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<sup>(1)</sup> 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<sup>(2)</sup>, 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<sup>(3)</sup>.
- (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<sup>(4)</sup>, Commission Decision 95/307/EC of 24 July 1995 determining the specimen animal health certificate for trade in semen of the equine species<sup>(5)</sup>, 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<sup>(6)</sup> and Commission Decision 95/483/EC of 9 November 1995 determining the specimen certificate for intra-Community trade in ova and embryos of swine<sup>(7)</sup> 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.

# [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.

# [F1ANNEX 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

EUF	ROPE	AN UNION	Intra trade certificate				
	l.1.	Consignor Name	I.2. Certificate reference No I.2.a. Local reference No				
		Address	I.3. Central competent authority				
ented		Postal code	I.4. Local competent authority				
nt prese	I.5.	Consignee Name Address	1.6.				
gnme		Postal code	1.7.				
Part I: Details of consignment presented	1.8.	Country of origin ISO code I.9. Region of origin Code	I.10. Country of destination ISO code I.11. Region of destination Code				
ails	l.12.	Place of origin	I.13. Place of destination				
Det		Semen Centre	Semen Centre Holding				
art I:		Name Approval number Address	Name Approval number Address				
•		Postal code	Postal code				
	l.14.		1.15.				
	I.16.	Means of transport	1.17.				
		Aeroplane Ship Railway wagon Shoad vehicle Other Identification:					
	l.18.	Description of commodity	I.19. Commodity code (HS code) 05 11 99 85				
			I.20. Quantity				
	I.21.	Temperature of products	I.22. Number of packages				
		Ambient ☐ Chilled ☐ Frozen [					
	1.23.	Seal/Container No	I.24. Type of packaging				
	1.25.	Commodities certified for: Artificial reproduction □					
	1.26.	Transit through third country	I.27. Transit through Member States				
		Third country ISO code	Member State ISO code				
		Exit point Code	Member State ISO code				
		Entry point BIP No	Member State ISO code				
	1.28.	Export	1.29.				
		Third country ISO code					
		Exit point Code					
	1.30.						
	I.31.	Identification of the commodities					
		Species Donor identity (Scientific name)	Date of collection Quantity				

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	EUROPEA	UNION		Equine semen – Part A						
	II.	Health information	II.a. Certificate reference No	II.b.						
	I, the unde	ersigned official veterinarian, hereby certify	that:							
	II.1.		the semen described above was collected a competent authority in accordance with Ch							
c	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:								
Part II: Certification	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);								
art II:	II.1.1.2.	fulfilled the conditions for a holding laid of	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 20 have been admitted onto the centre.										
	II.3.	The semen described above was collected	ed from donor stallions, which:							
	II.3.1.	did not show any clinical sign of an infe collection centre and on the day the sem	ctious or contagious disease at the time of nen was collected;	admission onto the semen						
	II.3.2.		o the date of semen collection in holdings whatagious equine metritis during that period;	ere no equine showed any						
	II.3.3.		period of at least 30 days prior to the date of to in point II.3.5.1, II.3.5.2 or II.3.5.3 until the							
	II.3.4.	Tests and Vaccines for Terrestrial Anima	t the requirements of the relevant Chapter of als of the OIE, carried out in a laboratory we referred to hereinafter included in its accredible (5), as follows:	which is recognised by the						
	II.3.4.1.		gar-gel immuno-diffusion test (AGID or Coggin infectious anaemia with a negative result;	s test) or an enzyme-linked						
	II.3.4.2.	for equine viral arteritis (EVA),								
	(1) either	[II.3.4.2.1. a serum neutralisation test w	ith a negative result at a serum dilution of o	ne in four;]						
	(1) and/or	[II.3.4.2.2. a virus isolation test, polymer aliquot of the entire semen o	rase chain reaction (PCR) or real-time PCR $\nu$ of the donor stallion;	vith a negative result on an						
	II.3.4.3.		agent identification test carried out on three with an interval of not less than 7 days at ledis;							
		antimicrobial treatment of the donor stallic	lier than 7 days (systemic treatment) or 21 or and were placed in transport medium with aboratory where they were subjected with n	activated charcoal, such as						
	( <sup>1</sup> ) either		uigenitalis after cultivation under microaerop rs after taking the specimens from the donor during transport;]							

EUROPEAN	UNION				Equine semen - Part			
II.	Health informa	ation	II.a.	Certificate reference No	II.b.			
( <sup>1</sup> ) and/or	[II.3.4.3.2. th	e detection of genome of 8 hours after taking the sp	f <i>Taylorella</i> ecimens fr	equigenitalis by PCR or real-tirom the donor animal;]	me PCR, carried out within			
II.3.5.		ed with the results specific ints II.3.5.1, II.3.5.2 and II.		II.3.4 in each case to at least o	one of the test programmes			
( <sup>6</sup> ) [II.3.5.1.	to the date of	the first collection and duri	ng the perio	semen collection centre for a per od of collection of the semen desc contact with equidae of lower he	ribed above and no equidae			
	year at the be chilled or froz	eginning of the breeding se	ason or pr an 14 day	t on samples taken (7) from the di ior to the first collection of semen s following the date of the comm semen collection.]	intended for trade in fresh,			
( <sup>6</sup> ) [II.3.5.2.	the first collect the responsib	tion and during the period illity of the centre veterinaria	of collection an for a co	ction centre for a period of at least n of the semen described above, ntinuous period of less than 14 da ct with equidae of lower health s	but has left the centre under lys, and/or other equidae on			
	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:							
	( <sup>1</sup> ) either			II.3.4.2 was last carried out on a e collection of the semen describ				
	( <sup>1</sup> ) or	the donor stallion taken (7 described above and a b	) not more lood samp esult in a	II.3.4.2.2 was carried out on an al than 6 months prior to the date of le taken (7) from the donor stallion serum neutralisation test for equi	f the collection of the semen n during the 6-month period			
		s (swabs) taken (7) not n		sts described in point II.3.4.3 wa 60 days prior to the date of the				
	(1) either	[on two occasions at leas	st 7 days a	apart;]				
	( <sup>1</sup> ) or	[on a single occasion and	d subjected	d to a PCR or real-time PCR.]]				
( <sup>6</sup> ) [II.3.5.3.		allion does not meet the o		set out in points 1.6(a) and (b) or trade in frozen semen.	of Chapter II of Annex D to			
		scribed in points II.3.4.1, II. st once a year at the begin		II.3.4.3 were carried out on samp e breeding season,	les taken (7) from the donor			
and	during the sto semen and be	orage period of the semen	of a minin d from the	vere carried out on samples take num period of 30 days from the semen collection centre, not less scribed above,	date of the collection of the			

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EUROPE	AN UNION						Equine s	emen – Part A
II.	Health inf	formation		II.a. Ce	rtificate refere	nce No	II.b.	
and	( <sup>1</sup> ) either	during the storage of the semen and	e period of the before the se	teritis described in point II.3.4.2 were carried out on samples taken (7) he semen of a minimum period of 30 days from the date of the collection semen is removed from the semen collection centre or used, not less than 30 days after the collection of the semen described above.]				
II.3.6.	(1) or	[the non-shedder isolation test, PCI entire semen of t donor stallion rea neutralisation test at the testing provid	R or real-time the donor stall acted with a part of the forequine vi	PCR carried of lion taken (7) to consitive result ral arteritis.]]	out with a nega wice a year a at a serum d	ative result on it an interval o lilution of at le	samples of an of at least 4 m east one in fo	aliquot of the onths and the
	underwen			T 11.0.5 011 5a11	·			
semer	e e		date ( <sup>7</sup> )			ampling for he	· · · ·	
Identification of semen	Test programme	Donor residence	Semen collection	EIA II.3.4.1.	Blood sample	Semen sample	1. sample	2. sample
	_							
( <sup>1</sup> ) either	[II.4. No	antibiotics were ad	ded to the se	men;]				
( <sup>1</sup> ) or		following antibiotic ted semen of not le						
II.5.	The semen	n described above	was:					
II.5.1.		processed, stored				hich comply	with the red	quirements of
II.5.2.	in the case	e of frozen semen,	stored for a	minimum perio	od of 30 days	from the date	of collection	of the semen;
II.5.3.		e place of loading 92/65/EEC and bear				point 1.4 of	Chapter III(I)	of Annex D to

EUROPEAN UNION	Equine semen - Part A

II.	Health information	II.a.	Certificate reference No	II.b.
1				

### 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

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.

	Jo u	Test programme	Start date (7)		Date of sampling for health tests (7)					
	catior		Donor residence		EVA II.3.4.2.		CEM	CEM II.3.4.3.		
	Identification semen			Semen collection	EIA II.3.4.1.	Blood sample	Semen sample	1. sample	2. sample	
Γ	Α	В	С	D	EIA-1	EVA-B1	EVA-S1	CEM-11	CEM-12	
	^		C		EIA-2	EVA-B2	EVA-S2	CEM-21	CEM-22	

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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

EUF	IOPEAN UNION				Equille Selliell - Part A
II.	Health information	II.a.	Certificate reference	No	II.b.
( <sup>1</sup> )	Delete as appropriate				
(2)	Only approved semen collection centres listed in website:	accordance	with Article 11(4) of D	irective 92/6	5/EEC on the Commission
	http://ec.europa.eu/food/animal/approved_establis	shments/est	ablishments_vet_field_d	en.htm	
(3)	OJ L 268, 14.9.1992, p. 54.				
( <sup>4</sup> )	OJ L 192, 23.7.2010, p. 1.				
( <sup>5</sup> )	OJ L 165, 30.4.2004, p. 1.				
( <sup>6</sup> )	Cross out the programme(s) that do(es) not app	ly to the co	nsignment.		
( <sup>7</sup> )	Insert date in table in point II.3.6 (follow Guidane	ce in Part I	of the Notes).		
(8)	Insert names and concentrations.				
_	The colour of the stamp and signature must be	different fro	m that of the other pa	rticulars in t	he certificate.
Off	icial veterinarian				
	Name (in capital letters):		Qua	alification an	d title:
	Local veterinary unit:		LVU	J No:	
	Date:		Sign	nature:	
	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

EUF	ROPE	AN UNION	Intra trade certificate				
	l.1.	Consignor Name	I.2. Certificate reference No I.2.a. Local reference No				
_		Address	I.3. Central competent authority				
ented		Postal code	I.4. Local competent authority				
Part I: Details of consignment presented	1.5.	Consignee	1.6.				
Ħ P		Name					
ner		Address	1.7.				
gur		Postal code					
	1.8.	Country of ISO code I.9. Region of code origin	I.10. Country of destination ISO code I.11. Region of destination destination				
18	1.12.	Place of origin	I.13. Place of destination				
etai		Semen Centre □	Semen Centre ☐ Holding ☐				
ă		Name Approval number	Name Approval number				
Ξ		Address	Address				
Ъ		Postal code	Postal code				
	l.14.		1.15.				
	I.16.	Means of transport	I.17.				
		Aeroplane Ship Railway wagon Shoad vehicle Other Identification:					
	I.18.	Description of commodity	I.19. Commodity code (HS code)				
		•	05 11 99 85				
			I.20. Quantity				
	I.21.	Temperature of products	I.22. Number of packages				
		Ambient ☐ Chilled ☐ Frozen [					
	1.23.	Seal/Container No	I.24. Type of packaging				
	1.25.	Commodities certified for:					
		Artificial reproduction					
	1.26.	Transit through third country	I.27. Transit through Member States				
		Third country ISO code	Member State ISO code				
		Exit point Code	Member State ISO code				
		Entry point BIP No	Member State ISO code				
	1.28.	Export	1.29.				
		Third country ISO code					
		Exit point Code					
	1.30.						
	I.31.	Identification of the commodities					
		Species Donor identity (Scientific name)	Date of collection Quantity				

	<u> </u>	V UNION	ormot!			Contificate	vofovono - N-		Equine semen - F	=	
II.		Health inf	ormation		II.a.	Certificate	reference No		II.b.		
I, th	e unde	ersigned off	ficial veterii	narian, hereby certify	/ that:						
II.1.		trade is ap	oproved an	n centre ( <sup>2</sup> ), in which d supervised by the tive 92/65/EEC;	h the sen competen	nen described t authority in a	above was coll ccordance with	lected, Chapter	processed and store r I(I)(1) and Chapter I(	d fo	
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.	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 equic	dae which were free	of clinica	l signs of equ	iine viral arteritis	s and co	ontagious equine me	triti	
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.									
11.3.		The seme	en describe	d above was collect	ted from (	donor stallions	, which:				
11.3.	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;										
11.3.2	2.			30 days prior to the e viral arteritis or co					no equine has shown	ı ar	
11.3.3	3.								semen collection and d of the collection pe		
11.3.4	4.	Diagnostic	c Tests and		trial Anim	als of the OIE	, carried out on s	samples	Chapter of the Manus taken in accordance petent authority:		
		( <sup>1</sup> ) either	[II.3.4.1.	an agar-gel immun negative result;]	no-diffusio	n test (Coggir	ns test) for equi	ne infed	ctious anaemia (EIA)	wi	
		( <sup>1</sup> ) or	[II.3.4.1.	an ELISA for equir	ne infectio	us anaemia (I	EIA) with negativ	ve resul	lt;]		
and	1	( <sup>1</sup> ) either	[11.3.4.2.	a serum neutralisa dilution of one in fo		for equine vira	al arteritis (EVA)	with n	egative result at a se	eru	
		( <sup>1</sup> ) or	[11.3.4.2.	a virus isolation te aliquot of the entire				ed out w	vith negative result o	n a	
and	•		II.3.4.3.	occasions on samp after a cultivation of	oles taken of 7 to 14 n at least	with an interval	al of 7 days by is re-ejaculatory flu	solation uid or a	EM) carried out on of <i>Taylorella equigen</i> semen sample and rethral fossa with neg	nita fro	

II.	Health in	formation	II.a.	Certificate reference No	II.b.	
II.3.5.		on subjected with the results spin points II.3.5.1, II.3.5.2 and II.		II.3.4 in each case to at least one ollows:	of the test programmes (4)	
II.3.5.1.	of the fire	st collection and during the per	iod of col	ne semen collection centre for at lea lection of the semen described about with equidae of lower health stat	ve and no equidae on the	
				ried out on samples taken (5) prior to commencement of the residence p		
II.3.5.2. The donor stallion was resident on the semen collection centre for at least 30 days prior to the date of collection and during the period of collection of the semen described above, but has left the centre un responsibility of the centre veterinarian for a continuous period of less than 14 days, and/or other equidae 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 beriod in the year the semen described above was collected and 14 days following the date of the commencement of the residence period of at least 30 days,						
and	(¹) either			.4.2 for equine viral arteritis was la the semen described above was co		
	(¹) or	entire semen of the donor stall was collected and a blood san	ion taken nple taken	teritis was carried out with negative (5) not more than 6 months before to on the same date (5) reacted positive dilution of more than one in four,]	he semen described above	
and				contagious equine metritis was la the semen described above was co		
II.3.5.3.				ed out on samples taken (5) prior to ear the semen described above was		
and		described in point II.3.4 were ladays after the collection of the		d out on samples taken (5) not less secribed above.	than 14 days and not more	

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

<b>EUROPEA</b>	n uni	ON						Equine se	emen – Part I	
II.	Heal	th informa	tion		II.a. Ce	rtificate referer	nce No	II.b.		
II.3.6.	have	undergon	e the testing p	provided for	in point II.3.5 on samples taken on the following dates:					
<b>J</b> o		ae	Start o	date ( <sup>5</sup> )	Date of sampling for health tests (5)					
itification		gram				EVA II.3.4.2.		CEM	II.3.4.3.	
Identification of semen		Test programme	Donor residence	Semen collection	EIA II.3.4.1.	Blood sample	Semen sample	1. sample	2. sample	
	+	<u> </u>								
	+									
	+									
	+									
	$\top$									
' '	[11.4.		iotics were add		•					
(1) or	[11.4.	diluted se	emen of not le	ss than ( <sup>6</sup> ):	ation of antibiot					
									;]	
II.5.			cribed above							
II.5.1.					sported under Directive 92/65/E		hich comply	with the rec	uirements of	
II.5.2.					container in acc		point 1.4 of	Chapter III(I) o	of Annex D to	
Notes										
Part I:										
Box 1.12:	place	e of origin	shall correspo	and to the se	emen collection	centre of origi	n of the seme	en.		
Box I.13:		e of destir ination.	nation shall co	orrespond to	the semen co	llection or sto	orage centre	or to the hold	ing of semen	
Box 1.23:	iden	tification of	f container and	d seal numb	er shall be indic	ated.				
Box I.31:	dono	or identity :	shall correspor	nd to the off	icial identificatio	n of the anima	al.			
	date	of collecti	on shall be in	dicated in th	e following form	at: dd/mm/yyy	y.			
			er of the centr		spond to the ap	proval number	of the semen	centre indicat	ed in Box I.12	

EUROPEAN UNION Equine semen – Part B

II. F	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.

Jo c	эше	Start o	date ( <sup>5</sup> )	Date of sampling for health tests (5)				
ication	programme	Donor	Semen		EVA II.3.4.2.		CEM II.3.4.3.	
Identification semen	Test p	residence	collection	EIA II.3.4.1.	Blood sample	Semen sample	1. sample	2. sample
A	В	С	D	EIA-1	EVA-B1	EVA-S1	CEM-11	CEM-12
				EIA-2	EVA-B2	EVA-S2	CEM-21	CEM-22

<sup>(1)</sup> Delete as appropriate.

 $http://ec.europa.eu/food/animal/approved\_establishments/establishments\_vet\_field\_en.htm$ 

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

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

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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

EUF	ROPEAN UNION			Equine semen – Part B							
II.	Health information	II.a.	Certificate reference No	II.b.							
(4)	(4) Cross out the programme(s) that do(es) not apply to the consignment.										
( <sup>5</sup> )	i) Insert date in table in point II.3.6 (follow Guidance in Part II of the Notes).										
( <sup>6</sup> )	Insert names and concentrations.										
_	The colour of the stamp and signature must be	different fr	om that of the other particulars in	the certificate.							
Off	icial veterinarian										
	Name (in capital letters):		Qualification an	nd title:							
	Local veterinary unit:		LVU No:								
	Date: Signature:										
	Stamp:										
l											

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

EUF	ROPE	AN UNION	Intra trade certificate			
	l.1.	Consignor Name	I.2. Certificate reference No I.2.a. Local reference No			
٠		Address	I.3. Central competent authority			
ente		Postal code	I.4. Local competent authority			
pres	1.5.	Consignee Name	1.6.			
턽		Address				
n me		Postal code	1.7.			
of consignment presented	1.8.	Country of ISO code I.9. Region of Code origin Code	I.10. Country of ISO code I.11. Region of destination destination			
<u>s</u>	112	Place of origin	I.13. Place of destination			
l: Details of		Semen Centre	Semen Centre Holding			
Part I: [		Name Approval number Address	Name Approval number Address			
4		Postal code	Postal code			
	1.14.		I.15.			
_	1.16.	Means of transport	1.17.			
		Aeroplane Ship Railway wagon IRoad vehicle Other Identification:				
	l.18.	Description of commodity	I.19. Commodity code (HS code)			
			05 11 99 85			
			I.20. Quantity			
	I.21.	Temperature of products	I.22. Number of packages			
		Frozen				
	1.23.	Seal/Container No	I.24. Type of packaging			
	1.25.	Commodities certified for:	·			
		Artificial reproduction				
	1.26.	Transit through third country	I.27. Transit through Member States			
		Third country ISO code	Member State ISO code			
		Exit point Code	Member State ISO code			
		Entry point BIP No	Member State ISO code			
	1.28.	Export	1.29.			
		Third country ISO code				
		Exit point Code				
	1.30.					
	I.31.	Identification of the commodities				
		Species Donor identity (Scientific name)	Date of collection Quantity			

	EUROPEA	N UNION						Equine semen – Part			
	II.	Health inforn	nation		II.a.	Certificate referer	nce No	II.b.			
	I, the und	lersigned officia	al veterinariar	n, hereby certify	that:						
	II.1. The semen collection centre (2), in which the semen described above was collected, processed and stored trade:										
	II.1.1.	II.1.1. is approved and supervised by the competent authority according to the conditions of Chapter I of Annex D Directive 92/65/EEC;									
Part II: Certification	II.1.2.	on the day so	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);								
Part II: Ce	II.1.3.	dispatched a	s fresh/chille		until the 30	days mandatory sto		ntil the date the semen war r frozen semen elapsed ( <sup>1</sup> )			
	II.1.4.	was dispatc	hed as fresh	/chilled (1) sem	nen or unt	il the 30 days mar	ndatory storage	on until the date the semele period for frozen semele contagious equine metritis			
	II.2.	All equidae 2009/156/EC		admitted onto	the cent	re under the prov	visions of Artic	cle 4 and 5 of Directive			
	II.3.	The semen	described ab	ove was collec	ted from d	onor stallions, which	n:				
	II.3.1.	on the day t	he semen wa	as collected hav	ve not sho	wn clinical signs of	an infectious o	r contagious disease,			
	II.3.2.	during at lea	st 30 days p	rior to collection	n of the se	emen have not been	used for natu	ral service,			
	II.3.3.		st 30 days p of equine vi		n of the se	emen have been ke	pt on holdings	where no equidae shower			
	II.3.4.			rior to collectio is equine metri		emen have been ke	pt on holdings	where no equidae shower			
	II.3.5.					scertain, have not be immediately prece		with equidae suffering from of the semen,			
	II.3.6.					carried out in a labo cified in point II.3.7.		sed by the competent auth			
			[II.3.6.1.	an agar-gel negative resi		fusion test (Coggin	s test) for equi	ne infectious anaemia witl			
	and	(1) either	[II.3.6.2.	a serum neu dilution of or			l arteritis with	negative result at a serun			
		( <sup>1</sup> ) or	[II.3.6.2.			or equine viral arteri men of the donor st		with negative result on a			

# EUROPEAN UNION Equine semen – Part C

						·
II.	Health info	rmation		II.a.	Certificate reference No	II.b.
and		II.3.6.3.	samples collected Taylorella equiger	d from the nitalis from east from the	donor stallion with an inter- pre-ejaculatory fluid or a se	arried out on two occasions on val of 7 days by isolation of men sample and from genital d urethral fossa with negative
II.3.7.	have been	subject to	the one of the follo	owing test p	rogrammes ( <sup>4</sup> ):	
II.3.7.1.	collection,	and during	the collection peri-	od, and no		st 30 days prior to the semen tre came during that time into
	in the case	e of contagi	ous equine metritis	on a seco	nd sample taken on	( <sup>5</sup> ) and( <sup>5</sup> ), being at at the beginning of the breeding
II.3.7.2.					he collection centre or other e is than the donor stallion.	equidae on the collection centre
	in the case	e of contagi	ous equine metritis	on a seco	nd sample taken on	( <sup>5</sup> ) and ( <sup>5</sup> ), being within nning of the breeding season,
and		blood take	escribed in point II. on onabove was collect		uine infectious anaemia was l ( <sup>5</sup> ), being not more tha	last carried out on a sample of an 120 days before the semen
and	( <sup>1</sup> ) either	collected of	e tests described ins collected,]	n point II.3.6 ( <sup>5</sup> )	5.2 for equine viral arteritis wa , being not more than 30 day	as last carried out on a sample ys before the semen described
	( <sup>1</sup> ) or	isolation to	est which was carri	ied out on a	in aliquot of the entire semen	eritis was confirmed by a virus of the donor stallion collected ar before the semen described
II.3.7.3.	semen a	nd not I	ess than 14 c	days after	the collection of the	datory storage period of frozen semen on samples taken imple taken on
II.4.					sed, stored and transported to D to Directive 92/65/EEC.	under conditions which comply
Notes						
Part I:						
Box I.12:	place	of origin s	hall correspond to	the semen	collection centre of origin of t	he semen.
Box I.13:		of destinat	tion shall correspor	nd to the se	men collection or storage cer	ntre or to the holding of semen
Box 1.23:			container and seal	number sha	all be indicated.	

EUROPEA	UROPEAN UNION Equine semen – Part								
II.	Health information	II.a.	Certificate reference No		II.b.				
Box I.31.:	Box I.31.: donor identity shall correspond to the official identification of the animal.								
	date of collection shall be indicated	d in the follo	owing format: dd/mm/yyyy.						
	approval number of the centre shall I.12 where the semen was collecte		to the approval number of t	the sem	en centre indicated in Box				
Part II:									
(1) Delete	as appropriate.								
(²) Only a websit	approved semen collection centres listed in te:	accordance	with Article 11(4) of Directi	tive 92/6	5/EEC on the Commission				
http://e	ec.europa.eu/food/animal/approved_establis	shments/est	ablishments_vet_field_en.ht	ıtm					
(3) OJ L	192, 23.7.2010, p. 1.								
(4) Cross	out the programme(s) that do(es) not app	ly to the co	onsignment.						
(5) Insert	date.								
— The co	plour of the stamp and signature must be	different fro	om that of the other particul	lars in t	he certificate.				
Official ve	terinarian or official inspector								
N:	Name (in capital letters): Qualification and title:								
Lo	Local veterinary unit: LVU No:								
D	ate:		Sign	nature:					
St	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

EUF	ROPE	AN UNION	Intra trade certificate
	l.1.	Name	I.2. Certificate reference No I.2.a. Local reference No
ъ		Address	I.3. Central competent authority
ente		Postal code	I.4. Local competent authority
consignment presented	I.5.	Consignee Name Address	I.6. No(s) of related original certificates No(s) of accompanying documents
Ĕ		Postal code	1.7.
of consig	1.8.	Country of origin ISO code I.9. Region of origin Code	I.10. Country of destination ISO code I.11. Region of destination Code
ils	l.12.	Place of origin	I.13. Place of destination
eta		Semen Centre □	Semen Centre ☐ Holding ☐
Part I: Details		Name Approval number Address	Name Approval number Address
4		Postal code	Postal code
	l.14.		1.15.
	I.16.	Means of transport	1.17.
		Aeroplane Ship Railway wagon IRoad vehicle Other Identification:	
	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
	1.23.	Seal/Container No	I.24. Type of packaging
	1.25.	Commodities certified for: Artificial reproduction □	'
	1.26.	Transit through third country	I.27. Transit through Member States
		Third country ISO code	Member State ISO code
		Exit point Code	Member State ISO code
		Entry point BIP No	Member State ISO code
	1.28.	Export	1.29.
		Third country ISO code	
		Exit point Code	
	1.30.		
	I.31.	Identification of the commodities	
		Species Donor identity (Scientific name)	Date of collection Quantity

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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** Equine semen - Part D II. II.a. Certificate reference No. II.b. Health information I, the undersigned official veterinarian, hereby certify that the semen described above was collected, processed and stored for a minimum period of 30 days immediately following collection in an approved semen collection centre $\binom{2}{2}$ 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 (1) either [II.1. 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 Part II: Certification (1) either [Part A of Annex I to Decision 2010/470/EU;] (1) or [Part B of Annex I to Decision 2010/470/EU;] [Part C of Annex I to Decision 2010/470/EU;] (1) or [Decision 95/307/EC;]] was collected, processed and stored for a minimum period of 30 days immediately following collection in (1) or []].1. an approved semen collection centre $\binom{2}{2}$ 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;] [Part D of Annex I to Decision 2010/470/EU;] (1) or [Decision 95/307/EC;]] (1) or was collected, processed and stored in an approved semen collection centre $\binom{2}{2}$ 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 (1) or []].1. 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:1] was stored in the approved semen storage centre $\binom{2}{2}$ 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; 11.2. 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. 11.3. 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.

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

EUR	OPEAN	UNION			Equine semen – Part D				
II.	Health information II.a. Certificate reference No II.b.								
Вох	Sox 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 of the semen.	to the ser	nen collection or storage centre o	r to the holding of destination				
Вох	1.23:	The identification of container and seal r	number sh	all be indicated.					
Вох	I.31:	The donor identity shall correspond to the The date of collection shall be indicated							
Part	t II:								
(¹)	Delete	e as appropriate							
(2)	92/65 http://	approved semen collection or storage cen /EEC on the Commission websites: ec.europa.eu/food/animal/approved_establ ec.europa.eu/food/animal/semen_ova/equir	ishments/e	establishments_vet_field_en.htm;	or Article 17(3)(b) of Directive				
(3)	OJ L	268, 14.9.1992, p. 54							
— ·	The co	plour of the stamp and signature must be	different fr	rom that of the other particulars i	n the certificate.				
Offic	cial vet	erinarian 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

	IOFE	AN UNION	Intra trade certificate			
	l.1.	Consignor Name Address	Certificate reference No     I.2.a. Local reference No     I.3. Central competent authority			
ted		Postal code	I.4. Local competent authority			
consignment presented	1.5.	Consignee Name Address	1.7.			
guu		Postal code				
75	1.8.	Country of ISO code I.9. Region of code origin Code	I.10. Country of destination ISO code I.11. Region of destination Code			
tails	I.12.	Place of origin	I.13. Place of destination			
<u>۾</u>		Embryo team □	Holding ☐ Embryo team ☐			
Part I: Details		Name Approval number Address	Name Approval number Address			
		Postal code	Postal code			
	l.14.		1.15.			
	I.16.	Means of transport	1.17.			
		Aeroplane Ship Railway wagon IRoad vehicle Other Identification:				
	I.18.	Description of commodity	I.19. Commodity code (HS code)			
			05 11 99 85			
			I.20. Quantity			
	I.21.	Temperature of products	I.22. Number of packages			
		Ambient ☐ Chilled ☐ Frozen [	<b>_</b>			
	1.23.	Seal/Container No	I.24. Type of packaging			
	1.25.	Commodities certified for:	'			
		Artificial reproduction				
	1.26.	Transit through third country	I.27. Transit through Member States			
		Third country ISO code	Member State ISO code			
		Exit point Code	Member State ISO code			
		Entry point BIP No	Member State ISO code			
	1.28.	Export	1.29.			
		Third country ISO code				
		Exit point Code				
	1.30.					
	I.31.	Identification of the commodities				
		Species Category Donor ide (Scientific name)	entity Date of collection Quantity			

	EUROPEA				Opulificate reference		ova and embryos - Part A
	II.	Health info	ormation	II.a.	Certificate reference	No	II.b.
	I, the und	ersigned off	icial veterinarian, hereby certify	that:			
	(1) either	[II.1.	the <i>in vivo</i> derived embryos/ stored by an embryo collection of Annex D to Directive 92/6	on team (²	) approved and supervi		
	( <sup>1</sup> ) or	[II.1.	the <i>in vivo</i> produced emb processed and stored by ar with Chapter I(III)(1) and (2)	embryo	production team (2), ap	proved and	
fication	(1) either	[II.2.	the <i>in vivo</i> derived embryos of Directive 92/65/EEC;]	described	above meet the requirer	ments of Ch	apter III(II)(1) of Annex D to
(1) either [II.2. the in vivo derived embryos described above meet the requirement Directive 92/65/EEC;]  (1) or [II.2. the in vivo derived ova described above meet the requirements Directive 92/65/EEC;]							oter III(II)(2) of Annex D to
ď	(¹) or	[II.2.	the <i>in vivo</i> produced embryos to Directive 92/65/EEC;]	s describe	d above meet the requi	rements of	Chapter III(II)(3) of Annex D
	(1) or	[II.2.	the micromanipulated embr Annex D to Directive 92/65/8		ribed above meet the	requireme	nts of Chapter III(II)(4) of
		II.3.	the ova or embryos describe	ed above	come from donor mares	s which:	
		II.3.1.	come from holdings fulfilling which only equidae satisfyin Directive 2009/156/EC were	g the con			
		II.3.2.	meet the requirements of Ch	napter IV(4	) of Annex D to Directi	ive 92/65/El	EC;
		II.3.3.	were not used for natural bre the ova or embryos and betv and the date of the collectio	veen the c	late of the first sample		
		II.3.4.	underwent the tests, which r Diagnostic Tests and Vaccin recognised by the compete accreditation in accordance	es for Ter ent authori	restrial Animals of the C ty and has the tests	DIE, carried referred to	out in a laboratory which is hereinafter included in its
		II.3.4.1.	for equine infectious anaemic enzyme-linked immunosorbe taken on	nt assay ( g not less II.3.3, and	ELISA) with a negative than 14 days following the test was last car	result carri- ng the date ried out on	ed out on a blood samples of commencement of the a sample of blood taken
		II.3.4.2.	for contagious equine metritis at least two specimens (swa mucosal surfaces of the clito	abs) taken	during the period refe	rred to in p	oint II.3.3 from at least the

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	UROPEA			Equine ova and embryos - Part A				
the case of isolation of <i>Taylorala equigentalias</i> after cultivation under microaerophilic conditions in a period of at least 7 days, set up within 24 hours after taking the specimens from the donor anim or 48 hours where the specimens are kept cool during transport.]  (1) and/or [II.3.4.2.2. on one cocasion on	II.	Health info	rmation	II.a.	Certificate reference No	II.b.		
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 da (systemic treatment) or 21 days (local treatment) after antimicrobial treatment of the donor stallion as were placed in transport medium with activated charcoal, such as Amies medium, before dispatch the laboratory;  (1) either [II.4. the embryos described above were conceived as a result of artificial insemination of the donor man with semen which was collected, processed, stored and transported under conditions which complying 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 <i>in vitro</i> fertilisation of ova complying with the conditions set out in point 2 of Chapter III(III) of Annex D to Directive 92/65/EEC.]  (1) or [II.4. the own have not been in contact with semen of the equine species.]  (1) or [II.4. the ova have not been in contact with semen of the equine species.]  (1) or [II.5. the ova or embryos described above were sent to the place of loading in a sealed container 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.13: The place of origin shall correspond to the embryo collection team or embryo production team or to the holding of ova/embryos destination shall correspond to the embryo collection team, embryo production team or to the indicated.  Box I.23: The identification of container and seal number shall be indicated.  The identification of container and seal number shall be indicated.  The date of collection shall be indicated in the following format: dd/mm/yyyyy.  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 of the Commission website:	( <sup>1</sup> ) either	[II.3.4.2.1.	the case of isolation of Tay a period of at least 7 days,	<i>rlorella equig</i> set up within	<i>enitalis</i> after cu <sup>l</sup> tivation under r 24 hours after taking the speci	nicroaerophilic conditions fo		
(systemic treatment) or 21 days (local treatment) after antimicrobial treatment of the donor stallion as were placed in transport medium with activated charcoal, such as Amies medium, before dispatch the laboratory;  (1) either [II.4. the embryos described above were conceived as a result of artificial insemination of the donor many with semen which was collected, processed, stored and transported under conditions which comp 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;]  (1) at the ova or embryos described above were sent to the place of loading in a sealed container 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 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 micromanipulated 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/yyyyy.  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 of the Commission website:	( <sup>1</sup> ) and/or	[II.3.4.2.2.	by a polymerase chain reac	tion (PCR) o	case of the detection of genor real-time PCR test, carried ou	me of <i>Taylorella equigenitali</i> it within 48 hours after takin		
with semin which was collected, processed, stored and transported under conditions which comp with the requirements of Chapters I(I), III(I) and III(I) of Annex D to Directive 92/65/EEC with semen which was collected, processed, stored and transported under conditions which comply 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 conditions which comply with the conditions which comply with the requirements of Chapters I(I), III(I) and III(I) of Annex D to Directive 92/65/EEC;  II.5. the ova or embryos described above were sent to the place of loading in a sealed container accordance with point 6 of Chapter III(II) of Annex D to Directive 92/65/EEC and bearing the number detailed in Box 1.23.  Notes  Part I:  Box I.12: The place of origin shall correspond to the embryo collection team or embryo production team or ova/embryo 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 micr 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 of the Commission website:			(systemic treatment) or 21 d were placed in transport me	ays (local tre	atment) after antimicrobial treatr	ment of the donor stallion an		
the conditions set out in point 2 of Chapter III(II) of Annex D to Directive 92/65/EEC with semen whis was collected, processed, stored and transported under conditions which comply with the requirements of Chapters I(I), III(I) and IIII(I) of Annex D to Directive 92/65/EEC;]  II.5. 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 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/embryo 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 micromanipulated 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:	( <sup>1</sup> ) either	[II.4.	with semen which was colle	cted, proces	sed, stored and transported ur	der conditions which compl		
II.5. the ova or embryos described above were sent to the place of loading in a sealed container 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/embryo 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: <i>in vivo</i> derived embryos, <i>in vivo</i> derived ova, <i>in vivo</i> produced embryos or micromanipulated 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 of the Commission website:	( <sup>1</sup> ) or	[II.4.	the conditions set out in poir was collected, processed,	nt 2 of Chapte stored and	er III(II) of Annex D to Directive of transported under condition	92/65/EEC with semen which sometimes which comply with the		
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/embryo 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 micromanipulated 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 of the Commission website:	( <sup>1</sup> ) or	[II.4.	the ova have not been in co	ontact with s	emen of the equine species;]			
Part I:  Box I.12: The place of origin shall correspond to the embryo collection team or embryo production team of ova/embryo 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 micromanipulated 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.  (c) Only approved embryo collection or production teams listed in accordance with Article 11(4) of Directive 92/65/EEC of the Commission website:		II.5.	accordance with point 6 of	f Chapter III				
Box I.12: The place of origin shall correspond to the embryo collection team or embryo production team of ova/embryo 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 micromanipulated 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 of the Commission website:	Notes							
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 micromanipulated 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 of the Commission website:	Part I:							
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 micr 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 of the Commission website:	Box I.12:			o the embryo	collection team or embryo pro	duction team of ova/embryo		
Box I.31: the category: specify if: in vivo derived embryos, in vivo derived ova, in vivo produced embryos or micr 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 of the Commission website:	Box I.13:			pond to the	embryo collection team, embry	o production team or to the		
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 of the Commission website:	Box 1.23:	The ide	ntification of container and se	al number sh	all be indicated.			
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 of the Commission website:	Box I.31:			red embryos,	in vivo derived ova, in vivo p	produced embryos or micro		
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 of the Commission website:		The dor	nor identity shall correspond to	the official	identification of the animal.			
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 of the Commission website:		The date	e of collection shall be indica	ted in the fol	lowing format: dd/mm/yyyy.			
(2) Only approved embryo collection or production teams listed in accordance with Article 11(4) of Directive 92/65/EEC of the Commission website:	Part II:				0			
the Commission website:	( <sup>1</sup> ) Dele	ete as appro	priate.					
http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm	(²) Only the			n teams listed	d in accordance with Article 11(	(4) of Directive 92/65/EEC o		
	http:	//ec.europa.e	eu/food/animal/approved_estal	blishments/es	stablishments_vet_field_en.htm			

EURC	OPEAN 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) (	( <sup>5</sup> ) OJ L 165, 30.4.2004, p. 1.								
( <sup>6</sup> ) I	(6) Insert date.								
<u> </u>	The colour of the stamp and signature must be	different fro	om that of the other particulars in	the certificate.					
Offic	ial veterinarian or official inspector								
	Name (in capital letters):		Qualification and t	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

EUF	UROPEAN UNION Intra trade certificate								
	l.1.	Consignor Name Address	I.2. Certificate reference No     I.2.a. Local reference No     I.3. Central competent authority						
sented		Postal code	I.4. Local competent authority						
consignment presented	I.5.	Consignee Name Address	1.6.						
ignm		Postal code	1.7.						
οę	1.8.	Country of ISO code I.9. Region of Code origin	I.10. Country of destination ISO code I.11. Region of destination Code						
I: Details	I.12.	Place of origin  Embryo team □	I.13. Place of destination  Holding ☐ Embryo team ☐						
Part		Name Approval number Address Postal code	Name Approval number Address Postal code						
	l.14.		I.15.						
	I.16.	Means of transport	1.17.						
		Aeroplane Ship Railway wagon IRoad vehicle Other IRoad vehicle Identification:							
	l.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						
	1.23.	Seal/Container No	I.24. Type of packaging						
		Commodities certified for:							
		Artificial reproduction							
	1.26.	Transit through third country	I.27. Transit through Member States						
		Third country ISO code Exit point Code Entry point BIP No	Member State ISO code  Member State ISO code  Member State ISO code						
	1.28.	Export	1.29.						
		Third country ISO code Exit point Code							
	1.30.								
	I.31.	Identification of the commodities							
		Species Category Donor id (Scientific name)	identity Date of collection Quantity						

II.	Health	information II.a. Certificate reference No II.b.										
I, the und	ersigned	official veterinarian, hereby certify that:										
( <sup>1</sup> ) either	[II.1.	the <i>in vivo</i> derived embryos/ <i>in vivo</i> derived ova (1) described above were collected, procestored by an embryo collection team (2) approved and supervised in accordance with Chap of Annex D to Directive 92/65/EEC;]										
( <sup>1</sup> ) or	[II.1.		ne <i>in vitro</i> produced embryos/micromanipulated embryos (1) described above were produced, rocessed and stored by an embryo production team (2), approved and supervised in accordance ith Chapter I(III)(1) and (2) of Annex D to Directive 92/65/EEC;]									
(1) either	[II.2.	ne in vivo derived embryos described above meet the requirements of Chapter III(II)(1) of Annex D to irective 92/65/EEC;]										
( <sup>1</sup> ) or	[II.2.	the <i>in vivo</i> derived ova described above meet the requirements of Chapter II Annex D to Directive 92/65/EEC;]	l(ll)(2) c									
( <sup>1</sup> ) or	[II.2.	the <i>in vitro</i> produced embryos described above meet the requirements of Chapter III(II)(3) of A Directive 92/65/EEC;]	nnex D t									
( <sup>1</sup> ) or	[II.2.	the micromanipulated embryos described above meet the requirements of Chapter III(II)(4) of to Directive 92/65/EEC;]	f Annex I									
	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/E which only equidae satisfying the conditions laid down in Articles 4 and 5 or Articles 12 to 16 o 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 embryos and between the date of the first sample referred to in points II.3.4 and II.3.5 and the collection of ova and embryos;										
	II.3.4.	have been subjected with negative result to an agar-gel immuno-diffusion test (Coggins to ELISA for equine infectious anaemia carried out on a blood sample on	s take of ova o d take									
	II.3.5.	have been subjected to an agent identification test for contagious equine metritis by is Taylorella equigenitalis after a cultivation of 7 to 14 days carried out with negative result case on samples taken during the past 30 days prior to the date of the first collection embryos from mucosal surfaces of the clitoral fossa and clitoral on two consecutives oestrus periods on	s in eac of ova o sinuse additiona									
( <sup>1</sup> ) either	[11.4.	the embryos described above were conceived as a result of artificial insemination of the dol with semen which was collected, processed, stored and transported under conditions which could be the requirements of Chapters I(I), II(I) and III(I) of Annex D to Directive 92/65/EEC;										

ELIBODEAN LINION

Document Generated: 2023-08-23

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

EUROPEA	OINU P	N	Eq	uine	ova and embryos – Part B				
II.	Health	information	II.a. Certificate reference No		II.b.				
( <sup>1</sup> ) or	[II.4.	conditions in point 2 of Chapter II	ere conceived as a result of <i>in vitro</i> fertilisation of ova complying with the II(II) of Annex D to Directive 92/65/EEC with semen which was collected, ed under conditions which comply with the requirements of Chapters I(I), ctive 92/65/EEC;]						
(1) or	[11.4.	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 accordar with point 6 of Chapter III(II) of Annex D to Directive 92/65/EEC and bearing the number detailed Box I.23.							
Notes									
Part I:									
Box I.12:		ace of origin shall correspond to to to to the conference of the c	he embryo collection team or embryo	produ	ction team of ova/embryos				
Box I.13:		ace of destination shall correspond ova/embryos destination.	to the embryo collection team, embryo	produ	ction team or to the holding				
Box 1.23:	id	entification of container and seal n	umber shall be indicated.						
Box I.31:	Box I.31: category: specify if: <i>in vivo</i> derived embryos, <i>in vivo</i> derived ova, <i>in vitro</i> produced embryos of manipulated embryos.								
	do	onor identity shall correspond to the	e official identification of the animal.						
	da	ate of collection shall be indicated	in the following format: dd/mm/yyyy.						
		oproval number of the team shall c va/embryos collection/production.	orrespond to the embryo collection tea	m or	embryo production team of				
Part II:									
(1) Delete	as app	propriate.							
		d embryo collection or production te vebsite:	ams listed in accordance with Article 11	(4) of	Directive 92/65/EEC on the				
http://e	c.europ	ea.eu/food/animal/approved_establis	hments/establishments_vet_field_en.htm	n					
(3) Insert	date.								
(4) OJ L	192, 23.	.7.2010, p. 1.							
— The co	— The colour of the stamp and signature must be different from that of the other particulars in the certificate.								
Official ve	terinaria	n or official inspector							
Nai	me (in c	capital letters):	Qualification	and	title:				
Loc	al veter	rinary unit:	LVU No:						
Dat	e:		Signature:						
Sta	mp:								
I									

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

EUF	ROPE	AN UNION	Intra trade certificate					
	l.1.	Consignor Name Address	I.2. Certificate reference No     I.2.a. Local reference No     I.3. Central competent authority					
ented		Postal code	I.4. Local competent authority					
consignment presented	I.5.	Consignee Name Address	1.6.					
gnn		Postal code	1.7.					
φ	1.8.	Country of origin ISO code I.9. Region of origin Code	I.10. Country of destination ISO code I.11. Region of destination Code					
Part I: Details	I.12.	Place of origin  Embryo team   □	I.13. Place of destination  Holding ☐ Embryo team ☐					
art		Name Approval number Address	Name Approval number Address					
ď		Postal code	Postal code					
	l.14.		1.15.					
	I.16.	Means of transport	1.17.					
		Aeroplane Ship Railway wagon Shoad vehicle Other Identification:						
	l.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					
	1.23.	Seal/Container No	I.24. Type of packaging					
	1.25.	Commodities certified for: Artificial reproduction □	'					
	1.26.	Transit through third country	I.27. Transit through Member States					
		Third country ISO code	Member State ISO code					
		Exit point Code Entry point BIP No	Member State ISO code Member State ISO code					
	1.28.	Export	1.29.					
		Third country ISO code Exit point Code						
	1.30.							
	I.31.	Identification of the commodities						
		Species Category Donor ide (Scientific name)	entity Date of collection Quantity					

Document Generated: 2023-08-23

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** Equine ova and embryos - Part C Health information Certificate reference No II.b. I, the undersigned official veterinarian, hereby certify that: Ova/embryos (1) described above were collected by a collection team (2) approved by the competent authority and II.1. processed in an appropriate laboratory; Ova/embryos (1) were collected from donor mares which: 11.2. on the day of collection have been located in premises situated on the territory or in the case of regionalisation in a II.2.1. part of the territory of a Member State which is not considered to be infected with African horse sickness in Part II: Certification accordance with Article 5(2)(a) and (b) of Directive 2009/156/EC (3), 11.2.2. 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, 11.2.3. have been kept prior to the collection in holdings free from clinical signs of contagious equine metritis for 60 days, 11.2.4. have not been used for natural breeding during the period of 30 days prior to the collection of ova/embryos (1), 11.2.5. 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), 11.2.6. have on the day of collection not shown clinical signs of an infectious or contagious disease: Ova/embryos (1) were collected, processed, stored and transported under conditions which comply with the 11.3. requirements of Annex D of Directive 92/65/EEC; 11.4. The semen used for the artificial insemination of the donor mares complies with the requirements of Directive 92/65/EEC (4)(1); The ova used for the in vitro production of embryos comply with the requirements of Directive 92/65/EEC (1). 11.5. Notes Part I: Box I.12: place of origin shall correspond to the embryo collection team of ova/embryos collection. place of destination shall correspond to the embryo collection team, embryo production team or to the holding of Box I.13: ova/embryos destination. identification of container and seal number shall be indicated. Box 1.23: Box I.31: category: specify if: in vivo derived embryos, in vivo derived ova, in vitro produced embryos or micromanipulated 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. Only approved embryo collection teams listed in accordance with Article 11(4) of Directive 92/65/EEC on the $(^{2})$ Commission website:

http://ec.europa.eu/food/animal/approved establishments/establishments vet field en.htm

EUR	DPEAN UNION		Equine	ova and embryos – Part C					
II.	Health information	II.a.	Certificate reference No	II.b.					
(3)	( <sup>3</sup> ) OJ L 192, 23.7.2010, p. 1.								
( <sup>4</sup> )	(4) Does not apply to ova.								
_	The colour of the stamp and signature must be	different fr	om that of the other particulars in	the certificate.					
Offic	cial veterinarian or official inspector								
	Name (in capital letters):		Qualification and t	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

# F2PART 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]

EUR	OPEAN	N UNION									Intra trade ce	rtificate		
	Name						1.2.	Certificate reference No I.2.a. Local reference No				erence		
							1.3.	Central co	mpetent auth	ority				
	Postal code							I.4. Local competent authority						
-	1.5.	5. Consignee Name Address						I.6.						
Part I: Details of consignment presented								1.7.						
ment p		Postal code												
onsign	1.8.	Country of origin	ISO code	1.9.	Region of origin	Code	I.10.	Country of destination		l.11	. Region of destination	Code		
ls of c														
Detai	I.12.	12. Place of origin Semen centre □					I.13. Place of destination							
Part I:							Semen centre ☐ Holding ☐							
_		Name Approval number				r	Name Approval number							
		Address						Address						
		Postal cod	е				Postal code							
	1.14.						1.15.							
	I.16.	Means of t	ransport				I.17.							
		Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification:												
	I.18. Description of commodity						I.19. Commodity code (CN code) 05 11 99 85					e)		
										1.20	). Quantity			

I.21.	Temperature of products					1.22.	Number of packages	
	Ambient	Chilled		Frozen				
I.23.	Seal/Container No					1.24.	Type of packaging	
1.25.	Commodities certified for	•						
	Artificial reproduction							
1.26.	Transit through third cour	ntry 🗆	1.27.	Transit through	Membe	er State	es 🗆	
	Third country	ISO code		Member State		ISO co	ode	
	Exit point	Code		Member State		ISO code		
	Entry point	BIP No		Member State		ISO code		
1.28.	Export		1.29.					
	Third country	ISO code						
	Exit point	Code						
I.30.								
I.31.	Identification of the comm	nodities						
	Species Breed entific name)	Donor identity		Date of collection	numbe	roval er of the ntre	Quantity	

### **European Union**

### Ovine and caprine semen — Part A

		<b></b>		O Timo dina daprinio domen					
	II.	Health in	formation	II.a. Certificate reference No	II.b.				
	I, the under	rsigned offic	cial veterinarian, hereby	certify that:					
		II.1. The semen described above:							
ation		SI		d and stored in a semen collection ent authority in accordance with Chap 1/65/EEC;					
of Annex D to Directive 92/65/EEC;  II.1.2. comes from donor animals which meet the requirements of Chapter II(II) of An Directive 92/65/EEC;  II.1.3. was collected, processed, stored and transported under conditions which comply requirements of Chapters II(II) and III(I) of Annex D to Directive 92/65/EEC;									
Part II:				, stored and transported under condi II(II) and III(I) of Annex D to Directive 9					
	(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	th be C w	e collection on a holdir efore the collection with hapter A of Annex VIII to ere kept at a semen col	s which have been kept continuously fing or holdings which has/have comp the requirements laid down in points a Regulation (EC) No 999/2001, except lection centre that complied during the of point 1.3(c)(iv) of that Section;]	lied for the last three years 1.3(a) to (f) of Section A of t during the period when they				
	(¹) or	or	zone of a Member Sta	als which have been kept continuously since birth in a Member State State listed in point 2.3 of Section A of Chapter A of Annex VIII to /2001 as having a negligible risk status for classical scrapie;]					
	(1) or	[II.1.4. w	as collected from ovine a	e animals of the ARR/ARR prion protein genotype;]					
		С		of loading in a sealed container in accordance with point 1.4 of D to Directive 92/65/EEC and bearing the number detailed in					
	(1) either	[II.2. N	o antibiotics or no mixtur	ure of antibiotics were added to the semen.]					
	(¹) or		he following antibiotic or te final diluted semen of	or combination of antibiotics was added to produce a concentration of not less than $(^3)$ :					
					]				
	Notes								
	Part I:								
	Box I.12:	Place of or	igin shall correspond to t	he semen collection centre of origin of	the semen.				
	Box I.13:	Place of de semen des		nd to the semen collection or storage	e centre or to the holding of				
	Box I.23:	Identification	on of container and seal i	number shall be indicated.					
	Box I.31:	Donor iden	tity shall correspond to the	ne official identification of the animal.					
		Date of col	lection shall be indicated	in the following format: dd/mm/yyyy.					
			umber of the centre shall where the semen was c	I correspond to the approval number o ollected.	of the semen centre indicated				

**European Union** 

Name (in capital letters):

Local veterinary unit:

Date:

Stamp:

Ovine and caprine semen — Part A

Qualification and title:

LVU No:

Signature:

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

II.	Health information	II.a. Certificate reference No	II.b.							
Part	Part II:									
( <sup>1</sup> )	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.									
Offic	cial veterinarian or official inspector									

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

EUR	OPEA	N UNION	Intra trade certificate
	l.1.	Consignor	I.2. Certificate reference No I.2.a. Local reference No
		Name Address	I.3. Central competent authority
		Postal code	
nted			I.4. Local competent authority
rese	1.5.	Consignee	1.6.
ᇦ		Name Address	
l me			1.7.
lsig	_	Postal code	
s of consignment presented	1.8.	Country ISO code I.9. Region of origin Code of origin	I.10. Country of ISO code I.11. Region of Code destination
etail	110	Place of origin	140 50 4 4 4 4 4 4
Part I: Details	1.12.	Semen centre	I.13. Place of destination  Semen centre Holding H
Part		_	
		Name Approval number Address	Name Approval number Address
		Postal code	Postal code
	1.14.		1.15.
_	1.16.	Means of transport	1.17.
		Aeroplane Ship Railway wagon	
		Road vehicle Other	
		Identification	
	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
	1.23.	Seal/Container No	I.24. Type of packaging
	1.25.	Commodities certified for:	
		Artificial reproduction	
	1.26.	Transit through third country	I.27. Transit through Member States
		Third country ISO code	Member State ISO code
		Exit point Code Entry point BIP No	Member State ISO code  Member State ISO code
	100		1.29.
	1.28.	Export  Third country ISO code Exit point Code	1.20.
	1.30.		
	1.31.	Identification of the commodities  Species Breed Donor identity (Scientific name)	Date of collection Approval number Quantity of the centre

-	EUROPE/	IN UNION		Ovine and caprine semen — Part B						
	II. I	Health info	rmation	II.a. Certificate reference No	II.b.					
_	I, the und	dersigned	official veterinarian, hereby certify that the semen des	scribed above:						
		II.1. was collected, processed and stored in a semen collection centre (°) 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;								
<u> </u>		II.2.	comes from the donor animals which meet the requirements of Chapter II(II) of Annex D to Directive 92/65/EEC;							
Part II: Certification		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;							
ਜ਼ੂ ਜ਼	( <sup>1</sup> ) either	[II.4.	meets the requirements of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001;]							
Pa 	(¹) or	[11.4.	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 ( <sup>3</sup> ) requested by the Member State of destination.]							
$\dashv$	Notes									
	Part I:									
	Box I.12:	Place of	origin shall correspond to the semen collection centre	e 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 1.23:	Identifica	tion of container and seal number shall be indicated.							
	Box I.31:	Donor ide	entity shall correspond to the official identification of t	the animal.						
		Date of o	collection shall be indicated in the following format: do	d/mm/yyyy.						
		Approval collected	number of the centre shall correspond to the approva	I number of the semen centre indicate	d in Box I.12 where the semen was					
	Part II:									
	(1) Delete	as appro	priate.							
			semen collection centres listed in accordance with A eu/food/animal/approved_establishments/establishmen		E/EEC on the Commission website:					
	(3) Additi	onal guara	intees as laid down in Article 2 of Regulation (EC) N	o 546/2006 (OJ L 94, 1.4.2006, p. 28	3).					
	Official v	eterinarian	or official inspector (*)							
	Name	(in capita	I letters):	Q	ualification and title:					
	Local	veterinary	unit:	L	/U No:					
	Date:			Si	gnature:					
	Stamp	):								
	(*) The c	olour 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

EUR	PEA	N UNION		Intra trade certificate				
	l.1.	•	I.2. Certificate reference No	I.2.a. Local reference No				
		Name Address	I.3. Central competent authority					
Part I: Details of consignment presented		Postal code	I.4. Local competent authority					
presen	1.5.	Consignee Name	I.6. No(s) of related original certificates	No(s) of accompanying documents				
ment		Address	1.7.					
sigr	_	Postal code						
of con	1.8.	Country ISO code I.9. Region of origin Code of origin	I.10. Country of ISO code destination	e I.11. Region of Code destination				
tails	1.12.	Place of origin	I.13. Place of destination					
: De		Semen centre	Semen centre	Holding				
Part		Name Approval number Address	Name Address	Approval number				
		Postal code	Postal code					
	l.14.		1.15.					
	I.16.	Means of transport	1.17.					
		Aeroplane ☐ Ship ☐ Railway wagon ☐						
		Road vehicle Other						
		Identification						
	I.18.	Description of commodity	I.19. Commodity code (HS code) 05 11 99 85					
				I.20. Quantity				
	I.21.	Temperature of products		LOO Nambers of sections				
		Ambient ☐ Chilled ☐	Frozen	22. Number of packages				
	1.23.	Seal/Container No		I.24. Type of packaging				
	1.25.	Commodities certified for:						
	100	Artificial reproduction   Transit through third country	I.27. Transit through Member S	States				
	1.20.	Transit through third country  Third country  ISO code	Member State	ISO code				
		Exit point Code	Member State	ISO code				
		Entry point BIP No	Member State	ISO code				
	1.28.	Export	1.29.					
		Third country ISO code Exit point Code						
	1.30.							
	I.31.	Identification of the commodities						
		Species Breed Donor identity (Scientific name)		oval number Quantity he centre				

EUROPEA	NOINU NA		Ovine and caprine semen — Part C				
II. I	Health inform	aation	II.a. Certificate reference No	II.b.			
I, the und	dersigned off	icial veterinarian, hereby certify that the semen de	scribed above:				
( <sup>1</sup> ) either	[II.1.	was collected, processed and stored for a mini semen collection centre (2) situated in the Me accordance with Chapter I(I)(1) and Chapter I(II) moved to the semen storage centre detailed in F animal health and veterinary certification at least	ember State of origin of the semen (1) of Annex D to Directive 92/65/EE0 Part I.12 situated in the same Member	and operated and supervised in C, and from where the semen was			
	(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 mini semen collection centre (²) situated in the Europ and Chapter ((II) of Annex D to Directive 92/65/E accordance with:	ean Union and operated and supervis	sed in accordance with Chapter I(I			
	(1) either	[Part A of Annex III to Decision 2010/470/EU;]					
1	(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 mini semen collection centre $\binom{2}{2}$ situated in a third col- operated and supervised in accordance with Cha imported into the European Union under the cor-	untry or part(s) thereof listed in Annex oter I(I)(1) and Chapter I(II)(1) of Annex	I to Decision 2010/472/EU which is D to Directive 92/65/EEC, and was			
	(1) either	[Section A of Part 2 of Annex II to Decision 201	)10/472/EU;]				
	(¹) or	[Section B of Part 2 of Annex II to Decision 201	)10/472/EU;]				
	(¹) or	[Annex II to Decision 2008/635/EC;]					
	II.2.	was stored in the approved semen storage centre with Chapter $I(I)(2)$ and Chapter $I(I)(2)$ of Annex		rated and supervised in accordance			
	II.3.	was sent to the place of loading in a sealed cont 92/65/EEC and bearing the number indicated in		Chapter III(I) of Annex D to Directive			
Notes							
Part I:							
Box 1.6:	panied the	spond to the serial number of the individual official semen described above from the approved semen of this/these document(s) or certificate(s), or the continuous co	collection centre of its origin to the desc	cribed above semen storage centre			
Box I.12:	Place of ori	igin shall correspond to the semen storage centre	of dispatch of the semen.				
Box I.13:	Place of de	estination shall correspond to the semen collection	or storage centre or to the holding of	destination of the semen.			
Box 1.23:	Identification	n of container and seal number shall be indicated.					
Box I.31:	Date of col	tity shall correspond to the official identification of the lection shall be indicated in the following format: describes of the contract of t	d/mm/yyyy.	ntre of the semen origin			
	Approval nu	umber of the centre shall correspond to the approx	ral number of the semen collection ce	ntre of the semen origin.			

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	Equi	ne 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/establishme http://ec.europa.eu/food/animal/semen_ova/ovine/index_en.htm	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):	Quali	fication 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 pa	articulars in the certificate.								

### ANNEX IV

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

### [F3PART 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

EURO	PEAN	UNION	Intra trade certificat						
	l.1.	Consignor Name	I.2. Certificate reference No I.2.a. Local reference No						
		Address	I.3. Central competent authority						
ted		Postal code	I.4. Local competent authority						
resen	1.5.	Consignee Name	1.6.						
ment p		Address	1.7.						
ign		Postal code							
Part I: Details of consignment presented	1.8.	Country of ISO I.9. Region of Code origin code origin	I.10. Country of destination ISO code I.11. Region of destination Code						
l: Detail	I.12.	Place of origin Embryo team □	I.13. Place of destination Holding ☐ Embryo team ☐						
Part		Name Approval number Address	Name Approval number Address						
		Postal code	Postal code						
	I.14.		1.15.						
	I.16.	Means of transport	1.17.						
		Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification							
	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						
	1.23.	Seal/Container No	I.24. Type of packaging						
	1.25.	Commodities certified for: Artificial reproduction							
	1.26.	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						
	1.28.	Export ISO code Exit point Code	1.29.						
	1.30.								
	I.31. (S	Identification of the commodities Species Category Donor identity Scientific name)	Date of collection Approval number of the team Quantity						

EUROPEA	N UNION				Ovine an	d caprine ova/embryos — Par				
II.	Health inf	ormation	II.a.	Certificate reference	e number	II.b.				
I, the unde	ersigned o	official veterinarian	, hereby certify t	that:						
( <sup>1</sup> ) either	[II.1.		bryo collection t	team (²) approved and		e were collected, processed a cordance with Chapter I(III)(1)				
( <sup>1</sup> ) or	[II.1.	processed and s	the <i>in vitro</i> 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.		ne in vivo derived embryos described above meet the requirements of Chapter III(II)(1) of Annex D to Directive 92/65/EEC;]							
( <sup>1</sup> ) or	[11.2.	the <i>in vivo</i> derive 92/65/EEC;]	ne in vivo derived ova described above meet the requirements of Chapter III(II)(2) of Annex D to Directive 2/65/EEC;]							
( <sup>1</sup> ) or	[II.2.	he <i>in vitro</i> produced embryos described above meet the requirements of Chapter III(II)(3) of Annex D to Directive 92/65/EEC;]								
( <sup>1</sup> ) or	[11.2.	the micromanipul Directive 92/65/E		described above meet	the requirements	of Chapter III(II)(4) of Annex D				
	( <sup>1</sup> ) [II.3.	the consignment	consists of emb	ryos of the ovine or ca	prine species whicl	h:				
	(¹) either	recognised as ha	aving a negligib		of classical scrap	e birth on a holding or holdin pie in accordance with point 1				
	( <sup>1</sup> ) or	collection on a h	olding or holding down in points	gs which have complie	ed for the last three	the last three years before to e years before collection with to pter A of Annex VIII to Regulation				
	( <sup>1</sup> ) or	Member State w	ith a negligible	risk status for class	ical scrapie appro	th in a Member State or zone or ved in accordance with the fi ulation (EC) No 999/2001;]]				
	(1) or	[were collected fr	om ovine anima	ls and						
		(1) either [are	of the ARR/ARR	R prion protein genotyp	e;]]					
		(1) or [carr	y at least one Al	RR allele and were col	lected after the dat	te of 1 January 2015;]]				
	II.4.			bove come from fema r IV(3) of Annex D to D		vine (¹)/caprine species (¹) whi				
(¹) either	[11.5.	semen which wa	s collected, pro		ansported under o	nination of the donor females w conditions which comply with t EEC;]				
( <sup>1</sup> ) or	[11.5.	conditions in Ch	apter III(II)(2) of and transported	of Annex D to Directed under conditions w	tive 92/65/EEC wi	isation of ova complying with t ith semen which was collecte the requirements of Chapters I				
(1) or	[11.5.	the ova have not	been in contact	with semen of the ovir	ne and caprine spe	cies;]				
	II.6.					a sealed container in accordan bearing the number detailed				
Notes										
Part I:										
Box I.12:		of origin shall con/production.	orrespond to the	ne embryo collection	team or embryo	o production team of embry				
Box I.13:		f destination shall bryos destination.	correspond to t	the embryo collection	team, embryo pro	duction team or to the holding				

**EUROPEAN UNION** 

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

EUROPEA	OPEAN UNION Ovine and caprine ova/embryos						
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 derive embryos.	ed embry	vos, in vivo derived ova, in vitro produ	iced embryos or micromanipulated			
Donor identity shall correspond to the official identification of the animal.							
	Date of collection shall be indicated	d in the fo	ollowing format: dd/mm/yyyy.				
	Approval number of the team sl ova/embryos collection/production.		espond to the embryo collection tea	m or embryo production team of			
Part II:							
(¹) Dele	ete as appropriate.						
	y approved embryo collection or prod nmission website:	duction to	eams listed in accordance with Article	11(4) of Directive 92/65/EEC on the			
http:	://ec.europa.eu/food/animal/approve	d_establi	shments/establishments_vet_field_en.h	ntm.			
— The	colour of the stamp and signature m	ust be di	ifferent from that of the other particulars	in the certificate.			
Official ve	eterinarian 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

EUR	JROPEAN UNION Intra trade certificate											
		Consignor Name				I.2. Ce	ertificate	reference No		I.2.a. L	ocal referen	ce No
		Address				I.3. Central competent authority						
nted		Postal code				I.4. Lo	cal com	petent authority				
ese	1.5.	Consignee				I.6.						
ğ	Name											
Jen		Address			ľ	1.7.						********************
signm		Postal code							******			
Part I: Details of consignment presented		Country of origin	ISO code	I.9. Region of origin C	ode		Country of destination		ISO code	l.11.	Region of destination	Code
stail	112	Place of origin			_	I 13 F	Place of	destination				
ä			Embryo tear	m 🔲			1400 01	Holding	ı 🗆		Embryo tea	am 🔲
Ĕ		Name	•	Approval number		١	Name	·	_		Approval n	
ď		Address		7,7		,	Address				,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	
		Postal code					Postal co	ode				
	1.14					l.15.						
	1.16	Means of transport				1.17.						
		Aeroplane	Ship	Railway wagon								
		Road vehicle	Other	, , , –								
		Identification	_									
	1.18	Description of commod	ditv					I.19. Commodity	/ code	(HS cod	e)	
			,							05 11		
									1.2	0. Quar	ntity	
	1.21	Temperature of produc	ets				_		1.2	2. Num	ber of packa	ges
		Ambient		Chilled		Frozen						
	1.23	Seal/Container No							1.2	4. Type	of packagin	ng
	1.25	Commodities certified	for:									
		Artificial reproduction [										
	1.26	Transit through third or	ountry		T I	I.27. <b>T</b>	ransit th	rough Member S	States			
		Third country		ISO code		N	1ember 8	State			ISO cod	le
		Exit point		Code		M	1ember 8	State			ISO cod	le
		Entry point		BIP No		N	1ember 8	State			ISO cod	le
	1.28	Export [				1.29.						
		Third country		ISO code								
		Exit point		Code								
	1.30											
	1.31	Identification of the cor	mmodities									
		Species	Breed	Category Do	nor ide	entity		Date of		val num		Quantity
		(Scientific name)					co	ollection	of	the tean	1	

	EUROPEA	N UNION		Ovine and caprine ova/embryos — Part						
	II.	Health in	formation	II.a. Certificate reference No	II.b.					
	I, the und	lersigned	official veterinarian, hereby certify that the ova/embryo	os (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 92/65/EEC;								
tification	( <sup>1</sup> ) 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.]							
The latest the requirements of Chapter III of Annex D to Directive 92/65/EEC and of Chapter A(I) of Annex VIII to find No 999/2001.]  meet the requirements of Chapter III of Annex D to Directive 92/65/EEC and of Chapter A(I) of Annex VIII to find No 999/2001 and are destined for a Member State which benefits, for all or part of its territory, from the provision point (b) or (c) of Chapter A(I) of Annex VIII to find find the donor animals comply rewith the guarantees provided for by the programmes referred to in that point and with the guarantees (2) rewith the guarantees (3) rewith the guarantees (4) rewith the guarantees (5) rewith the guarantees (6) rewith the guarantees (7) rewith the guarantees (8) rewith the guarantees (9) rewith the guarantees (9) rewith the guarantees (1) rewith the guarantees (2) rewith the guarantees (3) rewith the guarantees (4) rewith the guarantees (4) rewith the guarantees (4) rewith the guarantees (5) rewith the guarantees (6) rewith the guarantees (8) rewith the guarantees (1) rewith t					itory, from the provisions laid down in nor animals comply regarding scrapie					
	( <sup>1</sup> ) either	[II.4.	in the case of embryos, the semen used for fertilisat of Chapter A(I) of Annex VIII to Regulation (EC) No		tive 92/65/EEC and the requirements					
	(¹) or	[11.4.	in the case of embryos, the semen used for fertilisat of Chapter A(I) of Annex VIII to Regulation (EC) No 9 of its territory, from the provisions laid down in point and the donor animals comply regarding scrapie with and with the guarantees (2) requested by the Memb	99/2001 and is destined for a Memb (b) or (c) of Chapter A(l) of Annex to the guarantees provided for by the	er State which benefits, for all or part VIII to Regulation (EC) No 999/2001					
	Notes									
	Part I:									
	Box I.12:	Place of	origin shall correspond to the embryo collection team	n of ova/embryos collection.						
	Box I.13:	Place of destination	destination shall correspond to the embryo collection.	tion team, embryo production tean	n or to the holding of ova/embryos					
	Box 1.23:	Identifica	ation 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:										
		al veterina	ary unit:	LVU No:						
	Date			S	gnature:					
	Stamp:									
	(*) The col	our of the s	stamp and signature must be different from that of the other p	articulars in the certificate.						

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

### 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

	JFE,	AN UNION	Intra trade certifica					
		Consignor	I.2. Certificate reference No I.2.a. Local reference No					
		Name Address	I.3. Central competent authority					
Ited		Postal code	I.4. Local competent authority					
reser	1.5.	Consignee	l.6.					
ן בַּ		Name						
9		Address	1.7.					
ll l		Postal code						
ran i: Details of consignment presented		Country of ISO code I.9. Region of origin Code origin	I.10. Country of destination ISO I.11. Region of Code destination Code					
Detail	l.12.	Place of origin  Embryo team □	I.13. Place of destination  Holding   Embryo team   ■					
<u>:</u>		-	Name Approval number					
,au		Name Approval number						
۱ -		Address	Address					
		Postal code	Postal code					
	1.14		l.15.					
	I.16.	Means of transport	1.17.					
		Aeroplane Ship Railway wagon						
		Road vehicle Other						
		Identification						
ı	I.18.	Description of commodity	I.19. Commodity code (HS code) 05 11 99 85					
			I.20. Quantity					
ł	1 21	Temperature of products	I.22. Number of packages					
	1.21.	·	zen 🗆					
	1.23.	Seal/Container No	I.24. Type of packaging					
ŀ	125	Commodities certified for:						
	1.25.							
	1.00	Artificial reproduction	107 Tarak Maraha Sahara					
	1.26.	Transit through third country	I.27. Transit through Member States					
		Third country ISO code	Member State ISO code					
		Exit point Code	Member State ISO code					
		Entry point BIP No	Member State ISO code					
	1.28.	Export	1.29.					
		Third country ISO code						
		Exit point Code						
	1.30.							
	I.31.	Identification of the commodities Species Breed Category Donor iden (Scientific name)	tity Date of Approval number Quantity collection of the team					

	E	EUROPEAN UNION			Porcine ova/embryos — Part A						
		II.	Health inf	ormation	II.a. Certificate reference No	II.b.					
Г	4	I, the und	lersigned o	official veterinarian, hereby certify that the ova/embryo	os (1) described above:						
		( <sup>1</sup> ) either	II.1.	were produced/collected (1), processed and stored by accordance with Chapter I(III) of Annex D to Directive		team (2) approved and supervised in					
			II.2.	meet the requirements of Chapter III(II) of Annex D	to Directive 92/65/EEC;						
	II: Certification		II.3.	come from donor females of the porcine species which meet the requirements of Chapter IV(2) of Annex D to D 92/65/EEC;							
	: Cert		[II.4.	are in vivo derived embryos which:							
	Part		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:							
			( <sup>1</sup> ) either	[listed in Annex I to Decision 2008/185/EC and are de 2008/185/EC;]	estined for a Member State or region	thereof listed in Annex I to Decision					
L			(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 d 2008/185/EC and have been washed with trypsin;]	estined for a Member State or region	thereof listed in Annex I to Decision					
			( <sup>1</sup> ) or	[listed in Annex II to Decision 2008/185/EC and are do 2008/185/EC;]	estined for a Member State or region	thereof listed in Annex II to Decision					
			( <sup>1</sup> ) or	[not listed in Annex I or II to Decision 2008/185/EC art to Decision 2008/185/EC and have been washed with the control of the	nd are destined for a Member State or region thereof listed in Annex I or II lith trypsin;]						
			( <sup>1</sup> ) 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 v	d/micromanipulated (¹) embryos which:						
			II.4.1.	were conceived as a result of in vitro fertilisation wit	th semen meeting the requirements of Directive 90/429/EEC,						
			II.4.2.	originate from a Member State or region thereof:							
			( <sup>1</sup> ) either	[listed in Annex I to Decision 2008/185/EC and are do 2008/185/EC;]	estined for a Member State or region	thereof listed in Annex I to Decision					
			( <sup>1</sup> ) or	[listed in Annex I to Decision 2008/185/EC and are d Decision 2008/185/EC;]	estined for a Member State or region	thereof not listed in Annex I or II to					
			(¹) or		ecision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision donor females of the ova used for their production comply with the conditions of Article 1 of Decision						
			( <sup>1</sup> ) or	[listed in Annex II to Decision 2008/185/EC and are do 2008/185/EC;]	destined for a Member State or region thereof listed in Annex II to Decision						
			(¹) or		ed 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 sion 2008/185/EC and the donor females of the ova used for their production comply with the conditions of Article 1 of n 2008/185/EC;]						
			(¹) or	[not listed in Annex I or II to Decision 2008/185/EC ar or II to Decision 2008/185/EC;]]	and are destined for a Member State or region thereof not listed in Annex I						

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

EUROPE	UROPEAN UNION Porcine ova/embryos — P								
II.	Health info	rmation	II.a. Certificate reference No	II.b.					
(¹) or	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 De 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 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 De 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 a 2008/185/EC;]	destined for a Member State or region	n thereof listed in Annex II to Decision					
	( <sup>1</sup> ) or	[not listed in Annex I or II to Decision 2008/185/EC at Decision 2008/185/EC and which come from 2008/185/EC;]							
	(¹) or	[not listed in Annex I or II to Decision 2008/185/EC a or II to Decision 2008/185/EC;]]	and are destined for a Member State	or region thereof not listed in Annex I					
	II.5.	were sent to the place of loading in a sealed contain Directive 92/65/EEC and bearing the number details		point 6 of Chapter III(II) of Annex D to					
Notes									
Part I:									
Box I.12	2: place of	origin shall correspond to the embryo collection team	n or embryo production team of ova/	embryos collection/production.					
Box I.13	3: place of	destination shall correspond to the embryo collection t	team, embryo production team or to the	ne holding of ova/embryos destination.					
Box I.23	3: identifica	ation of container and seal number shall be indicated.							
Box I.3	1: category	r: specify if: in vivo derived embryos, in vivo derived	ova, in vitro produced embryos or mi	icromanipulated embryos.					
		entity shall correspond to the official identification of							
		collection shall be indicated in the following format: d I number of the team shall correspond to the embry	****	etion team of ova/embryos collection/					
		on indicated in Box I.12.	o concentration to an emplye product	Not tour of orwants, you concern					
Part II:									
1 '-'	ete as appro	•							
web	site:	embryo collection or production teams listed in accord	, ,	ective 92/65/EEC on the Commission					
http:	//ec.europa.	.eu/food/animal/approved_establishments/establishmen	nts_vet_field_en.htm						
Official	Official veterinarian or official inspector (*)								
Na	ame (in cap								
Local veterinary unit:			LVU No:						
Da	ate:		Signature:						
St	Stamp:								
(*) The c	*) 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

EUR	UROPEAN UNION Intra trade certificate											
		I.1. Consignor Name					I.2. Certificate reference No I.2.a. Local reference No					
		Address				I.3. Central competent authority						
ted	Postal code				1.4. L	ocal Cor	mpetent Authori	ity				
Part I: Details of consignment presented	1.5.	Consignee				1.6.						
		Name						***************************************				
	.	Address				1.7.						
signn	Postal code											
of con		Country of origin	ISO code	I.9. Region of origin	Code	I.10.	Country destination		ISO code		egion of estination	Code
ails	<u> </u>											
: Det	I.12. Place of origin Embryo team □					I.13.	Place of	destination Holdi	ing 🔲	E	mbryo team	
Έ		Name	,	Approval number		Name		•	A	pproval num	ber	
å		Address		,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,			Address					
		Postal code					Postal c					
	1.14					l.15.	1 00141 0					
	I.16.	Means of transport	~··		_	l.17.				_		
		Aeroplane	Ship 🔲	Railway wagon [	_							
		Road vehicle	Other									
		Identification										
	I.18.	I.18. Description of commodity				I.19. Commodity code (HS code) 05 11 99 85						
						I.20. Quantity						
	1.21.	Temperature of produc	ots			I.22. Number of packages						
		Ambient	Ch	illed 🗌	Fro	zen 🗆						
	1.23.	Seal/Container No				I.24. Type of packaging						
	1.25.	Commodities certified	for:									
		Artificial reproduction [										
	1.26.	Transit through third o	ountry			1.27.	Transit th	rough Member	States			
		Third country		ISO code			Member	State			ISO code	
		Exit point		Code			Member	State			ISO code	
	Entry point BIP No		BIP No			Member State						
	1.28.	Export [				1.29.						
		Third country		ISO code								
		Exit point		Code								
	1.30.											
	1.31.	I.31. Identification of the commodities										
		Species (Scientific name)	Breed	Category Do	onor ide	ntity		ate of ection		/al number ne team		Quantity

	EUROPEA	OPEAN UNION Porcine ova/embryos -								
	II.	Health in	formation	II.a. Certificate reference No	II.b.					
	I, the und	lersigned	official veterinarian, hereby certify that:							
		II.1.	The ova/embryos (1) described above:							
		II.1.1.	were collected, processed and stored under condition	ons which meet the requirements of	Directive 92/65/EEC;					
		II.1.2.	II.1.2. come from donor female swine which meet the requirements of Chapter IV of Annex D to Directive 92/65/EEC;							
tion		II.1.3.	meet the requirements of Chapter III of Annex D to	Directive 92/65/EEC.						
ertifica	(1) either	[II.2.	In the case of embryos,							
Part II: Certification		II.2.1.	I.2.1. the semen used for fertilisation meets the requirements of Directive 90/429/EEC;							
Ъ		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 dono 2008/185/EC $(^2)$ .]	r female swine which meets the o	conditions of Article 1 of Decision					
	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 collect destination.			ion team, embryo production team	or to the holding of ova/embryos					
	Box 1.23:	identifica	tion of container and seal number shall be indicated.							
	Box I.31:		: specify if: in vivo derived embryos, in vivo derived of		cromanipulated embryos.					
			entity shall correspond to the official identification of t collection shall be indicated in the following format: do							
			number of the team shall correspond to the embryo	****	ection indicated in Box I.12.					
	Part II:									
		as appro	•							
	2008/1 Memb	185/EC (O	oplies only to ova and embryos which originate in the N J L 59, 4.3.2008, p. 19) and destined to the Member or regions thereof listed in Annex II of Decision 2008/1	States or regions thereof so listed. It	shall also apply to movements from					
	Official veterinarian or official inspector (*)									
	Nan	ne (in cap	ital letters):	Qualification and title:						
	Local veterinary unit: Date:			LVU No:						
				Signature:						
	Star	np:								
	(*) The col	lour of the s	stamp and signature must be different from that of the other pa	articulars 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))