Commission Decision of 26 August 2010 laying down model health certificates for trade within the Union in semen, ova and embryos of animals of the equine, ovine and caprine species and in ova and embryos of animals of the porcine species (notified under document C(2010) 5779) (Text with EEA relevance) (2010/470/EU)

COMMISSION DECISION

of 26 August 2010

laying down model health certificates for trade within the Union in semen, ova and embryos of animals of the equine, ovine and caprine species and in ova and embryos of animals of the porcine species

(notified under document C(2010) 5779)

(Text with EEA relevance)

(2010/470/EU)

THE EUROPEAN COMMISSION,

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

Having regard to Council Directive 92/65/EEC of 13 July 1992 laying down animal health requirements governing trade in and imports into the Community of animals, semen, ova and embryos not subject to animal health requirements laid down in specific Community rules referred to in Annex A(I) to Directive 90/425/EEC[(1)] and in particular the fourth indent of Article 11(2) and the third indent of Article 11(3) thereof,

Whereas:

(1) Directive 92/65/EEC lays down the animal health requirements governing trade within the Union of animals, semen, ova and embryos not subject to the animal health requirements laid down in specific Union acts. It includes requirements for trade in semen, ova and embryos of animals of the equine, ovine and caprine species and in ova and embryos of animals of the porcine species (‘the commodities’). In addition, it provides for health certificates to be established for trade in the commodities within the Union.

(2) Annex D to Directive 92/65/EEC, as amended by Commission Regulation (EU) No 176/2010[(2)], sets out certain new requirements for the commodities which are to apply from 1 September 2010.

(3) Annex D to Directive 92/65/EEC, as thus amended by Regulation (EU) No 176/2010, introduces rules concerning semen storage centres and detailed conditions for their approval and supervision. It also sets out detailed conditions for the approval and supervision of embryo collection and production teams, for the collection and processing of *in vivo* derived embryos and the production and processing of *in vitro* fertilised embryos and micromanipulated embryos. Annex D, as thus amended, also

amended the conditions to be applied to the donor animals of semen, ova and embryos of animals of the equine, ovine and caprine species and of ova and embryos of porcine species.

(4) It is necessary to establish new model health certificates for trade within the Union of the commodities taking into account the animal health requirements set out in Annex D to Directive 92/65/EEC, as amended by Regulation (EU) No 176/2010.

(5) In addition, provision should be made for existing stocks of commodities in the Union that comply with the provisions of Directive 92/65/EEC established prior to the entry into force of the amendments introduced by Regulation (EU) No 176/2010. Accordingly, it is necessary to set out separate model health certificates for trade within the Union in semen, ova and embryos of animals of the equine, ovine and caprine species and trade in ova and embryos of animals of the porcine species collected or produced, processed and stored in accordance with Annex D to Directive 92/65/EEC prior to 1 September 2010.

(6) The long lasting stocking capabilities for such commodities make it impossible at present to fix a date for the exhaustion of the existing stocks. Therefore, it is not possible to fix a date for the termination of the use of those model health certificates for the existing stocks.

(7) In the interests of consistency and simplification of Union legislation, the model health certificates should be set out in a single decision and take account of Commission Regulation (EC) No 599/2004 of 30 March 2004 concerning the adoption of a harmonised model certificate and inspection report linked to intra-Community trade in animals and products of animal origin[(3)].

(8) In order to ensure full traceability of the commodities, model health certificates should be set out in this Decision for trade within the Union in semen of animals of the equine, ovine and caprine species collected in approved semen collection centres and dispatched from an approved semen storage centre, whether or not the latter constitutes part of a semen collection centre approved under a different approval number.

(9) In the interests of clarity of Union legislation, the Union acts setting out model health certificates for trade within the Union in the commodities concerned should be expressly repealed. Accordingly, Commission Decision 95/294/EC of 24 July 1995 determining the specimen animal health certificate for trade in ova and embryos of the equine species[(4)], Commission Decision 95/307/EC of 24 July 1995 determining the specimen animal health certificate for trade in semen of the equine species[(5)], Commission Decision 95/388/EC of 19 September 1995 determining the specimen certificate for intra-Community trade in semen, ova and embryos of the ovine and caprine species[(6)] and Commission Decision 95/483/EC of 9 November 1995 determining the specimen certificate for intra-Community trade in ova and embryos of swine[(7)] should be repealed.

(10) The measures provided for in this Decision are in accordance with the opinion of the Standing Committee on the Food Chain and Animal Health,

HAS ADOPTED THIS DECISION:

Article 1

Subject matter

This Decision lays down model health certificates for trade within the Union in the following commodities:

(a) semen of animals of the equine species;

(b) ova and embryos of animals of the equine species;

(c) semen of animals of the ovine and caprine species;

(d) ova and embryos of animals of the ovine and caprine species;

(e) ova and embryos of animals of the porcine species.

I[F1] Article 2

Trade in semen of animals of the equine species

A health certificate in accordance with one of the following models set out in Annex I shall accompany consignments of semen of animals of the equine species during transport from one Member State to another:

(a) model health certificate IA for trade in the Union in consignments of semen of animals of the equine species 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;

(b) model health certificate IB for trade in the Union in 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 and dispatched after 31 August 2010 from an approved semen collection centre of origin of the semen;

(c) model health certificate IC for trade in the Union in consignments of stocks of semen of animals of the equine species 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;

(d) model health certificate ID for trade in the Union in consignments of:

  (i) semen of animals of the equine species collected, processed and stored in accordance with Directive 92/65/EEC after 30 September 2014 and dispatched from an approved semen storage centre;

  (ii) 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.

Textual Amendments

F1 Substituted by Commission Implementing Decision (EU) 2015/261 of 6 February 2015 amending Decisions 2010/470/EU and 2010/471/EU as regards the animal health certification requirements for trade in and for imports into the Union of semen, ova and embryos of animals of the equine species (notified under document C(2015) 548) (Text with EEA relevance).

Article 3

Trade in ova and embryos of animals of the equine species

A health certificate in accordance with one of the following models set out in Annex II shall accompany consignments of ova and embryos of animals of the equine species during transport from one Member State to another:

(a) model health certificate IIA for trade in the Union in consignments of ova and embryos of animals of the equine species 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;

(b) model health certificate IIB for trade in the Union in consignments of stocks of ova and embryos 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 and dispatched after 31 August 2010 by an approved embryo collection or production team of origin of the ova or embryos;

(c) model health certificate IIC for trade in the Union in consignments of stocks of ova and embryos of animals of the equine species collected, processed and stored in accordance with Directive 92/65/EEC before 1 September 2010 and dispatched after 31 August 2010 by an approved embryo collection team of origin of the ova or embryos.]

Textual Amendments

F1 Substituted by Commission Implementing Decision (EU) 2015/261 of 6 February 2015 amending Decisions 2010/470/EU and 2010/471/EU as regards the animal health certification requirements for trade in and for imports into the Union of semen, ova and embryos of animals of the equine species (notified under document C(2015) 548) (Text with EEA relevance).

Article 4

Trade in semen of animals of the ovine and caprine species

A health certificate in accordance with one of the following models set out in Annex III shall accompany consignments of semen of animals of the ovine and caprine species during transport from one Member State to another:

(a) model health certificate IIIA as set out in Part A, for consignments of semen collected after 31 August 2010 and dispatched from an approved semen collection centre of origin of the semen;

(b) model health certificate IIIB as set out in Part B, for consignments of stocks of semen collected, processed and stored before 1 September 2010 and dispatched after 31 August 2010 from an approved semen collection centre of origin of the semen;

(c) model health certificate IIIC as set out in Part C, for consignments of semen and stocks of semen referred to in (a) and (b) dispatched from an approved semen storage centre.

Article 5

Trade in ova and embryos of animals of the ovine and caprine species

A health certificate in accordance with one of the following models set out in Annex IV shall accompany consignments of ova and embryos of animals of the ovine and caprine species during transport from one Member State to another:

(a) model health certificate IVA as set out in Part A, for consignments of ova and embryos collected or produced after 31 August 2010 and dispatched by an approved embryo collection or production team of origin of the ova or embryos;

(b) model health certificate IVB as set out in Part B, for consignments of stocks of ova and embryos collected, processed and stored before 1 September 2010 and dispatched after 31 August 2010 by an approved embryo collection team of origin of the ova or embryos.

Article 6

Trade in ova and embryos of the porcine species

A health certificate in accordance with one of the following models set out in Annex V shall accompany consignments of ova and embryos of animals of the porcine species during transport from one Member State to another:

(a) model health certificate VA as set out in Part A, for consignments of ova and embryos collected or produced after 31 August 2010 and dispatched by an approved embryo collection or production team of origin of the ova or embryos;

(b) model health certificate VB as set out in Part B, for consignments of stocks of ova and embryos collected, processed and stored before 1 September 2010 and dispatched after 31 August 2010 by an approved embryo collection team of origin of the ova or embryos.

Article 7

Repeals

Decisions 95/294/EC, 95/307/EC, 95/388/EC and 95/483/EC are repealed.

Article 8

Applicability

This Decision shall apply from 1 September 2010.

Article 9

Addressees

This Decision is addressed to the Member States.

[F1 ANNEX I

Model health certificates for trade in the union in consignments of semen of animals of the equine species

PART A

Model health certificate IA for trade in the Union in consignments of semen of animals of the equine species 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

EUROPEAN UNION **Intra trade certificate**

Part I: Details of consignment presented

I.1. Consignor Name Address Postal code	I.2. Certificate reference No	I.2.a. Local reference No
	I.3. Central competent authority	
	I.4. Local competent authority	
I.5. Consignee Name Address Postal code	I.6.	
	I.7.	
I.8. Country of origin ISO code I.9. Region of origin Code	I.10. Country of destination ISO code I.11. Region of destination Code	
I.12. Place of origin Semen Centre ☐ Name Approval number Address Postal code	I.13. Place of destination Semen Centre ☐ Holding ☐ Name Approval number Address Postal code	
I.14.	I.15.	
I.16. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification:	I.17.	
I.18. Description of commodity	I.19. Commodity code (HS code) **05 11 99 85**	
	I.20. Quantity	
I.21. Temperature of products Ambient ☐ Chilled ☐ Frozen ☐	I.22. Number of packages	
I.23. Seal/Container No	I.24. Type of packaging	
I.25. Commodities certified for: Artificial reproduction ☐		
I.26. Transit through third country ☐ Third country ISO code Exit point Code Entry point BIP No	I.27. Transit through Member States ☐ Member State ISO code Member State ISO code Member State ISO code	
I.28. Export ☐ Third country ISO code Exit point Code	I.29.	
I.30.		
I.31. Identification of the commodities Species (Scientific name) Donor identity Date of collection Quantity		

EUROPEAN UNION **Equine semen – Part A**

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

I, the undersigned official veterinarian, hereby certify that:

Part II: Certification

II.1. The semen collection centre ([2]), in which the semen described above was collected, processed and stored for trade is approved and supervised by the competent authority in accordance with Chapters I(I)(1) and I(II)(1) of Annex D to Directive 92/65/EEC ([3]);

II.1.1. during the period commencing 30 days prior to the date of first collection of the semen described above until the date the fresh or chilled semen was dispatched or until the 30 days minimum storage period for frozen semen elapsed, the semen collection centre:

II.1.1.1. was situated on the territory or in the case of regionalisation in a part of the territory ([1]) of a Member State which was not considered to be infected with African horse sickness in accordance with Article 5(2)(a) and(b) of Directive 2009/156/EC ([4]);

II.1.1.2. fulfilled the conditions for a holding laid down in Article 4(5) of Directive 2009/156/EC;

II.1.1.3. contained only equidae which were free of clinical signs of equine viral arteritis and contagious equine metritis;

II.2. Only equidae satisfying the conditions laid down in Articles 4 and 5 or Articles 12 to 16 of Directive 2009/156/EC have been admitted onto the centre.

II.3. The semen described above was collected from donor stallions, which:

II.3.1. did not show any clinical sign of an infectious or contagious disease at the time of admission onto the semen collection centre and on the day the semen was collected;

II.3.2. were kept for a period of 30 days prior to the date of semen collection in holdings where no equine showed any clinical sign of equine viral arteritis or contagious equine metritis during that period;

II.3.3. were not used for natural mating during a period of at least 30 days prior to the date of first semen collection and from the dates of the first sample referred to in point II.3.5.1, II.3.5.2 or II.3.5.3 until the end of the collection period;

II.3.4. underwent the tests, which meet at least the requirements of the relevant Chapter of the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals of the OIE, carried out in a laboratory which is recognised by the competent authority and has the tests referred to hereinafter included in its accreditation in accordance with Article 12 of Regulation (EC) No 882/2004 ([5]), as follows:

II.3.4.1. for equine infectious anaemia (EIA), an agar-gel immuno-diffusion test (AGID or Coggins test) or an enzyme-linked immunosorbent assay (ELISA) for equine infectious anaemia with a negative result;

II.3.4.2. for equine viral arteritis (EVA),

([1]) *either* [II.3.4.2.1. a serum neutralisation test with a negative result at a serum dilution of one in four;]

([1]) *and/or* [II.3.4.2.2. a virus isolation test, polymerase chain reaction (PCR) or real-time PCR with a negative result on an aliquot of the entire semen of the donor stallion;]

II.3.4.3. for contagious equine metritis (CEM), an agent identification test carried out on three specimens (swabs) taken from the donor stallion on two occasions with an interval of not less than 7 days at least from the penile sheath (prepuce), the urethra and the fossa glandis;

The samples were in no case taken earlier than 7 days (systemic treatment) or 21 days (local treatment) 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 where they were subjected with negative result to a test for:

([1]) *either* [II.3.4.3.1. the isolation of *Taylorella equigenitalis* after cultivation under microaerophilic conditions for at least 7 days, set up within 24 hours after taking the specimens from the donor animal, or 48 hours where the specimens are kept cool during transport;]

EUROPEAN UNION **Equine semen – Part A**

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

(1) *and/or* [II.3.4.3.2. the detection of genome of *Taylorella equigenitalis* by PCR or real-time PCR, carried out within 48 hours after taking the specimens from the donor animal;]

II.3.5. were subjected with the results specified in point II.3.4 in each case to at least one of the test programmes detailed in points II.3.5.1, II.3.5.2 and II.3.5.3, as follows:

(6) [II.3.5.1. The donor stallion was continuously resident on the semen collection centre for a period of 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 into direct contact with equidae of lower health status than the donor stallion.

The tests described in point II.3.4 were carried out on samples taken (7) from the donor stallion at least once a year at the beginning of the breeding season or prior to the first collection of semen intended for trade in fresh, chilled or frozen semen and not less than 14 days following the date of the commencement of the residence period of at least 30 days prior to the date of first semen collection.]

(6) [II.3.5.2. The donor stallion was resident on the semen collection centre for a period of 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, and/or other equidae on the semen collection centre came into direct contact with equidae of lower health status.

The tests described in point II.3.4 were carried out on samples taken (7) from the donor stallion at least once a year at the beginning of the breeding season or prior to the first collection of semen intended for trade in fresh, chilled or frozen semen and not less than 14 days following the date of the commencement of the residence period of at least 30 days prior to the date of first semen collection,

and during the period of collection of the semen intended for trade in fresh, chilled or frozen semen the donor stallion was subjected to the tests described in point II.3.4, as follows:

(a) for equine infectious anaemia, one of the tests described in point II.3.4.1. was last carried out on a sample of blood taken (7) not more than 90 days prior to the date of the collection of the semen described above;

(b) for equine viral arteritis:

(1) *either* [one of the tests described in point II.3.4.2 was last carried out on a sample taken (7) not more than 30 days prior to the date of the collection of the semen described above;]

(1) *or* [one of the tests described in point II.3.4.2.2 was carried out on an aliquot of the entire semen of the donor stallion taken (7) not more than 6 months prior to the date of the collection of the semen described above and a blood sample taken (7) from the donor stallion during the 6-month period reacted with a positive result in a serum neutralisation test for equine viral arteritis at a serum dilution of more than one in four;]

(c) for contagious equine metritis, one of the tests described in point II.3.4.3 was last carried out on three specimens (swabs) taken (7) not more than 60 days prior to the date of the collection of the semen described above

(1) *either* [on two occasions at least 7 days apart;]

(1) *or* [on a single occasion and subjected to a PCR or real-time PCR.]]

(6) [II.3.5.3. The donor stallion does not meet the conditions set out in points 1.6(a) and (b) of Chapter II of Annex D to Directive 92/65/EEC and the semen is collected for trade in frozen semen.

The tests described in points II.3.4.1, II.3.4.2 and II.3.4.3 were carried out on samples taken (7) from the donor stallion at least once a year at the beginning of the breeding season,

and the tests described in points II.3.4.1 and II.3.4.3 were carried out on samples taken (7) from the donor stallion during the storage period of the semen of a minimum period of 30 days from the date of the collection of the semen and before the semen is removed from the semen collection centre, not less than 14 days and not more than 90 days after the collection of the semen described above,

EUROPEAN UNION **Equine semen – Part A**

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

and (1) *either* [the tests for equine viral arteritis described in point II.3.4.2 were carried out on samples taken (7) during the storage period of the semen of a minimum period of 30 days from the date of the collection of the semen and before the semen is removed from the semen collection centre or used, not less than 14 days and not more than 90 days after the collection of the semen described above.]

(1) *or* [the non-shedder state of a donor stallion seropositive for equine viral arteritis was confirmed by virus isolation test, PCR or real-time PCR carried out with a negative result on samples of an aliquot of the entire semen of the donor stallion taken (7) twice a year at an interval of at least 4 months and the donor stallion reacted with a positive result at a serum dilution of at least one in four in a serum neutralisation test for equine viral arteritis.]]

II.3.6. underwent the testing provided for in point II.3.5 on samples taken on the following dates

Identification of semen	Test programme	Start date (7)		Date of sampling for health tests (7)				
					EVA II.3.4.2.		CEM II.3.4.3.	
		Donor residence	Semen collection	EIA II.3.4.1.	Blood sample	Semen sample	1. sample	2. sample

(1) *either* [II.4. No antibiotics were added to the semen;]

(1) *or* [II.4. The following antibiotic or combination of antibiotics was added to produce a concentration in the final diluted semen of not less than (8): .....
..... ;]

II.5. The semen described above was:

II.5.1. collected, processed, stored and transported under conditions which comply with the requirements of Chapters II(I) (1) and III(I) of Annex D to Directive 92/65/EEC;

II.5.2. in the case of frozen semen, stored for a minimum period of 30 days from the date of collection of the semen;

II.5.3. sent to the place of loading in a sealed container in accordance with point 1.4 of Chapter III(I) of Annex D to Directive 92/65/EEC and bearing the number indicated in Box I.23.

EUROPEAN UNION **Equine semen – Part A**

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

Notes

Part I:

Box I.12: The place of origin shall correspond to the semen collection centre of origin of the semen.

Box I.13: The place of destination shall correspond to the semen collection or storage centre or to the holding of semen destination.

Box I.23: The identification of container and seal number shall be indicated.

Box I.31: The donor identity shall correspond to the official identification of the animal.

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

Part II:

Guidance for the completion of the table in point II.3.6:

Abbreviations:

EIA-1 Equine infectious anaemia (EIA) testing first occasion

EIA-2 EIA testing second occasion

EVA-B1 Equine viral arteritis (EVA) testing on blood sample first occasion

EVA-B2 EVA testing on blood sample second occasion

EVA-S1 EVA testing on semen sample first occasion

EVA-S2 EVA testing on semen sample second occasion

CEM-11 Contagious equine metritis (CEM) testing first occasion first sample

CEM-12 CEM testing first occasion second sample taken 7 days after CEM-11

CEM-21 CEM testing second occasion first sample

CEM-22 CEM testing second occasion second sample taken 7 days after CEM-21

Instructions:

For each semen identification in column A in the example below, the test programme (points II.3.5.1, II.3.5.2 and/or II.3.5.3) shall be described in column B and columns C and D shall be completed with the dates required.

The dates when samples were taken for laboratory testing prior to the first collection of the semen described above, as required in points II.3.5.1, II.3.5.2 and II.3.5.3, 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.3.5.2 or II.3.5.3 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.

Identification of semen	Test programme	Start date ([7])		Date of sampling for health tests ([7])				
		Donor residence	Semen collection	EIA II.3.4.1.	EVA II.3.4.2.		CEM II.3.4.3.	
					Blood sample	Semen sample	1. sample	2. sample
A	B	C	D	EIA-1	EVA-B1	EVA-S1	CEM-11	CEM-12
				EIA-2	EVA-B2	EVA-S2	CEM-21	CEM-22

EUROPEAN UNION **Equine semen – Part A**

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

(1) Delete as appropriate

(2) Only approved semen collection centres listed in accordance with Article 11(4) of Directive 92/65/EEC on the Commission website:

http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm

(3) OJ L 268, 14.9.1992, p. 54.

(4) OJ L 192, 23.7.2010, p. 1.

(5) OJ L 165, 30.4.2004, p. 1.

(6) Cross out the programme(s) that do(es) not apply to the consignment.

(7) Insert date in table in point II.3.6 (follow Guidance in Part II of the Notes).

(8) Insert names and concentrations.

— The colour of the stamp and signature must be different from that of the other particulars in the certificate.

Official veterinarian

Name (in capital letters): Qualification and title:

Local veterinary unit: LVU No:

Date: Signature:

Stamp:

PART B Model health certificate IB for trade in the Union in 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 and dispatched after 31 August 2010 from an approved semen collection centre of origin of the semen

EUROPEAN UNION **Intra trade certificate**

Part I: Details of consignment presented

I.1. Consignor Name Address Postal code		I.2. Certificate reference No	I.2.a. Local reference No
		I.3. Central competent authority	
		I.4. Local competent authority	
I.5. Consignee Name Address Postal code		I.6.	
		I.7.	
I.8. Country of origin ISO code	I.9. Region of origin Code	I.10. Country of destination ISO code	I.11. Region of destination Code
I.12. Place of origin Semen Centre ☐ Name Approval number Address Postal code		I.13. Place of destination Semen Centre ☐ Holding ☐ Name Approval number Address Postal code	
I.14.		I.15.	
I.16. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification:		I.17.	
I.18. Description of commodity		I.19. Commodity code (HS code) **05 11 99 85**	
		I.20. Quantity	
I.21. Temperature of products Ambient ☐ Chilled ☐ Frozen ☐		I.22. Number of packages	
I.23. Seal/Container No		I.24. Type of packaging	
I.25. Commodities certified for: Artificial reproduction ☐			
I.26. Transit through third country ☐ Third country ISO code Exit point Code Entry point BIP No		I.27. Transit through Member States ☐ Member State ISO code Member State ISO code Member State ISO code	
I.28. Export ☐ Third country ISO code Exit point Code		I.29.	
I.30.			
I.31. Identification of the commodities Species (Scientific name) Donor identity Date of collection Quantity			

EUROPEAN UNION **Equine semen – Part B**

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

I, the undersigned official veterinarian, hereby certify that:

Part II: Certification

II.1. The semen collection centre ([2]), in which the semen described above was collected, processed and stored for trade is approved and supervised by the competent authority in accordance with Chapter I(I)(1) and Chapter I(II)(1) of Annex D to Directive 92/65/EEC;

II.1.1. during the period commencing 30 days prior to the date of first collection of the semen described above until the date the fresh or chilled semen was dispatched or until the 30 days storage period for frozen semen elapsed, the semen collection centre:

II.1.1.1. was situated on the territory or in the case of regionalisation in a part of the territory ([1]) of a Member State which was not considered to be infected with African horse sickness in accordance with Article 5(2)(a) and(b) of Directive 2009/156/EC ([3]);

II.1.1.2. fulfilled the conditions for a holding laid down in Article 4(5) of Directive 2009/156/EC ([3]);

II.1.1.3. contained only equidae which were free of clinical signs of equine viral arteritis and contagious equine metritis;

II.2. Only equidae satisfying the conditions laid down in Articles 4 and 5 or Articles 12 to 16 of Directive 2009/156/EC ([3]) have been admitted onto the centre.

II.3. The semen described above was collected from donor stallions, which:

II.3.1. have not shown any clinical sign of an infectious or contagious disease at the time of admission onto the centre and on the day the semen was collected;

II.3.2. have been kept for 30 days prior to the date of semen collection in holdings where no equine has shown any clinical sign of equine viral arteritis or contagious equine metritis during that period;

II.3.3. have not been used for natural mating during at least 30 days prior to the date of first semen collection and from the dates of the first sample referred to in points II.3.5.1, II.3.5.2 or II.3.5.3 until the end of the collection period;

II.3.4. have undergone the following tests, which meet at least the requirements of the relevant Chapter of the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals of the OIE, carried out on samples taken in accordance with one of the programmes specified in point II.3.5 in a laboratory recognised by the competent authority:

([1]) *either* [II.3.4.1. an agar-gel immuno-diffusion test (Coggins test) for equine infectious anaemia (EIA) with negative result;]

([1]) *or* [II.3.4.1. an ELISA for equine infectious anaemia (EIA) with negative result;]

and ([1]) *either* [II.3.4.2. a serum neutralisation test for equine viral arteritis (EVA) with negative result at a serum dilution of one in four;]

([1]) *or* [II.3.4.2. a virus isolation test for equine viral arteritis (EVA) carried out with negative result on an aliquot of the entire semen of the donor stallion;]

and II.3.4.3. an agent identification test for contagious equine metritis (CEM) carried out on two occasions on samples taken with an interval of 7 days by isolation of *Taylorella equigenitalis* 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;

EUROPEAN UNION **Equine semen – Part B**

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

II.3.5.	have been subjected with the results specified in II.3.4 in each case to at least one of the test programmes ([4]) detailed in points II.3.5.1, II.3.5.2 and II.3.5.3 as follows:	
II.3.5.1.	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 into direct contact with equidae of lower health status than the donor stallion.	
	The tests described in point II.3.4 have been carried out on samples taken ([5]) 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.3.5.2.	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, and/or other equidae on the collection centre came into direct contact with equidae of lower health status.	
	The tests described in point II.3.4 have been carried out on samples taken ([5]) prior to 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.3.4.1 for equine infectious anaemia was last carried out on a sample of blood taken ([5]) not more than 90 days before the semen described above was collected,	
and	*([1]) either* [one of the tests described in point II.3.4.2 for equine viral arteritis was last carried out on a sample taken ([5]) 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 ([5]) not more than 6 months before the semen described above was collected and a blood sample taken on the same date ([5]) 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.3.4.3 for contagious equine metritis was last carried out on samples taken ([5]) not more than 60 days before the semen described above was collected.	
II.3.5.3.	The tests described in point II.3.4 have been carried out on samples taken ([5]) prior to 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.3.4 were last carried out on samples taken ([5]) not less than 14 days and not more than 90 days after the collection of the semen described above.	

EUROPEAN UNION **Equine semen – Part B**

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

II.3.6. have undergone the testing provided for in point II.3.5 on samples taken on the following dates:

Identification of semen	Test programme	Start date ([5])		Date of sampling for health tests ([5])				
		Donor residence	Semen collection	EIA II.3.4.1.	EVA II.3.4.2.		CEM II.3.4.3.	
					Blood sample	Semen sample	1. sample	2. sample

([1]) *either* [II.4. No antibiotics were added to the semen;]

([1]) *or* [II.4. The following antibiotic or combination of antibiotics was added to produce a concentration in the final diluted semen of not less than ([6]): ..... ;]

II.5. The semen described above was:

II.5.1. collected, processed, stored and transported under conditions which comply with the requirements of Chapters II(I)(1) and III(I) of Annex D to Directive 92/65/EEC;

II.5.2. sent to the place of loading in a sealed container in accordance with point 1.4 of Chapter III(I) of Annex D to Directive 92/65/EEC and bearing the number indicated in Box I.23.

Notes

Part I:

Box I.12: place of origin shall correspond to the semen collection centre of origin of the semen.

Box I.13: place of destination shall correspond to the semen collection or storage centre or to the holding of semen destination.

Box I.23: identification of container and seal number shall be indicated.

Box I.31: donor identity shall correspond to the official identification of the animal.

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

approval number of the centre shall correspond to the approval number of the semen centre indicated in Box I.12 where the semen was collected.

EUROPEAN UNION **Equine semen – Part B**

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

Part II:

Guidance for the completion of Table in II.3.6:

Abbreviations:

EIA-1 Equine infectious anaemia (EIA) testing first occasion

EIA-2 EIA testing second occasion

EVA-B1 Equine viral arteritis (EVA) testing on blood sample first occasion

EVA-B2 EVA testing on blood sample second occasion

EVA-S1 EVA testing on semen sample first occasion

EVA-S2 EVA testing on semen sample second occasion

CEM-11 Contagious equine metritis (CEM) testing first occasion first sample

CEM-12 CEM testing first occasion second sample taken 7 days after CEM-11

CEM-21 CEM testing second occasion first sample

CEM-22 CEM testing second occasion second sample taken 7 days after CEM-21

Instructions:

For each semen identification in column A in the example below, the test programme (II.3.5.1, II.3.5.2 and/or II.3.5.3) must be described in column B and columns C and D must be completed with the dates required.

The dates when samples were taken for laboratory testing prior to the first collection of the semen described above as required in II.3.5.1, II.3.5.2 and II.3.5.3, are entered in the upper line of columns 5 to 9 of the table, this being the boxes marked with EIA-1, EVA-B1 or EVA-S1 and CEM-11 and CEM-12 in the example below.

The dates when samples were taken for repeat laboratory testing as required in accordance with II.3.5.2 or II.3.5.3 are entered in the lower line of columns 5 to 9 in table, this being the boxes EIA-2, EVA-B2 or EVA-S2 and CEM-21 and CEM-22 in the example below.

Identification of semen	Test programme	Start date ([5])		Date of sampling for health tests ([5])				
		Donor residence	Semen collection	EIA II.3.4.1.	EVA II.3.4.2.		CEM II.3.4.3.	
					Blood sample	Semen sample	1. sample	2. sample
A	B	C	D	EIA-1	EVA-B1	EVA-S1	CEM-11	CEM-12
				EIA-2	EVA-B2	EVA-S2	CEM-21	CEM-22

([1]) Delete as appropriate.

([2]) Only approved semen collection centres listed in accordance with Article 11(4) of Council Directive 92/65/EEC on the Commission website:

http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm

([3]) OJ L 192, 23.7.2010, p. 1.

EUROPEAN UNION **Equine semen – Part B**

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

([4]) Cross out the programme(s) that do(es) not apply to the consignment.

([5]) Insert date in table in point II.3.6 (follow Guidance in Part II of the Notes).

([6]) Insert names and concentrations.

— The colour of the stamp and signature must be different from that of the other particulars in the certificate.

Official veterinarian

Name (in capital letters): Qualification and title:

Local veterinary unit: LVU No:

Date: Signature:

Stamp:

PART C Model health certificate IC for trade in the Union in consignments of stocks of semen of animals of the equine species 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

EUROPEAN UNION **Intra trade certificate**

Part I: Details of consignment presented

I.1. Consignor Name Address Postal code		I.2. Certificate reference No	I.2.a. Local reference No
		I.3. Central competent authority	
		I.4. Local competent authority	
I.5. Consignee Name Address Postal code		I.6.	
		I.7.	
I.8. Country of origin ISO code	I.9. Region of origin Code	I.10. Country of destination ISO code	I.11. Region of destination Code
I.12. Place of origin Semen Centre ☐ Name Address Postal code	Approval number	I.13. Place of destination Semen Centre ☐ Holding ☐ Name Address Postal code	Approval number
I.14.		I.15.	
I.16. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification:		I.17.	
I.18. Description of commodity		I.19. Commodity code (HS code) **05 11 99 85**	
		I.20. Quantity	
I.21. Temperature of products Frozen ☐		I.22. Number of packages	
I.23. Seal/Container No		I.24. Type of packaging	
I.25. Commodities certified for: Artificial reproduction ☐			
I.26. Transit through third country ☐ Third country ISO code Exit point Code Entry point BIP No		I.27. Transit through Member States ☐ Member State ISO code Member State ISO code Member State ISO code	
I.28. Export ☐ Third country ISO code Exit point Code		I.29.	
I.30.			
I.31. Identification of the commodities Species (Scientific name) Donor identity Date of collection Quantity			

EUROPEAN UNION **Equine semen – Part C**

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

I, the undersigned official veterinarian, hereby certify that:

Part II: Certification

II.1. The semen collection centre (2), in which the semen described above was collected, processed and stored for trade:

II.1.1. is approved and supervised by the competent authority according to the conditions of Chapter I of Annex D to Directive 92/65/EEC;

II.1.2. is situated on the territory or in the case of regionalisation in a part of the territory (1) of a Member State which was on the day semen was collected until the date the semen was dispatched as fresh/chilled (1) semen or until the 30 days mandatory storage period for frozen semen elapsed (1) not considered to be infected with African horse sickness in accordance with Article 5(2)(a) and(b) of Directive 2009/156/EC (3);

II.1.3. fulfilled during the period commencing 30 days prior to the date of semen collection until the date the semen was dispatched as fresh/chilled (1) semen or until the 30 days mandatory storage period for frozen semen elapsed (1), the conditions of Article 4 of Directive 2009/156/EC;

II.1.4. contained during the period commencing 30 days prior to the date of semen collection until the date the semen was dispatched as fresh/chilled (1) semen or until the 30 days mandatory storage period for frozen semen elapsed (1) only equidae which were free of clinical signs of equine viral arteritis and contagious equine metritis;

II.2. All equidae have been admitted onto the centre under the provisions of Article 4 and 5 of Directive 2009/156/EC (3);

II.3. The semen described above was collected from donor stallions, which:

II.3.1. on the day the semen was collected have not shown clinical signs of an infectious or contagious disease,

II.3.2. during at least 30 days prior to collection of the semen have not been used for natural service,

II.3.3. during the last 30 days prior to collection of the semen have been kept on holdings where no equidae showed clinical signs of equine viral arteritis,

II.3.4. during the last 60 days prior to collection of the semen have been kept on holdings where no equidae showed clinical signs of contagious equine metritis,

II.3.5. to the best of my knowledge and as far as I could ascertain, have not been in contact with equidae suffering from an infectious or contagious disease during 15 days immediately preceding collection of the semen,

II.3.6. have undergone the following animal health tests, carried out in a laboratory recognised by the competent authority, in accordance with a test programme as specified in point II.3.7.

[II.3.6.1. an agar-gel immuno-diffusion test (Coggins test) for equine infectious anaemia with negative result;]

and (1) *either* [II.3.6.2. a serum neutralisation test for equine viral arteritis with negative result at a serum dilution of one in four; and]

(1) *or* [II.3.6.2. a virus isolation test for equine viral arteritis carried out with negative result on an aliquot of the entire semen of the donor stallion;]

EUROPEAN UNION **Equine semen – Part C**

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

and II.3.6.3. an agent identification test for contagious equine metritis carried out on two occasions on samples collected from the donor stallion with an interval of 7 days by isolation of *Taylorella equigenitalis* 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.3.7. have been subject to the one of the following test programmes ([4]):

II.3.7.1. The donor stallion was continuously resident on the collection centre for at least 30 days prior to the semen collection, and during the collection period, and no equidae on the collection centre came during that time into direct contact with equidae of lower health status than the donor stallions.

The tests described in point II.3.6 have been carried out on samples taken on ..... ([5]) and in the case of contagious equine metritis on a second sample taken on ..... ([5]), being at least 14 days after the commencement of the above residence period and at least at the beginning of the breeding season;

II.3.7.2. The donor stallion was not continuously resident on the collection centre or other equidae on the collection centre came into contact with equidae of lower health status than the donor stallion.

The tests described in point II.3.6 have been carried out on samples taken on ..... ([5]) and in the case of contagious equine metritis on a second sample taken on ..... ([5]), being within the 14 days period before the first semen collection and at least at the beginning of the breeding season,

and the test described in point II.3.6.1 for equine infectious anaemia was last carried out on a sample of blood taken on ..... ([5]), being not more than 120 days before the semen described above was collected;

and ([1]) *either* [one of the tests described in point II.3.6.2 for equine viral arteritis was last carried out on a sample collected on ..... ([5]), being not more than 30 days before the semen described above was collected,]

([1]) *or* [the non-shedder state of the seropositive stallion for equine viral arteritis was confirmed by a virus isolation test which was carried out on an aliquot of the entire semen of the donor stallion collected on ..... ([5]), being not more than 1 year before the semen described above was collected;]

II.3.7.3. The tests described in point II.3.6 have been carried out during the 30 days mandatory storage period of frozen semen and not less than 14 days after the collection of the semen on samples taken on ..... ([5]) and in the case of contagious equine metritis on a second sample taken on..... ([5]);

II.4. The semen described above was collected, processed, stored and transported under conditions which comply with the requirements of Chapters II and III of Annex D to Directive 92/65/EEC.

Notes

Part I:

Box I.12: place of origin shall correspond to the semen collection centre of origin of the semen.

Box I.13: place of destination shall correspond to the semen collection or storage centre or to the holding of semen destination.

Box I.23: identification of container and seal number shall be indicated.

EUROPEAN UNION **Equine semen – Part C**

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

Box I.31.: donor identity shall correspond to the official identification of the animal.

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

approval number of the centre shall correspond to the approval number of the semen centre indicated in Box I.12 where the semen was collected.

Part II:

(1) Delete as appropriate.

(2) Only approved semen collection centres listed in accordance with Article 11(4) of Directive 92/65/EEC on the Commission website:

http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm

(3) OJ L 192, 23.7.2010, p. 1.

(4) Cross out the programme(s) that do(es) not apply to the consignment.

(5) Insert date.

— The colour of the stamp and signature must be different from that of the other particulars in the certificate.

Official veterinarian or official inspector

Name (in capital letters): Qualification and title:

Local veterinary unit: LVU No:

Date: Signature:

Stamp:

PART D Model health certificate ID for trade in the Union in consignments of semen of animals of the equine species collected, processed and stored in accordance with Directive 92/65/EEC after 30 September 2014 and in 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

EUROPEAN UNION **Intra trade certificate**

Part I: Details of consignment presented

I.1. Consignor Name Address Postal code	I.2. Certificate reference No	I.2.a. Local reference No
	I.3. Central competent authority	
	I.4. Local competent authority	
I.5. Consignee Name Address Postal code	I.6. No(s) of related original certificates	No(s) of accompanying documents
	1.7.	
I.8. Country of origin ISO code I.9. Region of origin Code	I.10. Country of destination ISO code	I.11. Region of destination Code
I.12. Place of origin Semen Centre ☐ Name Approval number Address Postal code	I.13. Place of destination Semen Centre ☐ Holding ☐ Name Approval number Address Postal code	
I.14.	I.15.	
I.16. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification:	I.17.	
I.18. Description of commodity	I.19. Commodity code (HS code) **05 11 99 85**	
	I.20. Quantity	
I.21. Temperature of products Frozen ☐	I.22. Number of packages	
I.23. Seal/Container No	I.24. Type of packaging	
I.25. Commodities certified for: Artificial reproduction ☐		
I.26. Transit through third country ☐ Third country ISO code Exit point Code Entry point BIP No	I.27. Transit through Member States ☐ Member State ISO code Member State ISO code Member State ISO code	
I.28. Export ☐ Third country ISO code Exit point Code	I.29.	
I.30.		
I.31. Identification of the commodities Species (Scientific name) Donor identity Date of collection Quantity		

EUROPEAN UNION **Equine semen – Part D**

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

Part II: Certification

I, the undersigned official veterinarian, hereby certify that the semen described above

(1) either [II.1. was collected, processed and stored for a minimum period of 30 days immediately following collection in an approved semen collection centre (2) situated in the Member State of origin of the semen and operated and supervised in accordance with Chapters I(I)(1) and I(II)(1) of Annex D to Directive 92/65/EEC (3), and from where the semen was moved to the semen storage centre detailed in Box I.12 situated in the same Member State of origin of the semen under animal health and veterinary certification conditions at least as strict as those provided for in

(1) either [Part A of Annex I to Decision 2010/470/EU;]

(1) or [Part B of Annex I to Decision 2010/470/EU;]

(1) or [Part C of Annex I to Decision 2010/470/EU;]

(1) or [Decision 95/307/EC;]]

(1) or [II.1. was collected, processed and stored for a minimum period of 30 days immediately following collection in an approved semen collection centre (2) situated in the Union and operated and supervised in accordance with Chapters I(I)(1) and I(II)(1) of Annex D to Directive 92/65/EEC, and was moved to the semen storage centre detailed in Box I.12 in accordance with:

(1) either [Part A of Annex I to Decision 2010/470/EU;]

(1) or [Part B of Annex I to Decision 2010/470/EU;]

(1) or [Part C of Annex I to Decision 2010/470/EU;]

(1) or [Part D of Annex I to Decision 2010/470/EU;]

(1) or [Decision 95/307/EC;]]

(1) or [II.1. was collected, processed and stored in an approved semen collection centre (2) situated in a third country or part(s) thereof listed in columns 2 and 4 of Annex I to Decision 2004/211/EC which is operated and supervised in accordance with Chapters I(I)(1) and I(II)(1) of Annex D to Directive 92/65/EEC, and was imported into the Union in accordance with Article 4 of Decision 2004/211/EC in accordance with:

(1) either [Section A of Part 2 of Annex II to Decision 2010/471/EU;]

(1) or [Section B of Part 2 of Annex II to Decision 2010/471/EU;]

(1) or [Section C of Part 2 of Annex II to Decision 2010/471/EU;]

(1) or [Section D of Part 2 of Annex II to Decision 2010/471/EU;]

(1) or [Decision 96/539/EC;]]

II.2. was stored in the approved semen storage centre (2) indicated in Box I.12, which is operated and supervised in accordance with Chapters I(I)(2) and I(II)(2) of Annex D to Directive 92/65/EEC;

II.3. was sent to the place of loading in a sealed container in accordance with point 1.4 of Chapter III(I) of Annex D to Directive 92/65/EEC and bearing the number indicated in Box I.23.

Notes

Part I:

Box I.6: No(s) of related original certificates or accompanying documents shall correspond to the serial number of the individual official document(s) or health certificate(s) [either INTRA or CVED] that accompanied the semen described above from the approved semen collection centre of its origin to the described above semen storage centre. The original(s) of this/these document(s) or certificate(s) or the officially endorsed copy/copies thereof must be attached to this certificate.

EUROPEAN UNION **Equine semen – Part D**

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

Box I.12: The place of origin shall correspond to the semen storage centre of dispatch of the semen.

Box I.13: The place of destination shall correspond to the semen collection or storage centre or to the holding of destination of the semen.

Box I.23: The identification of container and seal number shall be indicated.

Box I.31: 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 appropriate

(2) Only approved semen collection or storage centres listed in accordance with Article 11(4) or Article 17(3)(b) of Directive 92/65/EEC on the Commission websites:
http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm;
http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm

(3) OJ L 268, 14.9.1992, p. 54

— The colour of the stamp and signature must be different from that of the other particulars in the certificate.

Official veterinarian or official inspector

Name (in capital letters): Qualification and title:

Local veterinary unit: LVU No:

Date: Signature:

Stamp:

ANNEX II

Model health certificates for trade in the union in consignments of ova and embryos of animals of the equine species

PART A

Model health certificate IIA for trade in the Union in consignments of ova and embryos of animals of the equine species 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

EUROPEAN UNION **Intra trade certificate**

Part I: Details of consignment presented

I.1. Consignor
Name
Address
Postal code

I.2. Certificate reference No | I.2.a. Local reference No

I.3. Central competent authority

I.4. Local competent authority

I.5. Consignee
Name
Address
Postal code

I.6.

I.7.

I.8. Country of origin | ISO code | I.9. Region of origin | Code

I.10. Country of destination | ISO code | I.11. Region of destination | Code

I.12. Place of origin
Embryo team ☐
Name | Approval number
Address
Postal code

I.13. Place of destination
Holding ☐ Embryo team ☐
Name | Approval number
Address
Postal code

I.14.

I.15.

I.16. Means of transport
Aeroplane ☐ Ship ☐ Railway wagon ☐
Road vehicle ☐ Other ☐
Identification:

I.17.

I.18. Description of commodity

I.19. Commodity code (HS code)
05 11 99 85

I.20. Quantity

I.21. Temperature of products
Ambient ☐ Chilled ☐ Frozen ☐

I.22. Number of packages

I.23. Seal/Container No

I.24. Type of packaging

I.25. Commodities certified for:
Artificial reproduction ☐

I.26. Transit through third country ☐
Third country | ISO code
Exit point | Code
Entry point | BIP No

I.27. Transit through Member States ☐
Member State | ISO code
Member State | ISO code
Member State | ISO code

I.28. Export ☐
Third country | ISO code
Exit point | Code

I.29.

I.30.

I.31. Identification of the commodities

Species (Scientific name)	Category	Donor identity	Date of collection	Quantity

EUROPEAN UNION **Equine ova and embryos – Part A**

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

Part II: Certification

I, the undersigned official veterinarian, hereby certify that:

(1) *either*	[II.1.	the *in vivo* derived embryos/*in vivo* derived ova (1) described above were collected, processed and stored by an embryo collection team (2) approved and supervised in accordance with Chapter I(III)(1) of Annex D to Directive 92/65/EEC (3);]
(1) *or*	[II.1.	the *in vivo* produced embryos/micromanipulated embryos (1) described above were produced, processed and stored by an embryo production team (2), approved and supervised in accordance with Chapter I(III)(1) and (2) of Annex D to Directive 92/65/EEC;]
(1) *either*	[II.2.	the *in vivo* derived embryos described above meet the requirements of Chapter III(II)(1) of Annex D to Directive 92/65/EEC;]
(1) *or*	[II.2.	the *in vivo* derived ova described above meet the requirements of Chapter III(II)(2) of Annex D to Directive 92/65/EEC;]
(1) *or*	[II.2.	the *in vivo* produced embryos described above meet the requirements of Chapter III(II)(3) of Annex D to Directive 92/65/EEC;]
(1) *or*	[II.2.	the micromanipulated embryos described above meet the requirements of Chapter III(II)(4) of Annex D to Directive 92/65/EEC;]
	II.3.	the ova or embryos described above come from donor mares which:
	II.3.1.	come from holdings fulfilling the conditions laid down in Article 4(5) of Directive 2009/156/EC (4) onto which only equidae satisfying the conditions laid down in Articles 4 and 5 or Articles 12 to 16 of Directive 2009/156/EC were admitted;
	II.3.2.	meet the requirements of Chapter IV(4) of Annex D to Directive 92/65/EEC;
	II.3.3.	were not used for natural breeding during a period of at least 30 days prior to the date of collection of the ova or embryos and between the date of the first sample referred to in points II.3.4.1 and II.3.4.2 and the date of the collection of the ova or embryos;
	II.3.4.	underwent the tests, which meet at least the requirements of the relevant Chapter of the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals of the OIE, carried out in a laboratory which is recognised by the competent authority and has the tests referred to hereinafter included in its accreditation in accordance with Article 12 of Regulation (EC) No 882/2004 (5), as follows:
	II.3.4.1.	for equine infectious anaemia (EIA), an agar-gel immuno-diffusion test (AGID or Coggins test) or an enzyme-linked immunosorbent assay (ELISA) with a negative result carried out on a blood samples taken on ..... (6), being not less than 14 days following the date of commencement of the period referred to in point II.3.3, and the test was last carried out on a sample of blood taken on ..... (6); being not more than 90 days prior to the date of the collection of the ova or embryos intended for trade;
	II.3.4.2.	for contagious equine metritis (CEM), an agent identification test carried out with a negative result on at least two specimens (swabs) taken during the period referred to in point II.3.3 from at least the mucosal surfaces of the clitoral fossa and the clitoral sinuses of the donor mare;

EUROPEAN UNION **Equine ova and embryos – Part A**

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

(1) *either* [II.3.4.2.1. on two occasions with an interval of not less than 7 days on ..... (6) and on ..... (6), in the case of isolation of *Taylorella equigenitalis* after cultivation under microaerophilic conditions for a period of at least 7 days, set up within 24 hours after taking the specimens from the donor animal, or 48 hours where the specimens are kept cool during transport;]

(1) *and/or* [II.3.4.2.2. on one occasion on ..... (6), in the case of the detection of genome of *Taylorella equigenitalis* by a polymerase chain reaction (PCR) or real-time PCR test, carried out within 48 hours after taking the specimens from the donor animal.]

The samples referred to in points II.3.4.2.1 and II.3.4.2.2 were in no case taken earlier than 7 days (systemic treatment) or 21 days (local treatment) 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;

(1) *either* [II.4. the embryos described above were conceived as a result of artificial insemination of the donor mares with semen which was collected, processed, stored and transported under conditions which comply with the requirements of Chapters I(I), II(I) and III(I) of Annex D to Directive 92/65/EEC;]

(1) *or* [II.4. the embryos described above were conceived as a result of *in vitro* fertilisation of ova complying with the conditions set out in point 2 of Chapter III(II) of Annex D to Directive 92/65/EEC with semen which was collected, processed, stored and transported under conditions which comply with the requirements of Chapters I(I), II(I) and III(I) of Annex D to Directive 92/65/EEC;]

(1) *or* [II.4. the ova have not been in contact with semen of the equine species;]

II.5. the ova or embryos described above were sent to the place of loading in a sealed container in accordance with point 6 of Chapter III(II) of Annex D to Directive 92/65/EEC and bearing the number detailed in Box I.23.

Notes

Part I:

Box I.12: The place of origin shall correspond to the embryo collection team or embryo production team of ova/embryos collection/production.

Box I.13: The place of destination shall correspond to the embryo collection team, embryo production team or to the holding of ova/embryos destination.

Box I.23: The identification of container and seal number shall be indicated.

Box I.31: the category: specify if: *in vivo* derived embryos, *in vivo* derived ova, *in vivo* produced embryos or micro-manipulated embryos.

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 appropriate.

(2) Only approved embryo collection or production teams listed in accordance with Article 11(4) of Directive 92/65/EEC on the Commission website:

http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm

EUROPEAN UNION **Equine ova and embryos – Part A**

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

([3]) OJ L 268, 14.9.1992, p. 54.

([4]) OJ L 192, 23.7.2010, p. 1.

([5]) OJ L 165, 30.4.2004, p. 1.

([6]) Insert date.

— The colour of the stamp and signature must be different from that of the other particulars in the certificate.

Official veterinarian or official inspector

Name (in capital letters): Qualification and title:

Local veterinary unit: LVU No:

Date: Signature:

Stamp:

PART B Model health certificate IIB for trade in the Union in consignments of stocks of ova and embryos 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 and dispatched after 31 August 2010 by an approved embryo collection or production team of origin of the ova or embryos

EUROPEAN UNION **Intra trade certificate**

Part I: Details of consignment presented

I.1. Consignor Name Address Postal code		I.2. Certificate reference No	I.2.a. Local reference No
		I.3. Central competent authority	
		I.4. Local competent authority	
I.5. Consignee Name Address Postal code		I.6.	
		I.7.	
I.8. Country of origin	ISO code	I.9. Region of origin	Code
I.10. Country of destination	ISO code	I.11. Region of destination	Code
I.12. Place of origin Embryo team ☐ Name Approval number Address Postal code		I.13. Place of destination Holding ☐ Embryo team ☐ Name Approval number Address Postal code	
I.14.		I.15.	
I.16. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification:		I.17.	
I.18. Description of commodity		I.19. Commodity code (HS code) **05 11 99 85**	
		I.20. Quantity	
I.21. Temperature of products Ambient ☐ Chilled ☐ Frozen ☐		I.22. Number of packages	
I.23. Seal/Container No		I.24. Type of packaging	
I.25. Commodities certified for: Artificial reproduction ☐			
I.26. Transit through third country ☐ Third country ISO code Exit point Code Entry point BIP No		I.27. Transit through Member States ☐ Member State ISO code Member State ISO code Member State ISO code	
I.28. Export ☐ Third country ISO code Exit point Code		I.29.	
I.30.			

I.31. Identification of the commodities

Species (Scientific name)	Category	Donor identity	Date of collection	Quantity

EUROPEAN UNION **Equine ova and embryos – Part B**

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

I, the undersigned official veterinarian, hereby certify that:

Part II: Certification

(1) *either* [II.1. the *in vivo* derived embryos/*in vivo* derived ova (1) described above were collected, processed and stored by an embryo collection team (2) approved and supervised in accordance with Chapter I(III)(1) of Annex D to Directive 92/65/EEC;]

(1) *or* [II.1. the *in vitro* produced embryos/micromanipulated embryos (1) described above were produced, processed and stored by an embryo production team (2), approved and supervised in accordance with Chapter I(III)(1) and (2) of Annex D to Directive 92/65/EEC;]

(1) *either* [II.2. the *in vivo* derived embryos described above meet the requirements of Chapter III(II)(1) of Annex D to Directive 92/65/EEC;]

(1) *or* [II.2. the *in vivo* derived ova described above meet the requirements of Chapter III(II)(2) of Annex D to Directive 92/65/EEC;]

(1) *or* [II.2. the *in vitro* produced embryos described above meet the requirements of Chapter III(II)(3) of Annex D to Directive 92/65/EEC;]

(1) *or* [II.2. the micromanipulated embryos described above meet the requirements of Chapter III(II)(4) of Annex D to Directive 92/65/EEC;]

II.3. the ova or embryos described above come from donor mares which:

II.3.1. coming from holdings fulfilling the conditions laid down in Article 4(5) of Directive 2009/156/EC (4) onto which only equidae satisfying the conditions laid down in Articles 4 and 5 or Articles 12 to 16 of Directive 2009/156/EC have been admitted;

II.3.2. meet the additional requirements of Chapter IV(4) of Annex D to Directive 92/65/EEC;

II.3.3. have not been used for natural breeding during at least 30 days prior to the date of collection of ova or embryos and between the date of the first sample referred to in points II.3.4 and II.3.5 and the date of the collection of ova and embryos;

II.3.4. have been subjected with negative result to an agar-gel immuno-diffusion test (Coggins test) or an ELISA for equine infectious anaemia carried out on a blood samples taken on ..... (3), being during the past 30 days prior to the date of the first collection of ova or embryos and the last test was carried out on a sample of blood taken on ..... (3); being not more than 90 days before the ova and embryos were collected;

II.3.5. have been subjected to an agent identification test for contagious equine metritis by isolation of *Taylorella equigenitalis* after a cultivation of 7 to 14 days carried out with negative results in each case on samples taken during the past 30 days prior to the date of the first collection of ova or embryos from mucosal surfaces of the clitoral fossa and clitoral sinuses on two consecutives oestrus periods on ..... (3) and on ..... (3), and on an additional culture specimen taken during one of the oestrus periods from the endometrial cervix on ..... (3);

(1) *either* [II.4. the embryos described above were conceived as a result of artificial insemination of the donor mares with semen which was collected, processed, stored and transported under conditions which comply with the requirements of Chapters I(I), II(I) and III(I) of Annex D to Directive 92/65/EEC;]

EUROPEAN UNION **Equine ova and embryos – Part B**

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

([1]) *or* [II.4. the embryos described above were conceived as a result of *in vitro* fertilisation of ova complying with the conditions in point 2 of Chapter III(II) of Annex D to Directive 92/65/EEC with semen which was collected, processed, stored and transported under conditions which comply with the requirements of Chapters I(I), II(I) and III(I) of Annex D to Directive 92/65/EEC;]

([1]) *or* [II.4. the ova have not been in contact with semen of the equine species;]

II.5. the ova or embryos described above were sent to the place of loading in a sealed container in accordance with point 6 of Chapter III(II) of Annex D to Directive 92/65/EEC and bearing the number detailed in Box I.23.

Notes

Part I:

Box I.12: place of origin shall correspond to the embryo collection team or embryo production team of ova/embryos collection/production.

Box I.13: place of destination shall correspond to the embryo collection team, embryo production team or to the holding of ova/embryos destination.

Box I.23: identification of container and seal number shall be indicated.

Box I.31: category: specify if: *in vivo* derived embryos, *in vivo* derived ova, *in vitro* produced embryos or micro-manipulated embryos.

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

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

approval number of the team shall correspond to the embryo collection team or embryo production team of ova/embryos collection/production.

Part II:

([1]) Delete as appropriate.

([2]) Only approved embryo collection or production teams listed in accordance with Article 11(4) of Directive 92/65/EEC on the Commission website:

http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm

([3]) Insert date.

([4]) OJ L 192, 23.7.2010, p. 1.

— The colour of the stamp and signature must be different from that of the other particulars in the certificate.

Official veterinarian or official inspector

Name (in capital letters): Qualification and title:

Local veterinary unit: LVU No:

Date: Signature:

Stamp:

PART C Model health certificate IIC for trade in the Union in consignments of stocks of ova and embryos of animals of the equine species collected, processed and stored in accordance with Directive 92/65/EEC before 1 September 2010 and dispatched after 31 August 2010 by an approved embryo collection team of origin of the ova or embryos**]**

EUROPEAN UNION **Intra trade certificate**

Part I: Details of consignment presented

I.1. Consignor
Name
Address
Postal code

I.2. Certificate reference No | I.2.a. Local reference No

I.3. Central competent authority

I.4. Local competent authority

I.5. Consignee
Name
Address
Postal code

I.6.

I.7.

I.8. Country of origin | ISO code | I.9. Region of origin | Code

I.10. Country of destination | ISO code | I.11. Region of destination | Code

I.12. Place of origin
Embryo team ☐
Name | Approval number
Address
Postal code

I.13. Place of destination
Holding ☐ Embryo team ☐
Name | Approval number
Address
Postal code

I.14.

I.15.

I.16. Means of transport
Aeroplane ☐ Ship ☐ Railway wagon ☐
Road vehicle ☐ Other ☐
Identification:

I.17.

I.18. Description of commodity

I.19. Commodity code (HS code)
05 11 99 85

I.20. Quantity

I.21. Temperature of products
Frozen ☐

I.22. Number of packages

I.23. Seal/Container No

I.24. Type of packaging

I.25. Commodities certified for:
Artificial reproduction ☐

I.26. Transit through third country ☐
Third country | ISO code
Exit point | Code
Entry point | BIP No

I.27. Transit through Member States ☐
Member State | ISO code
Member State | ISO code
Member State | ISO code

I.28. Export ☐
Third country | ISO code
Exit point | Code

I.29.

I.30.

I.31. Identification of the commodities

Species (Scientific name)	Category	Donor identity	Date of collection	Quantity

EUROPEAN UNION **Equine ova and embryos – Part C**

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

Part II: Certification

I, the undersigned official veterinarian, hereby certify that:

II.1. Ova/embryos (1) described above were collected by a collection team (2) approved by the competent authority and processed in an appropriate laboratory;

II.2. Ova/embryos (1) were collected from donor mares which:

II.2.1. on the day of collection have been located in premises situated on the territory or in the case of regionalisation in a part of the territory of a Member State which is not considered to be infected with African horse sickness in accordance with Article 5(2)(a) and (b) of Directive 2009/156/EC (3),

II.2.2. have been located in holdings under veterinary supervision which on the day of collection fulfilled the conditions of Article 4 of Directive 2009/156/EC,

II.2.3. have been kept prior to the collection in holdings free from clinical signs of contagious equine metritis for 60 days,

II.2.4. have not been used for natural breeding during the period of 30 days prior to the collection of ova/embryos (1),

II.2.5. to the best of my knowledge and as far as I could ascertain, have not been in contact with equidae suffering from an infectious or contagious disease during the 15 days immediately preceding the collection of ova/embryos (1),

II.2.6. have on the day of collection not shown clinical signs of an infectious or contagious disease;

II.3. Ova/embryos (1) were collected, processed, stored and transported under conditions which comply with the requirements of Annex D of Directive 92/65/EEC;

II.4. The semen used for the artificial insemination of the donor mares complies with the requirements of Directive 92/65/EEC (4)(1);

II.5. The ova used for the *in vitro* production of embryos comply with the requirements of Directive 92/65/EEC (1).

Notes

Part I:

Box I.12: place of origin shall correspond to the embryo collection team of ova/embryos collection.

Box I.13: place of destination shall correspond to the embryo collection team, embryo production team or to the holding of ova/embryos destination.

Box I.23: identification of container and seal number shall be indicated.

Box I.31: category: specify if: *in vivo* derived embryos, *in vivo* derived ova, *in vitro* produced embryos or micromanipulated embryos.

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

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

approval number of the team shall correspond to the embryo collection team of ova/embryos collection.

Part II:

(1) Delete as appropriate.

(2) Only approved embryo collection teams listed in accordance with Article 11(4) of Directive 92/65/EEC on the Commission website:

http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm

EUROPEAN UNION **Equine ova and embryos – Part C**

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

([3]) OJ L 192, 23.7.2010, p. 1.

([4]) Does not apply to ova.

— The colour of the stamp and signature must be different from that of the other particulars in the certificate.

Official veterinarian or official inspector

Name (in capital letters): Qualification and title:

Local veterinary unit: LVU No:

Date: Signature:

Stamp:'

ANNEX III

Model health certificates for trade in consigments of semen of animals of the ovine and caprine species

[[F2]PART A

Model health certificate IIIA for trade within the Union in consignments of semen of animals of the ovine and caprine species collected in accordance with Council Directive 92/65/EEC after 31 August 2010 and dispatched from an approved semen collection centre of origin of the semen**]**

EUROPEAN UNION **Intra trade certificate**

Part I: Details of consignment presented

I.1. Consignor Name Address Postal code		I.2. Certificate reference No	I.2.a. Local reference No
		I.3. Central competent authority	
		I.4. Local competent authority	
I.5. Consignee Name Address Postal code		I.6.	
		I.7.	
I.8. Country of origin	ISO code	I.9. Region of origin	Code
I.10. Country of destination	ISO code	I.11. Region of destination	Code
I.12. Place of origin Semen centre ☐ Name Approval number Address Postal code		I.13. Place of destination Semen centre ☐ Holding ☐ Name Approval number Address Postal code	
I.14.		I.15.	
I.16. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification:		I.17.	
I.18. Description of commodity		I.19. Commodity code (CN code) **05 11 99 85**	
			I.20. Quantity

I.21. Temperature of products Ambient ☐ Chilled ☐ Frozen ☐	I.22. Number of packages
I.23. Seal/Container No	I.24. Type of packaging
I.25. Commodities certified for: Artificial reproduction ☐	
I.26. Transit through third country ☐ Third country ISO code Exit point Code Entry point BIP No	I.27. Transit through Member States ☐ Member State ISO code Member State ISO code Member State ISO code
I.28. Export ☐ Third country ISO code Exit point Code	I.29.
I.30.	

I.31. Identification of the commodities

Species (Scientific name)	Breed	Donor identity	Date of collection	Approval number of the centre	Quantity

European Union **Ovine and caprine semen — Part A**

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

Part II: Certification

I, the undersigned official veterinarian, hereby certify that:

II.1. The semen described above:

II.1.1. was collected, processed and stored in a semen collection centre ([2]) approved and supervised by the competent authority in accordance with Chapter I(I)(1) and Chapter I(II)(1) of Annex D to Directive 92/65/EEC;

II.1.2. comes from donor animals which meet the requirements of Chapter II(II) of Annex D to Directive 92/65/EEC;

II.1.3. was collected, processed, stored and transported under conditions which comply with the requirements of Chapters II(II) and III(I) of Annex D to Directive 92/65/EEC;

([1]) *either* [II.1.4. was collected from animals which have been kept continuously since birth on a holding or holdings recognised as having a negligible or controlled risk of classical scrapie according to point 1 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001, except during the period when they were kept at a semen collection centre that complied during that period with the conditions set out in the four indents of point 1.3(c)(iv) of that Section;]

([1]) *or* [II.1.4. was collected from animals which have been kept continuously for the last three years before the collection on a holding or holdings which has/have complied for the last three years before the collection with the requirements laid down in points 1.3(a) to (f) of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001, except during the period when they were kept at a semen collection centre that complied during that period with the conditions set out in the four indents of point 1.3(c)(iv) of that Section;]

([1]) *or* [II.1.4. was collected from animals which have been kept continuously since birth in a Member State or zone of a Member State listed in point 2.3 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 as having a negligible risk status for classical scrapie;]

([1]) *or* [II.1.4. was collected from ovine animals of the ARR/ARR prion protein genotype;]

II.1.5. was sent to the place of loading in a sealed container in accordance with point 1.4 of Chapter III(I) of Annex D to Directive 92/65/EEC and bearing the number detailed in Box I.23.

([1]) *either* [II.2. No antibiotics or no mixture of antibiotics were added to the semen.]

([1]) *or* [II.2. The following antibiotic or combination of antibiotics was added to produce a concentration in the final diluted semen of not less than ([3]):

.....]

Notes

Part I:

Box I.12: *Place of origin* shall correspond to the semen collection centre of origin of the semen.

Box I.13: *Place of destination* shall correspond to the semen collection or storage centre or to the holding of semen destination.

Box I.23: Identification of container and seal number shall be indicated.

Box I.31: *Donor identity* shall correspond to the official identification of the animal.

Date of collection shall be indicated in the following format: dd/mm/yyyy.

Approval number of the centre shall correspond to the approval number of the semen centre indicated in Box I.12 where the semen was collected.

European Union **Ovine and caprine semen — Part A**

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

Part II:

(1) Delete as appropriate.

(2) Only approved semen collection centres listed in accordance with Article 11(4) of Directive 92/65/EEC on the Commission website:

http://ec.europa.eu/food/animals/live_animals/approved-establishments/index_en.htm].

(3) Insert names and concentrations.

— The colour of the stamp and signature must be different from that of the other particulars in the certificate.

Official veterinarian or official inspector

Name (in capital letters): Qualification and title:

Local veterinary unit: LVU No:

Date: Signature:

Stamp:

PART B Model health certificate IIIB for trade within the Union in consignments of stocks of semen of animals of the ovine and caprine species collected, processed and stored in accordance with Council 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

EUROPEAN UNION **Intra trade certificate**

Part I: Details of consignment presented

I.1. Consignor
Name
Address
Postal code

I.2. Certificate reference No | I.2.a. Local reference No

I.3. Central competent authority

I.4. Local competent authority

I.5. Consignee
Name
Address
Postal code

I.6.

I.7.

I.8. Country of origin | ISO code | I.9. Region of origin | Code

I.10. Country of destination | ISO code | I.11. Region of destination | Code

I.12. Place of origin
Semen centre ☐
Name | Approval number
Address
Postal code

I.13. Place of destination
Semen centre ☐ Holding ☐
Name | Approval number
Address
Postal code

I.14.

I.15.

I.16. Means of transport
Aeroplane ☐ Ship ☐ Railway wagon ☐
Road vehicle ☐ Other ☐
Identification

I.17.

I.18. Description of commodity

I.19. Commodity code (HS code)
05 11 99 85

I.20. Quantity

I.21. Temperature of products
Ambient ☐ Chilled ☐ Frozen ☐

I.22. Number of packages

I.23. Seal/Container No

I.24. Type of packaging

I.25. Commodities certified for:
Artificial reproduction ☐

I.26. Transit through third country ☐
Third country | ISO code
Exit point | Code
Entry point | BIP No

I.27. Transit through Member States ☐
Member State | ISO code
Member State | ISO code
Member State | ISO code

I.28. Export ☐
Third country | ISO code
Exit point | Code

I.29.

I.30.

I.31. Identification of the commodities

Species (Scientific name)	Breed	Donor identity	Date of collection	Approval number of the centre	Quantity

EUROPEAN UNION **Ovine and caprine semen — Part B**

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

Part II: Certification

I, the undersigned official veterinarian, hereby certify that the semen described above:

II.1. was collected, processed and stored in a semen collection centre (2) approved and supervised by the competent authority in accordance with Chapter I(I) and Chapter I(II) of Annex D to Directive 92/65/EEC;

II.2. comes from the donor animals which meet the requirements of Chapter II(II) of Annex D to Directive 92/65/EEC;

II.3. was collected, processed, stored and transported under conditions which comply with the requirements of Chapters II(II) and III of Annex D to Directive 92/65/EEC;

(1) either [II.4. meets the requirements of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001;]

(1) or [II.4. meets the requirements of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and is destined for a Member State which benefits, for all or part of its territory, from the provisions laid down in point (b) or (c) of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and the donor animals comply regarding scrapie with the guarantees provided for by the programmes referred to in that point and with the guarantees (3) requested by the Member State of destination.]

Notes

Part I:

Box I.12: Place of origin shall correspond to the semen collection centre of origin of the semen.

Box I.13: Place of destination shall correspond to the semen collection or storage centre or to the holding of semen destination.

Box I.23: Identification of container and seal number shall be indicated.

Box I.31: Donor identity shall correspond to the official identification of the animal.

Date of collection shall be indicated in the following format: dd/mm/yyyy.

Approval number of the centre shall correspond to the approval number of the semen centre indicated in Box I.12 where the semen was collected.

Part II:

(1) Delete as appropriate.

(2) Only approved semen collection centres listed in accordance with Article 11(4) of Council Directive 92/65/EEC on the Commission website: http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm

(3) Additional guarantees as laid down in Article 2 of Regulation (EC) No 546/2006 (OJ L 94, 1.4.2006, p. 28).

Official veterinarian or official inspector (*)

Name (in capital letters): Qualification and title:

Local veterinary unit: LVU No:

Date: Signature:

Stamp:

(*) The colour of the stamp and signature must be different from that of the other particulars in the certificate.

PART C Model health certificate IIIC for trade within the Union in consignments of semen of animals of the ovine and caprine species collected in accordance with Council Directive 92/65/EEC after 31 August 2010 and in consignments of stocks of semen of animals of the ovine and caprine species collected, processed and stored in accordance with Council Directive 92/65/EEC before 1 September 2010 and dispatched after 31 August 2010 from an approved semen storage centre

EUROPEAN UNION **Intra trade certificate**

Part I: Details of consignment presented

I.1. Consignor
Name
Address
Postal code

I.2. Certificate reference No | I.2.a. Local reference No

I.3. Central competent authority

I.4. Local competent authority

I.5. Consignee
Name
Address
Postal code

I.6. No(s) of related original certificates | No(s) of accompanying documents

I.7.

I.8. Country of origin | ISO code | I.9. Region of origin | Code

I.10. Country of destination | ISO code | I.11. Region of destination | Code

I.12. Place of origin
Semen centre ☐
Name | Approval number
Address
Postal code

I.13. Place of destination
Semen centre ☐ Holding ☐
Name | Approval number
Address
Postal code

I.14.

I.15.

I.16. Means of transport
Aeroplane ☐ Ship ☐ Railway wagon ☐
Road vehicle ☐ Other ☐
Identification

I.17.

I.18. Description of commodity

I.19. Commodity code (HS code)
05 11 99 85

I.20. Quantity

I.21. Temperature of products
Ambient ☐ Chilled ☐ Frozen ☐

I.22. Number of packages

I.23. Seal/Container No

I.24. Type of packaging

I.25. Commodities certified for:
Artificial reproduction ☐

I.26. Transit through third country ☐
Third country | ISO code
Exit point | Code
Entry point | BIP No

I.27. Transit through Member States ☐
Member State | ISO code
Member State | ISO code
Member State | ISO code

I.28. Export ☐
Third country | ISO code
Exit point | Code

I.29.

I.30.

I.31. Identification of the commodities

Species (Scientific name)	Breed	Donor identity	Date of collection	Approval number of the centre	Quantity

EUROPEAN UNION **Ovine and caprine semen — Part C**

Part II: Certification

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

I, the undersigned official veterinarian, hereby certify that the semen described above:

(1) *either* [II.1. was collected, processed and stored for a minimum period of 30 days immediately following collection in an approved semen collection centre (2) situated in the Member State of origin of the semen and operated and supervised in accordance with Chapter I(I)(1) and Chapter I(II)(1) of Annex D to Directive 92/65/EEC, and from where the semen was moved to the semen storage centre detailed in Part I.12 situated in the same Member State of origin of the semen under animal health and veterinary certification at least as strict as those provided for in:

(1) *either* [Part A of Annex III to Decision 2010/470/EU;]

(1) *or* [Part B of Annex III to Decision 2010/470/EU;]

(1) *or* [Decision 95/388/EC;]]

(1) *or* [II.1. was collected, processed and stored for a minimum period of 30 days immediately following collection in an approved semen collection centre (2) situated in the European Union and operated and supervised in accordance with Chapter I(I) and Chapter I(II) of Annex D to Directive 92/65/EEC and was moved to the semen storage centre detailed in Box I.12, in accordance with:

(1) *either* [Part A of Annex III to Decision 2010/470/EU;]

(1) *or* [Part B of Annex III to Decision 2010/470/EU;]

(1) *or* [Decision 95/388/EC;]]

(1) *or* [II.1. was collected, processed and stored for a minimum period of 30 days immediately following collection in an approved semen collection centre (2) situated in a third country or part(s) thereof listed in Annex I to Decision 2010/472/EU which is operated and supervised in accordance with Chapter I(I)(1) and Chapter I(II)(1) of Annex D to Directive 92/65/EEC, and was imported into the European Union under the conditions of Article 17(2) of Directive 92/65/EEC in accordance with:

(1) *either* [Section A of Part 2 of Annex II to Decision 2010/472/EU;]

(1) *or* [Section B of Part 2 of Annex II to Decision 2010/472/EU;]

(1) *or* [Annex II to Decision 2008/635/EC;]

II.2. was stored in the approved semen storage centre (2) indicated in Box I.12 which is operated and supervised in accordance with Chapter I(I)(2) and Chapter I(II)(2) of Annex D to Directive 92/65/EEC;

II.3. was sent to the place of loading in a sealed container in accordance with point 1.4 of Chapter III(I) of Annex D to Directive 92/65/EEC and bearing the number indicated in Box I.23.

Notes

Part I:

Box I.6: Shall correspond to the serial number of the individual official document(s) or health certificate(s) (either INTRA or CVED) that accompanied the semen described above from the approved semen collection centre of its origin to the described above semen storage centre. The original of this/these document(s) or certificate(s), or the officially endorsed copy/copies thereof must be attached to this certificate.

Box I.12: Place of origin shall correspond to the semen storage centre of dispatch of the semen.

Box I.13: Place of destination shall correspond to the semen collection or storage centre or to the holding of destination of the semen.

Box I.23: Identification of container and seal number shall be indicated.

Box I.31: Donor identity shall correspond to the official identification of the animal.
Date of collection shall be indicated in the following format: dd/mm/yyyy.
Approval number of the centre shall correspond to the approval number of the semen collection centre of the semen origin.

EUROPEAN UNION **Equine ova and embryos — Part C**

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

Part II:

(1) Delete as appropriate.

(2) Only approved semen collection or storage centres listed in accordance with Article 11(4) or Article 17(3)(b) of Council Directive 92/65/EEC on the Commission websites:

http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm
http://ec.europa.eu/food/animal/semen_ova/ovine/index_en.htm

Official veterinarian or official inspector (*)

Name (in capital letters): Qualification and title:

Local veterinary unit: LVU No:

Date: Signature:

Stamp:

(*) The colour of the stamp and signature must be different from that of the other particulars in the certificate.

ANNEX IV

Model health certificates for trade within the Union in consigments of ova/embryos of animals of the ovine and caprine species

[[F3]PART A

Model health certificate IVA for trade within the Union in consignments of ova and embryos of animals of the ovine and caprine species collected or produced in accordance with Council Directive 92/65/EEC after 31 August 2010 and dispatched by an approved embryo collection or production team of origin of the ova or embryos**]**

EUROPEAN UNION **Intra trade certificate**

Part I: Details of consignment presented

I.1. Consignor Name Address Postal code		I.2. Certificate reference No	I.2.a. Local reference No
		I.3. Central competent authority	
		I.4. Local competent authority	
I.5. Consignee Name Address Postal code		I.6.	
		I.7.	
I.8. Country of origin / ISO code	I.9. Region of origin / Code	I.10. Country of destination / ISO code	I.11. Region of destination / Code
I.12. Place of origin Embryo team ☐ Name / Approval number Address Postal code		I.13. Place of destination Holding ☐ Embryo team ☐ Name / Approval number Address Postal code	
I.14.		I.15.	
I.16. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification		I.17.	
I.18. Description of commodity		I.19. Commodity code (CN code) **05 11 99 85**	I.20. Quantity
I.21. Temperature of products Ambient ☐ Chilled ☐ Frozen ☐			I.22. Number of packages
I.23. Seal/Container No			I.24. Type of packaging
I.25. Commodities certified for: Artificial reproduction ☐			
I.26. Transit through third country ☐ Third country / ISO code Exit point / Code Entry point / BIP No		I.27. Transit through Member States ☐ Member State / ISO code Member State / ISO code Member State / ISO code	
I.28. Export ☐ Third country / ISO code Exit point / Code		I.29.	
I.30.			
I.31. Identification of the commodities Species (Scientific name) / Category / Donor identity / Date of collection / Approval number of the team / Quantity			

EUROPEAN UNION **Ovine and caprine ova/embryos — Part A**

II. Health information	II.a. Certificate reference number	II.b.

Part II: Certification

I, the undersigned official veterinarian, hereby certify that:

([1]) *either* [II.1. the *in vivo* derived embryos ([1])/*in vivo* derived ova ([1]) described above were collected, processed and stored by an embryo collection team ([2]) approved and supervised in accordance with Chapter I(III)(1) of Annex D to Directive 92/65/EEC;]

([1]) *or* [II.1. the *in vitro* produced embryos ([1])/micromanipulated embryos ([1]) described above were produced, processed and stored by an embryo production team ([2]) approved and supervised in accordance with Chapter I(III)(1) and (2) of Annex D to Directive 92/65/EEC;]

([1]) *either* [II.2. the *in vivo* derived embryos described above meet the requirements of Chapter III(II)(1) of Annex D to Directive 92/65/EEC;]

([1]) *or* [II.2. the *in vivo* derived ova described above meet the requirements of Chapter III(II)(2) of Annex D to Directive 92/65/EEC;]

([1]) *or* [II.2. the *in vitro* produced embryos described above meet the requirements of Chapter III(II)(3) of Annex D to Directive 92/65/EEC;]

([1]) *or* [II.2. the micromanipulated embryos described above meet the requirements of Chapter III(II)(4) of Annex D to Directive 92/65/EEC;]

([1]) [II.3. the consignment consists of embryos of the ovine or caprine species which:

([1]) *either* [were collected from animals which have been kept continuously since birth on a holding or holdings recognised as having a negligible or a controlled risk of classical scrapie in accordance with point 1 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001;]]

([1]) *or* [were collected from animals which have been kept continuously for the last three years before the collection on a holding or holdings which have complied for the last three years before collection with the requirements laid down in points (a) to (f) of point 1.3. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001;]]

([1]) *or* [were collected from animals which have been kept continuously since birth in a Member State or zone of a Member State with a negligible risk status for classical scrapie approved in accordance with the first subparagraph of point 2.2. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001;]]

([1]) *or* [were collected from ovine animals and

([1]) *either* [are of the ARR/ARR prion protein genotype;]]

([1]) *or* [carry at least one ARR allele and were collected after the date of 1 January 2015;]]

II.4. the ova or embryos described above come from female donors of the ovine ([1])/caprine species ([1]) which meet the requirements of Chapter IV(3) of Annex D to Directive 92/65/EEC;

([1]) *either* [II.5. the embryos described above were conceived as a result of artificial insemination of the donor females with semen which was collected, processed, stored and transported under conditions which comply with the requirements of Chapters I(I), II(I) and III(I) of Annex D to Directive 92/65/EEC;]

([1]) *or* [II.5. the embryos described above were conceived as a result of *in vitro* fertilisation of ova complying with the conditions in Chapter III(II)(2) of Annex D to Directive 92/65/EEC with semen which was collected, processed, stored and transported under conditions which comply with the requirements of Chapters I(I), II(I) and III(I) of Annex D to Directive 92/65/EEC;]

([1]) *or* [II.5. the ova have not been in contact with semen of the ovine and caprine species;]

II.6. the ova or embryos described above were sent to the place of loading in a sealed container in accordance with point 6 of Chapter III(II) of Annex D to Directive 92/65/EEC and bearing the number detailed in Box I.23.

Notes

Part I:

Box I.12: *Place of origin* shall correspond to the embryo collection team or embryo production team of embryos collection/production.

Box I.13: *Place of destination* shall correspond to the embryo collection team, embryo production team or to the holding of ova/embryos destination.

EUROPEAN UNION **Ovine and caprine ova/embryos — Part A**

II. Health information	II.a. Certificate reference number	II.b.

Box I.23: Identification of container and seal number shall be indicated.

Box I.31: *Category*: specify if: *in vivo* derived embryos, *in vivo* derived ova, *in vitro* produced embryos or micromanipulated embryos.

Donor identity shall correspond to the official identification of the animal.

Date of collection shall be indicated in the following format: dd/mm/yyyy.

Approval number of the team shall correspond to the embryo collection team or embryo production team of ova/embryos collection/production.

Part II:

([1]) Delete as appropriate.

([2]) Only approved embryo collection or production teams listed in accordance with Article 11(4) of Directive 92/65/EEC on the Commission website:

http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm.

— The colour of the stamp and signature must be different from that of the other particulars in the certificate.

Official veterinarian or official inspector

Name (in capital letters): Qualification and title:

Local veterinary unit: LVU No:

Date: Signature:

Stamp:'

PART B Model health certificate IVB for trade within the Union in consignments of stocks of ova and embryos of animals of the ovine and caprine species collected, processed and stored in accordance with Council Directive 92/65/EEC before 1 September 2010 and dispatched after 31 August 2010 by an approved embryo collection team of origin of the ova or embryos

EUROPEAN UNION **Intra trade certificate**

Part I: Details of consignment presented

I.1. Consignor Name Address Postal code		I.2. Certificate reference No	I.2.a. Local reference No
		I.3. Central competent authority	
		I.4. Local competent authority	
I.5. Consignee Name Address Postal code		I.6.	
		I.7.	
I.8. Country of origin ISO code	I.9. Region of origin Code	I.10. Country of destination ISO code	I.11. Region of destination Code
I.12. Place of origin Embryo team ☐ Name Approval number Address Postal code		I.13. Place of destination Holding ☐ Embryo team ☐ Name Approval number Address Postal code	
1.14.		I.15.	
I.16. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification		I.17.	
I.18. Description of commodity		I.19. Commodity code (HS code) **05 11 99 85**	I.20. Quantity
I.21. Temperature of products Ambient ☐ Chilled ☐ Frozen ☐			I.22. Number of packages
I.23. Seal/Container No			I.24. Type of packaging
I.25. Commodities certified for: Artificial reproduction ☐			
I.26. Transit through third country ☐ Third country ISO code Exit point Code Entry point BIP No		I.27. Transit through Member States ☐ Member State ISO code Member State ISO code Member State ISO code	
I.28. Export ☐ Third country ISO code Exit point Code		I.29.	
I.30.			

I.31. Identification of the commodities

Species (Scientific name)	Breed	Category	Donor identity	Date of collection	Approval number of the team	Quantity

EUROPEAN UNION **Ovine and caprine ova/embryos — Part B**

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

Part II: Certification

I, the undersigned official veterinarian, hereby certify that the ova/embryos (1) described above:

II.1. were collected, processed and stored under conditions which meet the requirements of Directive 92/65/EEC;

II.2. come from female donors of the ovine/caprine species (1) which meet the requirements of Chapter IV of Annex D to Directive 92/65/EEC;

(1) either [II.3. meet the requirements of Chapter III of Annex D to Directive 92/65/EEC and of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001.]

(1) or [II.3. meet the requirements of Chapter III of Annex D to Directive 92/65/EEC and of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and are destined for a Member State which benefits, for all or part of its territory, from the provisions laid down in point (b) or (c) of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and the donor animals comply regarding scrapie with the guarantees provided for by the programmes referred to in that point and with the guarantees (2) requested by the Member State of destination.]

(1) either [II.4. in the case of embryos, the semen used for fertilisation meets the requirements of Directive 92/65/EEC and the requirements of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001.]

(1) or [II.4. in the case of embryos, the semen used for fertilisation meets the requirements of Directive 92/65/EEC and the requirements of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and is destined for a Member State which benefits, for all or part of its territory, from the provisions laid down in point (b) or (c) of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and the donor animals comply regarding scrapie with the guarantees provided for by the programmes referred to in that point and with the guarantees (2) requested by the Member State of destination.]

Notes

Part I:

Box I.12: Place of origin shall correspond to the embryo collection team of ova/embryos collection.

Box I.13: Place of destination shall correspond to the embryo collection team, embryo production team or to the holding of ova/embryos destination.

Box I.23: Identification of container and seal number shall be indicated.

Box I.31: Category: specify if: *in vivo* derived embryos, *in vivo* derived ova, *in vitro* produced embryos or micromanipulated embryos.
Donor identity shall correspond to the official identification of the animal.
Date of collection shall be indicated in the following format: dd/mm/yyyy.
Approval number of the team shall correspond to the embryo collection team of ova/embryos collection indicated in Box I.12.

Part II:

(1) Delete as appropriate.
(2) Additional guarantees as laid down in Article 2 of Regulation (EC) No 546/2006 (OJ L 94, 1.4.2006, p. 28).

Official veterinarian or official inspector (*)

Name (in capital letters):	Qualification and title:
Local veterinary unit:	LVU No:
Date:	Signature:
Stamp:	

(*) The colour of the stamp and signature must be different from that of the other particulars in the certificate.

ANNEX V

Model health certificates for trade within the Union in consigments of ova/embryos of animals of the porcine species

PART A

Model health certificate VA for trade within the Union in consignments of ova and embryos of animals of the porcine species collected or produced in accordance with Council Directive 92/65/EEC after 31 August 2010 and dispatched by an approved embryo collection or production team of origin of the ova or embryos

EUROPEAN UNION — **Intra trade certificate**

Part I: Details of consignment presented

I.1. Consignor Name Address Postal code	I.2. Certificate reference No	I.2.a. Local reference No
	I.3. Central competent authority	
	I.4. Local competent authority	
I.5. Consignee Name Address Postal code	I.6.	
	I.7.	
I.8. Country of origin ISO code I.9. Region of origin Code	I.10. Country of destination ISO code I.11. Region of destination Code	
I.12. Place of origin Embryo team ☐ Name Approval number Address Postal code	I.13. Place of destination Holding ☐ Embryo team ☐ Name Approval number Address Postal code	
I.14.	I.15.	
I.16. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification	I.17.	
I.18. Description of commodity	I.19. Commodity code (HS code) **05 11 99 85**	
	I.20. Quantity	
I.21. Temperature of products Ambient ☐ Chilled ☐ Frozen ☐	I.22. Number of packages	
I.23. Seal/Container No	I.24. Type of packaging	
I.25. Commodities certified for: Artificial reproduction ☐		
I.26. Transit through third country ☐ Third country ISO code Exit point Code Entry point BIP No	I.27. Transit through Member States ☐ Member State ISO code Member State ISO code Member State ISO code	
I.28. Export ☐ Third country ISO code Exit point Code	I.29.	
I.30.		
I.31. Identification of the commodities Species (Scientific name) Breed Category Donor identity Date of collection Approval number of the team Quantity		

EUROPEAN UNION **Porcine ova/embryos — Part A**

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

Part II: Certification

I, the undersigned official veterinarian, hereby certify that the ova/embryos (1) described above:

II.1. were produced/collected (1), processed and stored by an embryo collection/production (1) team (2) approved and supervised in accordance with Chapter I(III) of Annex D to Directive 92/65/EEC;

II.2. meet the requirements of Chapter III(II) of Annex D to Directive 92/65/EEC;

II.3. come from donor females of the porcine species which meet the requirements of Chapter IV(2) of Annex D to Directive 92/65/EEC;

(1) either [II.4. are *in vivo* derived embryos which:

II.4.1. were conceived as a result of artificial insemination with semen meeting the requirements of Directive 90/429/EEC,

II.4.2. originate from a Member State or region thereof:

(1) either [listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC;]

(1) or [listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;]

(1) or [listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC and have been washed with trypsin;]

(1) or [listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex II to Decision 2008/185/EC;]

(1) or [not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I or II to Decision 2008/185/EC and have been washed with trypsin;]

(1) or [not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;]]

(1) or [II.4. are *in vitro* produced/micromanipulated (1) embryos which:

II.4.1. were conceived as a result of *in vitro* fertilisation with semen meeting the requirements of Directive 90/429/EEC,

II.4.2. originate from a Member State or region thereof:

(1) either [listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC;]

(1) or [listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;]

(1) or [listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC and the donor females of the ova used for their production comply with the conditions of Article 1 of Decision 2008/185/EC;]

(1) or [listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex II to Decision 2008/185/EC;]

(1) or [not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I or II to Decision 2008/185/EC and the donor females of the ova used for their production comply with the conditions of Article 1 of Decision 2008/185/EC;]

(1) or [not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;]]

EUROPEAN UNION **Porcine ova/embryos — Part A**

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

(1) *or* [II.4. are *in vivo* derived ova which originate from a Member State or region thereof:

(1) *either* [listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC;]

(1) *or* [listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;]

(1) *or* [listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC and which come from donor females complying with the conditions of Article 1 of Decision 2008/185/EC;]

(1) *or* [listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex II to Decision 2008/185/EC;]

(1) *or* [not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I or II to Decision 2008/185/EC and which come from donor females complying with the conditions of Article 1 of Decision 2008/185/EC;]

(1) *or* [not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;]]

II.5. were sent to the place of loading in a sealed container under conditions complying with point 6 of Chapter III(II) of Annex D to Directive 92/65/EEC and bearing the number detailed in Box I.23.

Notes

Part I:

Box I.12: place of origin shall correspond to the embryo collection team or embryo production team of ova/embryos collection/production.

Box I.13: place of destination shall correspond to the embryo collection team, embryo production team or to the holding of ova/embryos destination.

Box I.23: identification of container and seal number shall be indicated.

Box I.31: category: specify if: *in vivo* derived embryos, *in vivo* derived ova, *in vitro* produced embryos or micromanipulated embryos.

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

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

approval number of the team shall correspond to the embryo collection team or embryo production team of ova/embryos collection/production indicated in Box I.12.

Part II:

(1) Delete as appropriate.

(2) Only approved embryo collection or production teams listed in accordance with Article 11(4) of Council Directive 92/65/EEC on the Commission website:
http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm

Official veterinarian or official inspector (*)

Name (in capital letters): Qualification and title:

Local veterinary unit: LVU No:

Date: Signature:

Stamp:

(*) The colour of the stamp and signature must be different from that of the other particulars in the certificate.

PART B Model health certificate VB for trade within the Union in consignments of stocks of ova and embryos of animals of the porcine species collected, processed and stored in accordance with Council Directive 92/65/EEC before 1 September 2010 and dispatched after 31 August 2010 by an approved embryo collection team of origin of the ova or embryos

***Changes to legislation:** There are outstanding changes not yet made to Commission Decision of 26 August 2010 laying down model health certificates for trade within the Union in semen, ova and embryos of animals of the equine, ovine and caprine species and in ova and embryos of animals of the porcine species (notified under document C(2010) 5779) (Text with EEA relevance) (2010/470/EU). Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes*

EUROPEAN UNION **Intra trade certificate**

Part I: Details of consignment presented

I.1. Consignor Name Address Postal code		I.2. Certificate reference No	I.2.a. Local reference No
		I.3. Central competent authority	
		I.4. Local Competent Authority	
I.5. Consignee Name Address Postal code		I.6.	
		I.7.	
I.8. Country of origin ISO code	I.9. Region of origin Code	I.10. Country of destination ISO code	I.11. Region of destination Code
I.12. Place of origin Embryo team ☐ Name Approval number Address Postal code		I.13. Place of destination Holding ☐ Embryo team ☐ Name Approval number Address Postal code	
I.14.		I.15.	
I.16. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification		I.17.	
I.18. Description of commodity		I.19. Commodity code (HS code) **05 11 99 85**	
		I.20. Quantity	
I.21. Temperature of products Ambient ☐ Chilled ☐ Frozen ☐		I.22. Number of packages	
I.23. Seal/Container No		I.24. Type of packaging	
I.25. Commodities certified for: Artificial reproduction ☐			
I.26. Transit through third country ☐ Third country ISO code Exit point Code Entry point BIP No		I.27. Transit through Member States ☐ Member State ISO code Member State ISO code Member State ISO code	
I.28. Export ☐ Third country ISO code Exit point Code		I.29.	
I.30.			

I.31. Identification of the commodities

Species (Scientific name)	Breed	Category	Donor identity	Date of collection	Approval number of the team	Quantity

EUROPEAN UNION **Porcine ova/embryos — Part B**

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

Part II: Certification

I, the undersigned official veterinarian, hereby certify that:

II.1. The ova/embryos (1) described above:

II.1.1. were collected, processed and stored under conditions which meet the requirements of Directive 92/65/EEC;

II.1.2. come from donor female swine which meet the requirements of Chapter IV of Annex D to Directive 92/65/EEC;

II.1.3. meet the requirements of Chapter III of Annex D to Directive 92/65/EEC.

(1) either [II.2. In the case of embryos,

II.2.1. the semen used for fertilisation meets the requirements of Directive 90/429/EEC;

II.2.2. the embryos have been washed with trypsin (2).]

(1) or [II.2. In the case of ova, the ova comes from a donor female swine which meets the conditions of Article 1 of Decision 2008/185/EC (2).]

Notes

Part I:

Box I.12: place of origin shall correspond to the embryo collection team of ova/embryos collection.

Box I.13: place of destination shall correspond to the embryo collection team, embryo production team or to the holding of ova/embryos destination.

Box I.23: identification of container and seal number shall be indicated.

Box I.31: category: specify if: *in vivo* derived embryos, *in vivo* derived ova, *in vitro* produced embryos or micromanipulated embryos.
donor identity shall correspond to the official identification of the animal.
date of collection shall be indicated in the following format: dd/mm/yyyy.
approval number of the team shall correspond to the embryo collection team of ova/embryos collection indicated in Box I.12.

Part II:

(1) Delete as appropriate.

(2) This condition applies only to ova and embryos which originate in the Member States or regions thereof not listed in Annexes I and II to Decision 2008/185/EC (OJ L 59, 4.3.2008, p. 19) and destined to the Member States or regions thereof so listed. It shall also apply to movements from Member States or regions thereof listed in Annex II of Decision 2008/185/EC to Member States or regions thereof listed in Annex I of Decision 2008/185/EC.

Official veterinarian or official inspector (*)

Name (in capital letters):	Qualification and title:
Local veterinary unit:	LVU No:
Date:	Signature:
Stamp:	

(*) The colour of the stamp and signature must be different from that of the other particulars in the certificate.

(**1**) OJ L 268, 14.9.1992, p. 54.

(**2**) OJ L 52, 3.3.2010, p. 14.

(**3**) OJ L 94, 31.3.2004, p. 44.

(**4**) OJ L 182, 2.8.1995, p. 27.

(**5**) OJ L 185, 4.8.1995, p. 58.

(**6**) OJ L 234, 3.10.1995, p. 30.

(**7**) OJ L 275, 18.11.1995, p. 30.

Changes to legislation:
There are outstanding changes not yet made to Commission Decision of 26 August 2010 laying down model health certificates for trade within the Union in semen, ova and embryos of animals of the equine, ovine and caprine species and in ova and embryos of animals of the porcine species (notified under document C(2010) 5779) (Text with EEA relevance) (2010/470/EU). Any changes that have already been made to the legislation appear in the content and are referenced with annotations.
View outstanding changes

Changes and effects yet to be applied to :

- Annexes omitted by S.I. 2019/795 reg. 33(9) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Decision revoked by S.I. 2020/1462 Sch. para. 6
- Annex 3 Page A replacement by EUDN 2016/2002 Decision
- Art. 2 heading substituted by S.I. 2019/795 reg. 33(3)(a) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 2 words substituted by S.I. 2019/795 reg. 33(3)(b) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 3 heading substituted by S.I. 2019/795 reg. 33(4)(a) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 3 words substituted by S.I. 2019/795 reg. 33(4)(b) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 4 heading substituted by S.I. 2019/795 reg. 33(5)(a) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 4 words substituted by S.I. 2019/795 reg. 33(5)(b) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 5 heading substituted by S.I. 2019/795 reg. 33(6)(a) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 5 words substituted by S.I. 2019/795 reg. 33(6)(b) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 6 heading substituted by S.I. 2019/795 reg. 33(7)(a) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 6 words substituted by S.I. 2019/795 reg. 33(7)(b) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 9 omitted by S.I. 2019/795 reg. 33(8) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))

Changes and effects yet to be applied to the whole legislation item and associated provisions

- Art. 1(1) Art. 1 renumbered as Art. 1(1) by S.I. 2019/795 reg. 33(2)(a) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))

- Art. 1(1) words substituted by S.I. 2019/795 reg. 33(2)(b) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 1(2) inserted by S.I. 2019/795 reg. 33(2)(c) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 2(a) words substituted by S.I. 2019/795 reg. 33(3)(c) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 2(b) words substituted by S.I. 2019/795 reg. 33(3)(c) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 2(c) words substituted by S.I. 2019/795 reg. 33(3)(c) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 2(d) words substituted by S.I. 2019/795 reg. 33(3)(c) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 3(a) words substituted by S.I. 2019/795 reg. 33(4)(c) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 3(b) words substituted by S.I. 2019/795 reg. 33(4)(c) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 3(c) words substituted by S.I. 2019/795 reg. 33(4)(c) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 4(a) words substituted by S.I. 2019/795 reg. 33(5)(c) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 4(b) words substituted by S.I. 2019/795 reg. 33(5)(d) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 4(c) words substituted by S.I. 2019/795 reg. 33(5)(e) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 5(a) words substituted by S.I. 2019/795 reg. 33(6)(c) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 5(b) words substituted by S.I. 2019/795 reg. 33(6)(d) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 6(a) words substituted by S.I. 2019/795 reg. 33(7)(c) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))
- Art. 6(b) words substituted by S.I. 2019/795 reg. 33(7)(d) (This amendment not applied to legislation.gov.uk. S.I. 2019/795 revoked (31.12.2020) by virtue of S.I. 2020/1462, regs. 1(2)(b), 73(1))