Asian Games in			(3) and/or	[in Miami, Unites States of America;]	
a country (¹) other than those of the same Sanitary Group, and resident on holdings under veterins supervision, accommodated in separated stables without coming into contact with equidae of low health status except during competition and has taken part in or was stabled together with hors participating in  (²) either [the Asian Games in			(3) or	[in Shanghai, China;]]	
(3) or [the American Games in	(	( <sup>3</sup> ) or	a country supervision health sta	(¹) other than those of the same Sanitary Group, and on, accommodated in separated stables without com- tus except during competition and has taken part in	fresident on holdings under veterinary ing into contact with equidae of lower
(3) or [the Endurance World Cup in United Arab Emirates.]]  (3) or [less than 90 days ago, and since exit from the Union was never in a country, or part of the territory a country (1) other than those of the same Sanitary Group, and resident on holdings under veterins supervision, accommodated in separated stables without coming into contact with equidae of low health status except during competition and has taken part in or was stabled together with hors participating in  (3) either [the Test event for the Olympic Games in			(3) either	[the Asian Games in	(insert place).]]
(3) or [less than 90 days ago, and since exit from the Union was never in a country, or part of the territory a country (1) other than those of the same Sanitary Group, and resident on holdings under veterins supervision, accommodated in separated stables without coming into contact with equidae of low health status except during competition and has taken part in or was stabled together with hors participating in  (3) either [the Test event for the Olympic Games in			(3) or	[the American Games in	(insert place).]]
a country (¹) other than those of the same Sanitary Group, and resident on holdings under vetering supervision, accommodated in separated stables without coming into contact with equidae of low health status except during competition and has taken part in or was stabled together with hors participating in  (³) either [the Test event for the Olympic Games in			(3) or	[the Endurance World Cup in United Arab Emirates.]	
(3) or [the Olympic Games in	(	(³) or	a country supervision health sta	(¹) other than those of the same Sanitary Group, and on, accommodated in separated stables without com- tus except during competition and has taken part in	fresident on holdings under veterinary ing into contact with equidae of lower
(3) or [the Paralympics in			(³) either	[the Test event for the Olympic Games in	(insert place).]]
(3) or [the World Equestrian Games in			(3) or	[the Olympic Games in	(insert place).]]
The animal described in Box I.28 was examined today (¹) and found fit to be transported on the intended journ 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 Commission Implementing Regulation (EU) 2018/659.  Box I.15.: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) as information is to be provided. In case of unloading and reloading, the consignor must inform the Border Inspection Post of entry into the Union.			(3) or	[the Paralympics in	(insert place).]]
The animal described in Box I.28 was examined today (¹) and found fit to be transported on the intended journand 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 Commission Implementing Regulation (EU) 2018/659.  Box I.15.: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) as information is to be provided. In case of unloading and reloading, the consignor must inform the Border Inspection of the Union.			(3) or	[the World Equestrian Games in	(insert place).]]
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 Commission Implementing Regulation (EU) 2018/659.  Box I.15.: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) as information is to be provided. In case of unloading and reloading, the consignor must inform the Border Inspection Post of entry into the Union.	II.3.	Attestation	of animal v	welfare	
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 Commission Implementing Regulation (EU) 2018/659.  Box I.15.: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) a 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.8.: Provide the code of the country or part of the territory of the country as appearing in column 3 of Annex I Commission Implementing Regulation (EU) 2018/659.  Box I.15.: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) as information is to be provided. In case of unloading and reloading, the consignor must inform the Border Inspection Post of entry into the Union.	Notes:				
Commission Implementing Regulation (EU) 2018/659.  Box I.15.: Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) as information is to be provided. In case of unloading and reloading, the consignor must inform the Border Inspection Post of entry into the Union.	Part I:				
information is to be provided. In case of unloading and reloading, the consignor must inform the Border inspection Post of entry into the Union.				, , , , , , , , , , , , , , , , , , , ,	appearing in column 3 of Annex I to
Box I.23.: The container number and the seal number (if applicable) should be included.	i	nformation	is to be pr	ovided. In case of unloading and reloading, the consi	
	Box I.23.:	The contair	ner number	r and the seal number (if applicable) should be include	ed.

EURO	PEAN UNION	Registered equidae, equidae for breeding a	ind production equidae for slaughter
		II.a. Certificate reference number	II.b. Local reference number
Box I	identification documen the identification syste	The animal must bear an individual identifier which it as defined in Article 2(b) of Commission Implemention (such as ear tag, tattoo, brand, transponder) and the companying passport must be stated and the narrow	ng Regulation (EU) 2018/659. Specify a natomic place used on the animal.
	Age: Date of birth (dd/	mm/yyyy).	
	Sex (M = male, F = fer	male, C = castrated).	
Part	II:		
( <sup>1</sup> )	The certificate must be issued the Member State of destination	on the day of loading or on the last working day befor on in the Union.	re loading of the animal for dispatch to
	to the date of authorisation for referred to in point II.1.1, or du	export of this registered horse shall not be allowed what re-entry into the Union from the respective country ouring a period where restrictive measures have been at this part of the territory of the country of dispatch.	the part of the territory of the country
( <sup>2</sup> )		f the territory of the country, and the Sanitary Group menting Regulation (EU) 2018/659.	as appearing in columns 3 and 5 of
(3)	Delete as appropriate.		
This	health certificate shall:		
(a)		guage understood by the certifying officer and one of the Member State where the registered horse will e	
(b)	be made out to a single consig	gnee;	
(c)	be signed and stamped in a co	olour different to the colour of the printing;	
(d)		paper or all sheets of paper required are part of an otal number of pages, and each page shall bear the c are stapled and stamped.	
Offici	al veterinarian		
	Name (in capital letters):		Qualification and title:
	Date:		Signature:
	Stamp:		

Status: This is the original version (as it was originally adopted).

	Declaration by the owner or representative of the owner for the re-entry after temporary export of a registered horse for competition								
Ider	ntification o	f the animal (1	)						
Sp	ecies (Scie	entific name)	Identification system	Identification number	Age	Sex			
	Equus caballus								
I, th	e undersig	ned owner (²)	or representative of the owne	er (2) of the registered horse descr	ibed above, hereby decl	are, that:			
-	the horse	•							
	(²) either		arily exported from the Union days (²) or 90 days (²) prior t	to the country of dispatch on o this declaration;]		(insert date)			
	(²) or		country of dispatch on where horse entered country	(insert date) from y of dispatch);]		(insert name of			
_	the horse	has been tem	porarily exported from the U	nion to take part in					
	(²) either	[the Asian Ga	ames in	(insert p	lace);]				
	(2) or	[the America	n Games in	(insert p	lace);]				
	(2) or	[the Enduran	ce World Cup in United Arab	Emirates;]					
	(2) or	[the Test eve	nt for the Olympic Games in	(insert p	lace);]				
	(2) or	[the Olympic	Games in	(insert p	lace);]				
	(2) or	[the Paralym	pics in	(insert p	lace);]				
	(2) or	[the World Ed	questrian Games in	(insert p	lace);]				
	(2) or	[the LG Glob	al Champions Tour in						
		(2) either [ti	he Metropolitan area of Mexi	ico City, Mexico;]					
		(2) and/or [N	liami, Unites States of Amer	rica;]					
		(3) or [S	Shanghai, China;]						
_			days prior to the date of disp transmissible to equidae;	eatch the horse has not been in co	ntact with animals suffer	ing from infectious			
_				on as applicable in accordance v e country of dispatch are fulfilled;	vith point II.2 of the acc	companying health			
_	the trans		e effected in such a way tha	t health and well-being of the hors	e can be protected effec	ctively at all stages			
Nan	ne and add	fress of the ow	ner (²) or representative (²):						
Date	ə:		(dd/mm/yyyy)						
(1)	Article 2(b) of Commission Implementing Regulation (EU) 2018/659. Specify the identification system (such as ear tag, tattoo, brand, transponder) and the anatomic place used on the animal.  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).								

# Chapter 2

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

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

COUNTRY: Veterinary certificate to EU Certificate reference No I.1. Consignor I.2.a. Name Address I.3. Central competent authority Tel. I.4. Local competent authority Part I: Details of dispatched consignment Consignee 1.6. Name Address Postcode Country of ISO code I.8. Region of Code ISO code Code 1.9. Country of I.10. Region of origin destination destination origin I.11. Place of origin I.12. Place of destination Approval number Name Name 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 🔲 Aeroplane Railway wagon  $\square$ Road vehicle Other  $\square$ 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 I.21. I.22. Number of packages I.23. Seal/Container No 1.24. I.25. Animal certified for: Registered horse 1.26. I.27. For import or admission into EU  $\Box$ I.28. Identification of the animal Species (Scientific Identification system Identification number Age Sex name) Equus caballus

EUROPEAN UNION				Registered equidae, equidae for bree	ding and production equidae for slaughter					
				II.a. Certificate reference number	II.b. Local reference number					
	II.	Attestation	n of anima	ıl health and welfare						
	I, the unders	signed officia	l veterinari	an, hereby certify, that the animal described in B	30x I.28:					
	_	is a registe	red horse	as defined in Article 2(c) of Commission Implement	enting Regulation (EU) 2018/659;					
Part II: Certification	_	was examined today (1) and found free of clinical signs of diseases and of obvious signs of ectoparasite infestation;								
: Certif	_	is not intended for slaughter under a national programme of infectious or contagious disease eradication;								
Part	_	meets the r	equiremer	nts attested in points II.1 to II.3 of this certificate;						
	_	is accompa	nied by th	e written declaration, signed by the owner of the	horse or the representative of the owner.					
	II.1.	Attestation	on country	or part of the territory of the country and holding	g of dispatch					
	II.1.1.	country), a	country	ned from(insert name of country or part of the territory of a country which at	the date of issuing this certificate has the					
	(Trypanosoma equipe			spatch the following diseases are compulsorily erdum), glanders ( <i>Burkholderia mallei</i> ), equinoncephalomyelitis), equine infectious anaemia, ve	e encephalomyelitis (of all types including					
	II.1.3.	the animal	is dispatch	ned from a country or part of the territory of a cou	intry:					
		a)	which the African he	considered free from African horse sickness in accordance with Directive 2009/156/EC and in ere has been no clinical, serological (in unvaccinated equidae) or epidemiological evidence of orse sickness during the period of 2 years prior to the date of dispatch and in which there en no vaccinations against the disease during the period of 12 months prior to the date of						
		b)		Venezuelan equine encephalomyelitis has not o of dispatch;	occurred during the period of 2 years prior to					
		c)	in which o	dourine has not occurred during the period of 6 n	nonths prior to the date of dispatch;					
		d)	in which o	glanders has not occurred during the period of 6	months prior to the date of dispatch;					
	II.1.4.	points II.1.4	1.1 to II.1.4	come from a holding, and to the best of my kr 1.7 was not in contact with animals from holdings to in points II.1.4.1 to II.1.4.7 and which last for:						
		II.1.4.1.	in the cas	se of equidae suspected of having contracted do	urine,					
			( <sup>3</sup> ) either	[6 months beginning on the date of the last suspected of having contracted dourine or infections.]						
			( <sup>3</sup> ) or	[in the case of a stallion, until the animal is cast	trated;]					
			( <sup>3</sup> ) or	[30 days following the date of completion of th after all animals of susceptible species have be						

UROPEAN	UNION	Registered equidae, equidae for breeding and production equidae for slaugh	
		II.a. Certificate reference number II.b. Local reference number	
	II.1.4.2.	in the case of glanders,	
		(3) either [6 months beginning on the day on which the equidae suffering from the disease of subjected with positive results to a test for the detection of the causative pathoge Burkholderia mallei or antibodies to that pathogen, were killed and destroyed;]	
		or [30 days following the date of completion of the cleansing and disinfection of the premis after all animals of susceptible species have been killed and destroyed;]	
	II.1.4.3.	the case of equine encephalomyelitis of any type,	
		(3) either [6 months beginning on the day on which the equidae suffering from the disease have bee slaughtered;]	
		(3) or [6 months beginning on the day on which the equidae infected with the virus causing We Nile Fever, Eastern equine encephalomyelitis or Western equine encephalomyelitis have died, been removed from the holding or fully recovered;]	
		(3) or [30 days following the date of completion of the cleansing and disinfection of the premise 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 bee slaughtered, the remaining equine animals on the holding have shown a negative reaction in an aggel immunodiffusion test (AGID or Coggins test) carried out on blood samples collected two occasions 3 months apart;	
	II.1.4.5.	in the case of vesicular stomatitis,	
		(3) either [6 months following the last case;]	
		(3) or [30 days following the date of completion of the cleansing and disinfection of the premise 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 cleansing ar disinfection of the premises; $\frac{1}{2}$	
	II.1.4.7.	in the case of anthrax, 15 days following the last case and the date of completion of the cleansing ardisinfection of the premises;	
II.1.5.		of my knowledge, during the period of 15 days prior to the date of dispatch the animal has not been a equidae infected or suspected of an infectious or contagious disease.	
1.2.	Attestation	of residence and pre-export isolation	
I.2.1.		al was imported into the country or part of the territory of the country of dispatch o	
	(³) either	[directly from the EU Member State	
		(³) either [The Japan Cup;]	
		(³) or [The Melbourne Cup;]	
		(³) or [The Dubai Racing World-Cup;]	
		(³) or [The Hong Kong International Races;]	

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### **EUROPEAN UNION**

### Registered equidae, equidae for breeding and production equidae for slaughter

II.b. Local reference number

( <sup>3</sup> ) or	[from Australia (³), Canada (³), the United States of America (³), Hong Kong (³), Japan (³), Singapore (³), United Arab Emirates (³) or Qatar (³) for the participation in International Group/Grade
	meetings in the country of dispatch;]

II.a. Certificate reference number

- II.2.2. as far as can be ascertained and based on the declaration of the owner of the horse or representative of the owner (3) accompanying this certificate, the animal was:
  - not continuously outside the Union for more than 90 days, the date of scheduled return in accordance with this certificate included:
  - not outside the country of dispatch or in case of International Group/Grade meetings outside Australia,
     Canada, the United States of America, Hong Kong, Japan, Singapore, the United Arab Emirates or Qatar;
  - resident on holdings under veterinary supervision, accommodated in separated stables without coming into contact with equidae of lower health status except during racing;
- II.2.3. the animal entered the country of dispatch under animal health conditions at least as strict as those laid down in this health certificate.
- II.3. Attestation of animal welfare

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

### Notes:

### Part I:

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

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

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

### Part II:

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

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

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

EUROPEAN UNION		Registered equidae, equidae for bree	ding and production equidae for slaughter			
		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	olour different to the colour of the printing;				
(d)	consist of a single sheet of paper or all sheets of paper required are part of an integrated whole and indivisible by inserting page numbers and total number of pages, and each page shall bear the certificate reference number at the top of the page and those pages are stapled and stamped.					
Offic	ial veterinarian					
	Name (in capital letters):		Qualification and title:			
	Date: Signature:					
	Stamp:					

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	Declaration by the owner or representative of the owner for the re-entry after temporary export of a registered horse for racing						
Ider	Identification of the animal (¹)						
Spe	Species (Scientific name) Identification system Identification number Age Sex						
	Equus caballus						
I, the	e undersig	ned owner (2) or representative of the owner (2) of the registered horse described above, hereby declare, that:					
_	the horse						
	(²) either	[was temporarily exported from the Union to the country of dispatch on					
	(²) or	[entered the country of dispatch on					
_	the horse	has been temporarily exported from the Union to take part in					
	(²) either	[The Japan Cup;]					
	(²) or	[The Melbourne Cup;]					
	(²) or	[The Dubai Racing World-Cup;]					
	(²) or	[The Hong Kong International Races;]					
	(²) or	[International Group/Grade meetings in Australia $(^2)$ , Canada $(^2)$ , the United States of America $(^2)$ , Hong Kong $(^2)$ , Japan $(^2)$ , Singapore $(^2)$ , United Arab Emirates $(^2)$ or Qatar $(^2)$ ;]					
_		period of 15 days prior to the date of dispatch the horse has not been in contact with animals suffering from infectious ous diseases transmissible to equidae;					
_	<ul> <li>the conditions for residence and pre-export isolation as applicable in accordance with point II.2 of the accompanying health certificate for the country or part of the territory of the country of dispatch are fulfilled;</li> </ul>						
_	<ul> <li>the transportation will be effected in such a way that health and well-being of the horse can be protected effectively at all stages of the journey.</li> </ul>						
Nan	Name and address of the owner (²) or representative (²):						
Date	Date: (dd/mm/yyyy)						
(1)	Article 2(b) of Commission Implementing Regulation (EU) 2018/659. Specify the identification system (such as ear tag, tattoo, brand, transponder) and the anatomic place used on the animal.						
	Age: Date	rt accompanies the animal, its number should be stated and the name of the competent authority which validated it. of birth (dd/mm/yyyy).					
(2)	Sex (M = male, F = female, C = castrated).  Delete as appropriate.						

# PART 3

# **Imports**

# Section A

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

name)

Status: This is the original version (as it was originally adopted).

COUNTRY: Veterinary certificate to EU I.2.a. I.1. Consignor 1.2. Certificate reference No Name 1.3. Central competent authority Address 1.4. Local competent authority Tel. Part I: Details of dispatched consignment 1.6. Consignee Name Address Postcode Tel. Country of ISO code 1.8. Region of Code 1.9. Country of ISO code I.10. Region of Code origin origin destination destination I.11. Place of origin I.12. Place of destination Approval number Name Name Address Address Postcode I.13. Place of loading I.14. Date of departure I.15. Means of transport I.16. Entry BIP in EU Aeroplane  $\square$ Ship 🗖 Railway wagon  $\square$ Road vehicle Other  $\square$ 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 1.23. 1.24. Seal/Container No I.25. Animal certified for: Registered horse registered equine animal breeding and production 1.26. I.27. For import or admission into EU I.28. Identification of the animal Species (Scientific Identification system Identification number Sex Age

### **EUROPEAN UNION**

			registered equidae, equidae for breedi	<del> </del>
			II.a. Certificate reference number	II.b. Local reference number
II. Attestation of anir		n of anima	al health and welfare	
I, the undersigned official veterina		al veterinar	ian, hereby certify, that the animal described in Box	(1.28:
— (1) either [is a reg			stered equine animal, other than horse, as defined i	in Article 2(c) of Directive 2009/156/EC;]
				mmission Implementing Regulation (EU)
— comes from a count			uine animal for breeding and production as defined	in Article 2(e) of Directive 2009/156/EC;]
				thorised for imports into the Union of the
<ul> <li>was examined toda infestation;</li> </ul>			y (2) and found free of clinical signs of disease	es and of obvious signs of ectoparasite
_	is not inter	nded for sla	aughter under a national programme of infectious o	r contagious disease eradication;
_	meets the	requireme	nts attested in points II.1 to II.5 of this certificate;	
_	is accomp	anied by th	e written declaration, signed by the owner of the ar	nimal 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.	country or part of the territory of a country, w			the date of issuing this certificate
(Trypanosoma equip		oma equip	perdum), glanders (Burkholderia mallei), equine	encephalomyelitis (of all types including
II.1.3.	the animal	is dispatch	ned from a country or part of the territory of country	
	a)	which the African h have bee	ere has been no clinical, serological (in unvaccinate orse sickness during the period of 2 years prior to en no vaccinations against the disease during the	ed equidae) or epidemiological evidence of o the date of dispatch and in which there
	b)			curred during the period of 2 years prior to
	c)	in which	dourine has not occurred during the period of 6 mor	nths prior to the date of dispatch;
	d)	in which	glanders has not occurred during the period of 6 mo	onths prior to the date of dispatch;
				period of 6 months prior to the date of
( <sup>1</sup> ) or	[e)	and a blo of 21 day	od sample taken from the animal onvs prior to the date of dispatch, was tested with ne	(insert date), within a period
		(1) either	[in a virus neutralisation test at a serum dilution of	f 1 in 32;]]
		(¹) or	[in an ELISA in accordance with the relevant C and Vaccines for Terrestrial Animals of the OIE;]]	hapter of the Manual of Diagnostic Tests
	, the unders	He undersigned official  (1) either (1) or  (1) or  (1) or  (2) comes fro category of cate	., the undersigned official veterinari  — (1) either [is a regis 2018/658  (1) or [is an equivalent of section of the country of equidates of the country of part of the country of part of the country of part of the country of the country of the country of part of the country	II. Attestation of animal health and welfare  I, the undersigned official veterinarian, hereby certify, that the animal described in Boy  — (1) either [is a registered equine animal, other than horse, as defined (1) or [is an equistered horse as defined in Article 2(c) of Col 2018/659.]  — (1) or [is an equine animal for breeding and production as defined — comes from a country or part of the territory of a country which is au category of equidae specified in the first indent above;  — was examined today (2) and found free of clinical signs of disease infestation;  — is not intended for slaughter under a national programme of infectious one 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 is dispatched from

Status: This is the original version (as it was originally adopted).

EUROPEAN	UNION		Registered equidae, equidae for breeding	ng and production equidae for slaughte
			II.a. Certificate reference number	II.b. Local reference number
points II.1.4.1 to II.1.4		.4.1 to II.1.4	come from a holding, and to the best of my know 4.7 was not in contact with animals from holdings, w to in points II.1.4.1 to II.1.4.7 and which last for:	
	II.1.4.1.	in the cas	se of equidae suspected of having contracted douring	ne,
		(¹) either	[6 months beginning on the date of the last ac suspected of having contracted dourine or infected	
		(¹) or	[in the case of a stallion, until the animal is castrate	ed;]
		(¹) or	[30 days following the date of completion of the cafter all animals of susceptible species have been	
	II.1.4.2.	in the cas	se of glanders,	
		(¹) either	[6 months beginning on the day on which the subjected with positive results to a test for th <i>Burkholderia mallei</i> or antibodies to that pathogen	ne detection of the causative pathogen
		(¹) or	[30 days following the date of completion of the cafter all animals of susceptible species have been	•
	II.1.4.3.	in the cas	se of equine encephalomyelitis of any type,	
	(¹) eithe		[6 months beginning on the day on which the equi slaughtered;]	dae suffering from the disease have been
		(¹) or	[6 months beginning on the day on which the equ Nile Fever, Eastern equine encephalomyelitis or died, been removed from the holding or fully recov	Western equine encephalomyelitis have
		(¹) or	[30 days following the date of completion of the cafter all animals of susceptible species have been	
	II.1.4.4.	slaughter gel immi	se of equine infectious anaemia, until the date on red, the remaining equine animals on the holding ha unodiffusion test (AGID or Coggins test) carrie sions 3 months apart;	ave shown a negative reaction in an agar
	II.1.4.5.	in the cas	se of vesicular stomatitis,	
		(¹) either	[6 months following the last case;]	
		(¹) or	[30 days following the date of completion of the cafter all animals of susceptible species have been	
	II.1.4.6.		se of rabies, 30 days following the last case and tho on of the premises;	e date of completion of the cleansing and
	II.1.4.7.		se of anthrax, 15 days following the last case and thon of the premises;	ne date of completion of the cleansing and
II.1.5.	to the bes	st of my kno	wledge, during the period of 15 days prior to the day	ate of dispatch the animal has not been in

to the best of my knowledge, during the period of 15 days prior to the date of dispatch the animal has not been in contact with equidae infected or suspected of an infectious or contagious disease. II.1.5.

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# EUROPEAN UNION

II.a. Certificate reference number	
ii.a. Certificate reference number	II.b. Local reference number
sidence and pre-export isolation	
n 90 days old, or since entry if the animal was imported of days prior to the date of dispatch, the animal has bee	directly from the Union during a period of en resident on holdings under veterinary
ispatch, it was kept in pre-export isolation under veterinar	
	ved isolation centre described as place of
either [during the period of at least 40 days prior to the d	ate of dispatch;]]
or [during the period of at least 30 days prior to the Emirates;]]	e date of dispatch from the United Arab
igned to Sanitary Group F, and during the period of at le since birth if the animal is less than 90 days old, it wa ervision and was kept during the period of at least 60 day if it was imported directly from the Union during the patch, in the part of the territory described in point II.1.3	east 90 days prior to the date of dispatch, as resident on holdings under veterinary ays prior to the date of dispatch, or since e period of 60 days prior to the date of which is considered free of African horse
quarantine station) during the period of at least 40 n	O days prior to the date of dispatch tate), confined to the vector-protected o hours after sunrise and exercise was the application of insect repellents in prior to the removal from the stables, and ort under conditions at least as strict as
ert name of quarantine station) during the period of at le constant monitoring of the vector protection has prove	east 14 days prior to the date of dispatch
accination and health tests	
	ess in the country of dispatch and there is
animal was vaccinated against African horse sickness, a	and this vaccination was carried out:
re than 12 months prior to the date of dispatch;]]	
itary Group F and was vaccinated against African horse s e) not more than 24 months and at least 40 days prior to rantine by administration of a registered vaccine according	sickness on (insert to the date of entry in the vector-protected ng to manufacturer's instructions which is
Durinhard assistance of the control	,

**EUROPEAN UNION** 

		II.a. Certificate reference number	II.b. Local reference number		
II.3.2.		al was not vaccinated against Venezuelan equine prior to the date of dispatch from	e encephalomyelitis during the period of		
(¹) either		a country of which all parts of the territory are free of Venezuelan equine encephalomy operiod of at least 2 years prior to the date of dispatch;]			
(¹) (⁴) or	Venezue Venezue	[a part of the territory of a country which is assigned to Sanitary Group C or D, which is free Venezuelan equine encephalomyelitis for a period of at least 2 years prior to the date of dispatch a Venezuelan equine encephalomyelitis occurs in the remaining parts of the territory of the country dispatch, and			
	(¹) either	[is vaccinated against Venezuelan equine encourse and revaccinated according to manufac 60 days and no more than 12 months prior to the protected quarantine for a period of at least 21 during that period remained clinically healthy, remained within the normal physiological range holding which showed a rise in body temperature, for virus isolation for Venezuelan equine encephal	cturer's recommendations not less than date of dispatch, and was kept in vector-days prior to the date of dispatch, and and its body temperature, taken daily, and any equine animal on the same taken daily, was subjected to a blood test		
	( <sup>1</sup> ) or	[is not vaccinated against Venezuelan equine er protected quarantine for a period of at least 21 clinically healthy, and its body temperature, ta physiological range, and any equine animal on the body temperature, taken daily, was subjected Venezuelan equine encephalomyelitis with ne dispatched was subjected to a diagnostic test for with negative result conducted on a sample taken entry into the vector protected quarantine and rendispatch;]]	days, and during that period remained liken daily, remained within the normal he same holding which showed a rise in to a blood test for virus isolation for gative results, and the animal to be or Venezuelan equine encephalomyelitis in not less than 14 days after the date of		
	( <sup>1</sup> ) or	[was subjected to a haemagglutination intercephalomyelitis carried out by the same samples taken on two occasions with an interval of date) and on	e laboratory on the same day on f 21 days on		
(¹) [II.3.3.	the anima	al is an uncastrated male equine animal older than 1	80 days, and		
(¹) either		ched from a country in which equine viral arteritis ( not been officially reported during the period of 6 mo			
(¹) or	21 days	ed on a blood sample taken onprior to the date of dispatch, by virus neutralisation of 1 in 4;]]			
(¹) or	a period	ed on an aliquot of its entire semen taken on			
(¹) or	supervisi	cinated against EVA onon, and re-vaccinated at regular intervals according approved by the competent authority, and the initial value.	to the manufacturer's instructions, with a		

Status: This is the original version (as it was originally adopted).

### **EUROPEAN UNION**

				<del>, , , , , , , , , , , , , , , , , , , </del>
			II.a. Certificate reference number	II.b. Local reference number
		(¹) either	[before 31 December 2017, on the day a blood tested in a virus neutralisation test for EVA v of 1 in 4;]]]	
		(¹) or	[before 31 December 2017, during a period of i official veterinary supervision, commencing on the was tested during that isolation period in a virus result at a serum dilution of 1 in 4;]]]	he day a blood sample was taken which
		(¹) or	[at the age of 180 to 270 days, during a per supervision, during which the animal was subject carried out with negative result at a serum dilution by the same laboratory with stable or declining the 10 days apart;]]]	cted to a virus neutralisation test for EVA n of 1 in 4, or carried out on the same day
		(¹) or	[after the animal was subjected to a virus neutral a serum dilution of 1 in 4, carried out on a blood commencing a period of uninterrupted isolation vaccination;]]]	sample taken not earlier than 7 days after
		(¹) or	[at the age of 180 to 250 days, after the animal v for EVA carried out with negative result at a sen same day by the same laboratory with stable or d at least 14 days apart;]]]	um dilution of 1 in 4 or carried out on the
	(¹) or	carried of sample of to the date	ected to a virus isolation test, polymerase chain ut with negative result on an aliquot of its entire f that animal taken on	e semen collected after the date a blood rt date), within a period of 6 months prior
( <sup>1</sup> ) ( <sup>4</sup> ) either	[II.3.4.	anaemia,	al is dispatched from Iceland, which is certified where it was continuously resident since birth an ve entered Iceland from other countries;]	
(¹) or	[11.3.4.	Coggins	al was subjected with negative result to an a test) or to an ELISA for equine infectious anaemia 	a carried out on a blood sample taken on
	( <sup>1</sup> ) [II.3.5.	Sanitary of during a test for gl	al is dispatched from a country or part of the te Group B, D or E, or from China or Thailand, or from period of 3 years prior to the date of dispatch, and anders carried out with negative result at a serum (insert date), within a period of 30 da	a a country in which glanders was reported d was subjected to a complement fixation dilution of 1 in 5 on a blood sample taken
	( <sup>1</sup> ) [II.3.6.	country of China or the date negative (insert date	al is an uncastrated male or a female equine anim r part of the territory of a country which is assigne Thailand, or from a country in which dourine was r of dispatch, and was subjected to a complement result at a serum dilution of 1 in 5 on a blood samp to the date during the period of at least 30 days prior to and af	ed to Sanitary Group B, D, E or F, or from reported during a period of 2 years prior to a fixation test for dourine carried out with le taken on
	(¹) [II.3.7.		al is dispatched from a country or part of the te Group C or D, and	rritory of a country which is assigned to
		(¹) either	[Western and Eastern equine encephalomyelitis country or part of the territory of the country of diprior to the date of dispatch;]]	, ,

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EUROPEAN UNION	- Ne	gistereu eqt	iluae, equiuae for breedi	ng and production equidae for slaughte
	II.a. Certif	icate referen	ce number	II.b. Local reference number
(¹)	manufactu date of	rer's instruct dispatch wi	ions within a period of 6 th inactivated vaccine	ary course and revaccinated according to months and at least 30 days prior to the against Western and Eastern equine and on
(1)	protected	quarantine, a	and during this period subj	ays prior to the date of dispatch in a vector ected to haemagglutination inhibition tests is carried out by the same laboratory
	(¹) either			date of dispatch, with negative result;]]]
	(¹) or	21 days on (insert date the date o	e), the second of which wa	o occasions with an interval of at least rt date) and on
Sai	nitary Group G, o	nal is dispatched from a country or part of the territory of a country which is assigned Group G, or from a country in which Japanese encephalitis has been officially reporteduring the past 2 years, and the animal		
(1)	holding wh	nere there ha		n area of at least 30 km radius around that ese encephalitis during a period of at least
(1)	date of dis	patch, and d		g a period of at least 21 days prior to the temperature, taken daily, remained within d
	(¹) either	Japanese of samples of second of dispatch, v	encephalitis carried out by blood taken on two occas (insert date) and o which was taken within vithout a more than four-f	on or virus neutralisation test for the same laboratory on the same day on ions with an interval of at least 14 days on in
	(¹) or	Japanese sample tak	encephalitis virus with ren not earlier than 7 days (insert date), an	or the detection of antibodies against negative result, carried out on a blood after the date the isolation commenced on d remained protected from vector insects
(1)	revaccinat	ed according		s with a complete primary course and nendations during a period of not less than date of dispatch;]]
Sai	nitary Group E, ai	nd was subje	cted to a serological test	erritory of a country which is assigned to for African horse sickness as described in y the same laboratory on the same day
(1)	-	(i		an interval of between 21 and 30 days, on
	(¹) either	[with negat	ive results in each case;]]]	
	(¹) or	[with positive	ve result in the first sample	e, and
		(¹) either		subsequently tested with negative result in st as described in Annex IV to Directive

Status: This is the original version (as it was originally adopted).

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	rtegister	ca equiade, equiade for bre	eding and production equidae for slaughter
	II.a. Certificate re	eference number	II.b. Local reference number
	(¹) or	in antibody titre in a	tested without more than a two-fold increase virus neutralisation test as described in 2.5.1 of the OIE Terrestrial Manual for accines;]]]
to th	e date of dispatch, and le OIE as officially free	the country or part of the ter	(insert date), within a period of 21 days prior ritory of the country of dispatch is recognised d is not adjacent to a country in which African []
	animal is dispatched fi tary Group F and	rom a country or part of the	territory of a country which is assigned to
(¹) e	(1) 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 laboratory same day on blood samples taken on two occasions with an interval of between 30 days, on		as carried out by the same laboratory on the casions with an interval of between 21 and d on (insert date), the first duction into the vector-protected quarantine,
	(1) either [with	negative results in each case	:111
	(1) or [with	positive result in the first sam	ple, and
	(¹) ei		is subsequently tested with negative result in test as described in Annex IV to Directive
	(¹) or	in antibody titre in a	tested without more than a two-fold increase virus neutralisation test as described in 2.5.1 of the OIE Terrestrial Manual for accines;]]]]
(1) 0	as described in A case on a blood s 28 days after th	Annex IV to Directive 2009/15 sample taken on	identification test for African horse sickness 6/EC, carried out with negative result in each
(1) 0	Annex IV to Dire	ective 2009/156/EC, carried (insert date) not les	for African horse sickness as described in out with negative result on a blood sample so than 14 days after the date of introduction ore than 72 hours before dispatch;]]
II.4. Attestation of the	transport conditions		
Sani mark	tary Group A, B, C, D,	E or G and is transported dir	e territory of a country which is assigned to ectly to the Union, without passing through a oming into contact with other equidae of a
Sani com	(¹) (⁴) or [II.4.1. The animal is dispatched from a country or part of the territory of a country which is a Sanitary Group F and is transported directly from the vector-protected quarantine stat coming into contact with other equidae not accompanied by a health certificate either for im temporary admission into the Union		vector-protected quarantine station without
(¹) e.	the aircraft beir	ng cleansed and disinfected	ns and arrangements have been made that d in advance with a disinfectant officially d sprayed against vector insects just prior to

### **EUROPEAN UNION**

### Registered equidae, equidae for breeding and production equidae for slaughter

	II.a. Certificate reference number	II.b. Local reference number
(¹) or	[to a sea port in that country or part of the territor conditions and arrangements have been made scheduled directly to a port in the Union without or part of the territory of a country not approved for stalls which were cleansed and disinfected in recognised in the third country of dispatch and spaceparture.]]	e to transport it on a vessel which is calling into a port situated in a country or r the entry into the Union of equidae, in a advance with a disinfectant officially
11.4.2		

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

### II.5. Attestation of animal welfare

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

### Notes:

### Part I:

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

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

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

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

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

### Part II:

- (1) 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.

EUROPEAN UNION		Registered equidae, equidae for breedin	ng and production equidae for slaughte						
		II.a. Certificate reference number	II.b. Local reference number						
( <sup>3</sup> )	Code of the country or part of the territory of the country and the Sanitary Group as appearing in columns 3 ar respectively of Annex I to Commission Implementing Regulation (EU) 2018/659.								
(4)	Statements that relate entirely and exclusively to a Sanitary Group different from the Sanitary Group to which the coun of dispatch, or part of its territory, is assigned, may be left out, provided that the numbering of the subsequent statemer is maintained.								
This	health certificate shall:								
(a)	be drawn up in at least a language understood by the certifying officer and one of the official languages of the Memb State of destination and of the Member State where the animal will enter Union territory and undergo the veterina border checks;								
(b)	be made out to a single cons	ignee;							
(c)	be signed and stamped in a	colour different to the colour of the printing;							
(d)		paper or all sheets of paper required are part o total number of pages, and each page shall bear the are stapled and stamped.							
Offic	ial veterinarian								
	Name (in capital letters):		Qualification and title:						
	Date:		Signature:						
	Stamp:								

Declaration by the owner or representative of the owner for entry into the Union of an equine animal									
Iden	tification o	f the animal (1)							
Species (Scientific name) Identification system Identification number Age Sex									
I, the	e undersigi	ned owner (²) o	r representative of the owne	er (²) of the animal described abov	e, hereby declare, that:				
_	the anima	al							
	(²) either			e territory of the country of dispat a animal is less than 90 days of ag		ast 90 days prior			
	(²) or			y of the country of dispatch durin a Member State of the Union;]	g the required residence p	period of at least			
_			days prior to the date of diseases transmissible to ed	dispatch the animal has not bequidae;	en in contact with animal	s suffering from			
_				on as applicable in accordance vecountry of dispatch are fulfilled;	vith point II.2 of the accord	mpanying health			
_			nsport as applicable in acco	ordance with point II.4 of the according	mpanying health certificat	e for the country			
_		portation will be the journey;	e effected in such a way t	hat health and well-being of the	animal can be protected	effectively at all			
Nan	ne and add	ress of the owr	ner (2) or representative (2):						
Date	∋:		(dd/mm/yyyy)						
(¹)	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 Commissio		EU) 2018/659. Specify the identifica					
		ort accompanies to		e stated and the name of the competer	nt authority which validated it.				
0	•	nale, F = female,	C = castrated).						
(2)	<sup>2</sup> ) Delete as appropriate.								

# Section B

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

JUN	ITRY	•					Veterinary certific	tate to Et		
	l.1.	Consignor Name Address	I.2. Certificate reference No I.2.a.							
				1.4.	Local competent a	uthority				
nment	1.5.	Tel.  Consignee Name		1.6.						
d consi		Address								
tche		Tel.								
Part I : Details of dispatched consignment	1.7.	Country of ISO code I.8. Region of origin origin	Code	1.9.	Country of destination	ISO code	I.10. Region of destination	Code		
: Detai	l.11.	Place of origin		1.12.	Place of destinatio	n				
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 w	agon 🛘							
		Road vehicle Other I Identification Documentary references		I.17. No(s) of CITES						
	118	Description of animals				I 10 Commo	dity code (HS code)			
	1. 10.	Description of animals				1.13. Commo	01 01	'		
					•		I.20. Quantity			
	I.21.						I.22. Number of p	ackages		
	1.23.	Seal/Container No					1.24.			
	1.25.	Animals certified for:								
	Slaughter									
	1.26.				I.27. For import or	admission into	EU 🗆			
	1.28.	Identification of the animals								
	s	Species (Scientific Identification system name)	Identif	ication	number	Age	Sex			

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			II.a. Certificate reference number	II.b. Local reference number							
	II.	Attestation of anim	al health, animal welfare and public health								
	I, the undersigned official veterinarian, hereby certify, that the animals described in Box I.28:										
	_	are equidae for slau	ghter as defined in Article 2(d) of Directive 2009/156/E	EC;							
	_	were examined tod infestation;	(1) and found free of clinical signs of diseases and of obvious signs of ectoparasite								
	<ul> <li>are not intended for slaughter under a national programme of infectious or contagious disease eradical</li> </ul>										
	<ul> <li>meet the requirements attested in points II.1 to II.5 of this certificate;</li> </ul>										
_	_	<ul> <li>are accompanied by the written declaration, signed by the owner of the animals or the representative of owner.</li> </ul>									
catior	II.1.	Attestation on third country or part of the territory of third country and holding of dispatch									
Part II: Certification	II.1.1.	The animals are dispatched from									
Pa	II.1.2.	in the country of dispatch the following diseases are compulsorily notifiable: African horse sickness, dourine ( <i>Trypanosoma equiperdum</i> ), glanders ( <i>Burkholderia mallei</i> ), equine encephalomyelitis (of all types including Venezuelan equine encephalomyelitis), equine infectious anaemia, vesicular stomatitis, rabies and anthrax;									
	II.1.3.	the animals are disp	atched from a country or part of the territory of country	у							
		which th African h have be	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;								
			Venezuelan equine encephalomyelitis has not occur of dispatch;	rred during the period of 2 years prior to							
		c) in which	dourine has not occurred during the period of 6 mont	hs prior to the date of dispatch;							
		d) in which	glanders has not occurred during the period of 6 mor	nths prior to the date of dispatch;							
	(3) either	[e) in which dispatch	vesicular stomatitis has not occurred during the p	period of 6 months prior to the date of							
	(³) or	and a blo	vesicular stomatitis has occurred during the period of cood sample taken from each of the animals on	(insert date), within a							
		(³) either	[in a virus neutralisation test at a serum dilution of 1	I in 32;]]							
		(³) or	[in an ELISA in accordance with the relevant Cha and Vaccines for Terrestrial Animals of the OIE;]]	apter of the Manual of Diagnostic Tests							
	II.1.4.	points II.1.4.1 to II.1	come from holdings, and to the best of my knowle .4.7 have not been in contact with animals from hol is referred to in points II.1.4.1 to II.1.4.7 and which las	dings, which were subject to prohibition							

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			,						
			II.a. Certificate reference number	II.b. Local reference number					
	II.1.4.1.	in the cas	e of equidae suspected of having contracted dourine	<b>&gt;</b> ,					
		(³) either	[6 months beginning on the date of the last act suspected of having contracted dourine or infected						
		(3) or	[in the case of a stallion, until the animal is castrate	d;]					
		(³) or	[30 days following the date of completion of the cleansing and disinfection of the after all animals of susceptible species have been slaughtered;]						
	II.1.4.2.	in the cas	e of glanders,						
		(³) either	er [6 months beginning on the day on which the equidae suffering from the dis subjected with positive results to a test for the detection of the causative p Burkholderia mallei or antibodies to that pathogen, were killed and destroyed;]						
		(³) or	[30 days following the date of completion of the cleafter all animals of susceptible species have been keeper and the clear of the clea						
	II.1.4.3.	in the cas	e of equine encephalomyelitis of any type,						
		(³) either	[6 months beginning on the day on which the equid slaughtered;]	ae suffering from the disease have been					
		( <sup>3</sup> ) or	[6 months beginning on the day on which the equidae infected with the virus of Nile Fever, Eastern equine encephalomyelitis or Western equine encephalo died, been removed from the holding or fully recovered;]						
		(³) or	[30 days following the date of completion of the cleafter all animals of susceptible species have been s						
	II.1.4.4.	slaughter agar gel	in the case of equine infectious anaemia, until the date on which, the infected animals having slaughtered, the remaining equine animals on the holding have shown a negative reaction agar gel immunodiffusion test (AGID or Coggins test) carried out on blood samples collecte two occasions 3 months apart;						
	II.1.4.5.	in the cas	e of vesicular stomatitis,						
		(³) either	[6 months following the last case;]						
		( <sup>3</sup> ) or	[30 days following the date of completion of the cleafter all animals of susceptible species have been s						
	II.1.4.6.		se of rabies, 30 days following the last case and the on of the premises;	date of completion of the cleansing and					
	II.1.4.7.		ee of anthrax, 15 days following the last case and the on of the premises;	date of completion of the cleansing and					
II.1.5.			wledge, during the period of 15 days prior to the dat se infected or suspected of an infectious or contagiou						
II.2.	Attestation	n of residen	ce and pre-export isolation						
II.2.1.	of 90 days	prior to the	en resident in the country or part of the territory of t e date of dispatch, or since birth if the animals are le n, and they are dispatched from a country or part of t	ess than 90 days old, on holdings under					
	( <sup>3</sup> ) either		to Sanitary Group A and during the period of at lea kept apart from equidae not of equivalent health sta						

# EUROPEAN UNION

			II.a. Certificate reference number	II.b. Local reference number							
	(³) or	dispatch	It to Sanitary Groups B, C or D and during the period they were kept in pre-export isolation under vete iith equidae not of equivalent health status;]								
	(³) or		[assigned to Sanitary Group E and for the period of at least 40 days prior to the date of dispatch they were kept in the approved isolation centre described in Box I.11, protected from vector insects.]								
II.3.	Attestation	n of vaccina	ation and health tests								
(3) either	[II.3.1.		nals were not vaccinated against African horse sickn rmation suggesting previous vaccination;]	ess in the country of dispatch and there							
(3) or	[II.3.1.		nals were vaccinated against African horse sicknes n 12 months prior to dispatch;]]	is, and this vaccination was carried out							
	II.3.2.		als were not vaccinated against Venezuelan equino ispatch from	e encephalomyelitis during the 60 days							
	(³) either	•	y of which all parts of the territory are free of Ven at least 2 years prior to the date of dispatch;]	ezuelan equine encephalomyelitis for a							
	( <sup>3</sup> ) ( <sup>4</sup> ) or	Venezue	f the territory of a country which is assigned to S lan equine encephalomyelitis for a period of at least lan equine encephalomyelitis occurs in the remainin and	2 years prior to the date of dispatch and							
		(3) either [were vaccinated against Venezuelan equine encephalomyelitis with a concourse and revaccinated according to manufacturer's recommendations 60 days and not more than 12 months prior to the date of dispatch, and were protected quarantine for a period of at least 21 days prior to the date of during that period remained clinically healthy, and their body temperatur remained within the normal physiological range, and any equine animal holding which showed a rise in body temperature, taken daily, was subjected for virus isolation for Venezuelan equine encephalomyelitis with negative res									
		( <sup>3</sup> ) or	[were not vaccinated against Venezuelan equinvector-protected quarantine for a period of at leas and during that period remained clinically healthy. remained within the normal physiological range, holding which showed a rise in body temperature. It for virus isolation for Venezuelan equine encepha animals to be dispatched were subjected to a encephalomyelitis with negative result conducted after the date of entry into the vector-protected quector insects until dispatch;]]	at 21 days prior to the date of dispatch. and their body temperature, taken daily, and any equine animal on the same aken daily, was subjected to a blood test lomyelitis with negative results, and the diagnostic test for Venezuelan equine on a sample taken not less than 14 days							
( <sup>3</sup> ) ( <sup>4</sup> ) either	[II.3.3.	.3. the animals are dispatched from Iceland, which is certified as officially free from equine infection anaemia, where they have been continuously resident since birth and did not come into contact equidae which have entered Iceland from other countries;]									
( <sup>3</sup> ) or	or [II.3.3. the animals were subjected to an agar gel immunodiffusion test (AGID or Coggins test) or to a for equine infectious anaemia carried out with negative result in each case on blood samples										
(3)	[[II.3.4.	the animals are dispatched from a country or part of the territory of a country which is assigned to Sanitary Group B, D or E, or from a country in which glanders was reported during the period of 3 years prior to the date of dispatch, and were subjected to a complement fixation test for glanders carried out with negative result in each case at a serum dilution of 1 in 5 on blood samples taken on									

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		II.a. Certif	ficate refere	nce number	II.b.	Local	reference	number	
( <sup>3</sup> ) [II.3.5.	country o country ir were subj at a serur	r part of the which dou ected to a d n dilution of	e territory of rine was re complement 1 in 5 on bl	les or female equine animals f a country which is assigned ported during the period of 2 t fixation test for dourine carriood samples taken ons prior to the date of dispatch	I to Sa years ed ou	anitary prior t with	Group B, to the date negative re	D or E or e of dispa esult in ea	r from a tch, and ach case
(³) (⁴) [II.3.6.		als are disp Group C or l		n a country or part of the ter	ritory	of a c	ountry wh	ich is ass	igned to
	(³) either	country or		n equine encephalomyelitis herritory of the country of disp					
	( <sup>3</sup> ) or	to manufactorial date of	cturer's instr dispatch w	ccinated with a complete prin ructions within the period of 6 vith inactivated vaccine ag last vaccination was applied	montl gainst	ns and Wes	l at least 3 tern and	0 days pri Eastern	or to the equine
	(³) or	period sub	jected to h	ot for at least 21 days protect naemagglutination inhibition (insert	tests	for W	estern an		
		(³) either	on	e of blood taken from eac (insert date), spatch, with negative result in	withir	n the p	eriod of 1		
		( <sup>3</sup> ) or	occasions and on period of	of blood taken from each of with an interval of at least 21(insert date), the 10 days prior to the date of di nimals were vaccinated more	days secor spatch	on nd of n, with	which was	( <i>inse</i> s taken w se in antib	rt date) ithin the ody titre
(³) (⁴) [II.3.7.	Sanitary (	Group E, an	nd were sub	n a country of part of the ter jected to a serological test fo EC, which was carried out by	r Afric	an ho	rse sickne	ss as des	cribed in
	(³) either	with an inte	erval of betve) and on	ken from each of the animal ween 21 and 30 days, on(inse days prior to the date of disp	rt date				
		(³) either	[with nega	tive result in each case;]]]					
		(3) or	[with posit	ive results in the first sample,	and				
			(3) either	[the second samples were sin each case in an agent ide to Directive 2009/156/EC;]]]]	entifica				
			( <sup>3</sup> ) or	[the two samples of each a without more than a two-for neutralisation test as descri OIE Terrestrial Manual for D	old inc bed in	rease	in antibo	dy titre in apter 2.5	a virus .1 of the
	( <sup>3</sup> ) or	consignment dispatch, a the OIE as	ent on and the cou officially fre	n each case on a blood samp (insert date), within the ntry or part of the territory of ee of African horse sickness as a has occurred during the pre-	the pe the co and is	riod o untry not a	f 10 days p of dispatch djacent to	orior to the	e date of nised by

### **EUROPEAN UNION**

			II.a. Certificate reference number	II.b. Local reference number						
II.4.	Attestation of the transport conditions									
(³) either	[II.4.1. 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 contact with other equidae not authorised for the entry into the Union.]									
(³) or	[II.4.1.	Arrangements were made and verified to ensure that before the animals are transported to a slaughterhouse on the territory of the Union they pass only through a single approved market, marshalling or assembly centre referred to in Article 7(1) of Directive 2009/156/EC situated in the same Member State, from where they are transferred directly to the slaughterhouse without coming into contact with other equidae not authorised for the entry into the Union.]								
	II.4.2.	least the	nents were made and verified to prevent any contact same health requirements as described in this hon until dispatch to the Union.							
	II.4.3.	disinfecte	port vehicles or containers in which the animals ar d before loading with a disinfectant officially recogn so constructed that faeces, urine, litter or fodder can	ised in the third country of dispatch and						
II.5.	Attestation	of animal	welfare							
			ed in Box I.28 were examined today (1) and foun nents have been made to protect their health and v							
II.6.	Attestation	of public h	ealth							
	androgeni	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.								
			ring live equidae provided by the residue plan sub- 96/23/EC are fulfilled.	mitted and approved in accordance with						
Notes:										
Part I:										
Box I.8.:			the country or part of the territory of the country a enting Regulation (EU) 2018/659.	as appearing in column 3 of Annex I to						
Box I.15.:	information	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 contai	ner numbe	r and the seal number (if applicable) should be included	ded.						
Box I.28.:	Species: S	elect amor	ngst: "Equus caballus", "Equus asinus" or "Equus cal	ballus x Equus asinus".						
	identification	on docume	Each of the animals must bear an individual identifient. Specify the identification system (such as ear toon the animal.							
	Age: Date	of birth (dd	/mm/yyyy).							
	Sex (M = male, F = female, C = castrated).									

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### **EUROPEAN UNION**

### Registered equidae, equidae for breeding and production equidae for slaughter

		II.a. Certificate reference number	II.b. Local reference number					
Part	II:							
(1)	The certificate must be issued Union.	d on the day of loading of the animals for dispatch to	the Member State of destination in the					
	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.							
(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)	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.							
	health certificate shall:							
(a)		guage understood by the certifying officer and one he Member State where the animals will enter Uni						
(b)	be made out to a single consi	gnee;						
(c)	be signed and stamped in a c	colour different to the colour of the printing;						
(d)		paper or all sheets of paper required are part of total number of pages, and each page shall bear the are stapled and stamped.						
Offici	al veterinarian							
	Name (in capital letters):		Qualification and title:					
	Date:		Signature:					
	Stamp:							
Offici	al veterinarian  Name (in capital letters):  Date:		-					

(2) Delete as appropriate.

				owner or representative of the ov consignments of live equidae fo				
Ider	ntification o	f the animal (1	)					
Sp	ecies (Scie	ntific name)	Identification system	Identification number	Age			
I, th	e undersigi	ned owner (²)	or representative of the owne	er (²) of the animals described abov	e, hereby declare, that:			
-	the anima	als have remai	ned in the country or part of t	the territory of the country of dispat	ch for at least 90 days	prior to the date of		
-			days prior to the date of d diseases transmissible to ed	lispatch the animals have not bee quidae;	en in contact with anim	als suffering from		
-				on as applicable in accordance wi e country of dispatch are fulfilled;	th point II.2 of the acc	ompanying health		
-			ansport as applicable in according the country of dispatch are f	ordance with point II.4 of the accordulfilled;	npanying health certific	ate for the country		
-		portation will the journey;	pe effected in such a way the	hat health and well-being of the a	nimal can be protecte	d effectively at all		
_	the anima	als will be sent						
	(²) either		the premises of dispatch to of the same health status;]	o the slaughterhouse of destination	n without coming into	contact with other		
	(²) or	marshalling of		claughterhouse of destination pas to in Article 7(1) of Directive 2009 status;]				
Nar	ne and add	ress of the ow	mer (²) or representative (²):					
Date	ə:		(dd/mm/yyyy)					
(¹)	Identification Article 2(b) transponde If a passpo	on system: The b) of Commission er) and the anato	animal must bear an individual in on Implementing Regulation (E omic place used on the animal. the animal, its number should be	r indicate any cross between those. identifier which permits to link the anim EU) 2018/659. Specify the identificati e stated and the name of the competent	on system (such as ear	tag, tattoo, brand		
	-	nale F = female	****					

#### PART 4

### **Explanatory notes for the certification**

- (a) The health certificates shall be issued by the competent authority of the exporting country, based on the model set out in Part 1, 2 or 3 of Annex II, according to the layout of the model that corresponds to the animals concerned.
  - They shall contain, in the numbered order that appears in the model, the attestations that are required for any country and, as the case may be, those supplementary guarantees that are required for the exporting country or part of the territory of the country.
- (b) Where the model health certificate states that certain statements shall be kept as appropriate, statements which are not relevant, may be crossed out and initialled and stamped by the certifying officer, or completely deleted from the health certificate.
- (c) A separate and unique health certificate shall be issued for animals that are exported from a single territory appearing in columns 2 and 4 of Annex I which are consigned to the same destination and transported in the same railway wagon, lorry, aircraft or ship.
- (d) The original of the health certificate shall consist of a single sheet of paper, or, where more text is required, it shall be in such a form that all sheets of paper required are part of an integrated whole and indivisible.
- (e) The health certificate shall be drawn up in at least one of the official languages of the EU Member State of the border inspection post of entry of the consignment into the Union and of the EU Member State of destination. However, those EU Member States may authorise the health certificate to be drawn up in the official language of another EU Member State, and accompanied, if necessary, by an official translation.
- (f) If for reasons of identification of the animals of the consignment (schedule in Box 1.28 of the model health certificate), additional sheets of paper are attached to the health certificate, those sheets of paper shall also be considered as forming part of the original of the health certificate by application of the signature and stamp of the certifying official veterinarian, on each of the pages.

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

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

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

<sup>(1)</sup> OJ L 13, 16.1.1997, p. 28.

#### 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

### 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

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COUN	TRY:					Veterinary certificate to EU		
	l.1.	Name		Certificate reference		I.2.a.		
		Address	1.3.	I.3. Central competent authority				
멑		Tel.	1.4.	I.4. Local competent authority				
Part I : Details of dispatched consignment	1.5.	Consignee Name Address	1.6.	Person responsible Name Address	for the load	in EU		
oatched c		Postal code Tel.		Postal code Tel.				
ls of disp	1.7.	Country of ISO code I.8. Region of code origin crigin	1.9.	Country of destination	ISO code	I.10. Region of Code destination		
l : Detai	l.11.	Place of origin Semen centre □	I.12.	Place of destination Semen centre	n	Holding		
Part		Name Approval number Address		Name Address	,	Approval number		
		Postal code		Postal code				
	I.13.	Place of loading	I.14. Date of departure					
	I.15.	Means of transport	1.16	Entry BIP in EU				
		Aeroplane  Ship  Railway wagon						
		Road vehicle Other Identification  Documentary references	l.17.					
	I.18.	Description of commodity			.19. Commod	dity 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			EU 🔲		
		Third country ISO code						
	1.28.	Identification of the commodities						
	S	pecies (Scientific name) Donor identity		Date of colle	ection	Quantity		

### COUNTRY

### Equine semen - Section A

	II. Health information		II.a. Certificate reference No	II.b.					
	I, the undersigned, official ve	terinarian, of the	e exporting country (²)(name of exporting						
	certify that:								
u.	export to the U	Inion is approve	(3), in which the semen described above wa ed and supervised by the competent authorit Annex D to Directive 92/65/EEC (4);						
Part II: Certification		or chilled seme	g 30 days prior to the date of first collection on was dispatched or until the 30 days storage						
Part II:	II.2.1. was situated in the exporting country or, in the case of regionalisation according to Article 13 of Directive 2009/156/EC ( <sup>5</sup> ), in that part of the territory of the exporting country which was:								
	_		ed to be infected with African horse sickne ective 2009/156/EC,	ss in accordance with Article 5(2)(a)					
	_	free from Ven	nezuelan equine encephalomyelitis for a perio	d of at least 2 years,					
	_	free from glar	nders and dourine for a period of at least 6 mo	onths;					
II.2.2. fulfilled the conditions for a holding laid down in Article 4(5) of Directive 2009/156/EC and									
	(¹) either [II.2		ng a case of a disease mentioned below not all the animals of species susceptible to sease located in the holding were slaughtered or killed and the holding has been free:						
		be	om any type of equine encephalomyelitis eginning on the day on which the equid laughtered,						
		ne or	om equine infectious anaemia (EIA) for at egative result in an agar gel immunodiffusion n samples taken after the infected animals months apart from each of the remaining anir	test (AGID or Coggins test) carried out were slaughtered on two occasions					
			om vesicular stomatitis (VS) for a period of at ase,	least 6 months from the last recorded					
		— fr	om rabies for a period of at least one month fr	om the last recorded case,					
		— fr	om anthrax for a period of at least 15 days fro	m the last recorded case,]					
	( <sup>1</sup> ) or [II.2	disease and the encepha the case	g a case of a disease mentioned below all the located in the holding have been slaughtered a holding was free for a period of at leas alomyelitis, equine infectious anaemia, vesicula of anthrax, beginning on the day on which forfection of the premises was satisfactorily com	or killed and the premises disinfected, it 30 days from any type of equine ilar stomatitis and rabies or 15 days in ollowing the destruction of the animals					
		ntained only equ tritis,	uidae which were free of clinical signs of equi	ne viral arteritis and contagious equine					
	II.3. Prior to entering	g the semen col	llection centre the donor stallions and any oth	er equidae located in the centre:					

### COUNTRY

### Equine semen - Section A

II. Health	n informatio	on	II.a. Certificate reference No	II.b.				
	II.3.1.	a Member State regionalisation i	y resident for a period of 3 months (or since of the Union during the 3 months period) in taccordance with Article 13 of Directive 2009 ontry which was during that period:	the exporting country or, in the case of				
			red to be infected with African horse sicknown irective 2009/156/EC,	ess in accordance with Article 5(2)(a)				
		— free from \( \lambda \)	enezuelan equine encephalomyelitis for a perio	od of at least 2 years,				
		— free from g	anders and dourine for a period of at least 6 m	nonths;				
(¹) either	[11.3.2.		he country of export which was on the day tis (VS) for a period of at least 6 months,]	of admission into the centre free from				
( <sup>1</sup> ) or	[II.3.2.	result at a serul with the relevan	o a virus neutralisation test for vesicular stor didiution of 1 in 32 or a VS ELISA carried of Chapter of the Manual of Diagnostic Tests a did sample taken (6) within 14 days prior to ente	ut with a negative result in accordance and Vaccines for Terrestrial Animals of				
	II.3.3.	originated from point II.2.2;	oldings which on the day of admission onto	the centre fulfilled the requirements of				
II.4.	The sen	nen described above	was collected from donor stallions which:					
	II.4.1.		clinical sign of an infectious or contagious disease at the time of admission onto the centre and on the day the semen was collected;					
	II.4.2.		period of at least 30 days prior to the date of semen collection in holdings where no is shown any clinical sign of equine viral arteritis or contagious equine metritis during					
	II.4.3.	collection and be	or natural mating during a period of at least 30 days prior to the date of first semen etween the dates of the first sample referred to in points II.4.5.1, II.4.5.2 and/or II.4.5.3 If of the collection period;					
	II.4.4.	Manual of Diagr which is recogni	ollowing tests, which meet at least the require ostic Tests and Vaccines for Terrestrial Animal ed by the competent authority and has the testivalent to that provided for in Article 12 of	Is of the OIE, carried out in a laboratory sts referred to hereinafter included in its				
		test)	uine infectious anaemia (EIA), an agar-gel im r an enzyme-linked immunosorbent assay (EL tive result;]					
		II.4.4.2. for eq	uine viral arteritis (EVA),					
		(¹) either [II.4.4	a serum neutralisation test with a negatin four;]	ative result at a serum dilution of one				
		( <sup>1</sup> ) and/or [II.4.4	2.2. a virus isolation test, polymerase chain negative result on an aliquot of the entire					
		three	entagious equine metritis (CEM), an ager specimens (swabs) taken from the donor stalli s than 7 days at least from the penile sheat s;	on on two occasions with an interval of				

#### COUNTRY Equine semen - Section A

cou	NTRY				Equine semen – Section A
II.	Health informatio	n		II.a. Certificate reference No	II.b.
			(local tre	aples were in no case taken earlier than 7 days (systematment) after antimicrobial treatment of the donor so to medium with activated charcoal, such as Amies med by where they were subjected with a negative result to a	tallion and were placed in ium, before dispatch to the
		(¹) either	[11.4.4.3.1	<ol> <li>the isolation of Taylorella equigenitalis after cultivic conditions for a period of at least 7 days, set up with specimens from the donor animal, or 48 hours who cool during transport;]</li> </ol>	nin 24 hours after taking the
		(1) and/or	[11.4.4.3.2	<ol> <li>the detection of genome of Taylorella equigenitalis carried out within 48 hours after taking the specimer</li> </ol>	
	II.4.5.	programn		th the results specified in point II.4.4 in each case end respectively in points 1.6(a), (b) and (c) of Chapter ws:	
		( <sup>9</sup> ) [II.4.5.1.	at least 3 the seme	or stallion was continuously resident on the semen colle 30 days prior to the date of the first collection and durin en described above, and no equidae on the semen co into direct contact with equidae of lower health status th	g the period of collection of lection centre came during
			stallion a collection and not l	s described in point II.4.4 were carried out on sample at least once a year at the beginning of the breeding n of semen intended for imports into the Union of free less than 14 days following the date of the commencen at 30 days prior to the first semen collection.]	season or prior to the first sh, chilled or frozen semen
		( <sup>9</sup> ) [II.4.5.2.	30 days semen d centre ve	or stallion was resident on the semen collection cen prior to the date of the first collection and during the escribed above, but left the semen collection centre un eterinarian for a continuous period of less than 14 day en collection centre came into direct contact with equida	period of collection of the ider the responsibility of the is, and/or other equidae on
			stallion a the first semen a	s described in point II.4.4 were carried out on sample the least once a year at the beginning of the breeding se collection of semen intended for imports into the Unio nd not less than 14 days following the date of the common at least 30 days prior to the first semen collection,	eason or prior to the date of n of fresh, chilled or frozen
		and	chilled o	ne period of collection of the semen intended for impo or frozen semen the donor stallion was subjected .4, 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 sampl 30 days prior to the date of the collection of the sem	
			(¹) or	[in point II.4.4.2.2 was carried out on an aliquot donor stallion taken (6) not more than 6 months collection of the semen described above and a blood donor 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

Health information

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II.a. Certificate reference No

### COUNTRY

#### Equine semen - Section A

II.b.

11.	Health Information		ii.a. Certificate reference No	11.0.
		(c)	for contagious equine metritis, the test described carried out on three specimens (swabs) taken (6) not the date of the collection of semen described above	
		(1) either	[on two occasions;]	
		(¹) or	[on a single occasion and subjected to a PCR or real	-time PCR.]]
	( <sup>9</sup> ) [II.4.5.3.		or stallion does not meet the conditions set out in points D to Directive 92/65/EEC and the semen is collected formen.	
			s described in points II.4.4.1, II.4.4.2 and II.4.4.3 wer from the donor stallion at least once a year at the	
	and	the dono from the semen of	described in points II.4.4.1 and II.4.4.3 were carried out r stallion during the storage period of the semen of a r date of the collection of the semen and before the sollection centre, not less than 14 days and not mor	ninimum period of 30 days emen is removed from the
	and	(1) either	[the tests for equine viral arteritis described in point samples taken (6) during the storage period of the s of 30 days from the date of the collection of the sem- removed from the semen collection centre or used, not more than 90 days after the date of the collection above.]	emen of a minimum period en and before the semen is not less than 14 days and
		(¹) or	[the non-shedder state of a donor stallion seroposit was confirmed by virus isolation test, PCR or real-ti negative result on samples of an aliquot of the estallion taken (6) twice a year at an interval of at leastallion has reacted with a positive result at a serun four in a serum neutralisation test for equine viral arter.	me PCR carried out with a entire semen of the donor st 4 months and the donor in dilution of at least one in
	II.4.6. underwer	nt the test	ing provided for in points II.3.2 (1) and II.4.5 on same	les taken on the following

II.4.6. underwent the testing provided for in points II.3.2 (1) and II.4.5 on samples taken on the following dates:

of		Start o	date (6)	Date of sampling for health tests (6)						
Identification of semen	Test gramme	Donor residence	Semen VS (¹) collection II.3.2	VS (¹)	EIA		A II. 1.2.	CEM II.4.4.3.		
Ident	bro			II.4.4.1.	Blood sample	Semen sample	1. sample	2. sample		

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#### COUNTRY

#### Equine semen - Section A

II. H	lealth info	rmation		II.a. Certificate reference No	II.b.
(1) eith	er [II.	5.	No antibiotics were	added to the semen;]	
( <sup>1</sup> ) or	[II.	5.	diluted semen of no	iotic or combination of antibiotics was added to produc ot less than (10):	
					;]
II.6.	Th	e semen	described above w	as:	
	II.6	5.1.		ed, stored and transported under conditions which com ad III(I) of Annex D to Directive 92/65/EEC;	ply with the requirements of
	11.6	3.2.		of loading in a sealed container in accordance with per e 92/65/EEC and bearing the number indicated in Box	
Notes					
Part I:					
Box I.1	I1.: Th	e place o	of origin shall corres	pond to the semen collection centre of the semen origin	1.
Box I.2	22.: Th	e numbe	r of packages shall	correspond to the number of containers.	
Box I.2	23.: Th	e identifi	cation of container	and seal number shall be indicated.	
Box I.2	28.: Th	e donor i	dentity shall corresp	oond to the official identification of the animal.	
	Th	e date of	collection shall be	indicated in the following format: dd/mm/yyyy.	
Part II:	:				
Guidar	nce for the	complet	ion of the table in p	oint II.4.6.	
Abbrev	viations:				
VS	S	Vesicul	ar stomatitis (VS) te	esting if required in accordance with point II.3.2	
El	A-1	Equine	infectious anaemia	(EIA) testing first occasion	
El	A-2	EIA tes	ting second occasio	n	
E۱	VA-B1	Equine	viral arteritis (EVA)	testing on blood sample first occasion	
E۱	VA-B2	EVA tes	sting on blood samp	ole second occasion	
E۱	VA-S1	EVA tes	sting on semen sam	ple first occasion	
E۱	VA-S2	EVA tes	sting on semen sam	ple second occasion	
CE	EM-11	Contag	ious equine metritis	(CEM) testing first occasion first sample	
CE	EM-12	CEM te	sting first occasion	second sample taken 7 days after CEM-11	
CE	EM-21	CEM te	sting second occas	ion first sample	
CE	EM-22	CEM te	sting second occas	ion second sample taken 7 days after CEM-21	
Inetrue	tions				

### Instructions:

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.

Qualification and title:

Signature:

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COUNTRY	Equine semen –	Section A

II.	I. Health information II.a. Certificate reference No								II.b.		
	required i	n points	amples were t II.4.5.1, II.4.5.2 with EIA-1, EV	2 and II.4.5.3,	shall be en	tered in the	upper line of	columns 5 to	9 of the tabl		
	shall be e	entered in	amples were to the lower line example belo	of columns 5							
	of		Start	date			Date of samp	ing for health te	ests		
	Identification of semen	Test programme	Jonor Donor	Semen	vs	EIA		VA .4.2.		EΜ .4.3.	
	Ident	pro	residence	collection	II.3.2.	II.4.4.1.	Blood sample	Semen sample	1. sample	2. sample	
		_			Ve	EIA-1	EVA-B1	EVA-S1	CEM-11	CEM-12	
	Α	ВС	С	D	vs	EIA-2	EVA-B2	EVA-S2	CEM-21	CEM-22	
(²) (³) (⁴) (⁵)	Regulation (EU) 2018/659 provided that the semen was collected in the part of the territory of the third country detailed in column 4 of that Annex from a donor stallion of the category of equidae indicated in columns 11, 12 or 13 of that Annex.  (3) Only approved semen collection centres listed in accordance with Article 17(3)(b) of Directive 92/65/EEC on the Commission website: http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm  (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).										
(7)	Regulation (EC) No 882/2004 of the European Parliament and of the Council of 29 April 2004 on official controls performed to ensure the verification of compliance with feed and food law, animal health and animal welfare rules (OJ L 165, 30.4.2004, p. 1).										
(8)	donor e	quidae wl infectious	nunodiffusion to hich have conto anaemia and nd during the p	inuously resid no equidae a	ed in Iceland and their se	d since birth men, ova ar	, provided tha	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.

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 certific	ate to EU
	l.1.	Consignor Name					1.2.	Certificate reference	ce No	I.2.a.	
		Address					1.3.	Central competent	authority		
		Tel.					I.4. Local competent authority				
Part I : Details of dispatched consignment	1.5.	Consignee Name Address					1.6.	Person responsible Name Address	e for the load	in EU	
tched co		Postal code Tel.						Postal code Tel.			
s of dispat	1.7.	Country of IS origin	SO code		Region of origin	Code	1.9.	Country of destination	ISO code	I.10. Region of destination	Code
: I : Detail	I.11. Place of origin Semen centre □						I.12.	Place of destinatio Semen centre		Holding 🗖	·
Part	Name Approval number Address							Name Address	Approva	al number	
		Postal code						Postal code			
	I.13.	1.13. Place of loading						I.14. Date of departure			
	I.15.	Means of trans	sport				I.16.	Entry BIP in EU			
		Aeroplane 🗖	Ship 🗆		Railway wagor						
		Road vehicle L Identification Documentary r		Othe	r 🚨		I.17.				
	I.18.	Description of	commodity						I.19. Commo	dity code (HS code) 05 11 99 85	
										I.20. Quantity	
	I.21.									I.22. Number of pack	kages
	1.23.	Seal/Container	r No							1.24.	
	1.25.	Commodities of Artificial reprod									
	1.26.	For transit thro	ough EU to		country [	]		I.27. For import or	admission into	EU 🗆	
		Identification of		oditie	es Donor ider	ntity		Date of colle	ection	Quantity	

### COUNTRY Equine semen - Section B Health information II.a. Certificate reference No I, the undersigned, official veterinarian, of the exporting country (2) ........... hereby (name of exporting country) certify that : The semen collection centre (3), in which the semen described above was collected, processed and stored for II.1. export to the European Union is approved and supervised by the competent authority in accordance with the conditions of Chapter I(I) (1) and Chapter I(II) (1) of Annex D to Directive 92/65/EEC, Part II: Certification II.2. during the period commencing 30 days prior to the date of first collection of the semen described above until the 30 days storage period for frozen semen elapsed, the semen collection centre: II.2.1. was situated in the exporting country or, in the case of regionalisation according to Article 13 of 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; 11.2.2. fulfilled the conditions for a holding laid down in Article 4(5) of Directive 2009/156/EC (8) and in (1) either [II.2.2.1. following a case of a disease mentioned below not all the animals of species susceptible to the disease located on the holding were slaughtered or killed and the holding has been from any type of equine encephalomyelitis for at least 6 months, beginning on the day on which the equidae suffering from the disease are slaughtered, from equine infectious anaemia for at least the period required to obtain a negative result in an agar gel immunodiffusion test (Coggins test) carried out on samples taken after the infected animals were slaughtered on two occasions 3 months apart from each of the remaining animals, from vesicular stomatitis for at least 6 months from the last recorded case, from rabies for at least one month from the last recorded case, from anthrax for at least 15 days from the last recorded case,] (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 premises disinfected, the holding has been free for at least 30 days from any type of equine encephalomyelitis, equine infectious anaemia, vesicular stomatitis and rabies or 15 days in the case of anthrax, beginning on the day on which following the destruction of the animals the disinfection of the premises was satisfactorily completed;] 11 2 3 contained only equidae which were free of clinical signs of equine viral arteritis and contagious equine

Prior to entering the semen collection centre the donor stallions and any other equidae located in the centre:

metritis.

11.3.

### COUNTRY

#### Equine semen - Section B

II. He	ealth information		II.a. Certificate reference No	II.b.
	II.3.1.	State of the Europeregionalisation acc	resident for 3 months (or since entry if they were direan Union during the 3 months period) in the exportording to Article 13 of Directive 2009/156/EC (8), in which was during that period	ting country or, in the case of
			ed to be infected with African horse sickness in acceptive 2009/156/EC (8),	ecordance with Article 5(2)(a)
		— free from Ver	nezuelan equine encephalomyelitis for at least 2 years	,
		<ul><li>free from glan</li></ul>	nders and dourine for at least 6 months;	
(¹) eithe	er [II.3.2.		e country of export which was on the day of adm (VS) for at least 6 months,]	ission into the centre free of
( <sup>1</sup> ) or	[II.3.2.		a virus neutralisation test for vesicular stomatitis ('dilution of 1 in 12 on a blood sample taken (4) within	
	II.3.3.	originated from hopoint II.2.2;	ldings which on the day of admission onto the centr	e fulfilled the requirements of
II.4.	The semer	n described above w	vas collected from donor stallions, which:	
	II.4.1.		ny clinical sign of an infectious or contagious disease the day the semen was collected;	at the time of admission onto
	II.4.2.		r 30 days prior to the date of semen collection on hol- nical sign of equine viral arteritis or contagious equine	
	II.4.3.		d for natural mating during at least 30 days prior to the lates of the first sample referred to in points II.4.5.1, I action period;	
	II.4.4.	the Manual of Disamples taken in	ne following tests, which meet at least the requireme agnostic Tests and Vaccines for Terrestrial Animals accordance with one of the programmes specified competent authority:	s of the OIE, carried out on
	( <sup>1</sup> ) ( <sup>5</sup> ) either		-gel immuno-diffusion test (Coggins test) for equine e result; ]	infectious anaemia (EIA) with
	(¹) (⁵) or	[II.4.4.1. an ELIS	A for equine infectious anaemia (EIA) with negative re	esult;]
and	(¹) either	•	n neutralisation test for equine viral arteritis (EVA) with of one in four;]	ith negative result at a serum
	(¹) or		isolation test for equine viral arteritis (EVA) carried of the entire semen of the donor stallion;	out with negative result on an

### COUNTRY Equine semen – Section B

II.	Health information			II.a.	Certificate reference No		II.b.
and		II.4.4.3.	occasion equigenia and from	on a <i>lis</i> a geni	tification test for contagious equine samples collected with an interval ter a cultivation of 7 to 14 days from pal swabs taken at least from the peresult in each case;	of 7 days ore-ejaculato	by isolation of <i>Taylorella</i> ry fluid or a semen sample
	II.4.5.				h the results specified in II.4.4. in $\epsilon$ n points II.4.5.1, II.4.5.2 and II.4.5.3 as		o at least one of the test
		II.4.5.1.	30 days semen d	orior escrib	lion was continuously resident on th o the date of the first collection and ed above, and no equidae on the ser contact with equidae of lower health s	I during the men collection	period of collection of the on centre came during that
			first sem	n co	ribed in point II.4.4 have been carried ection and at least 14 days following d of at least 30 days.		
		II.4.5.2.	the date above, b continuo	of the ut ha s per	ion was resident on the semen collection and during the period is left the centre under the responsion of of less than 14 days, or other equivith equidae of lower health status.	d of collection ibility of the	n of the semen described centre veterinarian for a
			date of the	e firs escril	ribed in point II.4.4 have been carried semen collection of the breeding sea ed above was collected and at lea t of the residence period of at least 30	ason or colle est 14 days	ction period in the year the
	and			bloo	ped in point II.4.4.1 for equine infection displays by taken (4) not more than 90 days by		
	and	(¹) either			s described in point II.4.4.2 for equine t) not more than 30 days before the se		
	aliquot of the entire semen of semen described above wa			on test for equine viral arteritis was on the semen of the donor stallion taken ed above was collected and a blood in a serum neutralisation test for equin four,]	n (⁴) not more d sample ta	e than 6 months before the ken on the same date (4)	
	and				ped in point II.4.4.3 for contagious e (4), not more than 60 days before the		
	II.4.5.3. The tests described in point II.4.4 have been carried out on samples taken (4) produced the first semen collection of the breeding season or collection period in the semen described above was collected,						
	and				ribed in point II.4.4 have been carric after the collection of the semen desc		

### COUNTRY

### Equine semen - Section B

II. Health i	nformation		II.a. (	Certificate re	ference No	II.b.				
	II.4.6.	have undergor following dates		ing provided	d for in poir	and II.4.5	on samples ta	ken on the		
of		Start date	e (4)	Date of sampling for health tests (4)						
Identification of semen	Test	Donor	Semen	VS (¹)	EIA		A II. 1.2.	CE II.4.		
Identif	Dud	residence	collection	II.3.2	II.4.4.1.	Blood sample	Semen sample	1. sample	2. sample	
(1) oither	[U. 5	No optibiotics	wara addad	to the same	n:1					
(1) either	[II.5.	No antibiotics v								
(1) or	[II.5.	The following a diluted semen			of antibiotic	s was added	to produce	a concentration	n in the final	
									;]	
II.6.	The seme	n described abov	ve was:							
	II.6.1.	collected, proc Chapters II(I) (					which comp	ly with the requ	uirements of	
	II.6.2.	sent to the pla Annex D to Dir							pter III(I) of	
Notes										
Part I:										
Box I.11.:	The place	of origin shall co	orrespond to	the semen	collection ce	entre of the se	emen origin.			
Box I.22.:	The numb	er of packages s	hall corresp	ond to the r	umber of co	ntainers.				
Box I.23.:	The identif	ication of contain	ner and sea	I number sh	all be indicat	ted.				
Box I.28.:	The donor	identity shall con	rrespond to	the official i	dentification	of the anima	l.			
	The date of	of collection shall	l be indicate	d in the follo	wing format	: dd/mm/yyyy	<i>1</i> .			

#### COUNTRY

#### Equine semen - Section B

II.	Health info	rmation	II.a. Certificate reference No	II.b.					
Par	Part II:								
Gui	Guidance for the completion of the table in point II.4.6.								
Abb	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 samp	le second occasion						
	EVA-S1	EVA testing on semen sam	ple first occasion						
	EVA-S2	EVA testing on semen sam	ple second occasion						
	CEM-11	Contagious equine metritis	(CEM) testing first occasion first sample						
	CEM-12	CEM testing first occasion s	second sample taken 7 days after CEM-11						
	CEM-21	CEM testing second occasi	on first sample						

#### 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.

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

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	Test programme	Start date		Date of sampling for health tests						
Identification		Donor residence	Semen collection	VS II.3.2.	EIA	EVA II.4.4.2.		CEM II.4.4.3.		
					II.4.4.1.	Blood sample	Semen sample	1. sample	2. sample	
Α	ь	ВС	D	vs	EIA-1	EVA-B1	EVA-S1	CEM-11	CEM-12	
	В				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) 2018/659 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

Status: This is the original version (as it was originally adopted).

II.	Health information	II.a. Certificate reference No	II.b.				
(3)		es listed in accordance with Article 17(3)(b) of Counc eu/food/animal/semen_ova/equine/index_en.htm	bil Directive 92/65/EEC on the				
( <sup>4</sup> )	Insert date in table in point II.4.6 (follow	Guidance in Part II of the Notes)					
( <sup>5</sup> )	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.						
( <sup>6</sup> )	Cross out the programmes that do not apply to the consignment.						
( <sup>7</sup> )	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:									V	eterinary certifica	te to EU
	l.1.	Name						Certificate reference		1.3	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					1.6.	Person responsible Name Address Postal code	for the load	in El	J	
hed		Tel.						Tel.				
of dispate	1.7.	Country of origin	ISO code	1.8.	Region of origin	Code	1.9.	Country of destination	ISO code	I.10.	Region of destination	Code
: Details	l.11.	Place of origin Semen centre				I.12.	Place of destination	_				
Part		Name Approval number Address						Name Address	Approval number			
		Postal code						Postal code				
	I.13.	I.13. Place of loading					I.14.	Date of departure				
	I.15. Means of transport				I.16.	Entry BIP in EU						
		Aeroplane ☐			Railway wago her □	on 🗆	1.17.					
		Identification Documentary					1.17.					
	I.18.	Description of	of commodity	1				I.	19. Commo	-	code (HS code) 5 11 99 85	
										1.20	). Quantity	
	I.21.									1.22	. Number of packa	ages
	1.23.	Seal/Contain	er No							1.24		
	1.25.	Commodities	certified for	:								
		Artificial repre	oduction									
	1.26.	For transit the	rough EU to	third	country			I.27. For import or a	dmission int	o EU		
		Third country	, 15	SO co	ode							
		Identification		noditi		antita -		Data of aclic	otion		0	
	8	pecies (Scient	unc name)		Donor ide	entity		Date of colle	CUON		Quantity	

COUNTRY Equine semen - Section C

	II. Health information		ation	II.a. Certificate reference No	II.b.						
	I, the unders	igned	, official veterinarian, of the exp	porting country (²)	-						
				(name of exporting co	untry)						
	certify that:										
ou	II.1.		semen collection centre in whi e European Union:	ch the semen described above was collected, pro	ocessed and stored for export						
rtificati	II.1.1.		oproved and supervised by the true 92/65/EEC,	ne competent authority according to the condition	ons of Chapter I, Annex D to						
Part II: Certification	II.1.2.	part		case of regionalisation according to Article 13 of of export which was on the day the semen w							
		_	<ul> <li>African horse sickness, in accordance with EU legislation,</li> </ul>								
		<ul> <li>Venezuelan equine encephalomyelitis for 2 years,</li> </ul>									
		<ul> <li>glanders and dourine for 6 months;</li> </ul>									
	II.1.3.	was during the period commencing 30 days prior to the date of collection of the semen until the day of its dispatch not subject to a prohibition order for animal health reasons which laid down one of the following conditions:									
	II.1.3.1.	if not all the animals of species susceptible to the disease located in the holding were slaughtered or killed, the prohibition lasted for:									
		<ul> <li>6 months, beginning on the day on which the equidae suffering from the disease are slaughtered, in the case of equine encephalomyelitis,</li> </ul>									
		-		out with negative result two Coggins tests 3 inimals have been slaughtered, in the case of infe							
		_	6 months, in the case of vesion	cular stomatitis,							
		_	one month from the last reco	rded case, in the case of rabies,							
		_	15 days from the last recorde	ed case, in the case of anthrax.							
	II.1.3.2.	the p	premises disinfected, the proh	otible to the disease located in the holding have be ibition lasted for 30 days, or 15 days in the case iion of the animals the disinfection of the premises	of anthrax, beginning on the						
	II.1.4.			mencing 30 days prior to semen collection and ree of clinical signs of equine viral arteritis and co							
	II.2.	Prior	to entering the semen collecti	on centre the donor stallions and any other equida	ae located in the centre:						
	II.2.1.	Unio		onths (or since entry if they were directly importe in the territory or in the case of regionalisation in that period free of:							
		-	African horse sickness, in acc	cordance with EU legislation,							
		-	Venezuelan equine encephal	omyelitis for 2 years,							
		_	glanders for 6 months,								
		_	dourine for 6 months;								

### COUNTRY

### Equine semen - Section C

					Equine Semen – Section 6
II. Health	information		II.a.	Certificate reference No	II.b.
(1) either	[II.2.2.	originated from the ten		of the country of export which was on the or 6 months,]	day of admission into the centre
(¹) or	[II.2.2.		(4), th	neutralisation test for vesicular stomat his being within 14 days prior to entering th	
II.2.3.	originated	from holdings which on	he da	ay of admission onto the centre fulfilled the	e requirements of point II.1.3;
II.3.	The seme	n described above was o	ollect	ted from donor stallions, which:	
II.3.1.	on the day	the semen was collecte	d hav	re not shown clinical signs of an infectious	or contagious disease,
II.3.2.	during at l	east 30 days prior to coll	ection	n of the semen have not been used for nat	ural service,
II.3.3.		e last 30 days prior to d inical signs of equine vira		ion of the semen have been kept on ho pritis,	oldings where no equine animal
II.3.4.		last 60 days prior to dinical signs of contagious		ion of the semen have been kept on ho ine metritis,	oldings where no equine animal
II.3.5.		,		as I could ascertain have not been in cor 15 days immediately preceding the collect	
II.3.6.				health tests carried out in a laboratory ramme as specified in point II.3.7:	recognised by the competent
II.3.6.1.	an agar-ge	el immuno-diffusion test	Cogg	ins test) for equine infectious anaemia wit	th negative result (3);
(1) either	[II.3.6.2.	a serum neutralisation	test fo	or 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 equi	ine viral arteritis carried out with negative	result on an aliquot of the entire
II.3.6.3.	Taylorella	equigenitalis from pre-e	jacula	carried out on two occasions with an in atory fluid or a semen sample and from g urethral fossa with negative result in each	enital swabs taken at least from
II.3.7.	have beer	subjected to one of the	follow	ving test programmes ( <sup>5</sup> ):	
II.3.7.1.	collection,	and during the collection	n per	esident on the collection centre for at leading and no equidae on the collection centre that the collection centre that the contract of the status than the donor stallions.	
		4 days after the comme		een carried out on samples taken on nent of the above residence period and	
II.3.7.2.				resident on the collection centre or other lower health status than the donor stallion	
				een carried out on samples taken on semen collection and at least at the begin	
		equired in point II.3.6.1 v was collected on		ast carried out on a sample of blood taker(4);	n not more than 120 days before
(1) either	-	required in point II.3.6.2	was la	ast carried out not more than 30 days bef	fore the semen was collected on

COUNTRY

Equine semen - Section C

II. Healt	h information	II.a. Certificate reference No	II.b.					
(¹) or		opositive stallion for equine viral arteritis was co an one year before the semen was collected on .						
II.3.7.3.		nave been carried out during the 30 days manufter the collection of the semen on samples take						
II.4.	The semen described above was collected, processed, stored and transported under conditions which comply with the requirements of Chapter II and III of Annex D to Directive 92/65/EEC.							
Notes								
Part I:								
Box I.11.:	The place of origin shall correspon	d to the semen collection centre of the semen or	igin.					
Box I.22.:	The number of packages shall core	respond to the number of containers.						
Box I.23.:	The identification of container and	seal number shall be indicated.						
Box I.28.:	The donor identity shall correspond	d to the official identification of the animal.						
	The date of collection shall be indic	cate in the following format: dd/mm/yyyy.						
Part II:								
(¹) Dele	ete as necessary.							
Reg	ulation (EU) 2018/659 provided the s	from a third country listed in column 2 of Annex demen was collected in the part of the territory on of the category of equidae indicated in column	of the third country detailed in					
equi equi	dae 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 been was collected.	has remained officially free of					
(4) Inse	rt date.							
(5) Cros	ss out the programmes that do not app	ly to the consignment.						
( <sup>6</sup> ) 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 vet	erinarian							
Nam	ne (in capital letters):		Qualification and title:					
Date	×		Signature:					
Stan	np:							

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### 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 El
	l.1.	Consignor Name	1.2.	Certificate reference No	I.2.a.
		Address	1.3.	Central competent authority	
		Tel.	1.4.	Local competent authority	
Part I : Details of dispatched consignment	1.5.	Consignee Name Address Postal code	1.6.	Person responsible for the load Name Address	in EU
tched		Tel.		Tel.	
s of dispa	1.7.	Country of ISO code I.8. Region of origin Code	1.9.	Country of ISO code destination	I.10. Region of Code destination
I : Detai	l.11.	Place of origin Semen centre □	I.12.	Place of destination Semen centre ☐ Ho	olding
Part		Name Approval number Address		Name Appro Address	val 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 Identification Documentary references	l.17.	No(s) of CITES	
	I.18.	Description of commodity	l	I.19. Commod	dity 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

Box I.17.:

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#### COUNTRY Equine semen - Section D Health information II.a. Certificate reference No I, the undersigned official veterinarian of the exporting country (2) ...... ..... hereby (name of exporting country) certify that: II.1. The centre (3) described in Box I.11 at which the semen to be exported to the Union was stored: Part II: Certification (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);] (1) or meets the conditions laid down in Chapter I(I) (2) and is operated and supervised in accordance with [II.1.1. the conditions laid down in Chapter I(II) (2) of Annex D to Directive 92/65/EEC;] 11.2. The semen to be exported to the Union: has been collected, processed and stored for a minimum period of 30 days immediately following collection in an II.2.1. approved semen collection centre (5) operated and supervised in accordance with Chapters I(I) (1) and I(II) (1) of Annex D to Directive 92/65/EEC, which is (1) either [located in the exporting country:] (1) or ....... (2), and has been imported to the exporting country under conditions at least as strict as for imports of semen of animals of the equine species into the Union in accordance with Directive 92/65/EEC;] 11.2.2. was moved to the centre described in Box I.11 under conditions at least as strict as described in: (1) either [Model 1 in Section A of Part 1 of Annex III to Regulation (EU) 2018/659 (6);] (1) or [Model 2 in Section B of Part 1 of Annex III to Regulation (EU) 2018/659 (6);] (1) or [Model 3 in Section C of Part 1 of Annex III to Regulation (EU) 2018/659 (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);] was stored under conditions which satisfy the terms of Annex D to Directive 92/65/EEC; 11.2.3. 11.2.4. sent to the place of loading in a sealed container in accordance with point 1.4 of Chapter III(I) of 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 storage centre of semen dispatch.

The serial number of the individual official document(s) or health certificate(s) that accompanied the semen 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 endorsed

copy/copies of thereof must be attached to this certificate.

Equine semen - Section D

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COUNTRY

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II.	Health i	nformation	II.a. Certificate reference No	II.b.					
Box	1.22.:	The number of packages shall corr	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 indic	cated in the following format: dd/mm/yyyy.						
Part	Part II:								
(1)	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) 2018/659 provided that the semen was collected in the part of the territory of the third country detailed in column 4 of that Annex from a donor stallion of the category of equidae indicated in column 11, 12 or 13 of that Annex.								
(3)	Only approved semen collection or storage centres listed in accordance with Article 17(3)(b) of Directive 92/65/EEC on the Commission website:								
	http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm								
(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).								
( <sup>5</sup> )		pproved semen collection centres lis Commission websites:	sted in accordance with Article 11(4) and Article	7(3)(b) of Directive 92/65/EEC					
		ec.europa.eu/food/animals/live_anir c.europa.eu/food/animal/semen_ova							
( <sup>6</sup> )	accom		the health certificate(s) or the officially endo from the approved semen collection centre of the must be attached to this certificate.						
_	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	:							

### PART 2

## Model health certificate for imports of ova and embryos

### Section A

COUN	TRY:					Veterinary certifica	ate to El
	l.1.	Consignor Name	1.2.	Certificate referen	ice No	I.2.a.	
		Address	1.3.	Central competen	t authority		
		Tel.	1.4.	Local competent a	authority		
Part I : Details of dispatched consignment	1.5.	Consignee Name Address	1.6.	Person responsible Name Address	le for the load i	in EU	
atched co		Postal code Tel.		Postal code Tel.			
ls of dispa	1.7.	Country of ISO code I.8. Region of Code origin	1.9.	Country of destination	ISO code I	l.10. Region of destination	Code
I : Detai	l.11.	Place of origin Embryo team □	I.12.	Place of destination	on Embryo team		
Part		Name Approval number Address		Name Address	Approval num	ber	
		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	I.17.				
	I.18.	Description of commodity			I.19. Commod	lity code (HS code) 05 11 99 85	
						I.20. Quantity	
	I.21.					I.22. Number of pack	ages
	1.23.	Seal/Container No				1.24.	
	1.25.	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		Third country ISO code					
	1.28.	Identification of the commodities					
	s	pecies (Scientific Category D	Oonor i	dentity D	ate of collection	n Quantity	′

	COUNTRY				Equine 6	ova/embryos			
	II. Health	information	,	II.a. Certificate reference No	II.b.				
	I, the under	signed, offic	ial veterinarian, of the e	exporting country (²)(/	name of exporting country)	hereby			
	certify that:	rtify that:							
II.1. The ova (¹)/embryos (¹) described above:									
Part II: Certification	II.1.2.	accordance		by the team (3) described in Box I.11, word Annex D to Directive 92/65/EEC (4) calendar year;					
Part	II.1.3.		ected (1)/produced (1), to Directive 92/65/EEC;	processed and stored in accordance	with the requirements of Cha	pter III(II) of			
	II.1.4.		ected at a place separa and disinfected prior to t	rated from other parts of the premises of the collection;	r holding which is in good rep	pair and was			
	II.1.5. were examined, processed and packed in laboratory facilities which are not situated in a zone subject to prohibition or quarantine measures as set out in Box II.1.6, in a section which is separated from the section for storing equipment and materials used in contact with donor animals and from the area where the donor animals are handled;								
	II.1.6.	come from	m donor mares which:						
	II.1.6.1. were continuously resident for a period of 3 months (or since entry if they were directly impor a Member State of the Union during the 3 months period) in the exporting country or, in the regionalisation in accordance with Article 13 of Directive 2009/156/EC(5), in that part of the te the exporting country which was during that period			the case of					
				d to be infected with African horse sictive 2009/156/EC,	ckness in accordance with A	rticle 5(2)(a)			
			<ul><li>free from Venez</li></ul>	ezuelan equine encephalomyelitis for a p	eriod of at least 2 years,				
			<ul> <li>free from glande</li> </ul>	ders and dourine for a period of at least	6 months;				
	(1) either [II.1.6.2. originated from a country of export which was on the day of collection free from vesicular (VS) for a period of at least 6 months;]				ar stomatitis				
	(¹) or [II.1.6.2. were subjected to a virus neutralisation test for vesicular stomatitis (VS) carried out with a ne result at a serum dilution of 1 in 32 or a VS ELISA carried out with a negative result in accor with the relevant Chapter of the Manual of Diagnostic Tests and Vaccines for Terrestrial Anim the OIE on a blood sample taken on				accordance al Animals of				
	(1) either	[II.1.6.3.	veterinary supervisio	he past 30 days prior to the date of the on which fulfilled from the day of the co th the conditions for a holding laid dow	llection of the ova (1)/embryos	s (1) until the			
	(¹) or	(¹) or [II.1.6.3. in the case of frozen ova (¹)/embryos (¹), during a period of the past 30 days prior to the date of the collection were kept in holdings under veterinary supervision which fulfilled, from the day of the collection of the ova (¹)/embryos (¹) until the end of the period of 30 days mandatory storage at approved premises, the conditions for a holding laid down in Article 4(5) of Directive 2009/156/EC, and in particular:]							

COUNTRY Equine ova/embryos

II.	Health information	1		II.a. Certificate reference No	II.b.
	(¹) either	[II.1.6.3.1.	suscep	g a case of a disease mentioned below n ible to that disease located in the holding wer has been free:	
			beg	n any type of equine encephalomyelitis for a jinning on the day on which the equidae s ughtered,	
			res sar	n equine infectious anaemia for at least the periult in an agar gel immunodiffusion test (AGID on ples taken after the infected animals were nonths apart from each of the remaining equidae	or Coggins tests) carried out on slaughtered on two occasions
			— from	n vesicular stomatitis for a period of at least 6 e,	months from the last recorded
			— fro	n rabies for a period of at least one month from t	he last recorded case,
			— fro	n anthrax for a period of at least 15 days from th	e last recorded case,]
	(¹) or	[II.1.6.3.1.	that disinfed enceph period followin	g a case of a disease mentioned below all the a sease located in the holding were slaughters ted, the holding was free for a period of at least alomyelitis, equine infectious anaemia, vesiculof at least 15 days in the case of anthrax, but to g the destruction of the animals the disinterily completed;	ed or killed and the premises 30 days from any type of equine lar stomatitis and rabies or a leginning on the day on which
	II.1.6.4.		e of the	past 30 days prior to the collection the ova (¹)/e equidae has shown clinical signs of contagious	
	II.1.6.5.	of the ova (1	)/embryc	ural breeding during a period of at least 30 days s (¹) and between the date of the first sample date of the collection of the ova (¹)/embryos (¹);	
	II.1.6.6.	Manual of Dia which is reco	agnostic gnised b	tests, which meet at least the requirements of Tests and Vaccines for Terrestrial Animals of the y the competent authority and has the tests refe ent to that provided for in Article 12 of Regul	e OIE, carried out in a laboratory rred to hereinafter included in its
		( <sup>8</sup> ) [II.1.6.6.1.	Coggin carried being referred	ine infectious anaemia (EIA), an agar-gel ir s test) or an enzyme-linked immunosorbent assaout on a blood sample taken on	y (ELISA) with a negative result
		II.1.6.6.2.	for con negativ point II	tagious equine metritis (CEM), an agent ident e result on at least two specimens (swabs) taker 1.6.5 from at least the mucosal surfaces of the of the donor mare	ification test carried out with a n during the period referred to in
		(¹) either	[II.1.6.6	on two occasions with an interval on	(6), in the case of isolation ivation under microaerophilic s, set up within 24 hours after inimal, or 48 hours where the

COUNTRY	Equine ova/embryos
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							Equilie ova/ellibi yos
II. Health	information		II.a.	Cert	ificate reference	No	II.b.
		(¹) and/or [II.1.	6.6.2.2.	gend or re	ome of <i>Taylorella</i>	a <i>equigenitalis</i> by a po arried out within 48 ho	, in the case of detection of olymerase chain reaction (PCR) ours after taking the specimens
		earli treat	er than ment of	7 days	s (systemic treat donor stallion a	tment) or 21 days (loc	6.6.2.2 were in no case taken cal treatment) after antimicrobial ansport medium with activated laboratory.
	II.1.6.7.						n contact with equidae suffering ays immediately preceding the
	II.1.6.8.	on the day of the contagious disease		n of t	he ova (¹)/embr	yos (¹) did not show o	clinical signs of an infectious or
II.1.7.		cted (¹)/produced (¹) as approved by the					production (1) team described in
II.1.8.	collection		and we	re tra	ansported under		30 days immediately after their atisfy the terms laid down in
II.2.	The embryos described above were conceived by artificial insemination(1)/as a result of <i>in vitro</i> fertilisation (¹) using semen meeting the requirements of Directive 92/65/EEC and coming from semen collection centres approved in accordance with Article 11(2) or 17(3)(b) of Directive 92/65/EEC (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) (¹¹);						
( <sup>12</sup> ) [II.3.						ed above comply with p in points II.1.1 to II.1	the requirements of Annex D to .8 of this certificate.]
Notes							
Part I:							
Box I.11.:	ova/embry		roduced	l, prod	cessed, stored a		production team by which the ordance with Article 17(3)(b) of
	http://ec.eu	uropa.eu/food/anima	l/semen	_ova/	equine/index_en	ı.htm	
Box I.22.:	The numb	er of packages shall	corresp	ond to	the number of	containers.	
Box I.23.:	The identif	ication of container	and seal	numb	ber shall be indic	cated.	
Box I.28.:		gory: specify if <i>in</i> pulated embryos.	<i>vivo</i> d	erived	d embryos, <i>in</i>	vivo derived ova, ir	n vitro produced embryos or
	The donor	identity shall corres	ond to	the of	ficial identificatio	n of the animal.	
	The date of	f collection shall be	indicate	in the	following forma	t: dd/mm/yyyy.	

COUNTRY Equine ova/embryos

II.	Health information	II.a. Certificate reference No	II.b.				
Part	11:						
( <sup>1</sup> )	Delete as appropriate.						
( <sup>2</sup> )	Implementing Regulation (EU) 2018/659,	itory of third countries listed in columns 2 and respectively from which imports of registered eq s indicated in column 14 of Annex I thereto.					
( <sup>3</sup> )	Only approved embryo collection teams and embryo production teams listed in accordance with Article 17(3)(b) of Directive 92/65/EEC on the Commission website:						
	http://ec.europa.eu/food/animal/semen_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).						
( <sup>5</sup> )	Council Directive 2009/156/EC of 30 N importation from third countries of equidae	November 2009 on animal health conditions (OJ L 192, 23.7.2010, p. 1).	governing the movement and				
( <sup>6</sup> )	Insert date. (follow Guidance in Part II of t	he Notes).					
( <sup>7</sup> )	Regulation (EC) No 882/2004 of the European Parliament and of the Council of 29 April 2004 on official controls performed to ensure the verification of compliance with feed and food law, animal health and animal welfare rules (OJ L 165, 30.4.2004, p. 1).						
(8)	The agar gel immunodiffusion test (AGID or Coggins test) or the ELISA for equine infectious anaemia are not required for donor equidae which have continuously resided in Iceland since birth, provided that Iceland has remained officially free 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.						
( <sup>9</sup> )	Only approved semen collection centres on the Commission websites:	isted in accordance with Article 11(4) or Article	17(3)(b) of Directive 92/65/EEC				
	https://ec.europa.eu/food/animals/live_animals/approved-establishments_en; http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm						
(10)	Imports of equine semen are authorised from third countries listed in column 2 of Annex 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.						
( <sup>12</sup> )	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						
	Name (in capital letters):		Qualification and title:				
	Date:		Signature:				
	Stamp:						

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### 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 certificat	te to EU
	l.1.	Consignor Name	1.2.	Certificate referen	ce No	I.2.a.	
				Central competen	t authority		
		Tel.	1.4.	Local competent a	authority		
ment	I.5. Consignee Name Address			Person responsible	le for the load i	in EU	
nsign				Address			
cp payo		Postal code Tel.		Postal code Tel.			
Part I : Details of dispatched consignment	1.7.	Country of ISO code I.8. Region of Code origin Code	1.9.	Country of destination	ISO code	I.10. Region of destination	Code
tails	1.11.	Place of origin	1.12	2. Place of destination	on		l
. De		Embryo team □		Holding	Embryo team		
Part		Name Approval number Address		Name Address	Approval num	ber	
		Postal code		Postal code			
	I.13.	Place of loading	1.14	I. Date of departure			
	I.15.	Means of transport	1.16	6. Entry BIP in EU			
		Aeroplane  Ship  Railway wagon					
		Road vehicle Other Identification	1.17	1.17.			
		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 packa	ges
	1.23.	Seal/Container No				1.24.	
	1.25.	Commodities certified for:  Artificial reproduction					
	I.26. For transit through EU to third country			I.27. For import or admission into EU			
		Third country ISO code					
	1.28.	Identification of the commodities					
	Species (Scientific Category Doname)			identity D	ate of collectio	n Quantity	

COUNTRY Equine ova/embryos

	II. Health i	information			II.a. Certificat	e reference No		II.b.
	I, the unders	igned, officia	al veterinari	an, of the exp	orting country (2	,	name of exporting	n country)
	certify that:							
	II.1.	The ova (1)	/embryos (	1) described a	bove:			
u.		II.1.2.	supervise	d in accordar	nce with Chapte		x D to Directive	hich has been approved and 92/65/EEC and is subject to
Part II: Certification		II.1.3.			duced (1), proc D to Directive 92		red in accordanc	e with the requirements of
Part II: (		II.1.4.				m other parts of to the collection;		olding which is in good repair
		II.1.5.	to prohibit section for	ion or quarant r storing equi	tine measures a	as set out in Box	II.1.6, in a section	not situated in a zone subject n which is separated from the r animals and from the area
		II.1.6.	come fron	n donor mares	which:			
			II.1.6.1.	Member Stat or, in the cas	e of the Europe e of regionalisa	ean Union during to	the 3 months pe	were directly imported from a riod) in the exporting country ctive 2009/156/EC (8), in that nat period
						e infected with of Directive 2009		ickness in accordance with
				— free fro	om Venezuelan e	equine encepha	lomyelitis for at le	ast 2 years,
				— free fro	m glanders and	I dourine for at le	east 6 months;	
		(1) either	[II.1.6.2.		om a country of at least 6 month		was on the day o	of collection free of vesicular
		(¹) or	[II.1.6.2.		( <sup>4</sup> ) w			itis on a blood sample taken ith negative result at a serum
		(¹) either	[II.1.6.3.	supervision v	which fulfilled from the conditions	om the day of c	ollection of ova (1)	in holdings under veterinary //embryos (¹) until the date of (5) of Directive 2009/156/EC,
		(¹) or	[II.1.6.3.	supervision w of frozen ov premises ela	vhich fulfilled fro a (1)/embryos	om the day of co (1), the period ditions for a h	ollection of ova (1), of 30 days man	in holdings under veterinary /embryos (¹) until, in the case datory storage at approved in Article 4(5) of Directive

II.b.

Health information

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

(¹) either	[II.1.6.3.1.	following a case of a disease mentioned below not all the animals of species susceptible to the disease located on the holding were slaughtered or killed and the holding has been free:
		<ul> <li>from any type of equine encephalomyelitis for at least 6 months, beginning on the day on which the equidae suffering from the disease are slaughtered,</li> </ul>
		<ul> <li>from equine infectious anaemia for at least the period required to obtain a negative result in an agar gel immunodiffusion test (Coggins tests) carried out on samples taken after the infected animals were slaughtered on two occasions 3 months apart from each of the remaining equidae;</li> </ul>
		<ul> <li>from vesicular stomatitis for at least 6 months from the last recorded case,</li> </ul>
		<ul> <li>from rabies for at least one month from the last recorded case,</li> </ul>
		<ul> <li>from anthrax for at least 15 days from the last recorded case,]</li> </ul>
(¹) or	[II.1.6.3.1.	following a case of a disease mentioned below all the animals of species susceptible to the disease located in the holding have been slaughtered or killed and the premises disinfected, the holding has been free for at least 30 days from any type of equine encephalomyelitis, equine infectious anaemia, vesicular stomatitis and rabies or 15 days in the case of anthrax, beginning on the day on which following the destruction of the animals the disinfection of the premises was satisfactorily completed;]
II.1.6.4.		past 30 days prior to collection have been kept in holdings each of them having rom clinical signs of contagious equine metritis for at least 60 days;
II.1.6.5.	collection	peen used for natural breeding during at least 30 days prior to the date of of ova or embryos and between the date of the first samples referred to in 6.6 and II.1.6.7 and the date of the collection of ova and embryos;
II.1.6.6.	test) or an on collection o	subjected with negative result to an agar-gel immuno-diffusion test (Coggins a ELISA for equine infectious anaemia carried out on a blood sample taken
II.1.6.7.	isolation of negative re the first col sinuses on and on an	in subjected to an agent identification test for contagious equine metritis by ff Taylorella equigenitalis after a cultivation of 7 to 14 days carried out with esults in each case on samples taken during the past 30 days prior to the date of election of ova or embryos from mucosal surfaces of the clitoral fossa and clitoral two consecutives oestrus periods on
II.1.6.8.	equidae su	of my knowledge and as far as I could ascertain, have not been in contact with iffering from an infectious or contagious disease during the 15 days immediately the collection;
II.1.6.9.		e day of collection of ova (¹)/embryos (¹) not shown clinical signs of an infectious ous disease;
		educed (¹) after the date on which the embryo collection (¹)/production (¹) team was approved by the competent authority of the exporting country;

COUNTRY Equine ova/embryos

II.	Health information	II.a. Certificate reference No	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 Chapter III(II) of Annex D to Directive 92/65/EEC;
- 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. (<sup>6</sup>) (<sup>7</sup>);
- 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:

- 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.

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COUN	ITRY		Equine ova/embryos			
II.	Health information	II.a. Certificate reference No	II.b.			
( <sup>6</sup> )	Only approved semen collection centres listed in accordance with Article 11(4) or Article 17(3)(b) of Council Directive 92/65/EEC on the Commission websites:					
	https://ec.europa.eu/food/animals/live_anim http://ec.europa.eu/food/animal/semen_ova					
( <sup>7</sup> )	Does not apply to ova.					
(8)	OJ L 192, 23.7.2010, p. 1.					
_	The signature and the stamp must be in a d	ifferent colour to that of the printing.				
Offic	cial veterinarian					
	Name (in capital letters):		Qualification and title:			
	Date:	5	Signature:			
	Stamp:					

#### PART 3

### **Explanatory notes for the certification**

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

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

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

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

#### ANNEX IV

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

- 1. 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.
- 2. 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)

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Status: This is the original version (as it was originally adopted).

### PART 3

# **Model Transhipment Manifest**

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

	Serial Number:
	Reference No of Air Cargo Transfer Manifest:(1)
Country where transhipment takes p	place:
Airport (2)/Port (2) of arrival:	
Date of arrival:	
Date of transhipment:	
Transferring Carrier:	
Receiving Carrier:	
Description of consignment:	Animal species: Total number of animals:
Serial No of Health Certificate	Remarks
took place under my supervision a	arian (²)/customs officer (²) at the above mentioned airport (²)/port (²) declare that the transhipment and in compliance with the following conditions:  ranshipment protected from attacks by insect vectors of diseases transmissible to equidae;
(a) the equidae were daring the th	anomphion protocola from attacke by mood vocation of allocated standings to equidate,
(b) the equidae did not come into	contact with equidae of a different health status;
	talls and the surrounding airspace in the transport compartment were sprayed with an appropriate n with an insecticide immediately after the closing of the doors of the aircraft (²)/vessel (²).
The consignment has been transh	nipped in full and apparent good order and conditions except as noted in the "Remarks" column.
Done at	on
	Stamp
(signature of the official veter	I
(name in capital le	etters and title)
(¹) Keep empty if transhipment from v (²) Delete as appropriate	/essel to vessel

- (1) OJ L 268, 24.9.1991, p. 56.
- (2) OJ L 268, 14.9.1992, p. 54.
- (**3**) OJ L 192, 23.7.2010, p. 1.
- (4) Commission Decision 2004/211/EC of 6 January 2004 establishing the list of third countries and parts of territory thereof from which Member States authorise imports of live equidae and semen, ova and embryos of the equine species, and amending Decisions 93/195/EEC and 94/63/EC (OJ L 73, 11.3.2004, p. 1).
- (5) Council Directive 2008/73/EC of 15 July 2008 simplifying procedures of listing and publishing information in the veterinary and zootechnical fields and amending Directives 64/432/EEC, 77/504/EEC, 88/407/EEC, 88/661/EEC, 89/361/EEC, 89/556/EEC, 90/426/EEC, 90/427/EEC, 90/428/EEC, 90/429/EEC, 90/539/EEC, 91/68/EEC, 91/496/EEC, 92/35/EEC, 92/65/EEC, 92/66/EEC, 92/119/EEC, 94/28/EC, 2000/75/EC, Decision 2000/258/EC and Directives 2001/89/EC, 2002/60/EC and 2005/94/EC (OJ L 219, 14.8.2008, p. 40).
- (6) http://ec.europa.eu/food/animals/semen/equine en
- (7) Commission Regulation (EU) No 176/2010 of 2 March 2010 amending Annex D to Council Directive 92/65/EEC as regards semen collection and storage centres, embryo collection and production teams, and conditions for donor animals of the equine, ovine and caprine species and for handling semen, ova and embryos of those species (OJ L 52, 3.3.2010, p. 14).
- (8) Commission Decision 2010/470/EU of 26 August 2010 laying down model health certificates for trade within the Union in semen, ova and embryos of animals of the equine, ovine and caprine species and in ova and embryos of animals of the porcine species (OJ L 228, 31.8.2010, p. 15).
- (9) Decision 2002/309/EC, Euratom of the Council, and of the Commission as regards the Agreement on Scientific and Technological Cooperation of 4 April 2002 on the conclusion of seven Agreements with the Swiss Confederation (OJ L 114, 30.4.2002, p. 1).
- (10) OJ L 71, 18.3.1999, p. 3.
- (11) Council Decision 1999/201/EC of 14 December 1998 on the conclusion of the Agreement between the European Community and the Government of Canada on sanitary measures to protect public and animal health in respect of trade in live animals and animal products (OJ L 71, 18.3.1999, p. 1).
- (12) OJ L 57, 26.2.1997, p. 5.
- (13) Council Decision 97/132/EC of 17 December 1996 on the conclusion of the Agreement between the European Community and New Zealand on sanitary measures applicable to trade in live animals and animal products (OJ L 57, 26.2.1997, p. 4).
- (14) Commission Regulation (EC) No 180/2008 of 28 February 2008 concerning the Community reference laboratory for equine diseases other than African horse sickness and amending Annex VII to Regulation (EC) No 882/2004 of the European Parliament and of the Council (OJ L 56, 29.2.2008, p. 4).
- (15) Commission Decision 97/794/EC of 12 November 1997 laying down certain detailed rules for the application of Council Directive 91/496/EEC as regards veterinary checks on live animals to be imported from third countries (OJ L 323, 26.11.1997, p. 31).
- (16) Report of the Scientific Veterinary Committee on Equine Viral Arteritis, 12 December 1994, VI/4994/94 Rev. 4.
- (17) Commission Decision 95/329/EC of 25 July 1995 defining the categories of male equidae to which the requirement regarding viral arteritis laid down in Article 15 (b) (ii) of Council Directive 90/426/EEC applies (OJ L 191, 12.8.1995, p. 36).
- (18) http://www.oie.int/index.php?id=169&L=0&htmfile=chapitre eav.htm
- (19) Commission Decision 2003/24/EC of 30 December 2002 concerning the development of an integrated computerised veterinary system (OJ L 8, 14.1.2003, p. 44).
- (20) Commission Decision 2004/292/EC of 30 March 2004 on the introduction of the Traces system and amending Decision 92/486/EEC (OJ L 94, 31.3.2004, p. 63).

- (21) Commission Decision 2009/821/EC of 28 September 2009 drawing up a list of approved border inspection posts, laying down certain rules on the inspections carried out by Commission veterinary experts and laying down the veterinary units in Traces (OJ L 296, 12.11.2009, p. 1).
- (22) Commission Decision 2007/240/EC of 16 April 2007 laying down new veterinary certificates for importing live animals, semen, embryos, ova and products of animal origin into the Community pursuant to Decisions 79/542/EEC, 92/260/EEC, 93/195/EEC, 93/196/EEC, 93/197/EEC, 95/328/EC, 96/333/EC, 96/539/EC, 96/540/EC, 2000/572/EC, 2000/585/EC, 2000/666/EC, 2002/613/EC, 2003/56/EC, 2003/779/EC, 2003/804/EC, 2003/858/EC, 2003/863/EC, 2003/881/EC, 2004/407/EC, 2004/438/EC, 2004/595/EC, 2004/639/EC and 2006/168/EC (OJ L 104, 21.4.2007, p. 37).
- (23) Commission Regulation (EC) No 282/2004 of 18 February 2004 introducing a document for the declaration of, and veterinary checks on, animals from third countries entering the Community (OJ L 49, 19.2.2004, p. 11).
- (24) Commission Decision 93/444/EEC of 2 July 1993 on detailed rules governing intra-Community trade in certain live animals and products intended for exportation to third countries (OJ L 208, 19.8.1993, p. 34).
- (25) Commission Decision 92/260/EEC of 10 April 1992 on animal health conditions and veterinary certification for temporary admission of registered horses (OJ L 130, 15.5.1992, p. 67).
- (26) Commission Decision 93/195/EEC of 2 February 1993 on animal health conditions and veterinary certification for the re-entry of registered horses for racing, competition and cultural events after temporary export (OJ L 86, 6.4.1993, p. 1).
- (27) Commission Decision 93/196/EEC of 5 February 1993 on animal health conditions and veterinary certification for imports of equidae for slaughter (OJ L 86, 6.4.1993, p. 7).
- (28) Commission Decision 93/197/EEC of 5 February 1993 on animal health conditions and veterinary certification for imports of registered equidae and equidae for breeding and production (OJ L 86, 6.4.1993, p. 16).
- (29) Commission Decision 94/699/EC of 19 October 1994 providing for less frequent identity and physical checks on the temporary admission of certain equidae from Sweden, Norway and Finland and repealing Decision 93/321/EEC (OJ L 280, 29.10.1994, p. 88).
- (30) Commission Decision 2003/13/EC of 10 January 2003 on the temporary admission of horses participating in the pre-Olympic test event in Greece in 2003 (OJ L 7, 11.1.2003, p. 86).
- (31) Commission Decision 2004/177/EC of 20 February 2004 on the temporary introduction of registered horses participating in the Olympic Games or the Paralympic Games in Greece in 2004 (OJ L 55, 24.2.2004, p. 64).
- (32) Commission Decision 2010/57/EU of 3 February 2010 laying down health guarantees for the transit of equidae being transported through the territories listed in Annex I to Council Directive 97/78/EC (OJ L 32, 4.2.2010, p. 9).
- (33) Commission Decision 2010/471/EU of 26 August 2010 on imports into the Union of semen, ova and embryos of animals of the equine species as regards lists of semen collection and storage centres and embryo collection and production teams and certification requirements (OJ L 228, 31.8.2010, p. 52).
- (34) 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).
- (35) 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).