This text is meant purely as a documentation tool and has no legal effect. The Union's institutions do not assume any liability for its contents. The authentic versions of the relevant acts, including their preambles, are those published in the Official Journal of the European Union and available in EUR-Lex. Those official texts are directly accessible through the links embedded in this document

COMMISSION IMPLEMENTING REGULATION (EU) 2018/659

of 12 April 2018

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

(Text with EEA relevance)

(OJ L 110, 30.4.2018, p. 1)

Amended by:

►<u>B</u>

Official Journal

						No	page	date
► <u>M1</u>	Commission Implementing 27 September 2018	Regulation	(EU)	2018/1301	of	L 244	10	28.9.2018
► <u>M2</u>	Commission Implementing 28 November 2019	Regulation	(EU)	2019/2147	of	L 325	99	16.12.2019
► <u>M3</u>	Commission Implementing Reg	gulation (EU)	2020/485	5 of 2 April 2	2020	L 103	10	3.4.2020

Corrected by:

▶<u>C1</u> Corrigendum, OJ L 228, 11.9.2018, p. 2 (2018/659)

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

COMMISSION IMPLEMENTING REGULATION (EU) 2018/659

of 12 April 2018

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

(Text with EEA relevance)

SECTION 1

Subject matter, scope and definitions

Article 1

Subject matter and scope

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

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

Article 2

Definitions

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

- (a) 'regionalisation' means the official recognition of a part of the territory of a third country with precise geographical delimitations, containing an equidae subpopulation with a distinct health status with respect to one or more specific diseases and subject to appropriate surveillance, disease control and biosecurity measures;
- (b) 'identification document' means any document which may be used to prove the identity of an equine animal and which includes at least the following information:
 - (i) a narrative describing the animal and recording its marks depicted in a completed outline diagram;
 - (ii) a reference to specific marks, characteristics or identifiers which establish an unambiguous link between the animal and the document;
 - (iii) the information set out in points 1, 2, 3 and 6 to 10 of Part A and in points 12 to 18 in Part B of Section 1 of Annex I to Commission Implementing Regulation (EU) 2015/262 (¹);

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

(c)	'registered horse'	means an animal of the species <i>Equus</i> caballus registered as defined in Council Directive $90/427/\text{EEC}$ (¹), identified by means of an identification document issued by:
		 (i) the breeding authority or any other competent authority of the country where the animal originated which manages the studbook or register for that breed of animal; or
		 (ii) any international association or organis- ation which manages horses for competi- tion or racing;
(d)	'entry'	means the action of moving equidae or their semen, ova or embryos into one of the territories listed in Annex I to Council Directive $97/78/EC$ (²);
(e)	'type of entry'	means respectively the temporary admission, the re-entry after temporary export, imports and transit;
(f)	'temporary admission'	means the status of a registered horse originat- ing in a third country and moved into the Union territory for a period of less than 90 days;
(g)	'temporary export'	means the movement of a registered horse out of the Union for a period of less than 90 days;
(h)	're-entry'	means the movement of a registered horse from a third country into the Union after temporary export from the Union;
(i)	'imports'	means the movement of a consignment of equidae or their semen, ova or embryos into the Union for an undetermined period;
(j)	'transit'	means the movement of a consignment of equidae across Union territory by road, rail or waterway transport from one third country to another or from one part of the territory of a third country to another part of the territory of the same third country;
(k)	'border inspection post'	means any inspection post as defined in Article $2(2)(f)$ of Directive $91/496/EEC$ and Article $2(2)(g)$ of Directive $97/78/EC$ and approved for the commodity concerned in accordance with Decision $2009/821/EC$;
(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;

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

^{(&}lt;sup>2</sup>) Council Directive 97/78/EC of 18 December 1997 laying down the principles governing the organisation of veterinary checks on products entering the Community from third countries (OJ L 24, 30.1.1998, p. 9).

(m)	'ova'	means the haploid stages of the ootidogenesis including secondary oocytes and ova;
(n)	'operator'	means any natural or legal person subject to one or more of the rules provided for in this Regulation who has equidae or their germinal products under its responsibility;
(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 biose- curity rules under the control of the veterinary authority;
(q)	'vector-protected quarantine'	means the quarantine of equidae which
		(i) is carried out on dedicated premises that are:
		 screened against the intrusion of relevant vectors,
		 included in a system of vector surveillance within the premises and of measures to limit the presence of relevant vectors around the premises;
		 (ii) may include exercise of the quarantined animal under official supervision during the vector-low period of the day and subject to application of insecticides and insect repellents and where possible 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 (¹),
		 in which a vector surveillance system is operated,
		 where Standard Operating Procedures, including description of back-up and alarm systems, are implemented for the operation of the quarantine and the transport of equidae to the place of loading,
(s)	'TRACES'	means the integrated computerised veterinary system provided for in Decisions 2003/24/EC and 2004/292/EC.

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

2. Where equidae are transported by sea, the period of 10 days referred to in paragraph 1 shall be extended by the time of transport on sea.

SECTION 4

Transport requirements for entry of equidae into the Union

Article 8

General animal health requirements

1. The operator responsible for a consignment of equidae intended for entry into the Union shall ensure that those equidae are transported in compliance with the following:

- (a) the equidae are transported by a means of transport carrying only equidae that are destined for the Union or alternatively are accompanied by a health certificate required for transit;
- (b) the equidae are transported by a means of transport carrying only equidae of the same certified health status, except where otherwise authorised in the specific animal health requirements set out in Sections A and B of Part 1 and in Section A of Part 3 of Annex II;
- (c) the equidae are transported by road or railway or moved on foot only in a third country or a part of the territory of a third country that is authorised for at least one type of entry of at least one category of equidae.

2. The operator responsible for a consignment of equidae intended for entry into the Union shall ensure compliance with the following:

- (a) the crates, containers, stalls or jet-stalls and the means of transport or the transport compartment of the means of transport in which equidae will be transported are cleansed and disinfected prior to loading of the animals with a disinfectant officially recognised in the country of dispatch;
- (b) the means of transport used for road or railway transport are designed, constructed and operated to prevent the escape of faeces, urine and fodder during the intended journey;
- (c) measures to protect the animals from attacks of insect vectors shall be applied in case of the occurrence of one of the following diseases:

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

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

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

Article 9

Specific animal health requirements for transport by air

1. The operator responsible for a consignment of equidae intended for entry into the Union by air shall ensure compliance with the following:

- (a) the crates, containers or jet-stalls and the surrounding airspace in the transport compartment are sprayed with an appropriate insect repellent in combination with an insecticide immediately after the closing of the doors of the aircraft;
- (b) the captain of the aircraft completes and signs the declaration set out in Part 1 of Annex V.

2. By way of derogation from paragraph 3 of Article 8, Member States may, on request of the operator of the consignment, authorise direct transhipment from one aircraft to another aircraft which takes place in a country not listed in Annex I, provided that the following requirements are satisfied:

- (a) the transhipment is carried out in the same airport within the area of the same customs office under direct supervision of an official veterinarian or the responsible customs officer;
- (b) during the transhipment the equidae are protected from attacks by insect vectors of diseases transmissible to equidae;
- (c) the equidae do not come into contact with equidae of a different health status;
- (d) the measures provided for in points (a) and (b) of paragraph 1 are applied in relation to the aircraft to be used for onward travel;
- (e) compliance with the conditions set out in point (a) of paragraph 1 and in points (a), (b) and (c) of this paragraph is certified by the official veterinarian or the responsible customs officer in the Transhipment Manifest drawn up in accordance with the model set out in Part 3 of Annex V.

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

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

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

2. The competent authority of the third country dispatching equidae which are destined for the Union shall ensure that the laboratory tests provided for in the health certificates set out in Annex II for African horse sickness are carried out in accordance with Annex IV to Directive 2009/156/EC.

3. The competent authority of the third country dispatching equidae or semen, ova or embryos of equidae, which are destined for the Union shall ensure compliance with the following:

- (a) the tests referred to in paragraphs 1 and 2 are carried out in a laboratory recognised by the competent authority in the third country of dispatch;
- (b) the details of sampling and the results of the tests are stated as required in the health certificate set out for the consignment concerned in Annex II or III based on the laboratory report made available to the certifying official veterinarian.

Article 12

Testing upon arrival in the Union

1. Where a test carried out, in or on behalf of the Member State of entry, on a sample taken in accordance with Article 4 of Decision 97/794/EC does not confirm the result of a laboratory test attested in a health certificate accompanying equidae or semen, ova or embryos of equidae arriving in the Union, as set out in Annex II or III to this Regulation, the competent authority of that Member State of entry shall ensure that the test is repeated in the national reference laboratory designated for the disease concerned in accordance with Article 4(1) of Regulation (EC) No 882/2004 of the European Parliament and of the Council (¹).

2. Where the measures provided for in paragraph 1 do not result in a conclusive outcome of the checks for compliance carried out in accordance with Article 4 of Decision 97/794/EC, the competent authority referred to in paragraph 1 shall ensure that the sample referred to in that paragraph is subjected to definitive testing as follows:

(a) for African horse sickness, in the European Union reference laboratory for African horse sickness designated in accordance with Council Directive 92/35/EEC (²);

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

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

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

Article 13

Application of vaccines and recording of vaccination

1. The competent authority of the third country dispatching equidae or semen, ova or embryos of equidae, which are destined for the Union, shall ensure that the vaccination attested in any of the certificates set out in Annexes II or III is carried out in compliance with the following:

- (a) the vaccination is carried out in accordance with the manufacturers' instructions or national legislation, whatever is stricter;
- (b) the vaccination is carried out using a licensed vaccine which meets at least the requirements for safety, sterility and efficacy set out for the vaccine concerned in the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals, latest edition, of the World Organisation for Animal Health (OIE).

2. Where the competent authority of a third country attests that a positive laboratory finding in a serological test for African horse sickness is related to previous vaccination, the vaccination shall be documented in the identification document accompanying the equine animal, where such identification document is available.

Article 14

Requirements relating to equine viral arteritis

1. Uncastrated male equidae intended for entry into the Union, with the exception of those listed in point 1 of Annex IV, shall be subject to tests for equine viral arteritis to ascertain that their semen is free of equine arteritis virus.

2. Vaccination against equine viral arteritis, including the testing required in accordance with point 1(a) of Annex IV, shall be carried out under official veterinary supervision.

3. Vaccination against equine viral arteritis shall be valid where there is documented proof accompanying the equine animal of an uninterrupted history of a primary course carried out in compliance with one of the vaccination protocols provided for in point 1(a) of Annex IV and regular revaccination according to manufacturers' recommendations and in any event at intervals of not more than 12 months.

SECTION 6

Identification of equidae intended for entry into the Union

Article 15

Identification of equidae intended for entry into the Union

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

That identification shall:

- (a) either comply with the requirements of Article 14 of Implementing Regulation (EU) 2015/262; or
- (b) provide at least the information detailed in points 1, 2, 3 and 6 to 10 of Part A and in points 12 to 18 of Part B of Section I of Part 1 of Annex I to that Regulation.

2. Equidae for slaughter to be imported into the Union shall be marked individually with an electronic transponder or an ear tag, the number of which shall be recorded in the health certificate accompanying the animals during transport.

3. Equidae for slaughter to be imported into the Union shall bear on their left front hoof a clear and indelible hot-branded 'S' of the size not less than half the length of the hoof wall, in the following cases:

- (a) if they are marked individually, by way of derogation from paragraph 2, by an alternative method indicated in the health certificate, in which case the animals must be consigned to the slaughterhouse of destination in accordance with Article 21(a);
- (b) if they are destined to be consigned to the slaughterhouse of destination in accordance with Article 21(b).

SECTION 7

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

Article 16

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

1. Provided compliance with the entry conditions has been established, the competent authority at the border inspection post of entry shall:

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

- (iii) the competent authorities responsible for the places of temporary residence indicated in the declaration by the owner or the representative of the owner of the registered horse accompanying the health certificate referred to in Article 3(1)(a);
- (c) deliver at least one copy of the CVED to the operator identified as 'operator responsible for the consignment' in Box I.7 of the CVED referred to in paragraph 1(b).

2. Where a registered horse is to be moved from one Member State to another Member State during its temporary admission, the competent authority of the place of dispatch shall:

- (a) provided the animal health conditions of Articles 4 and 5 of Directive 2009/156/EC are fulfilled, issue a health certificate in accordance with Annex III to Directive 2009/156/EC either for an individual registered horse or for a consignment of registered horses of the same origin and with the same destination, and enter in Box 1.6 of that certificate a reference to the health certificate referred to in Article 3(1)(a) of each temporarily admitted registered horse forming the consignment and a reference to the CVED referred to in point (i) of paragraph 1(b);
- (b) inform, through TRACES, the competent authority at the place of destination, of the movement of a registered horse to that Member State, and request the verification of arrival by completing a further Part III of the CVED referred to in point (i) of paragraph 1(b);
- (c) deliver to the operator, as identified in Box I.7 of the CVED referred to in point (i) of paragraph 1(b), a new print of the CVED displaying the Part III added in accordance with point (b) of this paragraph;
- (d) invalidate or withdraw any print of the CVED delivered to the operator in accordance with paragraph 1(c), or, if there had been a previous movement to another Member State, in accordance with point (c) of this paragraph.

3. The competent authority of the place of destination referred to in point (i) of paragraph 1(b) and in paragraph 2(b) shall acknowledge through TRACES the arrival of the registered horse and document the checks carried out by completing Part III of the CVED.

4. At the end of the temporary admission, the competent authority referred to in points (i) or (iii) of paragraph 1(b) which certifies the temporarily admitted registered horse to the third country of origin or to another third country, shall:

- (a) inform, through TRACES, the border inspection post of exit of the departure of the temporarily admitted registered horse from the Union, by completing a further Part III of the CVED referred to in point (i) of paragraph 1(b);
- (b) deliver to the operator, as identified in Box I.7 of the CVED referred to in point (i) of paragraph 1(b), a new print of the CVED displaying the Part III added in accordance with point (a) of this paragraph;

- (c) where the border inspection post of exit is situated in another Member State,
 - (i) issue, in accordance with Decision 93/444/EEC, a certificate in accordance with Annex III to Directive 2009/156/EC either for an individual registered horse or for a consignment of registered horses of the same origin and with the same destination;
 - (ii) enter in Box I.6 of the certificate referred to in point (i) a reference to the health certificate referred to in Article 3(1)(a) of each temporarily admitted registered horse forming the consignment and a reference to the CVED referred to in point (i) of paragraph 1(b).

5. The border inspection post of exit referred to in point (a) of paragraph 4 shall document the termination of the temporary admission of the registered horse by completing Part III of the CVED accordingly.

6. Where the temporary admission of a registered horse has not been terminated in accordance with paragraph 5 within a period of less than 90 days following the date of issue of the CVED referred to in point (i) of paragraph 1(b), an alert is sent automatically through TRACES to the border inspection post of entry and the competent authorities referred to in this Article until those competent authorities have determined the status of the registered horse.

Article 17

Operator responsibilities for temporarily admitted registered horses

1. The operator responsible for a registered horse temporarily admitted into the Union, as identified in Box I.7 of the CVED referred to in point (i) of Article 16(1)(b), shall ensure that the following conditions are met:

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

2. The operator referred to in paragraph 1 shall remain responsible for the movement of the registered horse during its temporary admission in the Union, and in particular shall inform:

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

Article 18

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

1. Registered horses temporarily admitted into the Union may be authorised for re-entry after temporary export to a third country or part of the territory of a third country authorised for the re-entry of registered horses to take part in specific races, competitions or cultural events for which model health certificates for re-entry into the Union are laid down in accordance with Article 20(3), provided that the re-entry into the Union takes place within a period of less than 90 days following the date of issuing of the CVED referred to in point (i) of Article 16(1)(b).

2. In order to allow the re-entry of a registered horse referred to in paragraph 1, the competent authority referred to in points (i) and (iii) of Article 16(1)(b) issuing the certificate for the temporary export shall:

- (a) apply the measures provided for in points (a), (b) and, where applicable, (c) of Article 16(4);
- (b) inform, through TRACES, the border inspection post of scheduled re-entry by completing Part III of the CVED;
- (c) deliver to the operator as identified in Box I.7 of the CVED referred to in point (i) of Article 16(1)(b), a new print of the CVED displaying the Part III added in accordance with point (b) of this paragraph;
- (d) invalidate or withdraw any print of the CVED delivered in accordance with Article 16(1)(c) or, if there had been a previous movement to another Member State, in accordance with Article 16(2)(c).
- 3. The border inspection post of re-entry shall:
- (a) retain the original of the health certificate referred to in Article 3(1)(c);

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

4. Following the re-entry after temporary export of a temporarily admitted registered horse in accordance with paragraph 1, the rules laid down in Article 16 apply for the remaining period of less than 90 days following the date of issuing of the CVED referred to in point (i) of Article 16(1)(b).

Article 19

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

1. Where the operator, as identified in Box I.7 of the CVED referred to in point (i) of Article 16(1)(b), submits an application to the competent authority referred to in point (i) or (iii) of Article 16(1)(b) or in Article 16(2)(b), to convert the temporary admission of a registered horse into a permanent entry, a Member State may authorise that conversion provided that the following requirements are met:

- (a) in accordance with Annex I, imports of registered horses are authorised from the third country or part of the territory of the third country concerned;
- (b) the competent authority responsible for the place of temporary residence has complied with the following conditions:
 - (i) that competent authority has carried out with satisfactory results the checks necessary to verify compliance with the test and vaccination requirements for imports of registered horses from the third country or part of the territory of the third country concerned set out in Part 3 of Annex II;

(ii) that competent authority has ensured that the registered horse remained under official veterinary supervision in that Member State until 3 months have elapsed from the date of its entry into the Union indicated on the CVED referred to in point (i) of Article 16(1)(b).

2. The competent authority referred to in paragraph 1, or a border inspection post designated for this task by the Member State, shall:

- (a) terminate the temporary admission in TRACES by choosing 'Conversion into permanent entry' in Part III of the CVED delivered to the operator in accordance with either Article 16(1)(c), or, if there had been a previous movement to another Member State, with Article 16(2)(c) or, if there had been a previous re-entry after temporary export, with Article 18(3)(c);
- (b) deliver to the operator identified in Box I.7 of the CVED referred to in point (i) of Article 16(1)(b), a new print of the CVED referred to in point (a), or a new CVED, in which 'For internal market' is checked in Box I.21;
- (c) invalidate or withdraw any print of the CVED delivered to the operator in accordance with either Article 16(1)(c), or, if there had been a previous movement to another Member State, with Article 16(2)(c) or, if there had been a previous re-entry after temporary export, with Article 18(3)(c);
- (d) invalidate or withdraw the original of the health certificate referred to in Article 3(1)(a).

3. During the period of conversion, the operator, as identified in Box I.7 of the CVED issued in accordance with point (i) of Article 16(1)(b) or Article 18(3)(b) of the registered horse shall take the following measures:

- (a) arrange regular visits carried out and recorded by a veterinarian to check the registered horse for clinical signs of possible infectious diseases;
- (b) keep records on the movement of the registered horse and on movements of equidae on and off the holding where it is kept;
- (c) complete the customs procedures, as referred to in Article 15 of Implementing Regulation (EU) 2015/262;
- (d) make an application in accordance with Article 15(1) of Implementing Regulation (EU) 2015/262 for the issuing of an identification document or the adaptation of an existing identification document.

4. In the case of death or loss of a registered horse temporarily admitted into the Union, the competent authority of the place of death or loss, where required by the Member State concerned in close collaboration with a border inspection post, shall:

 (a) terminate the temporary admission in TRACES by choosing 'Death/Loss' in Part III of the CVED referred to in point (i) of Article 16(1)(b) or Article 18(3)(b);

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

Article 20

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

1. Member States shall authorise the re-entry of registered horses subject to compliance with the following conditions:

- (a) the registered horse has remained outside the Union for not more than 30 days, unless specifically provided for in paragraph 3;
- (b) the registered horse has neither been resident in nor transited on land through any third country or part of the territory of a third country that is not assigned to the same sanitary group as the third country or part of the territory of a third country in which the health certificate in accordance with Section A of Part 2 of Annex II has been signed by the official veterinarian;
- (c) the health certificate for temporary export signed by the official veterinarian in the Member State of origin, or an authorised copy thereof, is presented on request of the border inspection post of reentry into the Union.

2. The competent authority certifying a registered horse for temporary export to a third country shall ensure that in application of Article 2(1) of Decision 93/444/EEC the registered horse is accompanied until the exit point in another Member State by a health certificate in accordance with Annex III to Directive 2009/156/EC.

3. The re-entry after temporary export for a period of more than 30 days of registered horses taking part in specific races, competitions or cultural events is subject to specific animal health requirements as contained in the corresponding model health certificates provided for Section B of Part 2 of Annex II in respect of the relevant event.

4. The operator, as identified in Box I.7 of the CVED, responsible for the consignment shall ensure that during the temporary export the registered horse neither has been resident in nor has transited on land through any third country or part of the territory of a third country that is not assigned to the same sanitary group as the third country or part of the territory of a third country in which the health certificate in accordance with Section A of Part 2 of Annex II has been signed by the official veterinarian.

Article 21

Specific animal health conditions regarding imports of equidae for slaughter

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

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

SECTION 8

Transitional and final provisions

Article 22

Transitional provisions

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

Article 23

Repeals

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

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

Article 24

Entry into force and applicability

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

It shall apply from 1 October 2018.

▼M2

However, Article 16(1)(b)(iii), Article 16(2)(b), (c) and (d), Article 16(3), Article 16(4)(a) and (b), Article 16(5), Article 17(1)(d) and Article 19(2)(a) shall apply from 14 December 2019.

▼<u>B</u>

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

ANNEX I

▼<u>M2</u>

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

		Code of the			TA	Re-En		Imports		Imports			Transit	Specific conditions	
ISO- Code	Third country	part of the territory of the third	Description of the part of the territory of the third country	SG	RH	RH	RH	ES	RE +		SEMEN		O/E	Equi- dae	
		country							EBP	RH	RE	EBP			
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16
AE	United Arab Emirates	AE-0	Whole country	Е	Х	X	Х	_	—	Х	_	_	Х	X	
AR	Argentina	AR-0	Whole country	D	Х	X	Х	x	Х	Х	X	Х	Х	X	
AU	Australia	AU-0	Whole country	А	Х	X	Х	X	Х	Х	X	Х	х	Х	
BA	Bosnia and Herze- govina	BA-0	Whole country	В	Х	X	Х	_	—	_	—	_	_	Х	
BB	Barbados	BB-0	Whole country	D	Х	X	Х	_		_				X	
BH	Bahrain	BH-0	Whole country	Е	Х	X	Х	_		_				Х	
BM	Bermuda	BM-0	Whole country	D	Х	X	Х	_						Х	
BO	Bolivia	BO-0	Whole country	D	Х	X	Х	_	_	_	_	_	_	Х	
BR	Brazil	BR-0	Whole country	_			_	_					_		
		BR-1	The states of:	_											
			Paraná and Rio de Janeiro	D	Х	X	Х	_						X	
BY	Belarus	BY-0	Whole country	В	Х	X	Х	х	Х					Х	
СА	Canada	CA-0	Whole country	C	Х	X	Х	x	Х	Х	X	Х		X	

02018R0659 - EN - 06.04.2020 - 003.001 - 20

▼	M2
---	----

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

▼	M2
---	-----------

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

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

2															
		Code of the			TA	Re-En		Imports			Imp	orts		Transit	Specific conditions
ISO- Code	Third country	part of the territory of the third	Description of the part of the territory of the third country	SG	RH	RH	RH	ES	RE +		SEMEN		O/E	Equi- dae	
		country							EBP	RH	RE	EBP			
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16
MX	Mexico	MX-0	Whole country	С	_	_	_	-	_	_	_	_	_	-	
		MX-1	Metropolitan area of Mexico- City	С		х								_	Only if certified in accordance with Chapter 1 of Section B of Part 2 of Annex II
		MX-2	The whole country except the States of Chiapas, Oaxaca, Tabasco, Campeche, Yucatan, Quintana Roo, Veracruz and Tamaulipas	С	Х	Х	Х	_	Х					_	
NZ	New Zealand	NZ-0	Whole country	А	Х	X	Х	X	X	_	_	_	_	X	
ОМ	Oman	OM-0	Whole country	Е	Х	X	Х	_	_	_	_	_	_	X	
PE	Peru	PE-0	Whole country	_	_	_	_	_	_	_	_	_	_	_	
		PE-1	Region of Lima	D	Х	Х	Х	_	_		_		_	X	
РМ	St Pierre & Miquelon	PM-0	Whole country	А	_		Х	_	X					Х	
PY	Paraguay	PY-0	Whole country	D	Х	X	Х	X	X	_	_	_	_	X	
QA	Qatar	QA-0	Whole country	Е	Х	X	Х	_	_	Х	_	_	_	X	
RS	Serbia (⁵)	RS-0	Whole country	В	Х	Х	Х	X	X	_	_	_	_	X	

02018R0659 - EN - 06.04.2020 - 003.001 - 24

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

V IVIZ																
			Code of the			TA	Re-En		Imports			Imp	orts		Transit	Specific conditions
	ISO- Code	Third country	part of the territory of the third	Description of the part of the territory of the third country	SG	RH	RH	RH	ES	RE +		SEMEN		O/E	Equi- dae	
			country							EBP	RH	RE	EBP			
	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16
			SA-2	Protection and surveillance zones in the provinces of Jizan, Asir and Najran as described in BOX 3		_								_		
	SG	Singapore	SG-0	Whole country	G	Х	X	Х	_	_					Х	
▼ <u>M3</u>																
	TH	Thailand	TH-0	Whole country	G											
▼ <u>M2</u>																
	TN	Tunisia	TN-0	Whole country	Е	X	X	Х	Х	Х	—	—	—	—	Х	
	TR	Turkey	TR-0	Whole country	_			_	—	—	—				—	
			TR-1	Provinces of Ankara, Edirne, Istanbul, Izmir, Kirklareli and Tekirdag	E	Х	Х	Х					_	_	Х	
	UA	Ukraine	UA-0	Whole country	В	Х	X	Х	Х	Х	X	X	Х		Х	
	US	United States of America	US-0	Whole country	С	Х	X	Х	X	X	X	X	Х	Х	Х	
	UY	Uruguay	UY-0	Whole country	D	Х	X	Х	Х	Х	Х	X	Х		Х	
	ZA	South Africa	ZA-0	Whole country	_	_		_	_	_				_		
			ZA-1	Metropolitan area of Cape- Town	F				_	_	_				_	Commission Decision 2008/698/ EC
				(see BOX 4 for details)												

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

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

(³) Hereinafter understood as the State of Israel, excluding the territories under Israeli administration since June 1967, namely the Golan Heights, the Gaza Strip, East Jerusalem and the rest of the West Bank. (⁴) Without prejudice to specific certification requirements provided for in Article 17 of the Agreement on the European Economic Area (OJ No L 1, 3.1.1994, p. 3).

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

▼M2

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	

Columns	Information / Description of commodity	Required Health Certificate	
12	Imports of semen collected from registered equidae	Annex III Part 1 Section A Annex III Part 1 Section B Annex III Part 1 Section C Annex III Part 1 Section D	
13	Imports of semen collected from equidae for breeding and production		
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 condi- tions/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
Α	equine infectious anaemia, equine viral arteritis
В	equine infectious anaemia, equine viral arteritis, glanders, dourine
С	equine infectious anaemia, equine viral arteritis, Eastern and Western equine encephalomyelitis, vesicular stomatitis
D	equine infectious anaemia, equine viral arteritis, glanders, dourine, Eastern and Western equine encephalomyelitis, Vene- zuelan equine encephalomyelitis, vesicular stomatitis
Е	equine infectious anaemia, equine viral arteritis, glanders, dourine, African horse sickness
F	equine infectious anaemia, dourine, African horse sickness
G	equine infectious anaemia, equine viral arteritis, glanders, dourine, Japanese encephalitis

▼<u>B</u>____

BOX	1
DOA	

_

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

BOX	BOX 2						
EG	Egypt	EG-1	The Equine Disease Free Zone (EDFZ) of about 0,1 km ² size, establishe around the Egyptian Armed Forces Veterinary Hospital at El-Nasr Road across Al Ahly Club, on the Eastern outskirts of Cairo, (localised 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-Shaik 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 wit the Passage Inside Armed Forces, turning right and following the Passag for about 100 m to the East, turning right again and following the Passag for 150 m to the South, turning left and following the Passage for 300 m t the East, turning right and following the Passage for 300 m t the East, turning right and following El-Nasr Road for 300 m t South-West until opposite of the junction of El-Nasr Road with Hassa Ma'moon Road, turning right and following the Passage for 100 m to th North, turning left and following the Passage for 120 m to the West, turnin left and following the Passage for 200 m to the South, turning right an following El-Nasr Road for 100 m to the West until the crossing of El-Nasr Road with El-Shaheed Ibrahim El-Shaikh Road.				
			(b) Delineation of the boundaries of the pre-export quarantine area within th EDFZ:				
			From the point opposite of the junction of El-Nasr Road with Hassa Ma'moon Road following the Passage for 100 m to the North, turnin 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, turnin 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-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 All Qahmah, and north of road No 10. Province of Asir Protection zone: the part of the province delineated by road No 10, between Ad Darb, Abha and Khamis-Mushayt to the north, except the equestrian clubs at their air and military bases, and the part of the province delineated to the north by road No 15 leading from Khamis-Mushayt through Jarash, Al Utfah and Dhahran Al Janoub to the border with the province of Najran, and, the part of the province delineated to the north by the road leading from Al Utfah through Al Fayd to Badr Al Janoub (Province of Najran);

 Surveillance zone: the equestrian clubs at their air and military bases, the part of the province between the border of the protection zone and road No 209 from Ash-Shuqaiq to the road control post Muhayil on road No 211, the part of the province between the control post on road No 10 south of Abha, the city of Abha and the road control post Ballasmer 65 km from Abha on road No 15 leading north, the part of the province between Khamis-Mushayt and the road control post at Yarah, 90 km from Abha, on road No 10 leading to Riyadh, and, the part of the province south of a virtual line between the road control post at Yarah on road No 10 and Khashm-Ghurab on road No 177 up to the border of the province of Najran.
 Province of Najran Protection zone: the part of the province delineated by the road from Al Utfah (province of Asir) to Badr Al Janoub and to As Sebt and from As Sebt along Wadi Habunah to the conjunction with road No 177 between Najran and Riyadh to the north and from this conjunction by road No 177 leading south to the conjunction with road No 15 from Najran to Sharourah, and, the part of the province south of road No 15 between Najran and Sharourah and the border with the Yemen. Surveillance zone: the part of the province south of a line between the road control post at Yarah, on road No 10, and Khashm-Ghurab, on road No 177, from the border of the province of Najran until the road control post at Khashm-Ghurab, 80 km from Najran, and west of road No 175 leading to Sharourah.

BOX 4

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

ANNEX II

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

▼<u>M2</u>

PART 1

Temporary admission and transit

Section A

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

COUNTRY:

Veterinary certificate to EU

	I.1. Consignor Name	I.2. Certificate reference No I.2.a.
	Address Tel.	I.3. Central competent authority
ŧ	10.	I.4. Local competent authority
Part I : Details of dispatched consignment	I.5. Consignee Name Address Postcode Tel.	1.6.
s of dispato	I.7. Country ISO I.8. Region of Code of origin code origin	I.9. Country of ISO I.10. Region of Cod destination code destination e
Part I: Detail:	I.11. Place of origin Name Approval number Address	I.12. Place of destination Name Address
	I.13. Place of loading	Postcode I.14. Date of departure
	I.15. Means of transport Aeroplane Ship Railway	I.16. Entry BIP in EU
	wagon Road vehicle Identification Documentary references	I.17. No(s) of CITES
	I.18. Description of animal	I.19. Commodity code (HS code) 01 01
		I.20. Quantity 1
	l.21.	I.22. Number of packages
	I.23. Seal/Container No	1.24.
	I.25. Animal certified for: Registered horse	
	1.26.	1.27. For import or admission into EU
	I.28. Identification of the animal	1
	Species (Scientific name) Identification system <i>Equus caballus</i>	Identification number Age Sex

	COUNTRY			Temporary admission - Registered horse			
				II.a.	Certificate reference number	II.b. Local reference number	
	н.	Attestatio	on of anima	al health and w	elfare		
	I the un	dersigned	official voto	inarian bereby	certify that the animal described in I	Box 1 28 ·	
 I, the undersigned official veterinarian, hereby certify, that the animal of is a registered horse as defined in Article 2(c) of Commission 					이 이 방법법은 정말할 수 있는 것 같아. 이 있 것 같아. 이 집 ? 이 집 ? 이 ? 이 집 ? 이 ? 이 집 ? 이 ? 이 ?		
	-	was examined today ⁽¹⁾ and found free of clinical signs of diseases and of obvious signs of ectoparasite infestation;					
	-	is not intended for slaughter under a national programme of infectious or contagious disease eradication;					
	-			a service of the service of the service of	oints II.1. to II.5. of this certificate;		
		 is accompanied by the written declaration, signed by the owner of the animal or the representative of the owner. 					
_							
	II.1.				the territory of third country and hol		
	II.1.1.	a country	or part of	the territory of a	a country, which on the date of issi ary Group ⁽²⁾ ;		
	II.1.2.	(Trypanos	soma equip	erdum), glander	g diseases are compulsorily notifiab s (<i>Burkholderia mallei</i>), equine ence s), equine infectious anaemia, vesic	phalomyelitis (of all types including	
	II.1.3.	Venezuelan equine encephalomyelitis), equine infectious anaemia, vesicular stomatitis, rabies and anthrax; the animal is dispatched from a country or part of the territory of a country:					
	a) wh an evi in			of African horse	from African horse sickness in accor- een no clinical, serological (in unvacc e sickness during the period of 2 yea n no vaccinations against the diseas ch:	sinated equidae) or epidemiological ars prior to the date of dispatch and	
		 b) in which Venezuelan equine encephalomyelitis has not occurred during the period of 2 years prior to the date of dispatch; 					
		c) in which dourine has not occurred during the period of 6 months prior to the date of dispatch;					
	(2)	d)			t occurred during the period of 6 mo		
	⁽³⁾ either	[e)	in which dispatch;		titis has not occurred during the peri	od of 6 months prior to the date of	
	⁽³⁾ 0r	[e)	in which dispatch, period of vesicular	vesicular stoma and a blood si 21 days prior to stomatitis virus	atitis has occurred during the period ample taken from the animal on the date of dispatch, was tested with	(insert date), within a negative result for antibody to the	
			⁽³⁾ either	[in a virus neut	ralisation test at a serum dilution of	1 in 32;]]	
			⁽³⁾ 0r		accordance with the relevant Chapte or Terrestrial Animals of the OIE;]]	er of the Manual of Diagnostic Tests	
	II.1.4. the animal does not come from a holding and to the best of my knowledge for the time periods referred to in points II.1.4.1. to II.1.4.7. was not in contact with animals from holdings, which were subject to prohibition orders for the reasons referred to in points II.1.4.1. to II.1.4.7. and which last for:						
	(⁽⁴⁾ [II.1.4.1.	in the cas	se of equidae su	spected of having contracted dourin	e,	
			⁽³⁾ either		nning on the date of the last actual aving contracted dourine or infected		
			⁽³⁾ and/or	[in the case of	a stallion, until the animal is castrate	ed;]	
			(3)and/or		ving the date of completion of the all animals of susceptible species ha		

COUNTRY

Temporary admission - Registered horse

COONT		remporary admission - negistered noise					
			II.a.	Certificate reference number	II.b.	Local reference number	
1	⁽⁴⁾ [II.1.4.2.	in the case of gla	inders.				
	•	(3)either [6 months beginning on the day on which the equidae suffering from the disease 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;]					
				ng the date of completion of the Il animals of susceptible species ha			
	II.1.4.3.	in the case of eq	uine ence	phalomyelitis of any type,			
		⁽³⁾ either [6 months beginning on the day on which the equidae suffering from the disease hav been slaughtered;]					
		West	Nile Fe	ning on the day on which the equid ever, Eastern equine encepha is have died, been removed from th	lomyeli	tis or Western equin	
				ng the date of completion of the Il animals of susceptible species ha			
	II.1.4.4.	been slaughtered	d, the rem nunodiffus	ctious anaemia, until the date on w aining equine animals on the holdin sion test (AGID or Coggins test) carr s apart;	g have	shown a negative reaction	
	II.1.4.5.	in the case of ve	sicular sto	matitis,			
		⁽³⁾ either [6 mor	ths follow	ing the last case;]			
				ng the date of completion of the Il animals of susceptible species ha			
	II.1.4.6.	in the case of rat		ays following the last case and the on nises;	late of o	completion of the cleansir	
	II.1.4.7.	in the case of and and disinfection		lays following the last case and the on nises;	date of	completion of the cleansir	
II.1.5.		st of my knowledge, during the period of 15 days prior to the date of dispatch the animal has no ontact with equidae infected or suspected of an infectious or contagious disease.					
II.2.	Attestation	n of residence and	pre-expo	rt isolation			
⁽³⁾ either	[II.2.1.	During a period of at least 40 days prior to the date of dispatch, the animal has been resident or holdings under veterinary supervision situated in the country or part of the territory of the country of dispatch which is assigned to Sanitary Group A, B, C, D, E or G, and					
	⁽³⁾ either	[in a Member State of the Union;]					
	⁽³⁾ and/or	[in a country or part of the territory of a country with Code: ⁽²⁾ that is authorise for temporary admission into the Union of registered horses, and from which it was imported in the country or part of the territory of the country of dispatch under conditions at least as strict a those required in accordance with the Union legislation for the temporary admission of registered horses from this country or part of the territory of the country directly to the Union, and which is ⁽³⁾ either [assigned to the same Sanitary Group ⁽²⁾ as the country or part of the territory the country of dispatch;]]					
				anitary Group A, B or C;]]			
			•	ng Kong, Japan, Korea, Macao,	Malavs	ia (Peninsula). Singapor	

		II.a. Certificate reference number II.b. Local reference number							
⁽³⁾⁽⁷⁾ Or	[II.2.1.	During a period of at least 60 days prior to the date of dispatch, the animal has been resident on holdings under veterinary supervision situated in the country or part of the territory of the country of dispatch which is assigned to Sanitary Group F, or was imported during the 60 days prior to the date of dispatch from a Member State of the Union before entering the vector–protected or vector proof guarantine station in accordance with point II.2.2.;]							
⁽³⁾⁽⁷⁾ either	[11.2.2.	the animal is dispatched from a country or part of the territory of a country which is assigned to Sanitary Group E and							
	⁽³⁾ either	[has been kept in isolation in the country or part of the territory of the country of dispatch protected from vector insects for a period of at least 40 days prior to the date of dispatch, or since entry into the country or part of the territory of the country of dispatch, if it was imported in accordance with point II.2.1 from a Member State of the Union or a country or part of the territory of a country which is assigned to Sanitary Group A, B, C, D, E or G;]]							
	⁽³⁾ 0r	[has been kept in designated premises under official veterinary supervision for a period of at leas 40 days prior to the date of dispatch, or since entry into the country or part of the territory of the country of dispatch, if it was imported in accordance with point II.2.1 from a Member State of the Union or a country or part of the territory of a country which is assigned to Sanitary Group A, B C, D, E or G, and the country or part of the territory of the country of dispatch is recognised by the OIE as officially free of African horse sickness;]]							
⁽³⁾⁽⁷⁾ or	[11.2.2.	the animal is dispatched from a country or part of the territory of country which is assigned to Sanitary Group F and was kept:							
	⁽³⁾ either	[in the approved vector-protected quarantine station of							
⁽³⁾ or	name of a monitoring	ently confined in the approved vector-proof quarantine station of							
II.3.	Attestatio	n of vaccination and health tests							
⁽³⁾ either	[II.3.1.	The animal was not vaccinated against African horse sickness in the country of dispatch and there is no information suggesting previous vaccination;]							
⁽³⁾ Or	[II.3.1. ⁽³⁾ either ⁽³⁾ or	The animal was vaccinated against African horse sickness, and this vaccination was carried out [more than 12 months prior to the date of dispatch;]] [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⁾⁽⁷⁾ or	[11.3.1.	The animal is dispatched from a country or part of the territory of a country which is assigned to Sanitary Group F and was vaccinated against African horse sickness on							
	II.3.2.	the animal was not vaccinated against Venezuelan equine encephalomyelitis during the period of 60 days prior to the date of dispatch from							
	⁽³⁾ either	[a country of which all parts of the territory are free of Venezuelan equine encephalomyelitis for a period of at least 2 years prior to the date of dispatch;]							
	⁽³⁾⁽⁷⁾ 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							

▼<u>M2</u>

COUNTRY

Temporary admission - Registered horse

		П.	a.	Certificate reference number	II.b.	Local reference number
	⁽³⁾ either	course and 60 days an vector-prote and during daily, remains same holding	reva d no ected that ined ng w	gainst Venezuelan equine enceph accinated according to manufacture o more than 12 months prior to the d quarantine for a period of at least is period remained clinically healthy within the normal physiological ra hich showed a rise in body temper virus isolation for Venezuelan equi	er's reco date of 21 days , and it nge, and ature, ta	mmendations not less thar f dispatch, and was kept ir prior to the date of dispatch s body temperature, taker d any equine animal on the ken daily, was subjected to
	⁽³⁾ or	vector-prote remained c normal phys a rise in boo for Venezue dispatched encephalon days after	ected linica siolo dy te elan wa nyeli the	ted against Venezuelan equine e d quarantine for a period of at lea ally healthy, and its body temperatu gical range, and any equine animal imperature, taken daily, was subjec equine encephalomyelitis with neg as subjected to a diagnostic tis with negative result conducted date of entry into of the vector-p vector insects until dispatch;]]	st 21 da re, take on the s ted to a gative re test on a san	ays, and during that period n daily, remained within the same holding which showed blood test for virus isolatior sults, and the animal to be for Venezuelan equine nple taken not less than 14
	⁽³⁾ or	encephalon taken on tw 	of d n-po nyeli burs ks fi	I to a haemagglutination inhibitis tits carried out by the same labora ccasions with an interval of 21 da date), the second of which was tak ispatch, without an increase in ant lymerase chain reaction) test for t tits virus genome, carried out with prior to dispatch, on (insert rom the moment of the RT-PCR sa of approved insect repellents an of the stable and the means in whi	tory on uys on . en durin body tite ne detect negative date), au mpling u d insect	the same day on samples (insert date) and or og a period of 10 days prior re, and a RT-PCR (reverse ction of Venezuelan equine e result on a sample taker nd has been protected from initi loading for dispatch, by ticides on the animal and
⁽³⁾ [II.3.3.	the anima	al is an unca	strat	ed male equine animal older than :	80 days	and
⁽³⁾ either	the animal is an uncastrated male equine animal older than 180 days, and [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 o dispatch;]]					
⁽³⁾ <i>or</i>	[was tested on a blood sample taken on					
⁽³⁾ <i>or</i>	[was tested on an aliquot of its entire semen taken on					
⁽³⁾ or	 [was vaccinated against EVA on					

COUNTRY

Temporary admission - Registered horse

COUNTRY		
		II.a. Certificate reference number II.b. Local reference number
	official which	1 October 2018, during a period of isolation of not more than 15 days und veterinary supervision, commencing on the day a blood sample was take was tested during that isolation period in a virus neutralisation test for EVA wire result at a serum dilution of 1 in 4;]]]
	superv EVA ca same o	age of 180 to 270 days, during a period of isolation under official veterina ision, during which the animal was subjected to a virus neutralisation test arried out with negative result at a serum dilution of 1 in 4, or carried out on t day by the same laboratory with stable or declining titres on two blood sampl at least 10 days apart;]]]
	result a 7 days	he animal was subjected to a virus neutralisation test for EVA with negati at a serum dilution of 1 in 4, carried out on a blood sample taken not earlier the after commencing a period of uninterrupted isolation which lasted until 21 da ng vaccination;]]]
	test for on the	age of 180 to 250 days, after the animal was subjected to a virus neutralisati EVA carried out with negative result at a serum dilution of 1 in 4, or carried c same day by the same laboratory with stable or declining titres on two blo taken at least 14 days apart;]]]
⁽³⁾ <i>or</i>	EVA carried out blood sample of	to a virus isolation test, polymerase chain reaction (PCR) or real-time PCR is with negative result on an aliquot of its entire semen collected after the date that animal taken on (insert date), within a period of 6 months pri patch, was tested in a virus neutralisation test for EVA with positive result at at least 1 in 4;]]
⁽³⁾ <i>or</i>	[has previously to vaccinated again	ested positive for antibodies against the equine arteritis virus or has been st EVA, and
	consecu prior to serologio	period of 6 months prior to the date of dispatch, was test mated, on the tive days, to at least two mares which were kept in isolation during the 7 dat and until at least 28 days after test mating and which were subjected to the cal tests for EVA with negative results at a serum dilution of 1 in 4 on bloc collected at the time of test mating and at least 28 days after the test mating
		ected to a virus neutralisation test for EVA carried out on a blood sample tak days prior to the date of dispatch on(insert date),
	⁽³⁾ either ⁽³⁾ or	[with positive result at a serum dilution of at least 1 in 4;]]]
⁽³⁾ <i>or</i>	[any requirement legislation that the animal is specified in that participating in s	[with negative result at a serum dilution of 1 in 4;]]] s for testing for EVA or vaccination against EVA have been waived by Unii
⁽³⁾⁽⁷⁾ either [II.3.4.	anaemia, where	patched from Iceland, which is certified as officially free from equine infectio it was continuously resident since birth, and did not come into contact w ave entered Iceland from other countries;]
⁽³⁾ or [11.3.4.	the animal was Coggins test) or on(ii ⁽³⁾ either [a perio	subjected with negative result to an agar gel immunodiffusion test (AGID to an ELISA for equine infectious anaemia carried out on a blood sample take asert date), this being within be of 90 days prior to the date of dispatch from a country or part of the territor untry which is assigned to Sanitary Group A, B, C or G;]]
	(3)or [a perio	od of 30 days prior to the date of dispatch from a country or part of the territor

_

DUNTRY			9	Temporal	y aum	ission - Registered horse
			II.a.	Certificate reference number	II.b.	Local reference number
⁽³⁾ [II.3.5.	Sanitary reported complem on a blog	Group B o during a nent fixatio	or E, or period on test f taken	from a country or part of the territory from Brazil, China or Thailand, or fro of 3 years prior to the date of d or glanders carried out with negative on	m a cou ispatch result	untry in which glanders was , and was subjected to a at a serum dilution of 1 in 5
⁽³⁾ [II.3.6.	from a co F, or from 2 years p carried of not been	ountry or p n China or prior to the put with n	e date o egative (insert breedir	ted male or a female equine anima he territory of a country which is assi ind, or from a country in which dourin f dispatch, and was subjected to a co e result at a serum dilution of 1 in date), within a period of 30 days pri- ng during the period of at least 30 d	gned to ne was omplem 5 on or to the	o Sanitary Group B, D, E o reported during a period o nent fixation test for douring a blood sample taken or e date of dispatch, and has
⁽³⁾⁽⁷⁾ [II.3.7.		al is dispa Group C d		from a country or part of the territory	of a co	ountry which is assigned to
	⁽³⁾ either	[Westerr country	n and E or part	astern equine encephalomyelitis hav of the territory of the country of disp the date of dispatch;]]	e not b batch d	een officially reported in the uring a period of at least 2
	⁽³⁾ or	accordin days prid	ig to m or to the	as vaccinated with a complete p anufacturer's instructions within a pr date of dispatch with inactivated vac alomyelitis, the last vaccination was	eriod of ccine ag	f 6 months and at least 30 gainst Western and Easterr
	⁽³⁾ 0r	vector-p	rotecte n tests	s kept for a period of at least 21 day d quarantine and during this period w for Western and Eastern equine en y on the same day	as subj	ected to haemagglutination
		⁽³⁾ either		sample of blood taken on ys prior to the date of dispatch, with		
		⁽³⁾ or	days secor	amples of blood taken on two occas on (<i>insert date</i>) and o nd of which was taken within a perio tch, without increase in antibody titre	n od of 1	(insert date), the
⁽³⁾ [II.3.8.	Sanitary	Group G, ae during a [comes f that hold	or from a period from a h ding wh	from a country or part of the territory a a country in which Japanese encep d of at least 2 years prior to the date holding situated in the centre of an ar ere there has been no case of Japa to the date of dispatch;]]	halitis l of dispa rea of a	has been officially reported atch and the animal: t least 30 km radius around
	⁽³⁾ or	[was kep the date	ot in a v of dispa	rector-protected quarantine during a atch and during that period the body t al physiological range, and was subj	empera	
		⁽³⁾ either	[to a encep of blo which witho	haemagglutination inhibition or virus obalitis carried out by the same labor- bod taken on two occasions with a 	s neutratory or n inten 	n the same day on samples val of at least 14 days or <i>insert date</i>), the second o or to the date of dispatch ody titre between the two

			II.a. (Certificate reference number	II.b. Local reference number
		⁽³⁾ Or	Japanes sample t	e encephalitis virus with negati aken not earlier than 7 days after (insert date), and remained p	e detection of antibodies again: ive result, carried out on a bloc r the date the isolation commence protected from vector insects un
	⁽³⁾ Or	revaccin	ated accor		th a complete primary course an indations during a period of not les the date of dispatch;]]
⁽³⁾⁽⁷⁾ either [11.3.9.	Sanitary	Group E, a	and was su	ubjected to a serological test for A	y of a country which is assigned a African horse sickness as describe by the same laboratory on the same
	⁽³⁾ either	on			nterval of between 21 and 30 day (insert date), the second the date of dispatch:
		⁽³⁾ either	[with neg	gative results in each case.]]]	
		⁽³⁾ or	[with a p	ositive result in the first sample,	and
			⁽³⁾ either		ubsequently tested with negative negative negative test as described in Annex IV
			⁽³⁾ or	increase in antibody titre	ted without more than a two-fo in a virus neutralisation test apter 2.5.1. of the OIE Terrestri and Vaccines.]]]]
	⁽³⁾ Or	prior to t	the date of		t date), within a period of 21 day art of the territory of the country of African horse sickness.]]
⁽³⁾⁽⁷⁾ or [11.3.9.		al is dispa Group F,		n a country or part of the territor	y of a country which is assigned
	⁽³⁾ either	IV to Dire day on b days, on sample	not taken ne, the se	0/156/EČ, which was carried out to bles taken on two occasions with (insert date) and on I less than 7 days after intro	se sickness as described in Anne by the same laboratory on the sam n an interval of between 21 and 3 (insert date), the fir duction into the vector-protecte iod of 10 days prior to the date
		⁽³⁾ either	[with neg	gative results in each case.]]]	
		⁽³⁾ or	[with a p	ositive result in the first sample,	and
			⁽³⁾ either		equently tested with negative rest s described in Annex IV to Directive
			⁽³⁾ or	increase in antibody titre in a vi	ed without more than a two-for rus neutralisation test as describe of the OIE Terrestrial Manual f

COUNTR	Y						22	Tempora	iry adn	nission	- Registered	horse
			II.a	l.	Certific	ate refer	ence nu	mber	II.b.	Local	reference nu	umber
		⁽³⁾ 0r	[was subjec sickness as result in eac than 28 day within a perio	desc h ca s aft	cribed in a lase on a later the date	Annex IN blood sa ate of in	to Direction troduction	ctive 200 ken on on into th	9/156/E	C, carrie	ed out with n insert date) r	egative not less
		⁽³⁾ or	[was subject in Annex IV sample take introduction dispatch.]]	to [n on	Directive	2009/15	6/EC, c (insert a	arried ou ate) not	ut with less tha	negative an 14 da	e result on a ays after the	a blood date of
II.4.	Attestatio	n of the tra	nsport conditio	ons								
⁽³⁾⁽⁷⁾ either	[II.4.1.	Sanitary the Unio into con	nal is dispatch Group A, B, C n, without pass tact with other ed in this health	C, D, sing t r equ	E or G a through a uidae not	and arra	ngemen marsha	ts have b alling or a	een ma ssembl	ade to tr y centre	ansport it dir and without	ectly to coming
⁽³⁾⁽⁷⁾ or	[11.4.1.	Sanitary protecte	nal is dispatch Group F and d quarantine s certificate eith [to the airpo for the aircr recognised i prior to take	arra tation er fo rt un aft to in the	angemen n without or imports der vecto be clea e third co	ts have t coming s or for te or-protect ansed an	been m into con emporar ted con d disinfe	ade to tr tact with admissi ditions a ected in a	ansport other e ion into nd arra	t it direct quidae r the Union ngemen with a c	tly from the not accompa on ts have been disinfectant of	vector- nied by m made officially
		⁽³⁾ 0r	[to a sea por conditions a scheduled d or part of the in stalls whic recognised i to departure	nd a irectle terri ch wa n the	arrangem ly to a po itory of a ere clear	ort in the country nsed and	ve been Union w not appr d disinfe	made to ithout cal oved for cted in a	transpling into the enti dvance	oort it or a port s ry into th with a c	n a vessel w situated in a le Union of e disinfectant o	which is country quidae, officially
	II.4.2.	complyin	ments have b ng with at least od from certific	the s	same hea	alth requ	irements	s as desc				
	II.4.3.	The tran disinfect	sport vehicles ed before load av are so co	or co	ontainers with a dis	s in which	n the ani t officiall	mal is go y recogn	ised in	the third	country of d	ispatch
II.5.	Attestatio	n of anima	l welfare									
			ed in Box I.28. ements were r									
Notes:												
Part I:												
Box I.8.:			of the country of mission Impler						dispate	h as ap	pearing in co	olumn 3
Box I.15.	Registration	ation numb tion is to b	er (railway wa e provided. In entry into the l	gons case	e of unlo	ainer and	l lorries)	, flight nu				

Temporary	admission -	Registered	horse
-----------	-------------	------------	-------

		II.a.	Certificate reference number	II.b. Local reference num
			L /// P 11 A L 11 L 1	
			number (if applicable) should be inc	
BOX	identification document	as defined in	ust bear an individual identifier whi n Article 2(b) of Commission Impler ch as ear tag, tattoo, brand, transp	menting Regulation (EU) 2018
	If a passport accompanie which validated it.	es the anima	al, its number should be stated and t	he name of the competent aut
	Age: Date of birth (dd/m	m/yyyy).		
	Sex (M = male, F = fema	ale, C = cas	trated).	
Part	0:			
(1)		ed on the da	ay of loading or on the last working	day before loading of the anim
	dispatch to the Member Sta	te of destina	ation in the Union.	
	to the date of authorisation territory of the country refe	for tempora rred to in p	red horse shall not be allowed when ry admission into the Union from th oint II.1.1., or during a period whe of equidae from this country or this	ne respective country or part or re restrictive measures have
(2)			ory of the country and the Sanitary	
(2)		I to Commi	ssion Implementing Regulation (EU	J) 2018/659.
(3) (4)	Delete as appropriate.	tation in no	int II 1.2 combine to the entire equat	n, of dispotch
(5)			int II.1.3. applies to the entire count I for temporary admission as appear	
			Regulation (EU) 2018/659.	ing in columns 5 and 6 respec
(6)			assigned to Sanitary Group G.	
(7)	Statements that relate entire	ely and excl	usively to a Sanitary Group differer	t from the Sanitary Group to v
			ritory, is assigned, may be left out,	provided that the numbering of
	subsequent statements is m	aintained.		
This	health certificate shall:			
(a)		and of the	erstood by the certifying officer and Member State where the registered	
(b)	be made out to a single con			
(c)			original throughout its temporary ad	mission in the Union;
(d)			rent to the colour of the printing;	
(e)			sheets of paper required are part of	
			umber of pages, and each page s e pages are stapled and stamped.	snan bear the certificate refer
Offic	ial veterinarian			
	Name (in capital letters):			Qualification and title:
	Date:			Signature:
				•

				ne owner or representative rary admission of a registe		
Ider	ntification o	f the anir	nal ⁽¹⁾			
Spe	cies (Scier ne)	ntific	Identification system	Identification number	Age	Sex
Equ	us caballu	ıs		S		
			2020 De 104 - 1020 PD	NAME FOR THE STOCK		an a a an air a
, th	e undersig the horse		er ⁽²⁾ or representative of the	e owner ⁽²⁾ of the registered ho	rse described above, he	reby declare, that:
	⁽²⁾ either			(insert name of country or rs prior to the date of dispatch		a country of dispatch
	⁽²⁾ or			rt name of country or part of t east 40 days prior to the date		f dispatch) during th
		(a)		rom (insert ritory of country of dispatch)	name of country from	where horse entere
		(b)	on(insert date) f	rom (insert ritory of country of dispatch)	name of country from	where horse entere
		(c)		rom (insert ritory of country of dispatch);		where horse entere
	infectious	s or conta	agious diseases transmissil			
	the trans stages of			vay that health and well-being	of the horse can be prot	ected effectively at a
				isolation as applicable in act territory of the country of dis		of the accompanyir
			the transport as applicable the territory of the country	in accordance with point II.4 of dispatch are fulfilled;	. of the accompanying he	ealth certificate for th
	during its premises		ce inside the Union for a pe	eriod of less than 90 days the	horse will be accommo	dated on the following
				n (place of hold		20~~~~ 200 COURT COURT
				n (place of hold		
				n (place of hold n (place of hold		
	I am awa outlined i	are that in n this dee	n the event that the horse claration, it must be accomp	moves from one Member Sta panied by a health certificate i st be notified to the Member	ate of the Union to anoth ssued by an official veter	ner Member State, a
				Union on I place of border post of exit)		the border post
			(2)	10		
				e ⁽²⁾ :		
Jat	e:		(dd/mm/yyyy)			
				(Signature)		
1)	document	as defin	ed in Article 2(b) of Comr	an individual identifier which nission Implementing Regula onder) and the anatomic plac	tion (EU) 2018/659. Sp	
	If a passp validated i		mpanies the animal, its n	umber should be stated and	I the name of the comp	etent authority which
	•		(dd/mm/yyyy).			
2)	Sex (M = I Delete as		= female, C = castrated).			
	Delete as	appropria	ale.			

Section B

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

COU	INTRY:	Veterinary certificate to EU
	I.1. Consignor Name	I.2. Certificate reference No I.2.a.
	Address	I.3. Central competent authority
ŧ	Tel.	I.4. Local competent authority
Part I : Details of dispatched consignment	I.5. Consignee Name Address Postcode Tel.	I.6. Person responsible for the load in EU Name Address Postcode Tel.
ils of dispat	I.7. Country of ISO code I.8. Region Code origin	I.9. Country of ISO code I.10. Region of Code destination
Part I : Deta	I.11. Place of origin Name Approval number Address	1.12.
	I.13. Place of loading	I.14. Date of departure
	I.15. Means of transport Aeroplane Ship Railway wagon Road vehicle Other I Identification	I.16. Entry BIP in EU I.17. No(s) of CITES
	Documentary references I.18. Description of animals	I.19. Commodity code (HS code)
	<i>.</i>	01 01 1.20. Quantity
		1.20. Quantity
	1.21.	I.22. Number of packages
	I.23. Seal/Container No	1.24.
	I.25. Animals certified for:	
	Registered equidae breeding and pro	duction 🗆 slaughter 🗆
	1.26. For transit through EU to third country X	1.27.
	Third country ISO code	
	I.28. Identification of the animal	
	Species (Scientific name) Identification system	Identification number Age Sex

COUNTR	RY				Transit - Equid
			II.a.	Certificate reference number	II.b. Local reference number
н.	Attestatio	on of anim	al healt	n and welfare	
I, the un	dersigned	official vete	rinarian,	hereby certify, that the equine anim	al described in Box I.28.:
-	was exan		(¹⁾ and f	ound free of clinical signs of diseas	es and of obvious signs of ectoparas
-	is not inte	nded for sl	aughter	under a national programme of infec	tious or contagious disease eradicatio
-				sted in points II.1. to II.5. of this certi	
	is accomp owner	panied by t	he writte	n declaration, signed by the owner o	of the animal or the representative of t
II.1.	Attestatio	n on third o	country o	r part of the territory of third country	and holding of dispatch
II.1.1.	country), Code:	a country o ⁽²⁾ , is a d horses o	or part of assigned	the territory of a country, which on to Sanitary Group ⁽²⁾ , and is	of country or part of the territory of he date of issuing this certificate has t authorised for temporary admission equidae and equidae for breeding a
II.1.2.	dourine (types incl	Trypanosol	ma equi	perdum), glanders (Burkholderia n	orily notifiable: African horse sicknes nallei), equine encephalomyelitis (of nfectious anaemia, vesicular stomatit
II.1.3.	the anima	al is dispato	hed fron	a country or part of the territory of	a country
	a)	2009/15 or epider date of d	6/EC and miologica lispatch a	d in which there has been no clinica al evidence of African horse sickness	ckness in accordance with Directi al, serological (in unvaccinated equida s during the period of 2 years prior to t ccinations against the disease during t
	b)			elan equine encephalomyelitis has n of dispatch;	not occurred during the period of 2 yea
	c)	in which	dourine	has not occurred during the period of	of 6 months prior to the date of dispate
1.0000000000000000000000000000000000000	d)		•	v 1	of 6 months prior to the date of dispate
⁽³⁾ either	[e)	in which of dispat		r stomatitis has not occurred during	the period of 6 months prior to the da
⁽³⁾ or	[e)	dispatch period of	, and a l f 21 days	blood sample taken from the animal	e period of 6 months prior to the date I on(insert date), within ested with negative result for antibody
		⁽³⁾ either	[in a vi	rus neutralisation test at a serum dil	ution of 1 in 32;]]
		⁽³⁾ 0r		ELISA in accordance with the releva and Vaccines for Terrestrial Animals	ant Chapter of the Manual of Diagnos

COUNTRY

Transit -	Equidae
-----------	---------

			II.a.	Certificate reference number	II.b.	Local reference number			
II.1.4.	to in point	ts II.1.4.1.	to II.1.4	m a holding, and to the best of my k 7. was not in contact with animals cons referred to in points II.1.4.1. to II	from hold	lings, which were subject			
(•	⁴⁾ [II.1.4.1.	in the cas ⁽³⁾ either	[6 mon suspec	uidae suspected of having contracted the beginning on the date of the last at ted of having contracted dourir rdum;]	ctual or po				
		(3)and/or	[in the	case of a stallion, until the animal is c	astrated;]				
		(3)and/or		rs following the date of completion or es after all animals of susceptible spe					
(•	⁴⁾ [II.1.4.2.	in the cas	e of gla	nders,					
		⁽³⁾ either	subjec	ths beginning on the day on which the ed with positive result to a test for the <i>Ideria mallei</i> or antibodies to that pat	e detectio	on of the causative patho			
		⁽³⁾ and/or		vs following the date of completion or es after all animals of susceptible spe					
	II.1.4.3.	in the cas	e of eq	ine encephalomyelitis of any type,					
		⁽³⁾ either		ths beginning on the day on which t een slaughtered;]	he equida	e suffering from the dise			
		⁽³⁾ and/or	^b and/or [6 months beginning on the day on which the equidae infected with the v causing West Nile Fever, Eastern equine encephalomyelitis or Western equ encephalomyelitis have died, been removed from the holding or fully recovered						
		(3)and/or		s following the date of completion o as after all animals of susceptible spe					
	II.1.4.4.	in the case of equine infectious anaemia, until the date on which, the infected animals havi been slaughtered, the remaining animals on the holding have shown a negative reaction an agar gel immunodiffusion test (AGID or Coggins test) carried out on blood sampl collected on two occasions 3 months apart;							
	II.1.4.5.	in the cas	e of ves	icular stomatitis,					
		⁽³⁾ either	[6 mon	ths following the last case;]					
		⁽³⁾ and/or		rs following the date of completion or es after all animals of susceptible spe					
	II.1.4.6.			bies, 30 days following the last cas infection of the premises;	se and th	e date of completion of			
	II.1.4.7.			hthrax, 15 days following the last ca infection of the premises;	se and th	ne date of completion of			
II.1.5.				during the period of 15 days prior to t infected or suspected of an infectious					
II.2.	Attestation	n of residen	ce and	pre-export isolation					
⁽³⁾ either	[II.2.1 <i>.</i>	on holdin	gs und	f at least 40 days prior to the date of er veterinary supervision situated in sh which is assigned to Sanitary Grou	a country	or part of the territory			
	⁽³⁾ either			te of the Union;]]					
	⁽³⁾ and/or	for tempo into the c strict as t of registe and which	rary adi ountry o hose re- red hors n is:	art of the territory of country with Coo nission into the Union of registered h r part of the territory of the country o quired in accordance with the Union I es from this country or part of the terri	orses, and f dispatch egislation tory of the	d from which it was impo under conditions at leas for the temporary admiss country directly to the Un			
		⁽³⁾ either	[assi	gned to the same Sanitary Group	.(2) as the	country or part of the terri			

COUNTRY

Transit - Equidae

		0					1.000	
			II.a.	Certificate	reference number		II.b.	Local reference num
		(3)and/or	[assi	gned to Sar	iitary Group A, B or	C;]]]		Depte on the community
		⁽³⁾ and/or						mal is a registered hor Regulation (EU) 2018/6
⁽³⁾⁽⁵⁾ or	[II.2.1.	on holding country of days prior	dispat to the	er veterinar ch which is date of dis	y supervision situat assigned to Sanita	ary Group ber State	F, or of the	he animal has been res or part of the territory was imported during the Union before enterin e with point II.2.2.;]
⁽³⁾⁽⁵⁾ either	[11.2.2.	the anima to Sanitar			n a country or part o	of the territ	ory of	a country which is ass
	⁽³⁾ either	protected since entry in accorda	from ve / into th ince wi	ector insects ne country o th point II.2	for a period of at le r part of the territory	east 40 da of the counce State of the	ys prio untry o e Unio	y of the country of dis or to the date of dispat of dispatch, if it was imp n or a country or part c, D, E or G;]]
	⁽³⁾ 0r	[has been least 40 da of the cou State of th Group A, I	kept in ays pric ntry of e Unior B, C, D	designated or to the date dispatch, if n or a countr b, E or G, an	I premises under off e of dispatch, or sinc it was imported in y or part of the territo	ficial veteri ce entry inte accordanc ory of a cou rt of the te	inary s o the o e with untry v rritory	supervision for a period country or part of the ter point II.2.1 from a Me which is assigned to Sa of the country of dispa
		⁽³⁾ either			a registered horse egulation (EU) 2018		d in a	Article 2(c) of Comm
		⁽³⁾ <i>or</i>						ntry in which African rior to the date of dispa
⁽³⁾⁽⁵⁾ 0r	[11.2.2.			batched from F and was		of the territ	ory of	a country which is ass
	⁽³⁾ either	of quarant date) to two hours veterinary insecticide isolation fr	prior to superve effect om equ	tion) during (insert sunset unti vision, follow ive against uidae not be	the 40 days prior to t date), confined to I two hours after sun ving the application <i>Culicoides</i> prior to	the date of the vector nrise and ex- of insect the remo- port to the	dispat -prote xercise repelle val fro Union	(insert ich from(cted premises at least e was provided under c ents in combination wi m the stables, and in under conditions at lea Union.]]
	⁽³⁾ or	(insert nar dispatch a	ne of q and cor	<i>quarantine</i> s nstant moni	tation) during the p	eriod of at protection	least	station of 14 days prior to the da proven absence of ve
II.3.	Attestation	n of vaccina	tion and	d health tes	ts			
⁽³⁾ either	[II.3.1.				ed against African h esting previous vaco		iess ir	the country of dispate
⁽³⁾ <i>or</i>	[II.3.1.					0.52	s, and	this vaccination was c
	⁽³⁾ either	[more than	n 12 ma	onths prior t	o the date of dispate	ch;]]		
	⁽³⁾ or				than 12 months price ad to in point II.1.3.(admission into the part is dispatched;]]
⁽³⁾⁽⁵⁾ or	[II.3.1.	to Sanitary date) not protected	Group more the quaran	F and was nan 24 mon ntine by adn	vaccinated against ths and at least 40 ninistration of a regi	African hor days prior istered vac	se sic to the ccine	a country which is ass kness on(date of entry in the v according to manufact of the African horse sick

Transit - Equidae

		II.a. Certificate reference number	II.b. Local reference number
II.3.2		nal was not vaccinated against Venezuelan eq	uine encephalomyelitis during the
⁽³⁾ eith		ry of which all parts of the territory are free of Ver riod of at least 2 years prior to the date of dispatch	
(3)(5) _O	Venezu dispatch	If the territory of a country which is assigned to Sa elan equine encephalomyelitis for a period of at and Venezuelan equine encephalomyelitis occ of the country of dispatch, and	least 2 years prior to the date of
	⁽³⁾ either	[is vaccinated against Venezuelan equine en primary course and revaccinated according to not less than 60 days and no more than 12 m and was kept in vector-protected quarantine for the date of dispatch, and during that period rema temperature, taken daily, remained within the n equine animal on the same holding which show daily, was subjected to a blood test for virus encephalomyelitis with negative results;]]	manufacturer's recommendations onths prior to the date of dispatch, a period of at least 21 days prior to ained clinically healthy, and its body ormal physiological range, and any ed a rise in body temperature, taken
	⁽³⁾ or	[is not vaccinated against Venezuelan equine vector-protected quarantine for a period of at le remained clinically healthy, and its body temper the normal physiological range, and any equine showed a rise in body temperature, taken daily virus isolation for Venezuelan equine encephal the animal to be dispatched was subjected to equine encephalomyelitis with negative result less than 14 days after the date of entry intr remained protected from vector insects until dis	east 21 days, and during that period rature, taken daily, remained within a animal on the same holding which y, was subjected to a blood test for omyelitis with negative results, and a diagnostic test for Venezuelan conducted on a sample taken not o vector-protected quarantine and
	⁽³⁾ or	[was subjected to a haemagglutination inhib encephalomyelitis carried out by the same labo taken on two occasions with an interval of 21 on (<i>insert date</i>), the second of which we prior to the date of dispatch, without an increa PCR (reverse transcription-polymerase chain Venezuelan equine encephalomyelitis virus g result on a sample taken within 48 hours prior t and has been protected from vector attacks sampling until loading for dispatch, by combine and insecticides on the horse and disinsectizat which it is transported;]]	ratory on the same day on samples days on (insert date) and as taken during a period of 10 days use in the antibody titre, and a RT- reaction) test for the detection of penome, carried out with negative o dispatch, on (insert date), from the moment of the RT-PCR d use of approved insect repellents
⁽³⁾⁽⁵⁾ either [II.3.3	anaemia	nal is dispatched from Iceland, which is certified as a, where it was continuously resident since birth which have entered Iceland from other countries;	and did not come into contact with
⁽³⁾ or [II.3.3	Coggins	hal was subjected with negative result to an agar s test) or to an ELISA for equine infectious anaei n	mia carried out on a blood sample
	⁽³⁾ or	[a period of 30 days prior to the date of dispatch of a country which is assigned to Sanitary Grou	

COUNTRY

Transit - Equidae

		II.a.	Certificate reference number	II.b. Local reference number
⁽³⁾ [II.3.4.	to Sanita was repo complem in 5 on a	ry Group I orted durin nent fixatio	tched from a country or part of the terr B or E, or from Brazil, China or Thailand, Ig a period of 3 years prior to the date of In test for glanders carried out with neg nple taken on	, or from a country in which glanders of dispatch, and was subjected to a gative result at a serum dilution of 1
⁽³⁾⁽⁵⁾ [II.3.5.			tched from a country or part of the terr C or D, and	ritory of a country which is assigned
	⁽³⁾ either	[Western the cour	n and Eastern equine encephalomyelitis htry or part of the territory of the count ears prior to the date of dispatch;]]	
	⁽³⁾ 0r	accordin 30 days and Eas	mal was vaccinated with a complete g to manufacturer's instructions within prior to the date of dispatch with ina tern equine encephalomyelitis, the 	a period of 6 months and at leas activated vaccine against Westerr
	⁽³⁾ or	a vecto haemag	nal was kept for a period of at least 21 or-protected quarantine and during glutination inhibition tests for lomyelitis carried out by the same labo	this period was subjected to Western and Eastern equine
		⁽³⁾ either	[on a sample of blood taken on of 10 days prior to the date of dispate	
		⁽³⁾ 01	[on samples of blood taken on two or 21 days on (insert date) the second of which was taken within of dispatch, without increase in antibo	and on (insert date) a period of 10 days prior to the date
⁽³⁾ [II.3.6.	to Sanita	ary Group	tched from a country or part of the terr G, or from a country in which Japane e during a period of at least 2 years pr	ese encephalitis has been officially
	⁽³⁾ either	around t	from a holding situated in the centre of hat holding where there has been no ca of 21 days prior to the date of dispatch	ase of Japanese encephalitis during
	⁽³⁾ 0r	to the da	t in a vector-protected quarantine duri ate of dispatch, and during that period d within the normal physiological range	the body temperature, taken daily
		⁽³⁾ either	[to a haemagglutination inhibition or viencephalitis carried out by the sams samples of blood taken on two occasedays on	he laboratory on the same day or sions with an interval of at least 14 d on(<i>insert date</i>), the beriod of 10 days prior to the date o ld increase in antibody titre between
		⁽³⁾ or	[to a Ig-M capture ELISA test for the Japanese encephalitis virus with neg sample taken not earlier than 7 of commenced on	gative result, carried out on a blood days after the date the isolatior

COUNTRY

COUNTRY						Transit - Equi
		II.a.	Certificate	reference number	II.b.	Local reference number
	⁽³⁾ 0r	revaccin	ated accor	ainst Japanese encephalitis rding to manufacturer's recor and not more than 12 months	nmend	ations during a period of
⁽³⁾⁽⁵⁾ either [II.3.7.	to Sanita	ary Group d in Annex ame day [on bloo days, or	E and wa IV to Direct d samples	a country or part of the terriss subjected to a serological ctive 2009/156/EC, which was taken on two occasions with <i>(insert date)</i> and or as taken within a period of 10	test fo s carrie	or African horse sickness ad out by the same labora terval of between 21 and (insert date),
		⁽³⁾ either	[with neg	ative results in each case.]]]		
		⁽³⁾ or	[with a p	ositive result in the first samp	le, and	i
			⁽³⁾ either	[the second sample was result in an agent identific to Directive 2009/156/EC	ation t	
			⁽³⁾ or	[the two samples were te increase in antibody titr described in point 2.4 of 0 Manual for Diagnostic Te	e in a Chapte	virus neutralisation test r 2.5.1. of the OIE Terres
	⁽³⁾ 0r	prior to t	he date of	taken on(inse dispatch, and the country or sed by the OIE as officially fr	part of	the territory of the countri
		⁽³⁾ either		nal is a registered horse as de nting Regulation (EU) 2018/6		n Article 2(c) of Commis
		⁽³⁾ or		ntry of dispatch is not adjacen has occurred during the pe []]]		
⁽³⁾⁽⁵⁾ or [II.3.7.		In group [was sub Annex IV on the of betwee on introduct	F, and bjected to V to Directi same day een 21 a	a country or part of the terri a serological test for Africa ve 2009/156/EC, which was on blood samples taken of and 30 days, on (insert date), the first samp e vector-protected quarantine prior to the date of dispatch,	n hors carried on two le not	e sickness as describe d out by the same labora occasions with an inte (insert date) taken less than 7 days a
		(3)either		ative results in each case.]]]		
		⁽³⁾ or		ositive result in the first samp	le, and	1
			⁽³⁾ either	[the second sample was s result in an agent identificat Directive 2009/156/EC.]]]]		
			⁽³⁾ or	[the two samples were test increase in antibody titre described in point 2.4 of C Manual for Diagnostic Test	in a hapter	virus neutralisation test 2.5.1. of the OIE Terres

▼<u>M2</u>

_

COUNTRY

Transit - Equidae

			II.a.	Certificate reference number	II.b. Local reference number
		⁽³⁾ or	sicknes negativ date) n	ss as described in Annex IV to D ve result in each case on a blood s	tidentification test for African ho birective 2009/156/EC, carried out v sample taken on(<i>ins</i> of introduction into the vector-protec prior to the date of dispatch.]]
		⁽³⁾ Or	describ a blood the dat	bed in Annex IV to Directive 2009/15 d sample taken on(on test for African horse sickness 6/EC, carried out with negative result (<i>insert date</i>) not less than 14 days at tected quarantine and not more than
II.4.	Attestatio	n of the tra	ansport co	onditions	
	er[II.4.1.	to Sanit to the U coming	ary Group Inion, with into con	A, B, C, D, E or G and arrangement hout passing through a market, mars	e territory of a country which is assigr ts have been made to transport it direc shalling or assembly centre and with plying with at least the same hea
⁽³⁾⁽⁵⁾ <i>or</i>	[II.4.1.	to Sanita protecte by a hea	ary Group ed quarant alth certific the Unior [to the made f official!	F and arrangements have been ma titine station without coming into cont cate either for imports or for tempora n airport under vector-protected con for the aircraft to be cleansed and di	e territory of a country which is assign de to transport it directly from the vect lact with other equidae not accompan ary admission into the Union or for tran nditions and arrangements have be isinfected in advance with a disinfect of dispatch, and sprayed against vect
		⁽³⁾ Or	protect which is in a co Union o disinfeo	ted conditions and arrangements have is scheduled directly to a port in the puntry or part of the territory of a cou- of equidae, in stalls which were clear	he territory of the country under vect ve been made to transport it on a ves Union without calling into a port situal untry not approved for the entry into ansed and disinfected in advance wit I country of dispatch and sprayed agai
	II.4.2.	complyi	ng with a		vent any contact with other equidae ts as described in this health certific ne Union.
	II.4.3.	The tran	nsport veh infected b h and they	hicles or containers in which the ani before loading with a disinfectant off	imal is going to be loaded were clear ficially recognised in the third country ne, litter or fodder cannot escape dur
	II.4.4	the Uni	on). Arrai		(insert country of destination outs he necessary animal health condition without delay.
II.5.	Attestatio	n of anima	al welfare		
					und fit to be transported on the intend well-being effectively at all stages of

		II.a.	Certificate reference number	II.b. Local reference number	
Notes:					
Part I:					
Box I.6.:	Person responsible	for the	load in Union.		
Box I.8.:			untry or part of the territory of the coun n Implementing Regulation (EU) 2018		
Box I.15.:	and information is t	o be pr	ray wagons or container and lorries), rovided. In case of unloading and rele entry into the Union.		
Box I.23.:	The container num	ber and	the seal number (if applicable) should	d be included.	
Box I.28.:	kiang, Equus quag	ga, Equ	Equus caballus, Equus asinus, Equu uus zebra, Equus grevyi, or indicate a	ny cross between those	
	<i>Identification system</i> : 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).				
	Sex (M = male, F =	female	e, C = castrated).		
da Th	y before loading of th e entry into the Union	e anima of thes	on the day of loading or in the case of al for dispatch to the Member State of se animals shall not be allowed when ansit through the Union from the respo	destination in the Union. he animals were loaded either prior	
 (1) Th da The the the the (2) Cc in (3) De (4) De (5) Sta wh 	y before loading of the e entry into the Unior e date of authorisatio e country referred to i e Union against the er ode of the country or columns 3 and 5 resp elete as appropriate. elete statement if the atements that relate nich the country of disp	e anima of thes n for tra n point ntry of e part of t pectively attestat entirely patch, o	al for dispatch to the Member State of se animals shall not be allowed when insit through the Union from the respo II.1.1., or during a period where restri- quidae from this country or this part of the territory of the country of dispatch, y of Annex I to Commission Implemen- tion in point II.1.3. applies to the entire and exclusively to a Sanitary Group or part of its territory, is assigned, may I	destination in the Union. he animals were loaded either prior active country or part of the territory ctive measures have been adopted the territory of the country of dispate and the Sanitary Group as appeari ting Regulation (EU) 2018/659. country of dispatch. different from the Sanitary Group	
 (1) Th da Th the the the (2) Cc (3) De (4) De (5) Sta wh 	y before loading of th e entry into the Union e date of authorisatio e country referred to i e Union against the ei ode of the country or columns 3 and 5 resp elete as appropriate. elete statement if the atements that relate	e anima of thes n for tra n point ntry of e part of t pectively attestat entirely patch, o	al for dispatch to the Member State of se animals shall not be allowed when insit through the Union from the respo II.1.1., or during a period where restri- quidae from this country or this part of the territory of the country of dispatch, y of Annex I to Commission Implemen- tion in point II.1.3. applies to the entire and exclusively to a Sanitary Group or part of its territory, is assigned, may I	destination in the Union. he animals were loaded either prior active country or part of the territory ctive measures have been adopted the territory of the country of dispate and the Sanitary Group as appeari ting Regulation (EU) 2018/659. country of dispatch. different from the Sanitary Group	
(1) Th da Th the the (2) CC (3) De (4) De (5) Stt wh of This health (a) be the and	y before loading of the e entry into the Unior e date of authorisatio e country referred to i e ounion against the er ode of the country or columns 3 and 5 resp elete as appropriate. elete statement if the atements that relate nich the country of disp the subsequent state in certificate shall: drawn up in at least e Member State of de d undergo the veterin	e anima of thes n for tra n point htry of e part of t port of t port of t port of t port of t port of t port of t attestat entirely patch, o ments i a langu sstinatic ary bor	al for dispatch to the Member State of se animals shall not be allowed when in ansit through the Union from the respo- II.1.1., or during a period where restri- quidae from this country of this part of the territory of the country of dispatch, y of Annex I to Commission Implemen- tion in point II.1.3. applies to the entire and exclusively to a Sanitary Group or part of its territory, is assigned, may b is maintained.	destination in the Union. he animals were loaded either prior active country or part of the territory ctive measures have been adopted the territory of the country of dispate and the Sanitary Group as appeari ting Regulation (EU) 2018/659. country of dispatch. o different from the Sanitary Group be left out, provided that the numberi er and one of the official languages	
(1) Th da Th the the (2) CC (3) De (4) De (5) Sti (6) Sti (1) be the the ann (b) be	y before loading of the e entry into the Union e date of authorisatio e country referred to i e Union against the er- ode of the country or columns 3 and 5 resp elete as appropriate. elete statement if the atements that relate nich the country of dis the subsequent state in certificate shall: drawn up in at least of Member State of de d undergo the veterin made out to a single	e anima of these n for tran n point htry of e part of t part of t part of t entirely patch, o ments i a langu estinatio ary bor consign	al for dispatch to the Member State of se animals shall not be allowed when in ansit through the Union from the respo- II.1.1., or during a period where restri- quidae from this country of this part of the territory of the country of dispatch, y of Annex I to Commission Implemen- tion in point II.1.3. applies to the entire and exclusively to a Sanitary Group or part of its territory, is assigned, may b is maintained.	destination in the Union. he animals were loaded either prior active country or part of the territory trive measures have been adopted the territory of the country of dispate and the Sanitary Group as appeari ting Regulation (EU) 2018/659. country of dispatch. different from the Sanitary Group be left out, provided that the numberi er and one of the official languages e animal will enter the Union territor	
(1) Th da Th the the (2) CC (3) De (4) De (5) Sta wh of This health (a) be the c) be (d) be (d) co (d) co	y before loading of the e entry into the Union e date of authorisatio e country referred to i e Union against the en- ode of the country or columns 3 and 5 resp elete as appropriate. Elete statement if the atements that relate inch the country of dis the subsequent state in certificate shall: drawn up in at least Member State of de d undergo the veterin made out to a single signed and stamped nsist of a single she divisible by inserting p	e anima of thes n for tra n point ntry of e part of t bectively attestat entirely patch, o ments i a langu estinatio ary bor consign in a col et of p bage n.	al for dispatch to the Member State of se animals shall not be allowed when it insit through the Union from the respo- II.1.1., or during a period where restri- quidae from this country of this part of the territory of the country of dispatch, y of Annex I to Commission Implemen- tion in point II.1.3. applies to the entire and exclusively to a Sanitary Group or part of its territory, is assigned, may be is maintained.	destination in the Union. he animals were loaded either prior active country or part of the territory ctive measures have been adopted the territory of the country of dispate and the Sanitary Group as appeari ting Regulation (EU) 2018/659. country of dispatch. o different from the Sanitary Group be left out, provided that the numberi er and one of the official languages e animal will enter the Union territor ng; are part of an integrated whole a id each page shall bear the certifica	
(1) Th da Th the the (2) CC (3) De (4) De (5) Sta wh of This health (a) be the c) be (d) be (d) co (d) co	y before loading of the e entry into the Unior e date of authorisatio e country referred to i e country referred to i e union against the er ode of the country or columns 3 and 5 resp elete as appropriate. elete statement if the atements that relate nich the country of disp the subsequent state in certificate shall: drawn up in at least e Member State of de d undergo the veterin made out to a single signed and stamped ivisible by inserting p ference number at the	e anima of thes n for tra n point ntry of e part of t bectively attestat entirely patch, o ments i a langu estinatio ary bor consign in a col et of p bage n.	al for dispatch to the Member State of se animals shall not be allowed when t ansit through the Union from the respo- ll.1.1., or during a period where restri- equidae from this country of the spart of the territory of the country of dispatch, y of Annex I to Commission Implemen- tion in point II.1.3. applies to the entire and exclusively to a Sanitary Group or part of its territory, is assigned, may b is maintained. age understood by the certifying offic on and of the Member State where the der checks; nee; lour different to the colour of the printi aper or all sheets of paper required umbers and total number of pages, ar	destination in the Union. he animals were loaded either prior active country or part of the territory ctive measures have been adopted the territory of the country of dispate and the Sanitary Group as appeari ting Regulation (EU) 2018/659. country of dispatch. o different from the Sanitary Group be left out, provided that the numberi er and one of the official languages e animal will enter the Union territor ng; are part of an integrated whole a id each page shall bear the certifica	
(1) Th da Th da Th the the (2) CC (3) De (4) De (5) Sta (4) De (5) Sta (6) Sta (7) De (10) De (10) De (10) CO (10) CO	y before loading of the e entry into the Union e date of authorisatio e country referred to i e union against the en- ode of the country of columns 3 and 5 resp lete as appropriate. elete statement if the atements that relate inch the country of disj the subsequent state in certificate shall: drawn up in at least e Member State of de d undergo the veterin made out to a single signed and stamped nsist of a single she divisible by inserting p ference number at the terinarian	e anima of thes n for tra n point ntry of e part of t bectively attestat entirely patch, o ments i a langu estinatio ary bor consign in a col et of p bage n.	al for dispatch to the Member State of se animals shall not be allowed when t ansit through the Union from the respo- ll.1.1., or during a period where restri- equidae from this country of the spart of the territory of the country of dispatch, y of Annex I to Commission Implemen- tion in point II.1.3. applies to the entire and exclusively to a Sanitary Group or part of its territory, is assigned, may b is maintained. age understood by the certifying offic on and of the Member State where the der checks; nee; lour different to the colour of the printi aper or all sheets of paper required umbers and total number of pages, ar	destination in the Union. he animals were loaded either prior active country or part of the territory trive measures have been adopted the territory of the country of dispate and the Sanitary Group as appeari- ting Regulation (EU) 2018/659. country of dispatch. o different from the Sanitary Group be left out, provided that the numberi- er and one of the official languages e animal will enter the Union territor ng; are part of an integrated whole a d each page shall bear the certificat d and stamped.	
(1) Th da Th da Th the the (2) CC (3) De (4) De (5) Sta (4) De (5) Sta (6) Sta (7) De (10) De (10) De (10) CO (10) CO	y before loading of the e entry into the Unior e date of authorisatio e country referred to i e ounion against the er- ode of the country or columns 3 and 5 resp elete as appropriate. elete statement if the atements that relate nich the country of dis the subsequent state in certificate shall: drawn up in at least d undergo the veterin made out to a single signed and stamped nsist of a single she divisible by inserting p ierence number at the terinarian ne (in capital letters):	e anima of thes n for tra n point ntry of e part of t bectively attestat entirely patch, o ments i a langu estinatio ary bor consign in a col et of p bage n.	al for dispatch to the Member State of se animals shall not be allowed when t ansit through the Union from the respo- ll.1.1., or during a period where restri- equidae from this country of the spart of the territory of the country of dispatch, y of Annex I to Commission Implemen- tion in point II.1.3. applies to the entire and exclusively to a Sanitary Group or part of its territory, is assigned, may b is maintained. age understood by the certifying offic on and of the Member State where the der checks; nee; lour different to the colour of the printi aper or all sheets of paper required umbers and total number of pages, ar	destination in the Union. he animals were loaded either prior active country or part of the territory ctive measures have been adopted the territory of the country of dispate and the Sanitary Group as appeari ting Regulation (EU) 2018/659. country of dispatch. o different from the Sanitary Group be left out, provided that the numberi er and one of the official languages e animal will enter the Union territor ng; are part of an integrated whole a id each page shall bear the certifica	

	Declaration by the owner or representative of the owner for transit through the Union of an equine animal					
Identification of the animal ⁽¹⁾						
Spe	ecies (Scient ne)	ific	Identification system	Identification number	Age	Sex
I, th			r ⁽²⁾ or representative of the	owner ⁽²⁾ of the animal descril	bed above, hereby declar	re, that:
-	the anima					
	⁽²⁾ either			(insert name of country or s prior to the date of dispatch		country of dispatch)
	⁽²⁾ or			name of country or part of that ast 40 days prior to the date		dispatch) during the
		(a)		om (insert r itory of country of dispatch)	name of country from w	here animal entered
		(b)	country or part of the terr	om (insert r itory of country of dispatch)		
		(c)		om (insert r itory of country of dispatch);]		here animal entered
-			of 15 days prior to the date gious diseases transmissib	of dispatch the animal has n le to equidae;	ot been in contact with a	nimals suffering from
-				solation as applicable in acc territory of the country of disp		of the accompanying
-			he transport as applicable he territory of the country of the cou	in accordance with point II.4. f dispatch are fulfilled;	of the accompanying hea	alth certificate for the
-	the transp all stages			ay that health and well-being	g of the animal can be pr	otected effectively at
ā				on on place of border post of exit);		the border post of
Nar	ne and addr	ess of th	e owner ⁽²⁾ or representative	9 ⁽²⁾ :		
Dat	e:		(dd/mm/yyyy)			
				(Signature)		
(1)	 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). 					
(2)	Delete as a	ppropria	te.			

PART 2

Re-entry after temporary export

Section A

Model health certificate and model declaration for the re-entry into the Union of registered horses for racing, competition and cultural events after temporary export for a period of not more than 30 days

COUNTRY:

Veterinary	certificate	to	EU
vecennary	ouriniouto		

	I.1.	Consignor Name	I.2.	Certificate reference No		l.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 Postcode Tel.	1.6.				
batch		10.					
s of disp	1.7.	Country of ISO code I.8. Region of Code origin origin	1.9.	Country of ISO coo destination	e I.	10. Region of destination	Code
etail	l.11.	Place of origin	I.12.	Place of destination			
Part I : E		Name Approval number Address		Name Address			
_				Postcode			
	I.13.	Place of loading	I.14.	Date of departure			
	I.15.	Means of transport	I.16.	Entry BIP in EU			
		Aeroplane Ship Railway wagon Road vehicle Other I Identification Documentary references	l.17.	No(s) of CITES			
	I.18.	Description of animal		I.19. Con	nmoc	dity code (HS code) 01 01	
						I.20. Quantity 1	
	I.21.					I.22. Number of pack	kages
	1.23.	Seal/Container No				1.24.	
	1.25.	Animal certified for:				-	
		Registered horse					
	1.26.			I.27. For import or admission	n into	EU 🗖	
	1.28.	Identification of the animal					
	· ·	Species Identification system Identi Scientific name) Equus caballus	ificatio	on number Age		Sex	

			II.a. Certificate reference number	II.b. Local reference number				
	П.	Attestation of ani	mal health and welfare					
	I, the unde	rsigned official veterin	arian, hereby certify, that the animal described	l in Box I.28.:				
	_	is a registered hors	e as defined in Article 2(c) of Commission Imp	plementing Regulation (EU) 2018/659;				
=	_	was examined too infestation;	lay (1) and found free of clinical signs of a	diseases and of obvious signs of ectoparasit				
	_	is not intended for	slaughter under a national programme of infec	tious or contagious disease eradication;				
	_	meets the requiren	nents attested in points II.1. to II.3. of this certif	ïcate;				
	_	is accompanied by	the written declaration, signed by the owner o	f the horse or the representative of the owner.				
	II.1.	Attestation on third	country or part of the territory of third country	and holding of dispatch				
	II.1.1.	The animal is dispatched from						
	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:						
		, African	here has been no clinical, serological (in unva horse sickness during the period of 2 years een no vaccinations against the disease duri	in accordance with Directive 2009/156/EC and iccinated equidae) or epidemiological evidence prior to the date of dispatch and in which the ing the period of 12 months prior to the date				
			h Venezuelan equine encephalomyelitis has r e of dispatch;	not occurred during the period of 2 years prior				
		c) in whic	h dourine has not occurred during the period o	of 6 months prior to the date of dispatch;				
		d) in whic	h glanders has not occurred during the period	of 6 months prior to the date of dispatch;				
	II.1.4.	points II.1.4.1. to II		ny knowledge for the time periods referred to ioldings, which were subject to prohibition orde last for:				
		(³) [II.1.4.1. in the c	d dourine,					
		(⁴) eithe	er [6 months beginning on the date of the suspected of having contracted dourine or	last actual or possible contact with an anim infected with <i>Trypanosoma equiperdum</i> ;]				
		(⁴) and,	/or [in the case of a stallion, until the animal is	castrated;]				
		(⁴) and	/or [30 days following the date of completion after all animals of susceptible species hav	of the cleansing and disinfection of the premise				

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

Registered horse

II.a. Certificate reference number II.b. Local reference number (³) [II.1.4.2. in the case of glanders, II.b. Local reference number (⁴) either [6 months beginning on the day on which the equidae suffering from the disease or subjected with positive results to a test for the detection of the causative pathogen *Burkholderia mallei* or antibodies to that pathogen, were killed and destroyed;] (⁴) and/or [30 days following the date of completion of the cleansing and disinfection of the premises after all animals of susceptible species have been killed and destroyed;]] II.1.4.3. in the case of equine encephalomyelitis of any type,

- (⁴) either [6 months beginning on the day on which the equidae suffering from the disease have been slaughtered;]
- (⁴) and/or [6 months beginning on the day on which the equidae infected with the virus causing West Nile Fever, Eastern equine encephalomyelitis or Western equine encephalomyelitis have died, been removed from the holding or fully recovered;]
- (4) and/or [30 days following the date of completion of the cleansing and disinfection of the premises after all animals of susceptible species have been slaughtered;]
- II.1.4.4. in the case of equine infectious anaemia, until the date on which, the infected animals having been slaughtered, the remaining equine animals on the holding have shown a negative reaction in an agar gel immunodiffusion test (AGID or Coggins test) carried out on blood samples collected on two occasions 3 months apart;
- II.1.4.5. in the case of vesicular stomatitis,
 - (⁴) *either* [6 months following the last case;]
 - (⁴) and/or [30 days following the date of completion of the cleansing and disinfection of the premises after all animals of susceptible species have been slaughtered;]
- II.1.4.6. in the case of rabies, 30 days following the last case and the date of completion of the cleansing and disinfection of the premises;
- II.1.4.7. in the case of anthrax, 15 days following the last case and the date of completion of the cleansing and disinfection of the premises;
- II.1.5. to the best of my knowledge, during the period of 15 days prior to the date of dispatch the animal has not been in contact with equidae infected or suspected of an infectious or contagious disease.
- II.2. Attestation of residence and pre-export isolation
- II.2.1. The animal was imported on (insert date)
 - (⁴) either [directly from the EU Member State (insert name of EU Member State);]
- II.2.2. the animal exited from the Union less than 30 days ago, and since exit from the Union it was never in a country or part of the territory of a country (¹) other than those of the same Sanitary Group, and resident on holdings under veterinary supervision, accommodated in separated stables without coming into contact with equidae of lower health status, except during racing, competition or the cultural event.
- II.3. Attestation of animal welfare

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

▼<u>M1</u>

COUNTRY

COUN	TRY		Re-entry after tempo	orary export of not more than 30 days Registered horse				
			II.a. Certificate reference number	II.b. Local reference number				
Note	s:			·				
Part	l:							
Box	x 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	l.15.:		railway wagons or container and lorries), flight n vided. In case of unloading and reloading, the consig nion.					
Вох	1.23.:	The container number	and the seal number (if applicable) should be include	d.				
Box	1.28.:	identification document the identification system	The animal must bear an individual identifier whi as defined in Article 2(b) of Commission Implementi n (such as ear tag, tattoo, brand, transponder) and th companying passport must be stated and the name	ing Regulation (EU) 2018/659. Specify ne anatomic place used on the animal.				
		Age: Date of birth (dd/r	nm/yyyy).					
		Sex (M = male, F = fem	nale, C = castrated).					
Part	11:							
(1)		rtificate must be issued mber State of destinatio	on the day of loading or on the last working day befo n in the Union.	re loading of the animal for dispatch to				
	to the referre	date of authorisation for d to in point II.1.1., or o	xport of this registered horse shall not be allowed wh re-entry into the Union from the respective country o during a period where restrictive measures have be country or this part of the territory of the country of dis	r the part of the territory of the country een adopted by the Union against the				
(2)			of the territory of the country and the Sanitary Gro mission Implementing Regulation (EU) 2018/659.	up as appearing in columns 3 and 5				
(³)	Delete	statement if the attestat	ion in point II.1.3. applies to the entire country of disp	atch.				
(4)	Delete	as appropriate.						
This	health c	ertificate shall:						
(a)	State of	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;						
(b)	be ma	de out to a single consig	nee;					
(c)	be sigr	e signed and stamped in a colour different to the colour of the printing;						
(d)	insertir	nsist of a single sheet of paper or all sheets of paper required are part of an integrated whole and indivisible by serting page numbers and total number of pages, and each page shall bear the certificate reference number at the top the page and those pages are stapled and stamped.						
Offic	ial veter	inarian						
	Name	(in capital letters):		Qualification and title:				
	Date:			Signature:				
	Stamp							

	Declaration by the owner or representative of the owner for the re-entry after temporary export of a registered horse for racing, competition and cultural events					
lder	ntification o	f the animal (¹)				
Sp	ecies (Scie Equus c a	ntific name) aballus	Identification system	Identification number	Age	Sex
I, th	e undersigi	ned owner (²) c	or representative of the owne	r (²) of the registered horse descri	bed above, hereby dec	lare, that:
-	the horse					
	(²) either		rily exported from the Union prior to this declaration;]	to the country of dispatch on		(insert date) less
	(²) or	[entered the c country from	country of dispatch on	, (insert date) fro of dispatch);]	om	(insert name of
-			lays prior to the date of dispatransmissible to equidae;	atch the horse has not been in co	ntact with animals suffe	ring from infectious
-	the transp of the jou		effected in such a way that	health and well-being of the hors	e can be protected effe	ctively at all stages
_	 the conditions for residence and pre-export isolation as applicable in accordance with point II.2. of the accompanying health certificate for the country or part of the territory of the country of dispatch are fulfilled. 					
Nan	ne and add	ress of the own	ner (²) or representative (²):			
Date	Date: (dd/mm/yyyy)					
(1)	Article 2(b) of Commissio		identifier which permits to link the ani U) 2018/659. Specify the identifica		
		ort accompanies the of birth (dd/mm/y		e stated and the name of the competer	t authority which validated	l it.
0		nale, F = female,	C = castrated).			
(2)	Delete as appropriate.					

Section B

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

▼<u>M2</u>

Chapter 1

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

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

COU	NTRY:	Veterinary certificate to EU			
	I.1. Consignor Name	I.2. Certificate reference No I.2.a.			
	Address Tel.	I.3. Central competent authority			
ment		I.4. Local competent authority			
Part I : Details of dispatched consignment	I.5. Consignee Name Address Postcode	1.6.			
pato	Tel.				
ails of dis	I.7. Country ISO code I.8. Region Code of origin of origin	I.9. Country of ISO code I.10. Region Code destination of destination			
Deta	I.11. Place of origin	I.12. Place of destination			
Ë					
Pai	Name Approval number Address	Name Address			
		Postcode			
	I.13. Place of loading	I.14. Date of departure			
	I.15. Means of transport	I.16. Entry BIP in EU			
	Aeroplane Ship Railway wagon Road vehicle Other				
	Identification Documentary references	I.17. No(s) of CITES			
	I.18. Description of animal	I.19. Commodity code (HS code) 01 01			
		I.20. Quantity 1			
	1.21.	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			
	I.28. Identification of the animal				
	Species (Scientific name) Identification system <i>Equus caballus</i>	Identification number Age Sex			

	COL	JNTRY					t of not more than 90 day atitions - Registered hors			
				II.a.	Certificate reference number	II.b.	Local reference numbe			
	II.			al health and		in Day 1 Of				
ç	I, the u				y certify, that the animal described Article 2(c) of Commission Impleme					
Part II: Certification			ined today		free of clinical signs of diseases					
Cer	-		not intended for slaughter under a national programme of infectious or contagious disease eradication; neets the requirements attested in points II.1. to II.3. of this certificate;							
≣	-				points II.1. to II.3. of this certificate claration, signed by the owner of the		or the representative of th			
Part	_	owner.	anieu by ti	le whiteh dec	aradon, signed by the owner of the	le noise,	or the representative of th			
	II.1.	Attestation	on third c	ountry or part	of the territory of third country and	holding of	f dispatch			
	11.1.1.	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.		he animal is dispatched from a country or part of the territory of a country:							
		a)	and in wh evidence in which	ich there has of African hor	e from African horse sickness in ac been no clinical, serological (in unv rse sickness during the period of 2 sen no vaccinations against the dis tatch;	accinated years prio	equidae) or epidemiologica r to the date of dispatch an			
		b)		in which Venezuelan equine encephalomyelitis has not occurred during the period of 2 years prior to the date of dispatch;						
		c)	in which o	dourine has no	not occurred during the period of 6 months prior to the date of dispatch;					
		 in which glanders has not occurred during the period of 6 months prior to the date of dispatch the animal does not come from a holding, and to the best of my knowledge for the time periods referred to 								
	II.1.4.	points II.1.	4.1. to II.1	.4.7. was not	in contact with animals from holdin points II.1.4.1. to II.1.4.7. and whi	ngs, which	were subject to prohibitio			
		⁽³⁾ [II.1.4.1.	II.1.4.1. in the case of equidae suspected of having contracted dourine,							
			⁽⁴⁾ either		eginning on the date of the last act having contracted dourine or infec					
			(4)and/or	[in the case of	of a stallion, until the animal is castr	rated;]				
			(4)and/or		lowing the date of completion of er all animals of susceptible species					
		⁽³⁾ [II.1.4.2.	in the cas	e of glanders,						
			⁽⁴⁾ either	subjected with	eginning on the day on which the of th positive results to a test for the mallei or antibodies to that pathog	detection	of the causative pathoge			
			(4)and/or		lowing the date of completion of er all animals of susceptible species					
		II.1.4.3.	in the cas ⁽⁴⁾ either		ncephalomyelitis of any type, ginning on the day on which the ec tered;]	luidae suff	ering from the disease hav			
			(4)and/or	West Nile	eginning on the day on which the e Fever, Eastern equine ence yelitis have died, been removed fro	phalomyel	litis or Western equin			
			(4)and/or		lowing the date of completion of er all animals of susceptible species					

COUNTRY Re-entry after temporary export of not more than 90 days Specific competitions - Registered horse II.a. Certificate reference number II.b. Local reference number II.1.4.4. in the case of equine infectious anaemia, until the date on which, the infected animals having

II.1. II.1.5. to t bee I.2. Atte I.2.1. The (4) of I.2.2. the	een in co testation ne anima either or	 in the case of vesicular stomatitis, ⁽⁴⁾either [6 months following the last case;] ⁽⁴⁾and/or [30 days following the date of completion of the cleansing and disinfection of the premises after all animals of susceptible species have been slaughtered;] in the case of rabies, 30 days following the last case and the date of completion of the cleansing and disinfection of the premises; in the case of anthrax, 15 days following the last case and the date of completion of the cleansing and disinfection of the premises; st of my knowledge, during the period of 15 days prior to the date of dispatch the animal has not ontact with equidae infected or suspected of an infectious or contagious disease. <i>n of residence and pre-export isolation</i> nal was imported into the country or part of the territory of the country of dispatch on <i>(insert date)</i> [directly from the EU Member State
II.1.5. to t bee I.2. Atte I.2.1. The (4) <i>ej</i> (4) <i>o</i> (4) <i>o</i>	1.4.7. the best pen in co testation ne anima either pr e animal	premises after all animals of susceptible species have been slaughtered;] in the case of rabies, 30 days following the last case and the date of completion of the cleansing and disinfection of the premises; in the case of anthrax, 15 days following the last case and the date of completion of the cleansing and disinfection of the premises; st of my knowledge, during the period of 15 days prior to the date of dispatch the animal has not ontact with equidae infected or suspected of an infectious or contagious disease. <i>n of residence and pre-export isolation</i> mal was imported into the country or part of the territory of the country of dispatch on <i>(insert date)</i> [directly from the EU Member State
II.1.5. to t bee I.2. Atte I.2.1. The (4) <i>ej</i> (4) <i>o</i> (4) <i>o</i>	1.4.7. the best pen in co testation ne anima either pr e animal	and disinfection of the premises; in the case of anthrax, 15 days following the last case and the date of completion of the cleansing and disinfection of the premises; st of my knowledge, during the period of 15 days prior to the date of dispatch the animal has not ontact with equidae infected or suspected of an infectious or contagious disease. <i>n of residence and pre-export isolation</i> nal was imported into the country or part of the territory of the country of dispatch on <i>(insert date)</i> [directly from the EU Member State
I.1.5. to t bee I.2. Atte (4) (4) (4) (4) (4) (4) (4) (4) (4) (4)	the best een in co testation ne anima either or e animal	and disinfection of the premises; st of my knowledge, during the period of 15 days prior to the date of dispatch the animal has not ontact with equidae infected or suspected of an infectious or contagious disease. <i>n of residence and pre-export isolation</i> nal was imported into the country or part of the territory of the country of dispatch on <i>(insert date)</i> [directly from the EU Member State
(4) <i>or</i>	een in co testation ne anima either or e animal	ontact with equidae infected or suspected of an infectious or contagious disease. <i>n of residence and pre-export isolation</i> nal was imported into the country or part of the territory of the country of dispatch on <i>(insert date)</i> [directly from the EU Member State
I.2.1. The (4) <i>ej</i> (4) <i>oI</i> I.2.2. the (4) <i>ej</i>	ne anima either or e animal	nal was imported into the country or part of the territory of the country of dispatch on (insert date) [directly from the EU Member State
(4) <i>ei</i> (4) <i>o</i> (4) <i>o</i> (4) <i>ei</i>	either or e animal	(insert date) [directly from the EU Member State
(4)01 1.2.2. the (4) ₀₁	or e animal	[from a country or part of the territory of a country
1.2.2. the ⁽⁴⁾ <i>ei</i>	e animal	under conditions at least as strict as those set out in this certificate;] al exited from the Union [less than 30 days ago, and since exit from the Union was never in a country, or part of the
(⁴⁾ <i>ei</i>		[less than 30 days ago, and since exit from the Union was never in a country, or part of the
⁽⁴⁾ 01	either	
		territory of the country of dispatch it was resident on holdings under veterinary supervision, accommodated in separated stables without coming into contact with equidae of lower health status except during competition, and it has taken part in or was stabled together with horses participating in the LG Global Champions Tour ⁽⁴⁾ either [in the Metropolitan area of Mexico City, Mexico;]]
		(4) and/or [in Miami, Unites States of America;]
		(4)or [in Shanghai, China;]]
(4)01	or	[less than 60 days ago, and since exit from the Union was never in a country, or part of the territory of a country other than those of the same Sanitary Group. In the country or part of the territory of the country of dispatch it was resident on holdings under veterinary supervision, accommodated in separated stables without coming into contact with equidae of lower health status except during competition and it has taken part in or was stabled together with horses participating in ⁽⁴⁾ either [the Asian Games in
(4)00		(4) or [the American Games ⁽⁵⁾ in
(4)01		(4) or [the Endurance World Cup in United Arab Emirates.]]
	or	[less than 90 days ago, and since exit from the Union was never in a country, or part of the territory of a country ⁽¹⁾ other than those of the same Sanitary Group. In the country or part of the territory of the country of dispatch it was resident on holdings under veterinary supervision, accommodated in separated stables without coming into contact with equidae of lower health status except during competition and it has taken part in or was stabled together with horses participating in ⁽⁴⁾ either [the Test event for the Olympic Games in
		(4) or [the Olympic Games in(insert place).]]

II.3. Notes: Part I: Box I.8.:	⁽⁴⁾ or [th Attestation of animal wel The animal described in journey and arrangeme journey. Provide the code of th	e World Ec fare Box 1.28.	Certificate reference number bics in questrian Games/World Champions was examined today ⁽¹⁾ and found nade to protect its health and well-	hips in fit to be ti	(insert place).]]
Notes: Part I:	⁽⁴⁾ or [th Attestation of animal wel The animal described in journey and arrangeme journey.	e World Ec fare Box 1.28.	uestrian Games/World Champions was examined today ⁽¹⁾ and found	hips in fit to be ti	(insert place).]]
Notes: Part I:	Attestation of animal wel The animal described in journey and arrangeme journey. Provide the code of th	fare Box I.28.	was examined today(1) and found	fit to be to	ransported on the intend
Notes: Part I:	The animal described in journey and arrangeme journey. Provide the code of th	Box 1.28.			
Part I:	journey and arrangeme journey. Provide the code of th				
Part I:					
Box I.8.:					
	to Commission Imple		or part of the territory of the country egulation (EU) 2018/659.	as appea	aring in column 3 of Anne
Box I.15.		rovided. In	gons or container and lorries), flight case of unloading and reloading, t Union.		
Box I.23.			eal number (if applicable) should be	e included	
Box I.28.	identification docume Specify the identifica	nt as define tion system The numb which valid /mm/yyyy)		lementing ransponde	Regulation (EU) 2018/65 er) and the anatomic pla
Part II:	3ex (w = male, $F = 1e$	emale, C =	castrated).		
(1) T d T e te a d (2) C (2) a (3) D (4) D	ispatch to the Member St he re-entry after tempora ither prior to the date of a erritory of the country ref dopted by the Union agai ispatch. Code of the country or pai nd 5 respectively of Anne Delete statement if the atte Delete as appropriate.	ate of dest ry export of uthorisatio erred to in nst the ent rt of the ter ex I to Com estation in	day of loading or on the last workin ination in the Union. of this registered horse shall not be n for re-entry into the Union from th point II.1.1., or during a period wi ry of equidae from this country or th rritory of the country, and the Sanita mission Implementing Regulation (I point II.1.3. applies to the entire cou South American Games, Central Am	allowed we e respectionere restri is part of t ary Group EU) 2018/0 intry of dis	when the animal was load ve country or the part of t ictive measures have be he territory of the country as appearing in columns 659. spatch.
	ē.				
(a) be M (b) be (c) be (d) c b	lember State of destination ndergo the veterinary born e made out to a single co e signed and stamped in onsist of a single sheet of y inserting page number	on and of the der checks nsignee; a colour dif paper or a s and tota	nderstood by the certifying officer ar the Member State where the register ; iferent to the colour of the printing; all sheets of paper required are part I number of pages, and each page ose pages are stapled and stamped	ed horse v of an integ e shall be	will enter Union territory a grated whole and indivisit
	eterinarian			Na na	
	me (in capital letters):			Qualific	cation and title:
Dat				Signatu	
1200	amp:			Signatt	

		Declaration by th for the re-entry after tempo	e owner or representative prary export of a registered		on
Ide	ntification of th	e animal ⁽¹⁾			
	ecies (Scientifi me)	c Identification system	Identification number	Age	Sex
Eq	uus caballus				
I, ti	the horse ⁽²⁾ either ⁽²⁾ or	d owner ⁽²⁾ or representative of the [was temporarily exported from than 60 days ⁽²⁾ or 90 days ⁽²⁾ pri [entered the country of dispatch where horse entered country o as been temporarily exported from [the Asian Games in	the Union to the country of o or to this declaration;] n on	dispatch on(ins (insert place);] (insert place);] (insert place);] (insert place);] (insert place);]	(insert date) less
-	infectious or the condition health certifi	⁽³⁾ or [Shanghai, China;] eriod of 15 days prior to the date contagious diseases transmissib ns for residence and pre-export icate for the country or part of the tation will be effected in such a w e journey.	le to equidae; isolation as applicable in act territory of the country of dis	cordance with point II	I.2. of the accompanying
		ss of the owner ⁽²⁾ or representative	B ⁽²⁾ :		¥1
			(Signature)		
(1)	document as system (such	system: The animal must bear a defined in Article 2(b) of Comm as ear tag, tattoo, brand, transpo	nission Implementing Regula onder) and the anatomic place	ation (EU) 2018/659. ce used on the animal	Specify the identification
	validated it.	accompanies the animal, its nu	umber should be stated and	the name of the co	mpetent authority which
(2) (3)	Sex (M = ma Delete as ap	birth (dd/mm/yyyy). le, F = female, C = castrated). propriate. PanAmerican Games, South Am	erican Games, Central Ame	rican and Caribbean (Games.

Chapter 2

Model health certificate and model declaration applicable to re-entry into the Union of registered horses for racing after temporary export for a period of not more than 90 days to participate in specific race events in Australia, Canada, the United States of America, Hong Kong, Japan, Singapore, the United Arab Emirates or Qatar

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

cou	NTR	ŕ :							Ve	terinary certif	icate to EU
	l.1.	Consignor Name				1.2.	Certificate reference	e No		l.2.a.	
		-			1.3.	Central competent	authority				
ŧ					1.4.	Local competent a	uthority				
Imei	1.5.	Consignee				1.6.					
Isigr		Name Address							_		
cor											
hed		Postcode Tel.									
patc		101.									
s of dis	1.7.	Country of origin	ISO code	I.8. Region of origin	Code	1.9.	Country of Is destination	SO code		Region of destination	Code
Details	l.11.	. Place of origin				I.12.	Place of destination	า			
Part I : Details of dispatched consignment		Name App Address	proval numb	ber			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			igon 🗖							
					1.17.	No(s) of CITES					
	1.18.	. Description of ani						I.19. Com	moditv	code (HS code	e)
		•								01 01	,
									1	.20. Quantity 1	
	I.21.								I	.22. Number o	f packages
	I.23. Seal/Container No I.25. Animal certified for:							1	.24.		
		Registered horse									
	1.26.						I.27. For import or	admission in	ito EU		
	1.28.	. Identification of th	e animal				<u> </u>				
		Species (Scientific name) Equus caballus	Identi	fication system	Identi	ificatio	on number	Age		Se	K

DUNT	ſRY			Re-entry aff	ter temporary export of not more than 90 day Specific races — Registered hor					
			II.a. Certificate	reference number	II.b. Local reference number					
	II.	Attestatio	n of animal health and welf	are						
	I, the unde	rsigned offici	ıl veterinarian, hereby certify,	that the animal described i	n Box I.28.:					
	_	is a regist	ered horse as defined in Articl	e 2(c) of Commission Impl	ementing Regulation (EU) 2018/659;					
	—	was examined today (1) and found free of clinical signs of diseases and of obvious signs of ectoparasite infestation;								
	— is not intended for slaughter under a national programme of infectious or contagious disease eradication;									
	— meets the requirements attested in points II.1. to II.3. of this certificate;									
	_	is accomp	anied by the written declaration	on, 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 of dispatch									
	II.1.1.	a country	a country or part of the territ	tory of a country which at t	of country or part of the territory of the date of issuing this certificate has the Code gned to Sanitary Group(2);					
	II.1.2.	(Trypano	oma equiperdum), glanders	(Burkholderia mallei), eq	orily notifiable: African horse sickness, dourin uine encephalomyelitis (of all types includin , vesicular stomatitis, rabies and anthrax;					
	II.1.3.	the anima	is dispatched from a country	or part of the territory of a	country:					
		a)	which there has been no cli African horse sickness duri	nical, serological (in unvac ng the period of 2 years p	n accordance with Directive 2009/156/EC and i cinated equidae) or epidemiological evidence of prior to the date of dispatch and in which ther ng the period of 12 months prior to the date of					
		b)	in which Venezuelan equine the date of dispatch;	e encephalomyelitis has no	ot occurred during the period of 2 years prior t					
		c)	in which dourine has not occ	curred during the period of	6 months prior to the date of dispatch;					
		d)	in which glanders has not o	ccurred during the period o	f 6 months prior to the date of dispatch;					
	II.1.4.	points II.1		ntact with animals from ho	y knowledge for the time periods referred to i Idings, which were subject to prohibition order ast for:					
		(³) [II.1.4.1.	in the case of equidae susp	ected of having contracted	dourine,					
					ast actual or possible contact with an anima nfected with <i>Trypanosoma equiperdum</i> ;]					
			(⁴) and/or [in the case of a s	stallion, until the animal is c	castrated;]					
			(⁴) and/or [30 days following after all animals o		f the cleansing and disinfection of the premise					

Re-entry after temporary export of not more than 90 days

		Re-entry after	temporary export of not more than 90 days Specific races — Registered horse		
		II.a. Certificate reference number	II.b. Local reference number		
(³) [II.1.4.2.	in the cas	e of glanders,			
	(⁴) either	subjected with positive results to a test for	hs beginning on the day on which the equidae suffering from the disease or d with positive results to a test for the detection of the causative pathogen deria mallei or antibodies to that pathogen, were killed and destroyed;]		
	(⁴) and/or		following the date of completion of the cleansing and disinfection of the premises animals of susceptible species have been killed and destroyed;]]		
II.1.4.3.	in the cas	case of equine encephalomyelitis of any type,			
	(⁴) either	[6 months beginning on the day on which the e slaughtered;]	quidae suffering from the disease have been		
	(⁴) and/or	[6 months beginning on the day on which the environment of the law	5		

Nile Fever, Eastern equine encephalomyelitis or Western equine encephalomyelitis have died, been removed from the holding or fully recovered;]
(⁴) and/or [30 days following the date of completion of the cleansing and disinfection of the premises

II.1.4.4. in the case of equine infectious anaemia, until the date on which, the infected animals having been slaughtered, the remaining equine animals on the holding have shown a negative reaction in an agar gel immunodiffusion test (AGID or Coggins test) carried out on blood samples collected on two occasions 3 months apart;

after all animals of susceptible species have been slaughtered;]

- II.1.4.5. in the case of vesicular stomatitis,
 - (⁴) either [6 months following the last case;]
 - (⁴) and/or [30 days following the date of completion of the cleansing and disinfection of the premises after all animals of susceptible species have been slaughtered;]
- II.1.4.6. in the case of rabies, 30 days following the last case and the date of completion of the cleansing and disinfection of the premises;
- II.1.4.7. in the case of anthrax, 15 days following the last case and the date of completion of the cleansing and disinfection of the premises;
- II.1.5. to the best of my knowledge, during the period of 15 days prior to the date of dispatch the animal has not been in contact with equidae infected or suspected of an infectious or contagious disease.
- II.2. Attestation of residence and pre-export isolation
- II.2.1. The animal was imported into the country or part of the territory of the country of dispatch on (insert date)
 - (⁴) *either* [directly from the EU Member State (*insert name of EU Member State*) for the participation in
 - (⁴) either [The Japan Cup;]
 - (⁴) or [The Melbourne Cup;]
 - (⁴) or [The Dubai Racing World-Cup;]
 - (⁴) or [The Hong Kong International Races;]

▼M1

COUNTRY

			a Certificate reference number	11 6	l ocal reference number
			.a. Certificate reference number	11.6	b. Local reference number
	S	Singapore (4			merica (⁴), Hong Kong (⁴), Japan (⁴) varticipation in International Group/Grade
II.2.2.			ined and based on the declaration this certificate, the animal was:	of the owner	of the horse (4) or representative of the
		itinuously o tificate inclu		ays, the date	e of scheduled return in accordance with
					Froup/Grade meetings outside Australia ore, the United Arab Emirates or Qatar;
			s under veterinary supervision, acc a of lower health status except durin		in separated stables without coming into
II.2.3.	the animal e this health ce		ountry of dispatch under animal hea	alth conditior	ns at least as strict as those laid down ir
II.3.	Attestation o	f animal wel	fare		
			Box I.28. was examined today (1) a made to protect its health and well-b		o be transported on the intended journe ely at all stages of the journey.
Notes:					
Part I:					
Box I.8.:			country or part of the territory of the Regulation (EU) 2018/659.	he country a	as appearing in column 3 of Annex I to
Box I.15.:		s to be prov	ded. In case of unloading and reload		number (aircraft) or name (ship) and signor must inform the Border Inspection
Box I.23.:	The containe	r number aı	nd the seal number (if applicable) sh	ould be inclu	ded.
Box I.28.:	identification the identifica	document a tion system	as defined in Article 2(b) of Commiss (such as ear tag, tattoo, brand, tran	sion Impleme sponder) and	which permits to link the animal to the enting Regulation (EU) 2018/659. Specify d the anatomic place used on the animal name of the competent authority which
	Age: Date of	birth (dd/mi	n/yyyy).		
	Sex (M = ma	le, F = fema	le, C = castrated).		
Part II:					
	certificate must lember State of			orking day be	efore loading of the animal for dispatch to
to the refer	e date of autho red to in point l	risation for I.1.1., or du	re-entry into the Union from the res	pective cour asures have	when the animal was loaded either prio try or part of the territory of the country been adopted by the Union against the dispatch.
			the territory of the country, and thission Implementing Regulation (EU		Group as appearing in columns 3 and 5
(³) Delet	e statement if tl	ne attestatio	n in point II.1.3. applies to the entire	country of d	ispatch.

COUN	NTRY	Re-entry after	temporary export of not more than 90 days Specific races — Registered horse				
		II.a. Certificate reference number	II.b. Local reference number				
This	s health certificate shall:						
(a)		guage understood by the certifying officer and he Member State where the registered horse					
(b)	be made out to a single consi	gnee;					
(c)	be signed and stamped in a c	olour different to the colour of the printing;					
(d)	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	cial veterinarian						
	Name (in capital letters):		Qualification and title:				
	Date:		Signature:				
	Stamp:						

				owner or representative of the ov orary export of a registered hors		
lder	ntification o	f the animal (¹)			
Sp	ecies (Scie	ntific name)	Identification system	Identification number	Age	Sex
	Equus ca	aballus				
l, th	e undersigi	ned owner (²)	or representative of the owne	er (²) of the registered horse descrit	oed above, hereby dec	lare, that:
_	the horse					
	(²) either		arily exported from the Union days prior to this declaration;	to the country of dispatch on]		(insert date
	(²) or		country of dispatch on	(insert date) from v of dispatch);]		(insert name c
_	the horse	has been tem	nporarily exported from the Ur	nion to take part in		
	(²) either	[The Japan (Cup;]			
	(²) or	[The Melbou	rne Cup;]			
	(²) or	[The Dubai F	Racing World-Cup;]			
	(²) or	The Hong K	ong International Races;]			
	(²) or		I Group/Grade meetings in Angapore (²), United Arab Emin	Australia (²), Canada (²), the Uniterates (²) or Qatar (²);]	ed States of America	(²), Hong Kong (²
_			days prior to the date of dispatransmissible to equidae;	atch the horse has not been in con	tact with animals suffe	ring from infectiou
_				n as applicable in accordance wite country of dispatch are fulfilled;	th point II.2. of the ac	companying healt
_	the transp of the jou		e effected in such a way that	health and well-being of the horse	e can be protected effe	ctively at all stage
Nan	ne and add	ress of the ow	vner (²) or representative (²):			
Date	e:		(dd/mm/yyyy)			
(1)	Article 2(b	 of Commissi 		identifier which permits to link the anin EU) 2018/659. Specify the identificati		
		ort accompanies of birth (dd/mm		e stated and the name of the competent	t authority which validated	l it.
	Sex (M = r	nale, F = female	e, C = castrated).			
(²)	Delete as a	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

CC	UN	TRY:

Veterinary certificate to EL	Veterinary	y certificate	to EU
------------------------------	------------	---------------	-------

	I.1. Consignor Name Address	I.2. Certificate reference No I.2.a.									
	Tel.	I.3. Central competent authority									
		I.4. Local competent authority									
Part I : Details of dispatched consignment	I.5. Consignee Name Address Postcode Tel.	1.6.									
tails of disp	I.7. Country ISO code I.8. Region Code of origin	I.9. Country of ISO code I.10. Region destination destinat									
: De	I.11. Place of origin	I.12. Place of destination									
Part I	Name Approval number Address	Name Address									
		Postcode									
	I.13. Place of loading	I.14. Date of departure									
	I.15. Means of transport	I.16. Entry BIP in EU									
	Aeroplane Ship Railway wagon Road vehicle Other										
	Identification Documentary references	I.17. No(s) of CITES									
	I.18. Description of animal	I.19. Commodity code (i 01 01	HS code)								
		I.20. Quantity									
	l.21.	I.22. Number	of packages								
1	I.23. Seal/Container No	1.24.									
	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	1									
	Species (Scientific name) Identification system Identification number Age										

COUNT	RY			Import - Registered horse, registered equine animal or equine animal for breeding and production						
			II.a.	Certificate	reference number	II.b.	Local reference number			
II.	Attestat	ion of anima	health an	d welfare						
- (1)e (1)o (1)o - con cat - wa - is r - me	 meets the requirements attested in points II.1. to II.5. of this certificate; 									
11.1.		1999-19 1			• • • • • • • •					
II.1.1.	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,									
II.1.2.	(Trypand	untry of dispatch the following diseases are compulsorily notifiable: African horse sickness, dourine <i>psoma equiperdum</i>), glanders (<i>Burkholderia mallei</i>), equine encephalomyelitis (of all types including elan equine encephalomyelitis), equine infectious anaemia, vesicular stomatitis, rabies and anthrax;								
II.1.3.	the anim	al is dispatched from a country or part of the territory of country								
	a)	and in whi evidence in which t	which is considered free from African horse sickness in accordance with Directive 2009/156/EC and in which there has been no clinical, serological (in unvaccinated equidae) or epidemiological evidence of African horse sickness during the period of 2 years prior to the date of dispatch and in which there have been no vaccinations against the disease during the period of 12 months prior to the date of dispatch;							
	b)		in which Venezuelan equine encephalomyelitis has not occurred during the period of 2 y to the date of dispatch;							
	c)						prior to the date of dispatch;			
	d)				• •		prior to the date of dispatch;			
⁽¹⁾ either	[e)	in which vesicular stomatitis has not occurred during the period of 6 months prior to the dispatch;]								
⁽¹⁾ or	[e)	dispatch, a of 21 day	in which vesicular stomatitis has occurred during the period of 6 months prior to the date o dispatch, and a blood sample taken from the animal on(insert date), within a period of 21 days prior to the date of dispatch, was tested with negative result for antibody to the vesicular stomatitis virus							
		⁽¹⁾ either	in a virus r	neutralisation t	test at a serum dilu	tion of 1 in 3	32;]]			
					ce with the relevant rial Animals of the		the Manual of Diagnostic Test			

COUNTRY

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

			II.a.	Certificate ref	erence number	II.b.	Local reference number			
II.1.4.	the animal does not come from a holding, and to the best of my knowledge for the time periods referred to points II.1.4.1. to II.1.4.7. was not in contact with animals from holdings, which were subject to prohibitic orders for the reasons referred to in points II.1.4.1. to II.1.4.7. and which last for:									
	⁽⁴⁾ [II.1.4.1.	.1.4.1. in the case of equidae suspected of having contracted dourine, (1) <i>either</i> [6 months beginning on the date of the last actual or possible contact with an anim suspected of having contracted dourine or infected with <i>Trypanosoma equiperdum</i>								
		(1)and/or	[in the case	the case of a stallion, until the animal is castrated;]						
		(1)and/or	[30 days following the date of completion of the cleansing and disinfection of the premises after all animals of susceptible species have been slaughtered;]]							
	⁽⁴⁾ [II.1.4.2.	.4.2. in the case of glanders,								
		⁽¹⁾ either	[6 months b subjected w	b) generics, and the sequence of the sequen						
		(1)and/or	[30 days following the date of completion of the cleansing and disinfection of the premises after all animals of susceptible species have been killed and destroyed;]]							
	II.1.4.3.	in the cas	se of equine e	ncephalomyelit	tis of any type,					
		⁽¹⁾ either		months beginning on the day on which the equidae suffering from the disease hav een slaughtered;]						
		⁽¹⁾ and/or	nd/or [6 months beginning on the day on which the equidae infected with the virus West Nile Fever, Eastern equine encephalomyelitis or Western encephalomyelitis have died, been removed from the holding or fully recover							
		(1)and/or					ansing and disinfection of to been slaughtered;]			
	II.1.4.4.	in the case of equine infectious anaemia, until the date on which, the infected animals h been slaughtered, the remaining equine animals on the holding have shown a negative re in an agar gel immunodiffusion test (AGID or Coggins test) carried out on blood samples col on two occasions 3 months apart;								
	II.1.4.5.	in the cas	se of vesicula	stomatitis,						
		(1)either	[6 months fo	llowing the last	case;]					
		⁽¹⁾ and/or					ansing and disinfection of t been slaughtered;]			
	II.1.4.6.			abies, 30 days following the last case and the date of completion of the cleansing of the premises;						
	II.1.4.7.	in the case of anthrax, 15 days following the last case and the date of completion of the cleans and disinfection of the premises;								
II.1.5.					f 15 days prior to d of an infectious		of dispatch the animal has i			

COUNTRY

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

			23						
			II.a	a.	Certificate reference numbe	r II.b		Local reference number	
II.2.	Attestation	of resider	nce and pr	e-e	export isolation				
⁽¹⁾ either	[II.2.1.	During a period of at least the 90 days prior to the date of dispatch, or since birth if the animal is less than 90 days old, or since entry if the animal was imported directly from the Union during a period of 90 days prior to the date of dispatch, the animal has been resident on holdings under veterinary supervision situated in a country or part of the territory of a country which is:							
	(1)(5)either	[assigned to Sanitary Group A, and during the period of at least 30 days prior to the date of dispatch, it was kept apart from equidae not of equivalent health status;]]							
	(1)(5) 0 r	[assigned to Sanitary Groups B, C, D or G, and during the period of at least 30 days prior to the date of dispatch, it was kept in pre-export isolation under veterinary supervision without coming into contact with equidae not of equivalent health status;]]							
	(1)(5) 0 <i>r</i>				Group E, and it was kept in I.11., protected from vector ins		ved	isolation centre described a	
		⁽¹⁾ either [during the period of at least 40 days prior to the date of dispatch;]]]							
		⁽¹⁾ or [during the period of at least 30 days prior to the date of dispatch from a co- dispatch which is recognised by the OIE as officially free of African horse sickr							
			⁽¹⁾ either		he animal is a registered hors nplementing Regulation (EU) 2			in Article 2(c) of Commission	
			⁽¹⁾ or	si	he country of dispatch is not a ickness has occurred during ispatch;]]]]				
(1)(5) or	[II.2.1.	The animal is dispatched from a country of which at least a part of the territory of the country is assigned to Sanitary Group F, and during the period of at least 90 days prior to the date of dispatch, or since birth if the animal is less than 90 days old, it was resident on holdings under veterinary supervision and was kept during the period of at least 60 days prior to the date of dispatch, or since entry if it was imported directly from the Union during the period of 60 days prior to the date of dispatch horse sickness in accordance with the Union legislation and underwent the pre export isolation							
	⁽¹⁾ either	[in the approved vector-protected quarantine station of							
	⁽¹⁾ or	[permanently confined in the approved vector-proof quarantine station of (<i>insert name of quarantine station</i>) during the period of at least 14 days prior to the date dispatch and constant monitoring of the vector protection has proven absence of vectors ins the vector-protected part of the quarantine station.]]							
II.3.	Attestation	on of vaccination and health tests							
⁽¹⁾ either	[11.3.1.	The animal was not vaccinated against African horse sickness in the country of dispatch and there is no information suggesting previous vaccination;]							
(1) or	[II.3.1.				nated against African horse sid		d thi	s vaccination was carried o	
	(1)either	[more than 12 months prior to the date of dispatch;]]							
	(1)or	Image the			and less than 12 months prior	-	1		

COUNTRY

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

		II.a. Certificate reference number II.b. Local reference number
⁽¹⁾⁽⁵⁾ Or	[II.3.1.	The animal is dispatched from a country or part of the territory of a country which is assigned Sanitary Group F and was vaccinated against African horse sickness on
	II.3.2.	the animal was not vaccinated against Venezuelan equine encephalomyelitis during the peric of 60 days prior to the date of dispatch from
	⁽¹⁾ 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;]
	⁽¹⁾⁽⁵⁾ Or	[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 dispate and Venezuelan equine encephalomyelitis occurs in the remaining parts of the territory of the country of dispatch, and
		(1)either [is vaccinated against Venezuelan equine encephalomyelitis with a complete prima course and revaccinated according to manufacturer's recommendations not less tha 60 days and no more than 12 months prior to the date of dispatch, and was kept vector-protected quarantine for a period of at least 21 days prior to the date of dispatc and during that period remained clinically healthy, and its body temperature, take daily, remained within the normal physiological range, and any equine animal on th same holding which showed a rise in body temperature, taken daily, was subjected a blood test for virus isolation for Venezuelan equine encephalomyelitis with negative results;]]
		(1)or [is not vaccinated against Venezuelan equine encephalomyelitis and was kept vector-protected quarantine for a period of at least 21 days, and during that perior remained clinically healthy, and its body temperature, taken daily, remained within th normal physiological range, and any equine animal on the same holding which shows a rise in body temperature, taken daily, was subjected to a blood test for virus isolatic for Venezuelan equine encephalomyelitis with negative results, and the animal to the dispatched was subjected to a diagnostic test for Venezuelan equine encephalomyeliti with negative result conducted on a sample taken not less than 14 days after the da of entry into the vector protected quarantine and remained protected from vect insects until dispatch;]]
		(1)or [was subjected to a haemagglutination inhibition test for Venezuelan equir encephalomyelitis carried out by the same laboratory on the same day on sample taken on two occasions with an interval of 21 days on (insert date) and o
	¹⁾ [II.3.3.	the animal is an uncastrated male equine animal older than 180 days, and
	⁽¹⁾ either	[is dispatched from a country in which equine viral arteritis (EVA) is a compulsorily notifiab disease and has not been officially reported during the period of 6 months prior to the date dispatch;]]
	⁽¹⁾ 0r	[was tested on a blood sample taken on (insert date), within a period of 21 days print to the date of dispatch, by virus neutralisation test for EVA with negative result at a serum dilution of 1 in 4;]]

	RY		Import - Registered horse, registered equine animal or equine animal for breeding and production					
			II.a. Certificate reference number II.b. Local reference number					
	⁽¹⁾ Or	21 days	sted on an aliquot of its entire semen taken on <i>(insert date)</i> , within a period c s prior to the date of dispatch, by virus isolation test , polymerase chain reaction (PCR) o he PCR for EVA with negative result.]]					
	⁽¹⁾ 0r	[was va and re-	accinated against EVA on					
		⁽¹⁾ either	 [before 1 October 2018, on the day a blood sample was taken that was subsequentl tested in a virus neutralisation test for EVA with negative result at a serum dilution of in 4;]]] 					
		⁽¹⁾ or	[before 1 October 2018, during a period of isolation of not more than 15 days under official veterinary supervision, commencing on the day a blood sample was taken whic was tested during that isolation period in a virus neutralisation test for EVA wit negative result at a serum dilution of 1 in 4;]]]					
		⁽¹⁾ Or	[at the age of 180 to 270 days, during a period of isolation under official veterinar supervision, during which the animal was subjected to a virus neutralisation test for EVA carried out with negative result at a serum dilution of 1 in 4, or carried out on th same day by the same laboratory with stable or declining titres on two blood sample taken at least 10 days apart;]]]					
		⁽¹⁾ or	[after the animal was subjected to a virus neutralisation test for EVA with negative resu at a serum dilution of 1 in 4, carried out on a blood sample taken not earlier than 7 day after commencing a period of uninterrupted isolation which lasted until 21 day following vaccination;]]]					
		⁽¹⁾ or	[at the age of 180 to 250 days, after the animal was subjected to a virus neutralisatio test for EVA carried out with negative result at a serum dilution of 1 in 4 or carried ou on the same day by the same laboratory with stable or declining titres on two bloo samples taken at least 14 days apart;]]]					
	⁽¹⁾ 0r	EVA ca blood s to the c	ubjected to a virus isolation test, polymerase chain reaction (PCR) or real-time PCR for arried out with negative result on an aliquot of its entire semen collected after the date sample of that animal taken on					
	⁽¹⁾ <i>or</i>		reviously tested positive for antibodies against the equine arteritis virus or has bee ated against EVA, and					
		a)	within a period of 6 months prior to the date of dispatch, was test mated, on tw consecutive days, to at least two mares which were kept in isolation during the 7 day prior to and until at least 28 days after test mating and which were subjected to tw serological tests for EVA with negative results at a serum dilution of 1 in 4 on bloo samples collected at the time of test mating and at least 28 days after the test mating and					
		b)	was subjected to a virus neutralisation test for EVA carried out on a blood sample take within 21 days prior to the date of dispatch on(insert date),					
			(1)either[with positive result at a serum dilution of at least 1 in 4;]]](1)or[with negative result at a serum dilution of 1 in 4;]]]					
⁽¹⁾ either	[11.3.4.	anaem	nimal is dispatched from Iceland, which is certified as officially free from equine infectiou nia, where it was continuously resident since birth and did not come into contact wit ae which have entered Iceland from other countries;]					
⁽¹⁾ or	[11.3.4.	the an Coggii	nimal was subjected with negative result to an agar gel immunodiffusion test (AGID on the stest) or to an ELISA for equine infectious anaemia carried out on a blood sample take					

COUNTRY

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

		11.6	a. Certificate reference number	II.b. Local reference number
⁽¹⁾ [II.3.5.	Sanitary reported complet 5 on a t	y Group B, d during a ment fixatio	D or E, or from China or Thailand period of 3 years prior to the d on test for glanders carried out with ole taken on	territory of a country which is assigned t , or from a country in which glanders wa ate of dispatch, and was subjected to n negative result at a serum dilution of 1 isert date), within a period of 30 days price
⁽¹⁾ [II.3.6.	from a of or F, or of 2 yea dourine not bee	from China ars prior to carried ou	part of the territory of a country who or Thailand, or from a country in we the date of dispatch, and was su twith negative result at a serum dil (<i>insert date</i>), within a period of 30 breeding during the period of at le	te animal older than 270 days dispatche ich is assigned to Sanitary Group B, D, /hich dourine was reported during a perio /bjected to a complement fixation test for ution of 1 in 5 on a blood sample taken o days prior to the date of dispatch, and ha ast 30 days prior to and after the date th
⁽¹⁾⁽⁵⁾ [II.3.7.	Sanitary	Western Country o	or D, and and Eastern equine encephalomye	territory of a country which is assigned t elitis have not been officially reported in th y of dispatch during a period of at least
	⁽¹⁾ or	to manufa the date	acturer's instructions within a perior of dispatch with inactivated vacc	orimary course and revaccinated accordin d of 6 months and at least 30 days prior t ine against Western and Eastern equin applied on
	⁽¹⁾ <i>or</i>	vector pr	otected quarantine, and during th tests for Western and Eastern eq	21 days prior to the date of dispatch in is period subjected to haemagglutinatio uine encephalomyelitis carried out by the
		⁽¹⁾ either	[on a sample of blood taken on 10 days prior to the date of dispat	
		⁽¹⁾ or	days on (insert date	o occasions with an interval of at least 2 and on (<i>insert date</i>), th n a period of 10 days prior to the date o lody titre;]]]
⁽¹⁾ [II.3.8.	Sanitary	Group G, o		territory of a country which is assigned to encephalitis has been officially reported
	⁽¹⁾ either	that holdi		of an area of at least 30 km radius arour of Japanese encephalitis during a period (;]]
	⁽¹⁾ 0r	the date		during a period of at least 21 days prior t riod the body temperature, taken daily nge, and was subjected
		⁽¹⁾ either	encephalitis carried out by the sam of blood taken on two occasions 	or virus neutralisation test for Japanes ne laboratory on the same day on sample s with an interval of at least 14 days of

(1)(5)or

COUNTRY					registered equine animal or or breeding and production
		II.a. Certi	icate reference number	II.b.	Local reference number
	⁽¹⁾ C	Japanes sample t	e encephalitis virus with aken not earlier than 7 da (insert date), and rem	n negative reason negative re	ection of antibodies against sult, carried out on a blood date the isolation commenced cted from vector insects until
	rev	accinated accor		commendati	complete primary course and ons during a period of not less late of dispatch;]]
⁽¹⁾⁽⁵⁾ either [II.3.9.	Sanitary Gi described ir on the same ⁽¹⁾ either [or on	oup E, and wa Annex IV to Dir day blood samples	s subjected to a serolo rective 2009/156/EC, whi taken on two occasions v	ogical test fo ch was carrie with an interva	a country which is assigned to or African horse sickness as ed out by the same laboratory al of between 21 and 30 days, (insert date), the second of ate of dispatch
	(1)6	ither [with neg	ative results in each case	e;]]]	
	⁽¹⁾ C	r [with pos	itive result in the first san	nple, and	
		⁽¹⁾ either		ntification tes	quently tested with negative st as described in Annex IV to

(1)or [the two samples were tested without more than a two-fold increase in antibody titre in a virus neutralisation test as described in point 2.4 of Chapter 2.5.1. of the OIE Terrestrial Manual for Diagnostic Tests and Vaccines;]]]]

⁽¹⁾ or	to the da	od sample taken on (insert date), within a period of 21 days prior te of dispatch, and the country or part of the territory of the country of dispatch nised by the OIE as officially free of African horse sickness and
	⁽¹⁾ either	[the animal is a registered horse as defined in Article 2(c) of Commission Implementing Regulation (EU) 2018/659;]]]
	⁽¹⁾ or	[the country of dispatch is not adjacent to a country in which African horse sickness has occurred during the period of 2 years prior to the date of dispatch;]]]

[II.3.9. the animal is dispatched from a country or part of the territory of a country which is assigned to Sanitary Group F and ⁽¹⁾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

⁽¹⁾either [with negative results in each case;]]]

⁽¹⁾or [with positive result in the first sample, and

(1)either [the second sample was subsequently tested with negative result in an agent identification test as described in Annex IV to Directive 2009/156/EC;]]]]

(1)or [the two samples were tested without more than a two-fold increase in antibody titre in a virus neutralisation test as described in point 2.4 of Chapter 2.5.1. of the OIE Terrestrial Manual for Diagnostic Tests and Vaccines;]]]]

COUNTRY Import - Registered horse, registered equine animal or equine animal for breeding and production II a. Certificate reference number ll b Local reference number (1)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 guarantine and within a period of 10 days prior to the date of dispatch;]] (1)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 11.4. Attestation of the transport conditions The animal is dispatched from a country or part of the territory of a country which is assigned to (1)either [1].4.1. Sanitary Group A, B, C, D, E or G and is transported directly to the Union, without passing through a market, marshalling or assembly centre and without coming into contact with other equidae not complying with at least the same health requirements as described in this health certificate.] (1)(5)or [1.4.1. The animal is dispatched from a country or part of the territory of a country which is assigned to Sanitary Group F and is transported 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 (1)either [to the airport under vector-protected conditions and arrangements have been made for the aircraft to be 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.]] (1)or [to a sea port in that country or part of the territory of the country under vector-protected conditions and arrangements have been made to transport it on a 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.]] 11.4.2. 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. 11.4.3. 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. 11.5. Attestation of animal welfare The animal described in Box I.28. was examined today⁽²⁾ and found fit to be transported on the intended journey and arrangements were made to protect its health and well-being effectively at all stages of the journey.

Notes:

Part I:	
Box I.8.:	Provide the code of the country or the part of the territory of the country as appearing in column 3 of Annex I to Commission Implementing Regulation (EU) 2018/659.
Box I.15.:	Registration number (railway wagons or container and lorries), flight number (aircraft) or name (ship) and information is to be provided. In case of unloading and reloading, the consignor must inform the BIP of entry into the EU.
Box I.23.:	The container number and the seal number (if applicable) should be included.

COL	JNTRY				e, registered equine animal of I for breeding and production
		II.a.	Certificate reference number	II.b.	Local reference number
Box I			caballus, Equus asinus, Equus quus grevyi, or indicate any cros		
	identification document	as defir	al must bear an individual ident ed in Article 2(b) of Commissior n (such as ear tag, tattoo, brand,	Implemen	ting Regulation (EU) 2018/65
	If a passport accompanie which validated it.	es the a	nimal, its number should be state	d and the r	name of the competent authori
	Age: Date of birth (dd/m	m/yyyy)			
	Sex (M = male, F = fema	ale, C =	castrated).		
Part	11:				
(1)	Delete as appropriate.				
(2)		ed on t	he day of loading or in the case	of a regist	tered horse on the last workin
			dispatch to the Member State of		
			all not be allowed when the an		
			dual registered equine animal or		
			country or part of the territory of		
			neasures have been adopted by		
			erritory of the country of dispatc		against the entry of live equida
(3)			erritory of the country and the S		oup as appearing in columns
			mmission Implementing Regulat		
(4)			point II.1.3. applies to the entire		
(5)			exclusively to a Sanitary Group		
	the country of dispatch or n	art of it	s territory, is assigned, may be l	off out pro	vided that the numbering of th
	subsequent statements is m			on out, pro	vided that the numbering of th
This	health certificate shall:				
(a)		anade	understood by the certifying offic	er and one	of the official languages of th
(4)	Member State of destination	and of	the Member State where the an	imal will on	ter Union territory and under
	the veterinary border checks		the member otate where the an		nor onion territory and underg
(b)	be made out to a single cons				
(c)			different to the colour of the print	ina:	
(d)			all sheets of paper required are		integrated whole and indivisib
/			al number of pages, and each		
			nose pages are stapled and star		
Offici	ial veterinarian				
	Name (in capital letters):			Qua	alification and title:
	Date:			Sig	nature:
	Stamp				
	oramp				

	Declaration by the owner or representative of the owner for entry into the Union of an equine animal							
Ide	ntification of	the anima	ll(1)					
	ecies (Scienti ne)	ific	Identification system	Identification number	Age	Sex		
I, th	ne undersigne	ed owner	²⁾ or representative of the	owner ⁽²⁾ of the animal descr	ibed above, hereby de	clare, that:		
-	the animal							
	⁽²⁾ either			part of the territory of the cou , or since birth if the animal is				
	⁽²⁾ or			e territory of the country of di of dispatch from a Member S		ired residence period of		
-			15 days prior to the date ous diseases transmissib	of dispatch the animal has r ble to equidae;	not been in contact with	n animals suffering from		
-	the conditi health cert	ons for re ificate for	esidence and pre-export the country or part of the	isolation as applicable in act territory of the country of dis	cordance with point II.: patch are fulfilled;	2. of the accompanying		
-			e transport as applicable e territory of the country of	in accordance with point II.4 of dispatch are fulfilled;	. of the accompanying	health certificate for the		
-	the transpo all stages			vay that health and well-bein	g of the animal can be	protected effectively at		
Na	me and addre	ess of the	owner ⁽²⁾ or representative	e ⁽²⁾ :				
Dat	e:		(dd/mm/yyyy)					
				(Signature)				
(1)	quagga, Eq Identificatio document a system (suc If a passpo validated it. Age: Date o	n system as defined thas ear ort accomposition of birth (do	a, Equus grevyi, or indica : The animal must bear a I in Article 2(b) of Comm tag, tattoo, brand, transpo panies the animal, its nu t/mm/yyyy).	Equus asinus, Equus africa te any cross between those, an individual identifier which hission Implementing Regula onder) and the anatomic plac umber should be stated and	permits to link the ani tion (EU) 2018/659. S se used on the animal.	mal to the identification pecify the identification		
(2)	Sex (M = m Delete as a		emale, C = castrated). e.					

Section B

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

COL	INTRY:	Veterinary certificate to B	EU
	I.1. Consignor Name	I.2. Certificate reference No I.2.a.	~
	Address Tel.	I.3. Central competent authority	
	ו ט ו.	I.4. Local competent authority	
Part I : Details of dispatched consignment	I.5. Consignee Name Address Postcode Tel.	1.6.	~
ails of disp	I.7. Country of ISO code I.8. Region Code origin of origin	I.9. Country of ISO code I.10.Region of Coordestination	le
: Deta	I.11. Place of origin	I.12. Place of destination	
Part I	Name Approval number Address	Name Address	
		Postcode	
	I.13. Place of loading	I.14. Date of departure	
υC	I.15. Means of transport Aeroplane Ship Railway wagon	I.16. Entry BIP in EU	
	Road vehicle Other Identification Documentary references	I.17. No(s) of CITES	
	I.18. Description of animals	I.19. Commodity code (HS code) 01 01	
		I.20. Quantity	
	I.21.	I.22. Number of package	es
	I.23. Seal/Container No	1.24.	~
	I.25. Animals certified for: Slaughter		
	1.26.	I.27. For import or admission into EU	2
	I.28. Identification of the animals	Identification number Are O	
	Species (Scientific name) Identification system	n Identification number Age S	ex

COUNTRY

Import - Equidae for slaughter

	2474224.1 00.249004	500 f								
				II.a.	Certificate reference number	II.b.	Local reference number			
	П.	Attestatio	on of animal h	ealth, an	imal welfare and public health					
Part II: Certification	- are - we info - are - me - are	e equidae for re examine estation; e not intend eet the requ	or slaughter as ed today ⁽¹⁾ and ed for slaughte irements attes	defined in d found fr er under a ted in poir	by certify, that the animals desc n Article 2(d) of Directive 2009/1 ee of clinical signs of diseases national programme of infectiou nts II.1. to II.5. of this certificate; ration, signed by the owner of the	56/EC; and of o is or conta	bvious signs of ectoparasit			
	II.1.		n on third cour	try or par	t of the territory of third country	and holdir	a of dispatch			
	2000/00/200 2000/2000/2000				t of the territory of third country					
	II.1.1.	country),	a country or pa	art of the t	erritory of a country, which on th anitary Group ⁽²⁾ ;					
	II.1.2.	dourine (Trypanosoma uding Venezu	equiperd	Ilowing diseases are compulso um), glanders (Burkholderia ma ne encephalomyelitis), equine in	allei), equ	ine encephalomyelitis (of a			
	II.1.3.	the anima	ls are dispatch	ned from a	country or part of the territory o	f country				
		a)	2009/156/E0 or epidemiol date of dispa	C and in v logical evid atch and in	free from African horse sic which there has been no clinical dence of African horse sickness in which there have been no vacc rior to the date of dispatch;	, serologic during the	cal (in unvaccinated equidae e period of 2 years prior to the			
		b)	in which Venezuelan equine encephalomyelitis has not occurred during the period of prior to the date of dispatch;							
		c)	in which dou	irine has r	not occurred during the period of	6 months	prior to the date of dispatch			
	And an and a second	d)	in which gla	nders has	not occurred during the period o	f 6 months	s prior to the date of dispatch			
	⁽³⁾ either	[e)	in which ves of dispatch;]		matitis has not occurred during t	he period	of 6 months prior to the date			
	⁽³⁾ or	[e)	dispatch, an within a per	id a blood iod of 21	matitis has occurred during the sample taken from each of the days prior to the date of dispato lar stomatitis virus	animals o	ninsert date)			
			⁽³⁾ either [in	n a virus n	eutralisation test at a serum dilu	tion of 1 ir	n 32;]]			
					A in accordance with the relevant accines for Terrestrial Animals of					
	II.1.4.	in points I	I.1.4.1. to II.1.	4.7. have	dings, and to the best of my known not been in contact with animal referred to in points II.1.4.1. to II	s from ho	Idings, which were subject to			
	(*	⁴⁾ [II.1.4.1.	in the case of	of equidae	suspected of having contracted	dourine,				
					eginning on the date of the last a f having contracted dourine or in					
			⁽³⁾ and/or [ir	the case	of a stallion, until the animal is o	astrated;]				
					llowing the date of completion of the all animals of susceptible spe					

COUNTRY

Import - Equidae for slaughter

			1	I.a.	Certificate reference number	II.b.	Local reference numbe	
	⁽⁴⁾ [II.1.4.2.	in the cas	e of glan	der	S.			
		⁽³⁾ either	[6 month subjecte	ns b d w	eginning on the day on which th ith positive results to a test for t <i>a mallei</i> or antibodies to that pa	he detec	tion of the causative pathoge	
		⁽³⁾ and/or			llowing the date of completion of ter all animals of susceptible sp			
	II.1.4.3.	in the cas	e of equi	ne e	encephalomyelitis of any type,			
		⁽³⁾ either			beginning on the day on which slaughtered;]	the equid	lae suffering from the diseas	
		⁽³⁾ and/or	West N	lile	eginning on the day on which the Fever, Eastern equine en nyelitis have died, been removed	cephalon	nyelitis or Western equir	
		⁽³⁾ and/or			llowing the date of completion of ter all animals of susceptible sp			
	II.1.4.4.	been slau reaction i	ughtered, n an aga	the ar g	infectious anaemia, until the dat e remaining equine animals or lel immunodiffusion test (AGID two occasions 3 months apart;	the hol	Iding have shown a negativ	
	II.1.4.5.			of vesicular stomatitis,				
		⁽³⁾ either	[6 month	ns fo	ollowing the last case;]			
		⁽³⁾ and/or			llowing the date of completion of ter all animals of susceptible sp			
	II.1.4.6.				, 30 days following the last ca tion of the premises;	se and t	the date of completion of th	
	II.1.4.7.				x, 15 days following the last cation of the premises;	ase and	the date of completion of th	
II.1.5.					ng the period of 15 days prior to nfected or suspected of an infect			
II.2.	Attestation	n of residen	ce and p	re-e	export isolation			
II.2.1.	period of	90 days prid under veteri	or to the o	date	n the country or part of the territu e of dispatch, or since birth if the sion, and they are dispatched fr	animals	are less than 90 days old, c	
	⁽³⁾ either				Group A and during the period pt apart from equidae not of equ			
	⁽³⁾ or	[assigned date of di	to Sanita	ary ney	Groups B, C or D and during th were kept in pre-export isolatic ith equidae not of equivalent he	e period n under	of at least 30 days prior to the veterinary supervision without	
	⁽³⁾ or				Group E and for the period of at le approved isolation centre descr			

COUNTR	ΥF				Imp	oort - Equidae for slaughte
			II.a.	Certificate reference number	II.b.	Local reference number
II.3.	Attestatio	n of vaccination a	nd hea	alth tests		
⁽³⁾ either	[11.3.1.			t vaccinated against African hors ation suggesting previous vaccina		ss in the country of dispate
⁽³⁾ or	[II.3.1.			cinated against African horse sick ths prior to dispatch;]	ness, and	d this vaccination was carrie
	II.3.2.	the animals we days prior to di		vaccinated against Venezuelan e	quine end	cephalomyelitis during the 6
	⁽³⁾ either			I parts of the territory are free of 2 years prior to the date of dispa		lan equine encephalomyelit
	⁽³⁾⁽⁵⁾ Or	Venezuelan ec dispatch and V	uine e /enezu	of a country which is assigned to s incephalomyelitis for a period of relan equine encephalomyelitis of of dispatch, and	at least	2 years prior to the date
		prima less were date temp equir daily	ary cou than 60 kept ir of disp erature ne anin was	nated against Venezuelan equin rse and revaccinated according to 0 days and not more than 12 moi n vector-protected quarantine for a atch, and during that period rema a, taken daily, remained within the nal on the same holding which sho subjected to a blood test for vi nyelitis with negative result;]]	manufac nths prior a period c ained clin a normal wwed a ris	cturer's recommendations n r to the date of dispatch, ar of at least 21 days prior to the ically healthy, and their boo physiological range, and a se in body temperature, take
		in ve dispa temp equir daily ence subje nega entry	ctor-pr ttch. a erature e anin was phalon ected t tive res into t	accinated against Venezuelan eq otected quarantine for a period o ind during that period remaine a taken daily. remained within the al on the same holding which sho subjected to a blood test for vi nyelitis with negative results, and o a diagnostic test for Venezu sult conducted on a sample taken he vector-protected quarantine a dispatch;]]	f at least d clinica a normal wed a ris rus isola d the ani uelan eq not less	t 21 days prior to the date lly healthy. and their boo physiological range, and ar se in body temperature. take tion for Venezuelan equin mals to be dispatched wer uine encephalomyelitis wii than 14 days after the date
³⁾⁽⁵⁾ eithei	r [II.3.3.	infectious anae	mia, w	atched from Iceland, which is c here they have been continuously dae which have entered Iceland fr	resident	t since birth and did not corr
⁽³⁾ or	[II.3.3.	an ELISA for ea	quine ir on	jected to an agar gel immunodiffunt nfectious anaemia carried out with <i>(insert date)</i> , this being w	negative	e result in each case on bloc
(3	³⁾ [II.3.4.	to Sanitary Gro period of 3 yea test for glander	oup B, rs prio s carrie taken	ched from a country or part of the D or E, or from a country in wh r to the date of dispatch, and we ed out with negative result in eacl on (insert dat of dispatch;]	ich gland re subjec n case at	ders was reported during the ted to a complement fixation a serum dilution of 1 in 5 c

Import - Equidae for slaughter

COUNTRY						ort - Equidae for slaughte
		1	.a.	Certificate reference number	II.b.	Local reference number
⁽³⁾ [II.3.5.	from a co E or from of dispat negative	ountry or p n a country tch, and w result in	in vere ea	trated males or female equine an of the territory of a country which which dourine was reported during e subjected to a complement fixe ach case at a serum dilution of (insert date), this being within the	is assign the period ation test of 1 in 5	ed to Sanitary Group B, D o od of 2 years prior to the dat for dourine carried out wit 5 on blood samples take
⁽³⁾⁽⁵⁾ [II.3.6.		als are dis ary Group (ched from a country or part of the r D, and	territory o	f a country which is assigne
	⁽³⁾ either	the coun	ntry	nd Eastern equine encephalomyel or part of the territory of the cour to the date of dispatch;]]		
	⁽³⁾ Or	accordin 30 days Eastern	g to prio eo	s were vaccinated with a completed or manufacturer's instructions with or to the date of dispatch with ina quine encephalomyelitis, the (insert date);]]	in the per ctivated v	iod of 6 months and at leas vaccine against Western an
	⁽³⁾ 0r	this period	od s	were kept for at least 21 days pro- subjected to haemagglutination in ephalomyelitis on	hibition te	ests for Western and Easter
		⁽³⁾ either	or	sample of blood taken from each (insert date), within dispatch, with negative result in each	the period	d of 10 days prior to the dat
		⁽³⁾ or	tw da w	amples of blood taken from each vo occasions with an interval of a ate) and on (insert ithin the period of 10 days prior to antibody titre;]]]	t least 21 date), the	days on (insel
⁽³⁾ [II.3.7.	to Sanita	ary Group	G,	ched from a country or part of the or from a country in which Japa uring the past 2 years, and the an	inese end	
	⁽³⁾ either	[come fro those ho	om oldir	holdings situated in the centre of a ngs where there has been no ca least 21 days prior to the date of	in area of se of Jap	anese encephalitis during
	⁽³⁾ 0r	the date	of d	n a vector-protected quarantine du ispatch, and during that period the b remained within the normal physiol	ody temp	erature of each of the animals
		⁽³⁾ either	er sa da se di th	o a haemagglutination inhibition on neephalitis carried out by the sa amples of blood taken on two occ ays on	ime labor casions w ind on period o fold increa	ratory on the same day o vith an interval of at least 1 <i>(insert date)</i> , th f 10 days prior to the date of ase in antibody titre betwee

COUNTRY

Import - Equidae for slaughter

						or - Equidae for slaught		
2				II.a.	Certific	ate reference number	II.b.	Local reference numbe
			⁽³⁾ 0r	Ja sa co	panese mple ta mmence	encephalitis virus with n ken not earlier than 7	egative re 7 days af	ction of antibodies again sult, carried out on a bloc ter the date the isolatic nained protected from vector
		⁽³⁾ or	revacc	inated	accordi		commenda	omplete primary course an ations during a period of no the date of dispatch;]]
(3)(5) [II.3.8 .	Sanitary (Group E, IV to D	and v	vere subj	ected to a serological test	for African	country which is assigned horse sickness as describe the same laboratory on th
		⁽³⁾ either	occasi date) a	ons w and or	ith an int	erval of between 21 and	30 days, o he second	n the consignment on two on(inse d of which was taken with
			⁽³⁾ eithei	r [w	ith negat	ive result in each case;]]]	
			⁽³⁾ or	[w	ith positi	ve results in the first san	nple, and	
				(3)6	either		n agent ide	quently tested with negativentification test as describe S/EC;]]]]
				(3)	or	tested without more that in a virus neutralisation	an a two-fo on test as OIE Terres	al of the consignment we old increase in antibody tit described in point 2.4 strial Manual for Diagnost
		⁽³⁾ 0r	in the o to the dispate not adj	consig date ch is re acent	nment o of dispa ecognise to a cou	n (insert of tch, and the country or ed by the OIE as officially	ate), within part of the free of A prse sickne	ten from each of the anima in the period of 10 days pri- e territory of the country frican horse sickness and ass has occurred during th
II.4.	Attestation	of the trai	nsport co	onditio	ons			
⁽³⁾ either	[II.4.1.	slaughte	rhouse o y centre	on the referr	territory ed to in	of the Union, without pas	ssing throu 2009/156/E	are transported directly to ugh a market, marshalling (EC, and without coming in hion.]
⁽³⁾ 0r	[II.4.1.	slaughter marshalli the same	rhouse o ing or as e Memb	on the semb er St	territory ly centre ate, from	of the Union they pass of referred to in Article 7(1 n where they are trans	nly throug) of Direct ferred dire	animals are transported to h a single approved market ive 2009/156/EC situated actly to the slaughterhous the entry into the Union.]
	II.4.2.	complyin	g with a	t leas	t the sa		as describ	ct with other equidae not obtain this health certification

COUNT	інү			Imp	ort - Equidae for slaughte
		II.a.	Certificate reference number	II.b.	Local reference number
	disinfe	cted before loading	or containers in which the animals a ng with a disinfectant officially recog that faeces, urine, litter or fodder c	nised in the	e third country of dispatch and
II.5.	Attestation of anim	nal welfare			
			were examined today ⁽¹⁾ and four een made to protect their health a		
II.6.	Attestation of pub	lic health			
	oestrogenic, andr zootechnical treat The guarantees c	ogenic, gestage ment as defined overing live equid	8. have not received any stilber nic or beta-agonist substances for in Article 1(2)(b) and(c) of Directi dae provided by the residue plan s	or purpose ve 96/22/I	es other than therapeutic o
	with Article 29 of	Directive 96/23/E	C are fulfilled.		
Notes:					
Part I:					
Box I.8			or part of the territory of the count equilation (EU) 2018/659.	try as app	earing in column 3 of Annex
Box I.1	5.: Registration nur and information	nber (railway wa	gons or container and lorries), f d. In case of unloading and relo		
Box I.2			eal number (if applicable) should	be include	ed.
			s caballus", "Equus asinus" or "E		
	Identification s animal to the id transponder) a	vstem: Each of t	he animals must bear an individument. Specify the identification splace used on the animal.	ual identif	ier which permits to link the
	Sex (M = male,	F = female, C =	castrated).		
Part II:					
(1)	The certificate must destination in the U The import of these prior to the date of country or part of the	nion. equine animals authorisation for e territory of a co en adopted by the	he day of loading of the animals for slaughter shall not be allowed imports of live equidae for slaugh puntry mentioned under point II.1. e Union against the entryof equid	when the ter into the 1., or duri	animals were loaded eithe The Union from the respective ng a period where restrictive
(2)	Code of the country	or part of the te	rritory of the country and the Sani nmission Implementing Regulatio		
(3)	Delete as appropria	ite.			
(4) (5)	Statements that rela	te entirely and ex	point II.1.3. applies to the entire cclusively to a Sanitary Group diffe s territory, is assigned, may be le	erent from	the Sanitary Group to which

		II.a.	Certificate reference number	II.b.	Local reference numb
This	health certificate shall:				
(a) (b) (c) (d)	the Member State of dest undergo the veterinary bo be made out to a single or be signed and stamped in consist of a single sheet indivisible by inserting pa	ination an rder check onsignee; a colour of of paper ge numbe		e animals ing; 1 are part nd each pa	will enter Union territory a of an integrated whole a age shall bear the certifica
Offic	ial veterinarian				
	Name (in capital letters):			Qua	alification and title:
	Date:			Sig	nature:

			e owner or representative o of consignments of live equ		
dentificatio	on of the anim	als ⁽¹⁾			
Species (S name)	cientific	Identification system	Identification number	Age	Sex
, the unde	rsigned owner	(2) or representative of the	owner ⁽²⁾ of the animals descri	bed above, hereby declar	e, that:
	nimals have re of dispatch;	mained in the country or p	art of the territory of the count	ry of dispatch for at least	90 days prior to th
		15 days prior to the date o jious diseases transmissibl	f dispatch the animals have ne e to equidae;	ot been in contact with an	imals suffering fror
			solation as applicable in acco territory of the country of disp		the accompanyin
		ne transport as applicable in the territory of the country of	n accordance with point II.4. dispatch are fulfilled;	of the accompanying hea	th certificate for th
	ansportation v ages of the jou		ay that health and well-being	of the animal can be pro	tected effectively a
the a	nimals will be	sent			
(2)eith		y from the premises of disp quidae not of the same hea	atch to the slaughterhouse of alth status;]	f destination without comi	ng into contact wit
⁽²⁾ or	marsha		he slaughterhouse of destination of ferred to in Article 7(1) of Dir he same health status;]		
Name and	address of the	owner ⁽²⁾ or representative	(2).		
Date:		(dd/mm/yyyy)			
			(Signature)		
Identif docum systen	<i>lication system</i> nent as define n (such as ear	n: The animal must bear a d in Article 2(b) of Commi tag, tattoo, brand, transpor	(Signature) us asinus, or indicate any cro n individual identifier which p ssion Implementing Regulati nder) and the anatomic place	ess between those. ermits to link the animal on (EU) 2018/659. Spec used on the animal.	ify the identification
Identif docum systen	<i>lication system</i> nent as define n (such as ear	n: The animal must bear a d in Article 2(b) of Commi tag, tattoo, brand, transpor	(Signature) us asinus, or indicate any cro n individual identifier which p ssion Implementing Regulati	ess between those. ermits to link the animal on (EU) 2018/659. Spec used on the animal.	ify the identification
Identif docum system If a pas it. Age: D	<i>ication system</i> nent as define n (such as ear ssport accomp Date of birth (d	 The animal must bear and d in Article 2(b) of Committing, tattoo, brand, transpondenties the animal, its number d/mm/yyyy). 	(Signature) us asinus, or indicate any cro n individual identifier which p ssion Implementing Regulati nder) and the anatomic place	ess between those. ermits to link the animal on (EU) 2018/659. Spec used on the animal.	ify the identification
Identif docum systen If a pas it. Age: E	<i>ication system</i> nent as define n (such as ear ssport accomp Date of birth (d	b: The animal must bear and d in Article 2(b) of Committing, tattoo, brand, transport anies the animal, its number	(Signature) us asinus, or indicate any cro n individual identifier which p ssion Implementing Regulati nder) and the anatomic place	ess between those. ermits to link the animal on (EU) 2018/659. Spec used on the animal.	ify the identification

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.	(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
	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	(h)	on the top of the pages. The original of the health certificate shall be completed and
	country or part of the territory of the country.		signed by an official veterinarian within 24 hours prior loading
(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.		the consignment, or in the case of registered horses on the last working day prior to loading, for exportation to the Union. The competent authorities of the exporting country shall ensure that the principles of certification equivalent to those laid down in Council Directive 96/93/EC (¹) are followed.
(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.		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.
(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.	(i)	The original of the health certificate shall accompany the consignment until it reaches the border inspection post of entry into the Union.
(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.	(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.
(f)	If for reasons of identification of the animals of the consignment (schedule in Box I.28 of the model health certificate), additional sheets of paper are attached to the health certificate, those sheets of paper shall also be considered as forming part of the original of the health certificate by application of the signature and stamp of the certifying official veterinarian, on each of the pages.		

▼<u>B</u>

(¹) 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

▼<u>C2</u>

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

COUN	TRY:								Veterinary certifi	cate to EU
	l.1.	Name				1.2.	Certificate reference	ce No	l.2.a.	
		Address				1.3.	Central competent	authority		
ent		Tel.				1.4.	Local competent a	uthority		
Part I : Details of dispatched consignment	1.5.	Consignee Name Address				1.6.	Person responsible Name Address	e for the load	in EU	
batched o		Postal code Tel.					Postal code Tel.			
s of disp	1.7.	Country of origin	ISO code	I.8. Region of origin	Code	1.9.	Country of destination	ISO code	I.10. Region of destination	Code
: Detail	1.11.	Place of orig Semen cent		I	1	I.12.	Place of destination Semen centre	n	Holding 🗖	
Part I		Name Address	Approval n	umber			Name Address		Approval number	
		Postal code					Postal code			
	I.13.	Place of load	ding			I.14.	Date of departure			
	l.15.	Means of tra	ansport			I.16.	Entry BIP in EU			
		Aeroplane C			agon 🗖					
		Road vehicle Identification Documentar		. —		l.17.				
	l.18.	Description	of commodity	/			1	I.19. Commo	dity code (HS code) 05 11 99 85	
									I.20. Quantity	
	I.21.								I.22. Number of pac	kages
	1.23.	Seal/Contair	ner No						1.24.	
	1.25.	Commoditie Artificial rep								
	1.26.	For transit th	nrough EU to	third country			I.27. For import or a	admission int	o EU 🔲	
		Third countr	y I	SO code						
	1.28.	Identification	n of the com	nodities						
	s	pecies (Scier	ntific name)	Donor i	dentity		Date of colle	ection	Quantity	

	COUNTRY					Equine semen – Section A
	II. Health	information			II.a. Certificate reference No	II.b.
	I, the unders	igned, offic	ial veterinar	ian, of th	e exporting country (²)	
					(name of exporting	j country)
	certify that:					
u	II.1.	export to	the Union is	s approv	(³), in which the semen described above was ed and supervised by the competent authority Annex D to Directive 92/65/EEC (⁴);	
Part II: Certification	11.2.	date the f		ed seme	g 30 days prior to the date of first collection of n was dispatched or until the 30 days storage	
Part II:		II.2.1.			ne exporting country or, in the case of regio S/EC (⁵), in that part of the territory of the export	
					ed to be infected with African horse sicknes rective 2009/156/EC,	s in accordance with Article 5(2)(a)
			— free	from Ver	nezuelan equine encephalomyelitis for a period	of at least 2 years,
	-		— free	from gla	nders and dourine for a period of at least 6 mor	iths;
		II.2.2.	fulfilled th	e conditi	ons for a holding laid down in Article 4(5) of Dir	ective 2009/156/EC and in particular:
	(1)) either	[11.2.2.1.		g a case of a disease mentioned below not all ease located in the holding were slaughtered or	
				b	rom any type of equine encephalomyelitis f eginning on the day on which the equida laughtered,	
				n o	rom equine infectious anaemia (EIA) for at le egative result in an agar gel immunodiffusion te n samples taken after the infected animals months apart from each of the remaining anim	est (AGID or Coggins test) carried out were slaughtered on two occasions
					rom vesicular stomatitis (VS) for a period of at ase,	east 6 months from the last recorded
				— fr	rom rabies for a period of at least one month fro	m the last recorded case,
				— fr	rom anthrax for a period of at least 15 days fron	the last recorded case,]
		(¹) or	[II.2.2.1.	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 least alomyelitis, equine infectious anaemia, vesicula e of anthrax, beginning on the day on which fo nfection of the premises was satisfactorily comp	or killed and the premises disinfected, 30 days from any type of equine ar stomatitis and rabies or 15 days in llowing the destruction of the animals
		II.2.3.	containec metritis,	l only equ	uidae which were free of clinical signs of equin	e viral arteritis and contagious equine
	II.3.	Prior to er	ntering the s	emen co	llection centre the donor stallions and any othe	r equidae located in the centre:

II. H	ealth informatio	n	1	I.a. Certificate reference No	II.b.					
	II.3.1.	a Membe regionalis	r State of t ation in ac	sident for a period of 3 months (or since e he Union during the 3 months period) in t cordance with Article 13 of Directive 2009 which was during that period:	he exporting country or, in the case c					
				to be infected with African horse sickne tive 2009/156/EC,	ess in accordance with Article 5(2)(a					
		— free	from Venez	zuelan equine encephalomyelitis for a peric	od of at least 2 years,					
		— free	from gland	ers and dourine for a period of at least 6 m	onths;					
(¹) eithe	er [II.3.2.			country of export which was on the day o /S) for a period of at least 6 months,]	of admission into the centre free fron					
(¹) or	[II.3.2.	result at a with the r	a serum dil elevant Ch	virus neutralisation test for vesicular stom ution of 1 in 32 or a VS ELISA carried ou apter of the Manual of Diagnostic Tests a ample taken (⁶) within 14 days prior to ente	ut with a negative result in accordance and Vaccines for Terrestrial Animals o					
	II.3.3.	originated point II.2.2		ngs which on the day of admission onto	the centre fulfilled the requirements c					
11.4.	The sem	The semen described above was collected from donor stallions which:								
	II.4.1.			ical sign of an infectious or contagious dis tre and on the day the semen was collecte						
	II.4.2.		imal has sl	od of at least 30 days prior to the date of a nown any clinical sign of equine viral arteri						
	II.4.3.	collection	and betwe	atural mating during a period of at least 3 en the dates of the first sample referred to ne collection period;						
	II. 4 .4.	Manual of which is r	Diagnostic	ving tests, which meet at least the require Tests and Vaccines for Terrestrial Animal by the competent authority and has the tes lent to that provided for in Article 12 of	s of the OIE, carried out in a laborator sts referred to hereinafter included in it					
				infectious anaemia (EIA), an agar-gel im enzyme-linked immunosorbent assay (ELI result;]						
		II.4.4.2.	for equine	viral arteritis (EVA),						
		(¹) either	[II.4.4.2.1.	a serum neutralisation test with a nega in four;]	ative result at a serum dilution of on					
		(¹) and/or	[11.4.4.2.2.	a virus isolation test, polymerase chain r negative result on an aliquot of the entire						
		II.4.4.3.	three spec	gious equine metritis (CEM), an ager imens (swabs) taken from the donor stallic an 7 days at least from the penile sheath	on on two occasions with an interval o					

COUNTRY

Equine semen – Section A

II.	Health information	n	11	.a. Certificate reference No	II.b.
			(local treat transport n	les were in no case taken earlier than 7 days (systement) after antimicrobial treatment of the donor s nedium with activated charcoal, such as Amies med where they were subjected with a negative result to a	tallion and were placed in lium, before dispatch to the
		(¹) either	[11.4.4.3.1.	the isolation of <i>Taylorella equigenitalis</i> after cultiv conditions for a period of at least 7 days, set up with specimens from the donor animal, or 48 hours who cool during transport;]	hin 24 hours after taking the
		(¹) and/or	[11.4.4.3.2.	the detection of genome of <i>Taylorella equigenitalis</i> carried out within 48 hours after taking the specimer	
	II.4.5.	programm		the results specified in point II.4.4 in each case I respectively in points 1.6(a), (b) and (c) of Chapte ::	
		(⁹) [II.4.5.1.	at least 30 the semen	stallion was continuously resident on the semen coll days prior to the date of the first collection and durin described above, and no equidae on the semen co to direct contact with equidae of lower health status t	ng the period of collection o Ilection centre came durin
			stallion at collection of and not less	described in point II.4.4 were carried out on sample least once a year at the beginning of the breeding of semen intended for imports into the Union of free so than 14 days following the date of the commencen 30 days prior to the first semen collection.]	season or prior to the firs
		(⁹) [II.4.5.2.	30 days pr semen des centre vete	stallion was resident on the semen collection cen rior to the date of the first collection and during the scribed above, but left the semen collection centre ur erinarian for a continuous period of less than 14 day collection centre came into direct contact with equida	e period of collection of the oder the responsibility of the rs, and/or other equidae o
			stallion at the first co semen and	described in point II.4.4 were carried out on sample least once a year at the beginning of the breeding se illection of semen intended for imports into the Unio d not less than 14 days following the date of the comr t least 30 days prior to the first semen collection,	eason or prior to the date on of fresh, chilled or froze
		and	chilled or	period of collection of the semen intended for impo frozen semen the donor stallion was subjected , as follows:	
			(a)	for equine infectious anaemia, one of the tests det last carried out on a sample of blood taken (⁶) not the collection of the semen described above;	
			(b)	for equine viral arteritis, one of the tests described	
			(¹) either	[in point II.4.4.2 was last carried out on a samp 30 days prior to the date of the collection of the sem	
			(¹) or	[in point II.4.4.2.2 was carried out on an aliquot donor stallion taken (⁶) not more than 6 month- collection of the semen described above and a bloo 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 th od sample taken (⁶) from th I with a positive result in

_	Health in	nformatior	۱ 		II.a. (Certificate re	ference No			II.b.	
				(c)	car	ried out on tl	nree specim		taken (⁶) not i	in point II.4.4 more than 60	
				(¹) either	· [on	two occasio	ns;]				
				(¹) or	[on	a single occ	asion and s	ubjected to a	PCR or real-	time PCR.]]	
			(⁹) [II.4.5.3.		c D to D					l.6(a) and (b) r imports into	
					e tests described in points II.4.4.1, II.4.4.2 and II.4.4.3 were carried out on samples ten $(^6)$ from the donor stallion at least once a year at the beginning of the breeding ason,						
			and	eriod of the s semen and b	emen of a m efore the se	on samples ta inimum perioo men is remov e than 90 da	d of 30 day /ed from th				
			and (¹) eiti		san of 3 rem not	nples taken 30 days from 10ved from t	(⁶) during th the date of he semen c	e storage per the collection collection cen	riod of the se n of the seme tre or used, r	.4.4.2 were ca men of a mini n and before t not less than n of the seme	imum perio the semen 14 days ar
				(¹) or			er state of a	a donor stallio	on seropositiv	ve for equine	viral arterit
					neg stal stal	ative result llion taken (^e llion has rea	on sample) twice a ye cted with a	s of an aliquer at an inter	CR or real-tin uot of the er val of at leas It at a serum	ne PCR carrie ntire semen c st 4 months ar dilution of at ritis.]	ed out with of the don nd the don
		II.4.6.	underwen dates:	t the test	neg stal stal four	pative result llion taken (⁶ llion has rea r in a serum	on sample) twice a ye cted with a neutralisatio	s of an alique ar at an inter positive resu on test for equ	CR or real-tin uot of the er val of at leas It at a serum uine viral arte	ntire semen c at 4 months ar dilution of at	ed out with of the don nd the don least one
			dates:	t the test	neg stal stal four	pative result llion taken (⁶ llion has rea r in a serum	on sample) twice a ye cted with a neutralisation points II.3.	s of an alique ar at an inter positive resu on test for equ	CR or real-tin uot of the er val of at leas It at a serum line viral arte	ntire semen c at 4 months ar dilution of at ritis.] es taken on t	ed out with of the don nd the don least one
			dates:	rt date (°)	neg stal stal four	pative result llion taken (⁶ llion has rea r in a serum	on sample) twice a ye cted with a neutralisation points II.3.	s of an aliqu ar at an inter positive resu on test for equ 2 (¹) and II.4 Date of samplin	CR or real-tin uot of the er val of at leas It at a serum line viral arte	tire semen c at 4 months ar dilution of at ritis.] es taken on t ts (⁶)	ed out with of the dong nd the dong least one
	Identification of semen	II.4.6. Lest broßramme	dates: Sta	t date (⁶) Ser	neg stal stal fou	pative result lion taken (⁶ lion has rea r in a serum	on sample) twice a yected with a neutralisation points II.3.	s of an aliqu ar at an inter positive resu on test for equ 2 (¹) and II.4 Date of samplin	CR or real-tin uot of the er val of at leas lt at a serum ine viral arte l.5 on sampl g for health tes A II.	tire semen c at 4 months ar dilution of at ritis.] es taken on t ts (⁶)	ed out with of the dono least one the followin
			dates: Sta	t date (⁶) Ser	neg stal stal fou ting pro	pative result lion taken (⁶ lion has rea r in a serum povided for in VS (¹)	on sample) twice a yected with a neutralisation points II.3.	s of an aliqu ar at an inter positive resu on test for equ 2 (¹) and II.4 Date of samplin EV. 4.4 Blood	CR or real-tin uot of the er val of at leas It at a serum uine viral arte I.5 on sampl g for health tes A II. I.2. Semen	tire semen c t 4 months ar dilution of at ritis.] es taken on t ts (⁶) CE II.4. 1.	ed out with of the dominant the dominant the dominant least one the followin EM .4.3.
			dates: Sta	t date (⁶) Ser	neg stal stal fou ting pro	pative result lion taken (⁶ lion has rea r in a serum povided for in VS (¹)	on sample) twice a yected with a neutralisation points II.3.	s of an aliqu ar at an inter positive resu on test for equ 2 (¹) and II.4 Date of samplin EV. 4.4 Blood	CR or real-tin uot of the er val of at leas It at a serum uine viral arte I.5 on sampl g for health tes A II. I.2. Semen	tire semen c t 4 months ar dilution of at ritis.] es taken on t ts (⁶) CE II.4. 1.	ed out with of the dono least one the followin EM .4.3.
			dates: Sta	t date (⁶) Ser	neg stal stal fou ting pro	pative result lion taken (⁶ lion has rea r in a serum povided for in VS (¹)	on sample) twice a yected with a neutralisation points II.3.	s of an aliqu ar at an inter positive resu on test for equ 2 (¹) and II.4 Date of samplin EV. 4.4 Blood	CR or real-tin uot of the er val of at leas It at a serum uine viral arte I.5 on sampl g for health tes A II. I.2. Semen	tire semen c t 4 months ar dilution of at ritis.] es taken on t ts (⁶) CE II.4. 1.	ed out with of the dono least one i the followin EM .4.3.
			dates: Sta	t date (⁶) Ser	neg stal stal fou ting pro	pative result lion taken (⁶ lion has rea r in a serum povided for in VS (¹)	on sample) twice a yected with a neutralisation points II.3.	s of an aliqu ar at an inter positive resu on test for equ 2 (¹) and II.4 Date of samplin EV. 4.4 Blood	CR or real-tin uot of the er val of at leas It at a serum uine viral arte I.5 on sampl g for health tes A II. I.2. Semen	tire semen c t 4 months ar dilution of at ritis.] es taken on t ts (⁶) CE II.4. 1.	ed out with of the dominant the dominant the dominant least one the followin EM .4.3.
			dates: Sta	t date (⁶) Ser	neg stal stal fou ting pro	pative result lion taken (⁶ lion has rea r in a serum povided for in VS (¹)	on sample) twice a yected with a neutralisation points II.3.	s of an aliqu ar at an inter positive resu on test for equ 2 (¹) and II.4 Date of samplin EV. 4.4 Blood	CR or real-tin uot of the er val of at leas It at a serum uine viral arte I.5 on sampl g for health tes A II. I.2. Semen	tire semen c t 4 months ar dilution of at ritis.] es taken on t ts (⁶) CE II.4. 1.	ed out with of the dono least one the followin EM .4.3.

_

II. Hea	Ith information		II.a. Certificate reference No	II.b.
				II.U.
(¹) either	[11.5.	No antibiotics we	re added to the semen;]	
(¹) or	[II.5.		ibiotic or combination of antibiotics was adde not less than (¹⁰):	d to produce a concentration in the fina
II.6.	The seme	en described above	was:	
	II.6.1.		sed, stored and transported under conditions and III(I) of Annex D to Directive 92/65/EEC;	s which comply with the requirements o
	II.6.2.		e of loading in a sealed container in accord tive 92/65/EEC and bearing the number indic	
Notes				
Part I:				
Box I.11.:	The place	of origin shall corre	espond to the semen collection centre of the s	semen origin.
Box I.22.:	The numb	per of packages sha	Il correspond to the number of containers.	
Box I.23.:	The identi	fication of containe	r and seal number shall be indicated.	
Box I.28.:	The dono	r identity shall corre	spond to the official identification of the anima	al.
	The date	of collection shall b	e indicated in the following format: dd/mm/yyy	/у.
Part II:				
Guidance	for the compl	etion of the table in	point II.4.6.	
Abbreviat	ions:			
VS	Vesic	ular stomatitis (VS)	testing if required in accordance with point II.	3.2
EIA-	1 Equin	e infectious anaem	a (EIA) testing first occasion	
EIA-2	2 EIA te	esting second occas	ion	
EVA-	-B1 Equin	e viral arteritis (EVA	A) testing on blood sample first occasion	
EVA-	-B2 EVA t	esting on blood sar	nple second occasion	
EVA-	-S1 EVA t	esting on semen sa	mple first occasion	
EVA-	-S2 EVA t	esting on semen sa	mple second occasion	
CEM	-11 Conta	igious equine metrit	is (CEM) testing first occasion first sample	
CEM	-12 CEM	testing first occasio	n second sample taken 7 days after CEM-11	
CEM	-21 CEM	testing second occa	asion first sample	
CEM	-22 CEM	testing second occa	asion second sample taken 7 days after CEM-	-21
Instructio	ns:			
For (each semen i	dentified in columr	A in correspondence with Box I.28, the to	est programme (points II.4.5.1, II.4.5

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.

Equine semen – Section A

COUNTRY

			-4
II.	Health information	II.a. Certificate reference No	II.b.

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

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

of				Date of sampling for health tests					
Identification semen	Test programme	Donor	Semen	VS	EIA	EVA II.4.4.2.		CEM II.4.4.3.	
Identi	bro	residence	collection	II.3.2.			Semen sample	1. sample	2. sample
۸	в	с	D	vs	EIA-1	EVA-B1	EVA-S1	CEM-11	CEM-12
A				v3	EIA-2	EVA-B2	EVA-S2	CEM-21	CEM-22

(¹) Delete as necessary.

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

(³) Only approved semen collection centres listed in accordance with Article 17(3)(b) of Directive 92/65/EEC on the Commission website: http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm

- (4) Council Directive 92/65/EEC of 13 July 1992 laying down animal health requirements governing trade in and imports into the Community of animals, semen, ova and embryos not subject to animal health requirements laid down in specific Community rules referred to in Annex A(I) to Directive 90/425/EEC (OJ L 268, 14.9.1992, p. 54).
- (⁵) Council Directive 2009/156/EC of 30 November 2009 on animal health conditions governing the movement and importation from third countries of equidae (OJ L 192, 23.7.2010, p. 1).
- (6) Insert date in table in point II.4.6 (follow Guidance in Part II of the Notes).
- (⁷) Regulation (EC) No 882/2004 of the European Parliament and of the Council of 29 April 2004 on official controls performed to ensure the verification of compliance with feed and food law, animal health and animal welfare rules (OJ L 165, 30.4.2004, p. 1).
- (⁸) The agar gel immunodiffusion test (AGID or Coggins test) or the ELISA for equine infectious anaemia are not required for donor equidae which have continuously resided in Iceland since birth, provided that Iceland has remained officially free of equine infectious anaemia and no equidae and their semen, ova and embryos have been introduced into Iceland from outside prior to and during the period the semen was collected.
- (⁹) Cross out the programmes that do not apply to the consignment.
- (¹⁰) Insert names and concentrations.
- The signature and the stamp must be in a different colour to that of the printing.

Official veterinarian

Name (in capital letters):

Date:

Stamp:

Qualification and title:

Signature:

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

ITRY:					Veterinary certifie	cate to E	
l.1.	Consignor Name	1.2.	Certificate reference	ce No	l.2.a.		
	Address	1.3.	I.3. Central competent authority				
	Tel.	1.4.	Local competent a	uthority			
1.5.	Consignee Name Address	1.6.	Person responsible Name Address	e for the load	in EU		
	Postal code Tel.		Postal code Tel.				
1.7.	Country of ISO code I.8. Region of Code origin	1.9.	Country of destination	ISO code	I.10. Region of destination	Code	
l.11.	Place of origin Semen centre	I.12.	Place of destinatio Semen centre 🗖		Holding 🗖		
	Name Approval number Address		Name Address	Approv	al number		
	Postal code		Postal code				
l.13.	Place of loading	I.14.	Date of departure				
l.15.	Means of transport	I.16.	Entry BIP in EU				
	Aeroplane Ship Railway wagon Road vehicle Other I Identification Documentary references	l.17.					
l.18.	Description of commodity			I.19. Commo	dity code (HS code) 05 11 99 85		
			L		I.20. Quantity		
I.21.					I.22. Number of pac	kages	
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 int	o EU 🛛		
	Third country ISO code						
128	Identification of the commodities						
1.20.					Quantity		

	COUNTRY				Equine semen – Section B
	II. Health	information		II.a. Certificate reference No	II.b.
	I, the unders	igned, officia	al veterinar	ian, of the exporting country (²)(name of exporting co	
	certify that :				
ų	II.1.	export to t	the Europe	n centre (³), in which the semen described above was collecte ean Union is approved and supervised by the competent author $I(I)(1)$ and Chapter $I(II)(1)$ of Annex D to Directive 92/65/EEC,	
Part II: Certification	II.2.			nmencing 30 days prior to the date of first collection of the sem od for frozen semen elapsed, the semen collection centre:	en described above until the
Part II: C		II.2.1.		ated in the exporting country or, in the case of regionalisation 2009/156/EC ($^{\circ}$), in that part of the territory of the exporting coun	
				considered to be infected with African horse sickness in acc (b) of Directive 2009/156/EC (8),	ordance with Article 5(2)(a)
			— free	from Venezuelan equine encephalomyelitis for 2 years,	
	-		— free	from glanders and dourine for 6 months;	
		II.2.2.	fulfilled tl particular	ne conditions for a holding laid down in Article 4(5) of Direct	ive 2009/156/EC (⁸) and in
		(¹) either	[11.2.2.1.	following a case of a disease mentioned below not all the anim the disease located on the holding were slaughtered or killed free:	
				 from any type of equine encephalomyelitis for at least day on which the equidae suffering from the disease are 	
				 from equine infectious anaemia for at least the period r result in an agar gel immunodiffusion test (Coggins te taken after the infected animals were slaughtered on tw from each of the remaining animals, 	est) carried out on samples
				 from vesicular stomatitis for at least 6 months from the la 	st 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 of 	ase,]
		(¹) or	[II.2.2.1.	following a case of a disease mentioned below all the animals disease located on the holding have been slaughtered of disinfected, the holding has been free for at least 30 days encephalomyelitis, equine infectious anaemia, vesicular stomat the case of anthrax, beginning on the day on which following the the disinfection of the premises was satisfactorily completed;]	r killed and the premises s from any type of equine titis and rabies or 15 days in
		II.2.3.	containeo metritis,	d only equidae which were free of clinical signs of equine viral an	teritis and contagious equine
	II.3.	Prior to en	tering the s	semen collection centre the donor stallions and any other equidat	e located in the centre:

II. F	lealth information		II.a. Certificate reference No	II.b.
	II.3.1.	State of the E regionalisation	uropean Union during the 3 months perio	if they were directly imported from a Membe d) in the exporting country or, in the case of /156/EC (⁸), in that part of the territory of the
			sidered to be infected with African horse of Directive 2009/156/EC (°),	sickness in accordance with Article 5(2)(a
		— free from	n Venezuelan equine encephalomyelitis for	at least 2 years,
		— free from	n glanders and dourine for at least 6 months	5;
(1) eith	ner [II.3.2.		m the country of export which was on th natitis (VS) for at least 6 months,]	ne day of admission into the centre free c
(1) or	[11.3.2.			lar stomatitis (VS) carried out with negative taken $(^4)$ within 14 days prior to entering the
	II.3.3.	originated from point II.2.2;	n holdings which on the day of admission	n onto the centre fulfilled the requirements o
II. 4 .	The seme	n described abo	we was collected from donor stallions, whic	sh:
	II.4.1.		vn any clinical sign of an infectious or cont d on the day the semen was collected;	tagious disease at the time of admission ont
	II.4.2.		pt for 30 days prior to the date of semen or y clinical sign of equine viral arteritis or con	ollection on holdings where no equine anima tagious equine metritis during that period;
	II.4.3.	and between		days prior to the date of first semen collectio points II.4.5.1, II.4.5.2 and/or II.4.5.3 and unt
	II.4.4.	the Manual c samples take	f Diagnostic Tests and Vaccines for Ter	t the requirements of the relevant Chapter or rrestrial Animals of the OIE, carried out o imes specified in point II.4.5 in a laborator
	(¹) (⁵) <i>either</i>	-	agar-gel immuno-diffusion test (Coggins te ative result;]	est) for equine infectious anaemia (EIA) wit
	(¹) (⁵) or	[II.4.4.1. an	ELISA for equine infectious anaemia (EIA)	with negative result;]
and	(¹) either	•	erum neutralisation test for equine viral ar tion of one in four;]	rteritis (EVA) with negative result at a serur
	(¹) or		irus isolation test for equine viral arteritis uot of the entire semen of the donor stallior	(EVA) carried out with negative result on a

COUNTRY

II. Health information

II.a.

	Equine semen – Section B
Certificate reference No	II.b.
entification test for contagious equine metritis ((CEM) carried out on two

and	II.4.4.3.	an agent identification test for contagious equine metritis (CEM) carried out on two occasions on samples collected with an interval of 7 days by isolation of <i>Taylorella equigenitalis</i> after a cultivation of 7 to 14 days from pre-ejaculatory fluid or a semen sample and from genital swabs taken at least from the penile sheath, urethra and urethral fossa with negative result in each case;
II.4.5.		en subjected with the results specified in II.4.4. in each case to at least one of the test $(^6)$ detailed in points II.4.5.1, II.4.5.2 and II.4.5.3 as follows:
	II.4.5.1.	The donor stallion was continuously resident on the semen collection centre for at least 30 days prior to the date of the first collection and during the period of collection of the semen described above, and no equidae on the semen collection centre came during that time into direct contact with equidae of lower health status than the donor stallion.
		The tests described in point II.4.4 have been carried out on samples taken (⁴) prior to the first semen collection and at least 14 days following the date of the commencement of the residence period of at least 30 days.
	II.4.5.2.	The donor stallion was resident on the semen collection centre for at least 30 days prior to the date of the first collection and during the period of collection of the semen described above, but has left the centre under the responsibility of the centre veterinarian for a continuous period of less than 14 days, or other equidae on the collection centre came into direct contact with equidae of lower health status.
		The tests described in point II.4.4 have been carried out on samples taken (⁴) prior to the date of the first semen collection of the breeding season or collection period in the year the semen described above was collected and at least 14 days following the date of the commencement of the residence period of at least 30 days,
and		the test described in point II.4.4.1 for equine infectious anaemia was last carried out on a sample of blood taken (⁴) not more than 90 days before the semen described above was collected;
and	(¹) either	[one of the tests described in point II.4.4.2 for equine viral arteritis was last carried out on a sample taken (4) not more than 30 days before the semen described above was collected,]
	(¹) or	[a virus isolation test for equine viral arteritis was carried out with negative result on an aliquot of the entire semen of the donor stallion taken (⁴) not more than 6 months before the semen described above was collected and a blood sample taken on the same date (⁴) reacted positive in a serum neutralisation test for equine viral arteritis at a serum dilution of more than one in four,]
and		the test described in point II.4.4.3 for contagious equine metritis was last carried out on samples taken (⁴), not more than 60 days before the semen described above was collected.
	II.4.5.3.	The tests described in point II.4.4 have been carried out on samples taken (⁴) prior to the date of the first semen collection of the breeding season or collection period in the year the semen described above was collected,
and		the tests described in point II.4.4 have been carried out on samples taken (⁴) between 14 and 90 days after the collection of the semen described above.

COUNTRY

Equine semen – Section B

II. Health	n information	1	II.a. (Certificate r	eference No			II.b.		
	II.4.6.	have under following da		ing provide	d for in poi	nts II.3.2 (¹)	and II.4.5	on samples ta	aken on th	
of	a	Start o	date (4)		[Date of samplin	g for health te	ests (4)		
Identification of semen	Test programme	Donor	Semen	VS (1)	EIA	EVA II. 4.4.2.			CEM II.4.4.3.	
	brog	residence	collection	11.3.2	II.4.4.1.	Blood sample	Semen sample	1. sample	2. sample	
(¹) either	[11.5.	No antibiotic	cs were added	to the seme	en;]					
(1) or	[11.5.	The followin	ig antibiotic or	combinatio	n of antibiotio	s was added	to produce	a concentratio	n in the fin	
		diluted com	en of not less t	han (7).						
		ulluled serie		nan ().						
11.6.	The seme	en described a							;	
II.6.	The seme	en described al collected, pr	bove was:	ed and tran	sported unde	er conditions		ly with the requ		
II.6.		en described al collected, pr Chapters II(sent to the	bove was: rocessed, store I)(1) and III(I) c	ed and tran of Annex D ng in a se	sported unde to Directive S aled contain	er conditions 92/65/EEC; er in accorda	which comp nce with po	ly with the requ	uirements	
	II.6.1.	en described al collected, pr Chapters II(sent to the	bove was: rocessed, store I)(1) and III(I) c place of loadi	ed and tran of Annex D ng in a se	sported unde to Directive S aled contain	er conditions 92/65/EEC; er in accorda	which comp nce with po	ly with the requ	uirements	
Notes	II.6.1.	en described al collected, pr Chapters II(sent to the	bove was: rocessed, store I)(1) and III(I) c place of loadi	ed and tran of Annex D ng in a se	sported unde to Directive S aled contain	er conditions 92/65/EEC; er in accorda	which comp nce with po	ly with the requ	uirements	
II.6. Notes Part I: Box I.11.:	II.6.1. II.6.2.	en described al collected, pr Chapters II(sent to the Annex D to	bove was: rocessed, store I)(1) and III(I) c place of loadi	ed and tran of Annex D ng in a sea J/EEC and I	sported unde to Directive S aled contain bearing the n	er conditions 92/65/EEC; er in accorda umber indica	which comp nce with po ted in Box I.	ly with the requ bint 1.4 of Cha 23.	uirements	
Notes Part I:	II.6.1. II.6.2. The place	en described al collected, pr Chapters II(sent to the Annex D to	bove was: rocessed, store I)(1) and III(I) c place of loadi Directive 92/65	ed and tran of Annex D ng in a se J/EEC and I the semen	sported under to Directive S aled contain- bearing the n	er conditions 12/65/EEC; er in accorda umber indica	which comp nce with po ted in Box I.	ly with the requ bint 1.4 of Cha 23.	uirements	
Notes Part I: Box I.11.:	II.6.1. II.6.2. The place	en described al collected, pr Chapters II(sent to the Annex D to e of origin shall per of package	bove was: rocessed, store I)(1) and III(I) c place of loadi Directive 92/65	ed and tran of Annex D ng in a se JEEC and I the semen wond to the r	sported under to Directive S aled contain- bearing the n collection ce number of cc	er conditions i2/65/EEC; er in accorda umber indica entre of the se intainers.	which comp nce with po ted in Box I.	ly with the requ bint 1.4 of Cha 23.	uirements	
Notes Part I: Box I.11.: Box I.22.:	II.6.1. II.6.2. The place The numb	en described al collected, pr Chapters II(sent to the Annex D to e of origin shall per of package ification of con	bove was: rocessed, store I)(1) and III(I) c place of loadi Directive 92/65	ed and tran of Annex D ng in a se b/EEC and I the semen bond to the i	sported under to Directive S aled contain bearing the n collection ce number of cc nall be indica	er conditions i2/65/EEC; er in accorda umber indica entre of the se ntainers. ted.	which comp ince with po ted in Box I.	ly with the requ bint 1.4 of Cha 23.	uirements	

	ricalar in	formation		II.a. (Sertificate r	eference No			II.b.				
Part	II:			I									
Guid	lance for t	he comple	etion of the tab	ble in point II.4	.6.								
Abbr	reviations:												
	VS	Vesicu	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 te	sting second o	occasion									
	EVA-B1	Equin	e viral arteritis	(EVA) testing	on blood sa	ample first o	ccasion						
	EVA-B2	EVA te	esting on bloo	d sample seco	nd occasio	n							
	EVA-S1	EVA te	esting on sem	en sample first	occasion								
	EVA-S2		esting on sem			on							
	CEM-11		gious equine r	·			st sample						
	CEM-12		esting first oc	. ,	-								
	CEM-21		testing second		·								
	CEM-22		testing second		·	taken 7 dav	s after CEM-1	21					
		CLINI	esting second	000031011 3600	ond sample	aken / day		- 1					
	uctions:												
			identified in st be specified										
	The dates	s when sa	amples were t	aken for labor	atory testir	na prior to th	ne first collec	tion of the se	men describe	ed above a			
	required in	n II.4.5.1.	, II.4.5.2. and EVA-B1 or E\	II.4.5.3., are e	ntered in th	ie upper line	of columns 5	to 9 of the ta					
	entered ir	n the low	amples were ta er line of col- example below	umns 5 to 9									
		22 11 110											
	on of	ле	Start	date				ing for health te					
	Identification of semen	Test programme	Donor	Semen	VS	EIA		/A 4.2.		EM .4.3.			
	s ut	2	residence	collection	II.3.2.	II.4.4.1.	Blood	Semen	1.	2.			
	Ide	-					sample	sample	sample	sample			

(¹) Delete as necessary.

Imports of equine semen are authorised from a third country listed in column 2 of Annex I to Commission Implementing Regulation (EU) 2018/657 provided 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. (²)

EIA-2

EVA-B2

EVA-S2

CEM-21

CEM-22

COUNTRY

COU	NTRY		Equine semen – Section E				
II.	Health information	II.a. Certificate reference No	II.b.				
(³)		es listed in accordance with Article 17(3) eu/food/animal/semen_ova/equine/index_	(b) of Council Directive 92/65/EEC on the _en.htm				
(4)	Insert date in table in point II.4.6 (follow Guidance in Part II of the Notes)						
(5)	The agar gel immunodiffusion test (Coggins test) or the ELISA for equine infectious anaemia are not required for donor equidae which have continuously resided in Iceland since birth, provided that Iceland has remained officially free of equine infectious anaemia and no equidae and their semen, ova and embryos have been introduced into Iceland from outside prior to and during the period the semen was collected.						
(6)	Cross out the programmes that do not	apply to the consignment.					
(7)	Insert names and concentrations.						
(8)	OJ L 192, 23.7.2010, p. 1.						
_	The signature and the stamp must be i	n a different colour to that of the printing.					
Offi	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

UNT	TRY:				Veterinary certificate to EL			
	l.1.	Consignor Name	1.2.	Certificate reference No	l.2.a.			
		Address	1.3.	I.3. Central competent authority				
		Tel.	1.4.	Local competent authority				
Part I : Details of dispatched consignment	1.5.	Consignee Name Address	1.6.	Person responsible for the loa Name Address	id in EU			
tched co		Postal code Tel.		Postal code Tel.				
of dispa	1.7.	Country of ISO code I.8. Region of Code origin origin	9 1.9.	Country of ISO code destination	I.10. Region of Code destination			
I : Details	l.11.	Place of origin Semen centre 🗖	1.12	Place of destination Semen centre ☐ H	olding 🗖			
Part		Name Approval number Address		Name Approva Address	al number			
		Postal code		Postal code				
	I.13.	Place of loading	1.14	. Date of departure				
	l.15.	Means of transport	1.16	6. Entry BIP in EU				
		Aeroplane Abip Railway wagon Acou Vehicle Other Identification	1.17	·				
		Documentary references						
	l.18.	Description of commodity		I.19. Comm	nodity 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 in	nto EU 🔲			
		Third country ISO code						
		Identification of the commodities pecies (Scientific name) Donor identity		Date of collection	Quantity			

	COUNTRY			Equine semen – Section C			
	II. Health	information	II.a. Certificate reference No	II.b.			
	I, the unders	igned, official veterinarian, of the ex	xporting country (²) (name of exporting co				
	certify that:						
Part II: Certification	II.1.	The semen collection centre in which the semen described above was collected, processed and stored for export to the European Union:					
	II.1.1.	is approved and supervised by the competent authority according to the conditions of Chapter I, Annex D to Directive 92/65/EEC,					
	II.1.2.	is situated in the territory or in the case of regionalisation according to Article 13 of Directive 2009/156/EC (⁶) in a part of the territory of the country of export which was on the day the semen was collected until the date of dispatch free of:					
		 African horse sickness, in accordance with EU legislation, 					
		 Venezuelan equine encepha 	alomyelitis for 2 years,				
		— glanders and dourine for 6 months;					
	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.	 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: 6 months, beginning on the day on which the equidae suffering from the disease are slaughtered, in the case of equine encephalomyelitis, 					
		 a period required to carry out with negative result two Coggins tests 3 months apart in the anima remaining after the infected animals have been slaughtered, in the case of infectious equine anaemia, 					
		 6 months, in the case of vesicular stomatitis, 					
		— one month from the last reco	orded case, in the case of rabies,				
		 — 15 days from the last record 	led case, in the case of anthrax.				
	II.1.3.2.	if all the animals of species susceptible to the disease located in the holding have been slaughtered or killed an the premises disinfected, the prohibition lasted for 30 days, or 15 days in the case of anthrax, beginning on th day on which following the destruction of the animals the disinfection of the premises was satisfactorily completed					
	II.1.4.	contained during the period commencing 30 days prior to semen collection and lasting until the date of its dispatch only equidae which were free of clinical signs of equine viral arteritis and contagious equine metritis,					
	II.2.	Prior to entering the semen collection centre the donor stallions and any other equidae located in the centre:					
	II.2.1.	were continuously resident for 3 months (or since entry if they were directly imported from a Member State of the Union during the 3 months period) in the territory or in the case of regionalisation in a part of the territory(1) of the country of export which was during that period free of:					
		 African horse sickness, in ad 	ccordance with EU legislation,				
		 Venezuelan equine encepha 	alomyelitis for 2 years,				
		— glanders for 6 months,					

				Equine semen – Section C
II. Health inf	ormation		II.a. Certificate reference No	II.b.
(¹) either [I	1.2.2.	originated from the ten free of vesicular stoma	itory of the country of export which was on the datitis for 6 months,]	ay of admission into the centre
(¹) or [l	1.2.2.		irus neutralisation test for vesicular stomatiti (⁴), this being within 14 days prior to entering the 12;]	
II.2.3. o	riginated f	rom holdings which on	the day of admission onto the centre fulfilled the r	requirements of point II.1.3;
II.3. Т	he semen	described above was o	collected from donor stallions, which:	
II.3.1. o	n the day	the semen was collecte	d have not shown clinical signs of an infectious o	r contagious disease,
II.3.2. d	luring at le	ast 30 days prior to coll	ection of the semen have not been used for natur	ral service,
		last 30 days prior to c nical signs of equine vira	collection of the semen have been kept on hold al arteritis,	lings where no equine animal
		last 60 days prior to c nical signs of contagious	collection of the semen have been kept on hold s equine metritis,	lings where no equine animal
			as far as I could ascertain have not been in conta e the 15 days immediately preceding the collectio	
			nimal health tests carried out in a laboratory in programme as specified in point II.3.7:	recognised by the competent
II.3.6.1. a	in agar-ge	I immuno-diffusion test	(Coggins test) for equine infectious anaemia with	negative result (³);
(¹) either [I	1.3.6.2.	a serum neutralisation	test for equine viral arteritis with negative result a	at a serum dilution of 1 in 4;]
(¹) or [l	1.3.6.2.	a virus isolation test fo semen;]	r equine viral arteritis carried out with negative re	esult on an aliquot of the entire
Т	aylorella e	equigenitalis from pre-e	ritis carried out on two occasions with an inter jaculatory fluid or a semen sample and from ger n the urethral fossa with negative result in each ca	nital swabs taken at least from
ll.3.7. h	ave been	subjected to one of the	following test programmes (⁵):	
с	ollection,	and during the collection	sly resident on the collection centre for at leas on period, and no equidae on the collection cen r health status than the donor stallions.	
a		days after the comme	eve been carried out on samples taken on encement of the above residence period and at	
			ously resident on the collection centre or other ea ae of lower health status than the donor stallions.	
			ive been carried out on samples taken on e first semen collection and at least at the beginn	
		quired in point II.3.6.1 was collected on	vas last carried out on a sample of blood taken r 	not more than 120 days before
(¹) either [equired in point II.3.6.2 	was last carried out not more than 30 days befor	re the semen was collected on

COUNTRY

	ITRY			Equine semen – Section		
II.	Health	information	II.a. Certificate reference No	II.b.		
(1) 0	r		eropositive stallion for equine viral arteriti than one year before the semen was colle			
II.3.7	7.3.		6 have been carried out during the 30 d rs after the collection of the semen on sam ;			
11.4.			vas collected, processed, stored and tran r II and III of Annex D to Directive 92/65/E			
Note	es					
Part	:1:					
Box	l.11.:	The place of origin shall corresp	oond to the semen collection centre of the	semen origin.		
Box	1.22.:	The number of packages shall o	correspond to the number of containers.			
Вох	1.23.:	The identification of container a	nd seal number shall be indicated.			
Вох	l.28.:	The donor identity shall corresp	ond to the official identification of the anim	al.		
		The date of collection shall be in	ndicate in the following format: dd/mm/yyyy	y.		
Part	: 11:					
(¹)	Delete	as necessary.				
(²)	Regula	orts of equine semen are authorised from a third country listed in column 2 of Annex I to Commission Implementing julation (EU) 2018/659 provided the semen was collected in the part of the territory of the third country detailed in umn 4 of that Annex from a donor stallion of the category of equidae indicated in columns 11, 12 or 13 of that Annex.				
(³)	equida equine	e agar gel immunodiffusion test (Coggins test) or the ELISA for equine infectious anaemia are not required for donor uidae which have continuously resided in Iceland since birth, provided that Iceland has remained officially free of uine infectious anaemia and no equidae and their semen, ova and embryos have been introduced into Iceland from tside prior to and during the period the semen was collected.				
(4)	Insert	date.				
(⁵)	Cross	out the programmes that do not a	apply to the consignment.			
(⁶)	OJ L 1	92, 23.7.2010, p. 1.				
_	The sig	gnature and the stamp must be ir	a different colour to that of the printing.			
Offic	cial veter	inarian				
	Name	(in capital letters):		Qualification and title:		
	Date:			Signature:		

Section D

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

COUNTRY:

Veterinary certificate to EU

	I.1.	Consignor Name		I.2. Certificate reference No I.2.a.				
		Address	1.3.	I.3. Central competent authority				
		Tel.	1.4.	Local competent au	thority			
Part I : Details of dispatched consignment	1.5.	Consignee Name Address Postal code Tel.	 Person responsible for the load in EU Name Address Postal code Tel. 					
patch								
s of dis	1.7.	Country of ISO code I.8. Region of Code origin origin	1.9.	Country of destination	ISO code	I.10. Region of Coo destination	le	
I : Details	1.11.	Place of origin Semen centre	I.12.	Place of destination Semen centre □		blding 🗖		
Part		Name Approval number Address		Name 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			I.17. No(s) of CITES				
	I.18.	Description of commodity				odity 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.	b. 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.	.28. Identification of the commodities						
Species (Scientific name) Donor identity				Date of collection Quantity				

COUNTRY Equine semen - Section D II. Health information II.a. Certificate reference No II.b 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: Certificatior (1) either meets the conditions laid down in Chapter I(I)(1) and is operated and supervised in accordance with [11.1.1. 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 []].1.1. the conditions laid down in Chapter I(II)(2) of Annex D to Directive 92/65/EEC;] II.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 least as strict as for imports of semen of animals of the equine species into the Union in accordance with Directive 92/65/EEC:1 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) 2018/657 (6);] (1) or [Model 2 in Section B of Part 1 of Annex III to Regulation (EU) 2018/657 (6);] [Model 3 in Section C of Part 1 of Annex III to Regulation (EU) 2018/657 (6);] (1) or (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);] 11.2.3. was stored under conditions which satisfy the terms of Annex D to Directive 92/65/EEC; sent to the place of loading in a sealed container in accordance with point 1.4 of Chapter III(I) of Annex D to 11.2.4. 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. Box I.17.: 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.

▼<u>C2</u>

-

COUN	TRY			Equine semen – Section D						
II.	Health i	nformation	II.a. Certificate reference No	II.b.						
Box	1.22.:	The number of packages shall corre	espond to the number of containers.							
Box	1.23.:	The identification of container and	seal number shall be indicated.							
Box	x I.28.: The donor identity shall correspond to the official identification of the animal.									
	The date of collection shall be indicated in the following format: dd/mm/yyyy.									
Part	II:									
(¹)	Delete	as necessary.								
(2)) Imports of equine semen are authorised from a third country listed in column 2 of Annex I to Commission Implementing Regulation (EU) 2018/657 provided that the semen was collected in the part of the territory of the third country detailed in column 4 of that Annex from a donor stallion of the category of equidae indicated in column 11, 12 or 13 of that Annex.									
(3)	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).									
(⁵)		oproved semen collection centres lis Commission websites:	sted in accordance with Article 11(4) and Article	17(3)(b) of Directive 92/65/EEC						
		ec.europa.eu/food/animals/live_anin c.europa.eu/food/animal/semen_ova								
(⁶)	accom		he health certificate(s) or the officially endo from the approved semen collection centre of th nust be attached to this certificate.							
—	The sig	nature and the stamp must be in a d	different colour to that of the printing.							
Offic	ial veteri	narian								
	Name	(in capital letters):		Qualification and title:						
	Date:			Signature:						
	Stamp									

▼<u>C2</u>

PART 2

Model health certificate for imports of ova and embryos

Section A

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

COUNTRY:

	l.1.	Consignor Name					1.2.	Certificate referen	ce No	1.2.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					I.6.	I.6. Person responsible for the load in EU Name Address						
atched c	Postal code Tel.							Postal code 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 destination	Code			
betail	l.11.	Place of orig				1	I.12.	Place of destination		_				
		Embryo tea	m 🗖					Holding 🗖	Embryo team					
Par		Name Approval number Address						Name Address	Approval nun	nber				
		Postal code						Postal code						
	1.13.	13. Place of loading						Date of departure						
	l.15.	Means of tra	ansport				l.16.	Entry BIP in EU						
		Aeroplane D			Railway wago	n 🗖								
		Road vehicl Identification Documenta					l.17.							
	l.18.	Description	of commodity	/					I.19. Commo	dity code (HS code) 05 11 99 85				
										I.20. Quantity				
	I.21.									I.22. Number of pa	ackages			
	1.23.	Seal/Contai	ner No				1.24.							
	1.25.	Commoditie Artificial rep	es certified for roduction											
	1.26.	For transit th	hrough EU to	third	country			I.27. For import or	admission int	o EU 🛛				
		Third countr	ry ISC) cod	e									
	1.28.	Identification	n of the comr	noditie	es									
	s	Species (Scie name)	ntific	C	ategory	۵)onor i	dentity D	ate of collection	on Quar	ntity			

	II. Health	information	i	II.a. Ce	ertificate reference No		II.b.							
_	I, the under	signed, offic	ial veterinarian, c	f the exporting	country (²)		borting country)	eby						
	certify that:						ooning country							
	II.1.	The ova (1)/embryos (1) described above:												
	II.1.2.	accordan		I(III) of Annex I	m (³) described in Box I.11, w D to Directive 92/65/EEC (⁴) /ear;									
	II.1.3.	.1.3. were collected (1)/produced (1), processed and stored in accordance with the requirements of Chapter III(II) of Annex D to Directive 92/65/EEC;												
	II.1.4.	II.1.4. were collected at a place separated from other parts of the premises or holding which is in good repair and was cleaned and disinfected prior to the collection;												
	II.1.5.	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 fror	n donor mares wi	nich:										
		II.1.6.1.	a Member Sta regionalisation	te of the Union in accordance	a period of 3 months (or sin during the 3 months period) with Article 13 of Directive 20 as during that period	in the expo	orting country or, in the	case o						
				dered to be in Directive 2009	fected with African horse s /156/EC,	ickness in	accordance with Article	5(2)(a						
			— free from	Venezuelan eq	uine encephalomyelitis for a	period of at	least 2 years,							
			— free from	glanders and d	ourine for a period of at least	6 months;								
	(1) either	[II.1.6.2.		a country of e od of at least 6 r	export which was on the day months;]	of collectio	n free from vesicular sto	omatitis						
	(¹) or	[II.1.6.2.	result at a service with the relevant	um dilution of 1 int Chapter of t lood sample tak	Itralisation test for vesicular in 32 or a VS ELISA carrie he Manual of Diagnostic Tes en on	d out with a sts and Vac	a negative result in acco cines for Terrestrial Ani	ordance mals o						
(¹) <i>either</i> [II.1.6.3. during a period of the past 30 days prior to the date of the collection were located in veterinary supervision which fulfilled from the day of the collection of the ova (¹)/emb date of their dispatch the conditions for a holding laid down in Article 4(5) of Direction and in particular:]								until the						
	(1) or	[II.1.6.3.	collection were collection of the	e kept in holdine ova (1)/embinises, the cond	embryos (¹), during a period ngs under veterinary super ryos (¹) until the end of the itions for a holding laid dow	vision which period of	n fulfilled, from the day 30 days mandatory stor	of the						

►⁽¹⁾ <u>M1</u>

COUNTRY

► ⁽¹⁾	Eq	uine	ova	emb	ryos	- ;	Sect	ion	А	Ì
------------------	----	------	-----	-----	------	-----	------	-----	---	---

II Lineth information			II a Cartificato raforanza No.
II.:	Health information		II.a. Certificate reference No II.b.
	(1) either	[II.1.6.3.1.	following a case of a disease mentioned below not all the animals of specie susceptible to that disease located in the holding were slaughtered or killed and the holding has been free:
			 from any type of equine encephalomyelitis for a period of at least 6 months beginning on the day on which the equidae suffering from the disease ar slaughtered,
			 from equine infectious anaemia for at least the period required to obtain a negative result in an agar gel immunodiffusion test (AGID or Coggins tests) carried out of samples taken after the infected animals were slaughtered on two occasion 3 months apart from each of the remaining equidae,
			 from vesicular stomatitis for a period of at least 6 months from the last recorder case,
			 from rabies for a period of at least one month from the last recorded case,
			 from anthrax for a period of at least 15 days from the last recorded case,]
	(¹) or	[II.1.6.3.1.	following a case of a disease mentioned below all the animals of species susceptible that disease located in the holding were slaughtered or killed and the premise disinfected, the holding was free for a period of at least 30 days from any type of equin encephalomyelitis, equine infectious anaemia, vesicular stomatitis and rabies or period of at least 15 days in the case of anthrax, beginning on the day on whic following the destruction of the animals the disinfection of the premises was satisfactorily completed;]
	II.1.6.4.		d of the past 30 days prior to the collection the ova (1)/embryos (1) were kept in holding of the equidae has shown clinical signs of contagious equine metritis for a period of
	II.1.6.5.	of the ova (d for natural breeding during a period of at least 30 days prior to the date of the collection /embryos (1) and between the date of the first samples referred to in points II.1.6.6 and the date of the collection of the ova (1)/embryos (1);
	II.1.6.6.	Manual of Di which is reco	one the tests, which meet at least the requirements of the relevant Chapters of the ignostic Tests and Vaccines for Terrestrial Animals of the OIE, carried out in a laborator gnised by the competent authority and has the tests referred to hereinafter included in i equivalent to that provided for in Article 12 of Regulation (EC) No 882/2004(7), a
	,	(⁸) [II.1.6.6.1.	for equine infectious anaemia (EIA), an agar-gel immuno-diffusion test (AGID of Coggins test) or an enzyme-linked immunosorbent assay (ELISA) with a negative result carried out on a blood sample taken on
		II.1.6.6.2.	for contagious equine metritis (CEM), an agent identification test carried out with negative result on at least two specimens (swabs) taken during the period referred to point II.1.6.5 from at least the mucosal surfaces of the clitoral fossa and the clitor sinuses of the donor mare
		(1) either	[II.1.6.6.2.1. on two occasions with an interval of not less than 7 day on

▶⁽ⁱ⁾ Equine ova/embryos - Section A -

COUNTRY

II.

II.1.7.

II.1.8.

II.2.

(12) [II.3.

Notes

Part I:

Box I.11.:

Box I.22.:

Box 1.23.:

Box 1.28 .:

 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; The embryos described above were conceived by artificial insemination(1)/as a result of <i>in vitro</i> fertilisation (1) using semen meeting the requirements of Directive 92/65/EEC and coming from semen collection centres approved in accordance with Article 11(2) or 17(3)(b) of Directive 92/65/EEC (⁹) and located respectively in a Member State of the Union or in a third country or parts of the territory of a third country listed in columns 2 and 4 of Annex I to Commission Implementing Regulation (EU) 2018/659 from which the import of equine semen collected from registered horses, registered equidae or equidae for breeding and production is authorised in 										
 genome of Taylorella equigenitalis by a polymerase chain greation (PCR) acried out within 48 hours after taking the specimens from the donor animal.] The samples referred to in points II.16.6.2.1 and II.16.6.2.2 were in no case taken earlier than 7 days (systemic treatment) or 21 days (local treatment) alter antimicrobial treatment of the donor stallon and were placed in transport medium with activated charcoal, such as Amies medium, before dispatch to the laboratory. II.16.7. to the best of my knowledge and as far as I could accertain, were not in contact with equidae suffering from an infectious or contagious disease during the period of 15 days immediately preceding the collection; II.16.8. on the day of the collection of the ova (1)/embryos (1) did not show clinical signs of an infectious or contagious disease. were collected (1)/produced (1) after the date on which the embryo collection (1)/production (1) team described in Box I.11 was approved by the competent authority of the exporting country; were processed and stored under approved conditions for a period of at least 30 days immediately after their collection (1)/production (1), and were transported under conditions which satisfy the terms laid down in Chapter HI(II) of Annex D to Directive 92/85/EEC; 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 the tricitoy of a third country isted in columns 2 and 4 of Annex I the Commission Implementing Regulation (EU) 2018/559 from which the import of equine semen collected from registered equidae regulation (EU) 2018/559 from which the import of equine sement ocliceted from registered ends of the terroly of a third country isted in columns 2 and 4 of Annex I thereto. (*) (*) (*) (*) (*) (*) (*) (*) (*) (*)	Health information				II.a.	Certificate reference No	II.b.			
 earlier than 7 days (systemic treatment) or 21 days (local reatment) after antimicrobial treatment of the donor stallion and were placed in transport medium with activated charcoal, such as Amies medium, before dispatch to the laboratory. II.16.7. to the best of my knowledge and as far as I could ascertain, were not in contact with equidae suffering from an infectious or contagious disease during the period of 15 days immediately preceding the collection; II.16.8. on the day of the collection of the ova (1)/embryos (1) did not show clinical signs of an infectious or contagious disease; 7. were collected (1)/produced (1) after the date on which the embryo collection (1)/production (1) team described in Box I.11 was approved by the competent authority of the exporting country; 8. were processed and stored under approved conditions for a period of 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(1) of Annex D to Directive S2/65/EEC; 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 S2/65/EEC and coming from semen collection centres approved in accordance with Article 11(2) or 17(3)(b) of DirectiveS9/65/650 from which the import of equine semen collected for mergistered horses, registered equidae for breeding and production is authorised in accordance with Article 4 of Commission Implementing Regulation (EU) 2018/659 and indicated in columns 2 and 4 of Annex 1 to Compute 4 or 90° (1); (II.3. The ova used for <i>in vitro</i> production of the embryos described above comply with the requirements of Annex D to Directive 4 of commission implementing Regulation (EU) 2018/659 and indicated in columns 11, 12 and 13 of Annex I thereeto, (1°) (1°); (II.3. The lace of origin shall correspond to the embryos collection team or embryo prod	(¹) and/or [II.1.6					genome of <i>Taylorella equigenitalis</i> by a p or real-time PCR , carried out within 48 h	olymerase chain reaction (PCR)			
from an infectious or contagious disease during the period of 15 days immediately preceding the collection; II.16.8. on the day of the collection of the ova (1)/embryos (1) did not show clinical signs of an infectious or contagious disease; 7. were collected (1)/produced (1) after the date on which the embryo collection (1)/production (1) team described in Box 1.11 was approved by the competent authority of the exporting country; 8. were processed and stored under approved conditions for a period of 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; 7. 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 seme collection centres approved in accordance with Article 117(3)(b) of Directive 92/65/EEC (2) and located respectively in a Member State of the Union or in a third country or parts of the territory of a third country of equine semen collected from registered horses, registered equidae or equidae for breeding and production is authorised in accordance with Article 4 17 Commission Implementing Regulation (EU) 2018/659 free and indicated in columns 11, 12 and 13 of Annex I to Commission Implementing Regulation (EU) 2018/659 and indicated in columns 11, 12 and 13 of Annex I thereto. (¹⁰) (¹¹); (II.3. The ova used for <i>in vitro</i> 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.] tes < t.t. < t.t. < t.t. < t.t. < t.t. < The place of origin shall correspond to the embryos collection team or embryo production team by which the ova/embryos were collect/0/roduced, processed, stored and approved in accordance with Article 17(3)(b) of Directive 92/65/EEC and listed on the Commissi	earlie treatn					7 days (systemic treatment) or 21 days (loc the donor stallion and were placed in tr	cal treatment) after antimicrobial ansport medium with activated			
 contagious disease; 7. were collected (1)/produced (1) after the date on which the embryo collection (1)/production (1) team described in Box I.11 was approved by the competent authority of the exporting country; 8. were processed and stored under approved conditions for a period of 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; 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 (2) and located respectively in a Member State of the Union or in a thride 11(2) or 17(3)(b) Directive 92/65/EEC (2) and located respectively in a Member State of the Union or in a thride equilation (2) 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. (19) (11); [II.3. The ova used for <i>in vitro</i> 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.] tes tt: (1.11: 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 Directive 92/65/EEC and listed on the Commission website: http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm (1.22: The number of packages shall correspond to the number of containers. (1.23: The identification of container and seal number shall be indicated.		II.1.6.7.	from an infect							
 Box 1.11 was approved by the competent authority of the exporting country; a. were processed and stored under approved conditions for a period of 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; 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 el do 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 Union or in a third country or parts of the territory of a third country listed in columns 2 and 4 of Annex 1 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 1 thereto. (¹0) (¹¹); [II.3. The ova used for <i>in vitro</i> 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.] tes (1.11: 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 Directive 92/65/EEC and listed on the Commission website: http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm (1.22: The number of packages shall correspond to the number of containers. (1.23: The identification of container and seal number shall be indicated. <		II.1.6.8.			lectio	n of the ova (1)/embryos (1) did not show	clinical signs of an infectious or			
 collection (¹)/production (¹), and we're transported under conditions which satisfy the terms laid down in Chapter III(II) of Annex D to Directive 92/65/EEC; The embryos described above were conceived by artificial insemination(1)/as a result of <i>in vitro</i> fertilisation (¹) using semen meeting the requirements of Directive 92/65/EEC and coming from semen collection centres approved in accordance with Article 11(2) or 17(3)(b) of Directive 92/65/EEC (⁹) and located respectively in a Member State of the Union or in a third country or parts of the territory of a third country listed in columns 2 and 4 of Annex I to Commission Implementing Regulation (EU) 2018/659 from which the import of equine semen collected from registered horses, registered equidae or equidae for breeding and production is authorised in accordance with Article 4 of Commission Implementing Regulation (EU) 2018/659 and indicated in columns 11, 12 and 13 of Annex I thereto. (¹⁰) (¹¹); [II.3. The ova used for <i>in vitro</i> 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.] tts (1.11: 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 Directive 92/65/EEC and listed on the Commission website: http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm (1.22: The number of packages shall correspond to the number of containers. (1.23: The category: specify if <i>in vivo</i> derived embryos, <i>in vivo</i> derived ova, <i>in vitro</i> produced embryos or micromanipulated embryos. 	.7.						production (1) team described in			
 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 Union or in a third country parts of the territory of a third country listed in columns 2 and 4 of Annex 1 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 1 and 13 of Annex 1 thereto. (¹⁰) (¹¹); [II.3. The ova used for <i>in vitro</i> 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.] tt: (I.1.: 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 Directive 92/65/EEC and listed on the Commission website: http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm (1.22.: The number of packages shall correspond to the number of containers. (1.23.: The identification of container and seal number shall be indicated. (1.23.: The category: specify if <i>in vivo</i> derived embryos, <i>in vivo</i> derived ova, <i>in vitro</i> produced embryos or micromanipulated embryos. 	.8.	were processed and stored under approved conditions for a period of 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;								
Directive 92/65/EEC and in particular the requirements set up in points II.1.1 to II.1.8 of this certificate.] tes t1: (1.11:: 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 Directive 92/65/EEC and listed on the Commission website: http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm (1.22:: The number of packages shall correspond to the number of containers. (1.23:: The identification of container and seal number shall be indicated. (1.28:: The category: specify if <i>in vivo</i> derived embryos, <i>in vivo</i> derived ova, <i>in vitro</i> produced embryos or micromanipulated embryos.		The embryos described above were conceived by artificial insemination(1)/as a result of <i>in vitro</i> fertilisation (¹) using semen meeting the requirements of Directive 92/65/EEC and coming from semen collection centres approved in accordance with Article 11(2) or 17(3)(b) of Directive 92/65/EEC (⁹) and located respectively in a Member State of the Union or in a third country or parts of the territory of a third country listed in columns 2 and 4 of Annex I to Commission Implementing Regulation (EU) 2018/659 from which the import of equine semen collected from registered horses, registered equidae or equidae for breeding and production is authorised in accordance with Article 4 of Commission Implementing Regulation (EU) 2018/659 and indicated in columns 11, 12 and 13 of Annex I thereto. (¹⁰) (¹¹);								
t1: x1.11.: 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 Directive 92/65/EEC and listed on the Commission website: http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm x1.22.: The number of packages shall correspond to the number of containers. x1.23.: The identification of container and seal number shall be indicated. x1.28.: The category: specify if <i>in vivo</i> derived embryos, <i>in vivo</i> derived ova, <i>in vitro</i> produced embryos or micromanipulated embryos.	[11.3.									
 k1.11.: 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 Directive 92/65/EEC and listed on the Commission website: http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm k1.22.: The number of packages shall correspond to the number of containers. k1.23.: The identification of container and seal number shall be indicated. k1.28.: The category: specify if <i>in vivo</i> derived embryos, <i>in vivo</i> derived ova, <i>in vitro</i> produced embryos or micromanipulated embryos. 	tes									
 ova/embryos were collected/produced, processed, stored and approved in accordance with Article 17(3)(b) of Directive 92/65/EEC and listed on the Commission website: http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm x 1.22.: The number of packages shall correspond to the number of containers. x 1.23.: The identification of container and seal number shall be indicated. x 1.28.: The category: specify if <i>in vivo</i> derived embryos, <i>in vivo</i> derived ova, <i>in vitro</i> produced embryos or micromanipulated embryos. 	rt I:									
 k 1.22.: The number of packages shall correspond to the number of containers. k 1.23.: The identification of container and seal number shall be indicated. k 1.28.: The category: specify if <i>in vivo</i> derived embryos, <i>in vivo</i> derived ova, <i>in vitro</i> produced embryos or micromanipulated embryos. 	c I.11.:	ova/embry	os were collec	ted/prod	luced	, processed, stored and approved in acc				
k1.23.: The identification of container and seal number shall be indicated. k1.28.: The category: specify if <i>in vivo</i> derived embryos, <i>in vivo</i> derived ova, <i>in vitro</i> produced embryos or micromanipulated embryos.		http://ec.eu	ropa.eu/food/a	inimal/se	emen	_ova/equine/index_en.htm				
k I.28.: The category: specify if <i>in vivo</i> derived embryos, <i>in vivo</i> derived ova, <i>in vitro</i> produced embryos or micromanipulated embryos.	k I.22.:	The numbe	er of packages	shall co	rresp	and to the number of containers.				
micromanipulated embryos.	k I.23.:	The identifi	cation of conta	iner and	l seal	number shall be indicated.				
The donor identity shall correspond to the official identification of the animal.	k I.28.:				vo de	erived embryos, <i>in vivo</i> derived ova, <i>i</i>	n vitro produced embryos or			
		The donor	identity shall c	orrespor	nd to t	he official identification of the animal.				

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

►(1) <u>M1</u>

	ITRY	•	Equine ova/embryos - Section A				
II.	Health information	II.a. Certificate reference No	II.b.				
Part	11:						
(¹)	Delete as appropriate.						
(2)	Implementing Regulation (EU) 2018/659	rritory of third countries listed in columns 2 , respectively from which imports of registere as indicated in column 14 of Annex I thereto.					
(³)	Only approved embryo collection team Directive 92/65/EEC on the Commission	ns and embryo production teams listed in website:	accordance with Article 17(3)(b) of				
	http://ec.europa.eu/food/animal/semen_o	ova/equine/index_en.htm					
(4)	the Community of animals, semen, ova	1992 laying down animal health requirements a and embryos not subject to animal health I) to Directive 90/425/EEC (OJ L 268, 14.9.19	requirements laid down in specific				
(5)	Council Directive 2009/156/EC of 30 importation from third countries of equida	November 2009 on animal health conditi ae (OJ L 192, 23.7.2010, p. 1).	ions governing the movement and				
(⁶)	Insert date. (follow Guidance in Part II of	the Notes).					
(7)	Regulation (EC) No 882/2004 of the European Parliament and of the Council of 29 April 2004 on official controls performed to ensure the verification of compliance with feed and food law, animal health and animal welfare rules (OJ L 165, 30.4.2004, p. 1).						
(⁸)	donor equidae which have continuously equine infectious anaemia and no equid	D or Coggins test) or the ELISA for equine inf resided in Iceland since birth, provided that le dae and their semen, ova and embryos hav e ova or embryos were collected and the sem	celand has remained officially free of the been introduced into Iceland from				
(⁹)	Only approved semen collection centres on the Commission websites:	s listed in accordance with Article 11(4) or Ar	ticle 17(3)(b) of Directive 92/65/EEC				
	https://ec.europa.eu/food/animals/live_ar http://ec.europa.eu/food/animal/semen_c						
(¹⁰)	Regulation (EU) 2018/659 provided that	d from third countries listed in column 2 of A the semen was collected in the part of the te egory of equidae positively indicated in colum	erritory of the third country detailed in				
(11)	Does not apply to ova.						
(12)	Delete if none of the embryos in the cons	signment was produced by in vitro fertilisation	n 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:						

►(1) <u>M1</u>

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:								Veterin	nary certifica	te to EU
	l.1.	Consignor Name				1.2.	Certificate reference	ce No	l.2.a.		
		Address				1.3.	Central competent	authority			
		Tel.				1.4.	Local competent a	uthority			
Part I : Details of dispatched consignment	1.5.	Consignee Name Address				1.6.	Person responsible Name Address	e for the load	in EU		
tched co		Postal code Tel.					Postal code Tel.				
of dispa	1.7.	Country of ISC origin	O code	I.8. Region of origin	Code	1.9.	Country of destination	ISO code		on of nation	Code
tails	I.11.	Place of origin				I.12.	Place of destinatio				1
I: De		Embryo team C]				Holding 🗖	Embryo team			
Part		Name Address	A	pproval number			Name Address	Approval nun	nber		
		Postal code					Postal code				
	l.13.	Place of loading	g			l.14.	Date of departure				
	l.15.	Means of transp	port			I.16.	Entry BIP in EU				
		Aeroplane 🗖 _	Ship 🗆		on 🗖						
		Road vehicle Identification Documentary re		er 🗖		1.17.					
	I.18.	Description of c						I.19. Commo	dity code (HS code)	
									05 11 9	9 85	
									1.20. Qua	antity	
	I.21.								1.22. Nur	nber of packa	iges
	1.23.	Seal/Container	No						1.24.		
	1.25.	Commodities co Artificial reprodu									
	1.26.	For transit throu	ugh EU to	third country			I.27. For import or	admission inte	o EU		
		Third country	IS	O code							
	1.28.	Identification of	the comm	nodities		I					
	s	pecies (Scientifi name)	с	Category	D	onor i	dentity Da	ate of collectio	on	Quantity	

	II. Health	information		II.a. Certificate reference No	II.b.								
	I, the unders	igned, offici	al veterina	an, of the exporting country (2)									
				(nai	me of exporting country)								
	certify that:												
	II.1.	The ova (¹)/embryos	1) described above:									
		II.1.2.	supervise	were collected (¹)/produced (¹) by the team (³) described in Box I.11, which has been approved and supervised in accordance with Chapter I(III) of Annex D to Directive 92/65/EEC and is subject to nspection by an official veterinarian at least once every calendar year;									
		II.1.3.		were collected (¹)/produced (¹), processed and stored in accordance with the requirements of Chapter III(II) of Annex D to Directive 92/65/EEC;									
		II.1.4.		were collected at a place separated from other parts of the premises or holding which is in good repair and was cleaned and disinfected prior to the collection;									
II.1.5. were examined, processed and packed in laboratory facilities which are not situated in a to prohibition or quarantine measures as set out in Box II.1.6, in a section which is separa section for storing equipment and materials used in contact with donor animals and fr where the donor animals are handled;													
		II.1.6.	come fro	n donor mares which:									
			II.1.6.1.	were continuously resident for 3 months (or sind Member State of the European Union during th or, in the case of regionalisation according to A part of the territory of the exporting country whic	e 3 months period) in the exporting countr rticle 13 of Directive 2009/156/EC (8), in the								
				 not considered to be infected with Af Article 5(2)(a) and (b) of Directive 2009/15 	rican horse sickness in accordance wit 56/EC,								
				 free from Venezuelan equine encephalom 	yelitis for at least 2 years,								
				 free from glanders and dourine for at leas 	t 6 months;								
		(¹) either	[II.1.6.2.	originated from a country of export which was stomatitis for at least 6 months;]	s on the day of collection free of vesicula								
		(1) or	[II.1.6.2.	were tested by a virus neutralisation test for vi on									
		(¹) either	[II.1.6.3.	during the past 30 days prior to collection have supervision which fulfilled from the day of colle their dispatch the conditions for a holding laid do and in particular:]	ction of ova (1)/embryos (1) until the date of								
		(1) or	[II.1.6.3.	during the past 30 days prior to collection have supervision which fulfilled from the day of collect of frozen ova (')/embryos ('), the period of premises elapsed, the conditions for a hold	ction of ova (1)/embryos (1) until, in the cas 30 days mandatory storage at approve								

►⁽¹⁾ <u>M1</u>

_

Health information	n		II.a. Certificate reference No	II.b.
	(1) either	[II.1.6.3.1.	following a case of a disease mentioned bell susceptible to the disease located on the hold the holding has been free:	
			 from any type of equine encephalor beginning on the day on which the equinor slaughtered, 	
			 from equine infectious anaemia for at lean egative result in an agar gel immunodif out on samples taken after the infect two occasions 3 months apart from each 	ffusion test (Coggins tests) carrie ed animals were slaughtered o
			 from vesicular stomatitis for at least 6 model 	onths from the last recorded case
			 from rabies for at least one month from t 	he last recorded case,
			 from anthrax for at least 15 days from the 	e last recorded case,]
	(¹) or	[11.1.6.3.1.	following a case of a disease mentioned b susceptible to the disease located in the hi killed and the premises disinfected, the ho 30 days from any type of equine encephalom vesicular stomatitis and rabies or 15 days in the day on which following the destruction of premises was satisfactorily completed;]	olding have been slaughtered of lding has been free for at leas yelitis, equine infectious anaemia the case of anthrax, beginning o
	II.1.6.4.		ast 30 days prior to collection have been kep om clinical signs of contagious equine metritis	
	II.1.6.5.	collection of	een used for natural breeding during at lea f ova or embryos and between the date of .6 and II.1.6.7 and the date of the collection of	f the first samples referred to i
	II.1.6.6.	test) or an on	subjected with negative result to an agar-ge ELISA for equine infectious anaemia carrie 	ed out on a blood sample take ays prior to the date of the firs out on a sample of blood taken o
	II.1.6.7.	isolation of negative re- the first coll sinuses on and on an	subjected to an agent identification test for <i>Taylorella equigenitalis</i> after a cultivation of sults in each case on samples taken during the ection of ova or embryos from mucosal surface wo consecutives oestrus periods on	of 7 to 14 days carried out wit e past 30 days prior to the date of eas of the clitoral fossa and clitora
	II.1.6.8.	equidae sut	of my knowledge and as far as I could ascerta fering from an infectious or contagious diseas ne collection;	
	II.1.6.9.	have on the or contagion	day of collection of ova (1)/embryos (1) not sh is disease;	nown clinical signs of an infectiou
II.1.7.	were colle	ected (1)/pro	duced (1) after the date on which the embryo	collection (1)/production (1) teal

)			
-			

II.	Health i	information		II.a. Certificate reference No	II.b.
		II.1.8.	collection (1)/production		at least 30 days immediately after their itions which satisfy the terms laid down
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 and located respectively in a Member State of the European Union or in a third country or parts of the territory of third country listed in columns 2 and 4 of Annex I to Commission Implementing Regulation (EU) 2018/659 from which the import of equine semen collected from registered horses, registered equidae or equidae for breeding and production is authorised in accordance with Article 4 of Commission Implementing Regulation (EU) 2018/659 and indicated in columns 11, 12 and 13 of Annex I thereto. (⁶) (⁷);				
11.3.		The ova used for <i>in vitro</i> 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).			
Note	es				
Part	l:				
Box	l.11.:	ova/embry	os were collected/produ		embryo production team by which the in accordance with Article 17(3)(b) of
		http://ec.eu	ropa.eu/food/animal/sen	nen_ova/equine/index_en.htm	
Box	1.22.:	The numbe	er of packages shall corre	espond to the number of containers.	
Box	1.23.:	The identifi	cation of container and s	eal number shall be indicated.	
Box	1.28.:		ory: specify if <i>in vivo</i> pulated embryos.	derived embryos, in vivo derived	ova, in vitro produced embryos or
		The donor	identity shall correspond	to the official identification of the anima	al.
		The date o	f collection shall be indic	ate in the following format: dd/mm/yyyy	l.
Part	11:				
(¹)	Delete	as appropri	ate.		
(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 Count Directive 92/65/EEC on the Commission website:				
	http://e	c.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 donc equidae which have continuously resided in Iceland since birth, provided that Iceland has remained officially free c equine infectious anaemia and no equidae and their semen, ova and embryos have been introduced into Iceland fror outside prior to and during the period the semen was collected.				

►⁽¹⁾ <u>M1</u>

NTRY	F [∞] Equ	ne ova/embryos - Section B			
Health information	II.a. Certificate reference No	II.b.			
Only approved semen collection centres listed in accordance with Article 11(4) or Article 17(3)(b) of Council Directive 92/65/EEC on the Commission websites:					
Does not apply to ova.					
OJ L 192, 23.7.2010, p. 1.					
The signature and the stamp must be in a different colour to that of the printing.					
cial veterinarian					
Name (in capital letters):	C	Qualification and title:			
Date:	s	Signature:			
Stamp:					
	Health information Only approved semen collection centres li 92/65/EEC on the Commission websites: https://ec.europa.eu/food/animals/live_anim http://ec.europa.eu/food/animals/live_anim loos not apply to ova. OJ L 192, 23.7.2010, p. 1. The signature and the stamp must be in a c cial veterinarian Name (in capital letters): Date:	Health information II.a. Certificate reference No Only approved semen collection centres listed in accordance with Article 11(4) or Article 92/65/EEC on the Commission websites: https://ec.europa.eu/food/animals/live_animals/approved-establishments_en; http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm Does not apply to ova. OJ L 192, 23.7.2010, p. 1. The signature and the stamp must be in a different colour to that of the printing. cial veterinarian Name (in capital letters): C Date: S			

►(1) <u>M1</u>

PART 3

Explanatory notes for the certification

(a)	The health certificates shall be issued by the competent authority of the exporting country, based on the model set out in Part 1 or 2 of Annex III, according to the layout of the model that corresponds to the commodity concerned. They shall contain, in the numbered order that appears in the model, the attestations that are required for any country and, as the case may	(g)	When the health certificate, including additional schedules referred to in (f), comprises more than one page, each page shall be numbered -(<i>page number</i>) of (<i>total number</i> of <i>pages</i>)-, 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.
	be, those supplementary guarantees that are required for the exporting country or part of the territory of the country.	(h)	The original of the health certificate shall be completed and signed by an official veterinarian within 24 hours prior to loading
(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 wadon, lorry, aircraft or ship.		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
(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.		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.
(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.	(i) (i)	The original of the health certificate shall accompany the consignment until it reaches the border inspection post of entry into the Union. The certificate reference number referred to in Box 1.2 and
(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.	0,	Box II.a of the model health certificate shall be provided by t competent authority of the exporting country.
(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.		

(¹) OJ L 13, 16.1.1997, p. 28.

ANNEX IV

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

- 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.
- 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), dec certificate No	remained on board the ship during the voyage
Done at on	
(Port of arrival)	(Date of arrival)
	(signature of master)
(stamp)	
	(name in capital letters and title)

PART 3

Model Transhipment Manifest

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

Serial Number:

Reference No of Air Cargo Transfer Manifest: (1)

Country where transhipment takes 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

I, the undersigned, official veterinarian (²)/customs officer (²) at the above mentioned airport (²)/port (²) declare that the transhipment took place under my supervision and in compliance with the following conditions:

(a) the equidae were during the transhipment protected from attacks by insect vectors of diseases transmissible to equidae;

(b) the equidae did not come into contact with equidae of a different health status;

(c) the crates, containers or jet-stalls and the surrounding airspace in the transport compartment were sprayed with an appropriate insect repellent in combination with an insecticide immediately after the closing of the doors of the aircraft (²)/vessel (²).

The consignment has been transhipped in full and apparent good order and conditions except as noted in the "Remarks" column.

Done at	on
(signature of the official veterinarian or customs officer)	Stamp
(name in capital letters and title)	
 (¹) Keep empty if transhipment from vessel to vessel (²) Delete as appropriate 	