DECISIONS

COMMISSION DECISION

of 26 August 2010

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

(notified under document C(2010) 5779)

(Text with EEA relevance)

(2010/470/EU)

THE EUROPEAN COMMISSION,

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

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

Whereas:

- (1) Directive 92/65/EEC lays down the animal health requirements governing trade within the Union of animals, semen, ova and embryos not subject to the animal health requirements laid down in specific Union acts. It includes requirements for trade in semen, ova and embryos of animals of the equine, ovine and caprine species and in ova and embryos of animals of the porcine species ('the commodities'). In addition, it provides for health certificates to be established for trade in the commodities within the Union.
- (2) Annex D to Directive 92/65/EEC, as amended by Commission Regulation (EU) No 176/2010 (²), sets out certain new requirements for the commodities which are to apply from 1 September 2010.
- (3) Annex D to Directive 92/65/EEC, as thus amended by Regulation (EU) No 176/2010, introduces rules
- (¹) OJ L 268, 14.9.1992, p. 54.
- ⁽²⁾ OJ L 52, 3.3.2010, p. 14.

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 (¹).
- (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.
- In the interests of clarity of Union legislation, the Union (9) 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 (2), Commission Decision 95/307/EC of 24 July 1995 determining the specimen animal health certificate for trade in semen of the equine species (3), 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 (4) and Commission Decision 95/483/EC of 9 November 1995 determining the specimen certificate for intra-Community trade in ova and embryos of swine (5) 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;
- (1) OJ L 94, 31.3.2004, p. 44.
- (²) OJ L 182, 2.8.1995, p. 27.
- (³) OJ L 185, 4.8.1995, p. 58.
- (⁴) OJ L 234, 3.10.1995, p. 30.
- ⁽⁵⁾ OJ L 275, 18.11.1995, p. 30.

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

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 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 IB 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 IC 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 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 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 IIB 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 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.

Done at Brussels, 26 August 2010.

For the Commission John DALLI Member of the Commission

Intra trade certificate

ANNEX I

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

PART A

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

EUROPEAN UNION

	l.1.	Consignor Name	I.2. Certificate reference No I.2.a. Local reference No
		Address	I.3. Central competent authority
Ited		Postal code	I.4. Local competent authority
presen	l.5.	Consignee Name	1.6.
nent		Address	1.7.
sign		Postal code	
Part I: Details of consignment presented	1.8.	Country of origin ISO I.9. Region of origin Code	I.10. Country of ISO I.11. Region of Code destination code destination
Detai	I.12.	Place of origin	I.13. Place of destination
<u>≓</u>		Semen centre	Semen centre
Par		Name Approval number	Name Approval number
		Address	Address
		Postal code	Postal code
	l.14.		l.15.
	l.16.	Means of transport	1.17.
		Aeroplane Ship Railway wagon	
		Road 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 product	I.22. Number of packages
	1.00	Ambient Chilled	Frozen
	1.23.	Seal/container No	I.24. Type of packaging
	1.25.	Commodities certified for: Artificial reproduction	
	I.26.	Transit through a 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
	128	Export	1.29.
	1.20.	Third country ISO code	
		Exit point Code	
	I.30.		
	I.31.	Identification of the commodities	
		Species Breed Donor identity Da (scientific name)	ate of collection Approval number Quantity of the team

EUROPEAN UNION Equine semen — Part A Ш. Health information II.b. II.a. Certificate reference No I, the undersigned official veterinarian, hereby certify that: II.1. The semen collection centre (2), in which the semen described above was collected, processed and stored for trade is approved and supervised by the competent authority in accordance with Chapter I(I)(1) and Chapter I(II)(1) of Annex D to Directive 92/65/EEC; 11.1.1 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: 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 (3); II.1.1.2. fulfilled the conditions for a holding laid down in Article 4(5) of Directive 2009/156/EC; Part II.1.1.3. contained only equidae which were free of clinical signs of equine viral arteritis and contagious equine metritis. 11.2. Only equidae satisfying the conditions laid down in Articles 4 and 5 or Articles 12 to 16 of Directive 2009/156/EC have been admitted into the centre. 11.3. The semen described above was collected from donor stallions, which: II.3.1. have not shown any clinical sign of an infectious or contagious disease at the time of admission into the centre and on the day the semen was collected: II.3.2. have been kept for 30 days prior to the date of semen collection in holdings where no equine has shown any clinical sign of equine viral arteritis or contagious equine metritis during that period; II.3.3. have not been used for natural mating during at least 30 days prior to the date of first semen collection and from the dates of the first sample referred to in points II.3.5.1, II.3.5.2 or II.3.5.3 until the end of the collection period; 11.3.4. have undergone the following tests, which meet at least the requirements of the relevant chapter of the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals of the OIE, carried out on samples taken in accordance with one of the programmes specified in point II.3.5 in a laboratory recognised by the competent authority: $(^{1})$ either [II.3.4.1. an agar-gel immuno-diffusion test (Coggins test) for equine infectious anaemia (EIA) with negative result;] (1) or [II.3.4.1. an ELISA for equine infectious anaemia (EIA) with negative result;] and (1) either [II.3.4.2. a serum neutralisation test for equine viral arteritis (EVA) with negative result at a serum dilution of one in four;] (¹) or [II.3.4.2. a virus isolation test for equine viral arteritis (EVA) carried out with negative result on an aliquot of the entire semen of the donor stallion;] an agent identification test for contagious equine metritis (CEM) carried out on two occasions on samples taken [11.3.4.3. and with an interval of seven days by isolation of Taylorella equigenitalis after a cultivation of 7 to 14 days from preejaculatory fluid or a semen sample and from genital swabs taken at least from the penile sheath, urethra and urethral fossa with negative result in each case; 11.3.5. have been subjected with the results specified in II.3.4 in each case to at least one of the test programmes (4) detailed in points II.3.5.1, II.3.5.2 and II.3.5.3 as follows: II.3.5.1. the donor stallion was continuously resident on the semen collection centre for at least 30 days prior to the date of the first collection and during the period of collection of the semen described above and no equidae on the semen collection centre came into direct contact with equidae of lower health status than the donor stallion; the tests described in point II.3.4 have been carried out on samples taken (5) prior to the first semen collection and at least 14 days following the date of the commencement of the residence period of at least 30 days;

EUROPEAN UNION

Equine semen — Part A II. Health information II.a. Certificate reference No II.b. II.3.5.2. the donor stallion was resident on the semen collection centre for at least 30 days prior to the date of the first collection and during the period of collection of the semen described above, but has left the centre under the responsibility of the centre veterinarian for a continuous period of less than 14 days, and/or other equidae on the collection centre came into direct contact with equidae of lower health status: the tests described in point II.3.4 have been carried out on samples taken (5) prior to the first semen collection of the breeding season or collection period in the year the semen described above was collected and at least 14 days following the date of the commencement of the residence period of at least 30 days; the test described in point II.3.4.1 for equine infectious anaemia was last carried out on a sample of blood taken (5) not more than 90 days and before the semen described above was collected; and $(^{1})$ either [one of the tests described in point II.3.4.2 for equine viral arteritis was last carried out on a sample taken (5) not more than 30 days before the semen described above was collected;] (1) or [a virus isolation test for equine viral arteritis was carried out with negative result on an aliquot of the entire semen of the donor stallion taken (5) not more than six months before the semen described above was collected and a blood sample taken on the same date (5) reacted positive in a serum neutralisation test for equine viral arteritis at a serum dilution of more than one in four:1 and the test described in point II.3.4.3 for contagious equine metritis was last carried out on samples taken (5) not more than 60 days before the semen described above was collected. The tests described in point II.3.4 have been carried out on samples taken (5) prior to the first semen collection of the breeding season or II.3.5.3. collection period in the year the semen described above was collected, the tests described in point II.3.4 were last carried out on samples taken (5) not less than 14 days and not more than 90 days after the and collection of the semen described above. II.3.6. Have undergone the testing provided for in point II.3.5 on samples taken on the following dates: Start date (5) Date of sampling for health tests (5) Identification of semen Test programme Donor **FVA** CEM FIA Semen 11.3.4.3 residence collection II.3.4.1 11.3.4.2 Blood Semen 1. sample 2. sample sample sample (1) either [1].4 No antibiotics were added to the semen;] The following antibiotic or combination of antibiotics was added to produce a concentration in the final diluted semen of not less (¹) or [1].4 than (⁶): II.5. The semen described above was: collected, processed, stored and transported under conditions which comply with the requirements of Chapters II(I)(1) and III(I) of Annex II.5.1. D to Directive 92/65/EEC; II.5.2. sent to the place of loading in a sealed container in accordance with point 1.4 of Chapter III(I) of Annex D to Directive 92/65/EEC and bearing the number indicated in Box I.23.

EUROPEAN UNION

Equine semen — Part A II. Health information II.a. Certificate reference No II.b. Notes Part I: Box I.12: place of origin shall correspond to the semen collection centre of origin of the semen. Box I.13: place of destination shall correspond to the semen collection or storage centre or to the holding of semen destination. Box I.23: identification of container and seal number shall be indicated. Box I.31: donor identity shall correspond to the official identification of the animal. date of collection shall be indicated in the following format: dd/mm/yyyy. approval number of the centre shall correspond to the approval number of the semen centre indicated in Box I.12 where the semen was collected Part II: 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 where 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.

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Equine semen — Part A

П.	II. Health information					tificate reference No	II.b.	
_	0	Start date	· (⁵)		D	ate of sampling for he	ealth tests (⁵)	
Identification of semen	Test orogramme	Donor Semen residence collection		EIA II.3.4.1		EVA II.3.4.2		EM .4.3
Ident of s	Drog				Blood sample	Semen sample	1. sample	2. sample
			EIA-1	EVA-B1	EVA-S1	CEM-11	CEM-12	
A	В	С	D	EIA-2	EVA-B2	EVA-S2	CEM-21	CEM-22
(¹) Dele	(¹) Delete as appropriate.							
1		red semen collection opa.eu/food/animal/ap			,	l) of Council Directive	92/65/EEC on the 0	Commission website:

(³) OJ L 192, 23.7.2010, p. 1.

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

- (5) Insert date in table in point II.3.6 (follow guidance in part II of the Notes).
- (⁶) Insert names and concentrations.

Official veterinarian (*)

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

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

PART B

Model health certificate IB for trade within the Union in consignments of stocks of semen of animals of the equine 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

	AN UNION						Intra tr	ade certificat
I.1.	Consignor			I.2. Certificate	reference No		I.2.a. Local refe	rence No
	Name							
	Address			I.3. Central competent authority				
	Postal code			I.4. Local con	npetent author	ity		
1.5.	Consignee			1.6.				
	Name							
	Address	1.7.						
	Postal code			1.7.				
I.5. I.8.	Country of origin ISO code	I.9. Region of origin	Code	I.10. Country destination		ISO code	I.11. Region of destination	Code
1.12	. Place of origin			I.13. Place of	destination			
	Semen ce	entre		Se	emen centre		Holding	
	Name	Approval number		Name			Approval num	her
	Address			Address			Approvar nam	501
	Postal code			Postal co	do			
					Jue			
1.14				1.15.				
1.16	. Means of transport			1.17.				
		_	_					
	Aeroplane 🗌 Ship		י 🗋			_		
	Road vehicle Othe	er 🔲						
	Identification							
1.18	. Description of commodity				I.19. Commo	ditv coo	de (HS code)	
				05 11 99 85				
							Quantity	
1.21	. Temperature of product							
	Ambient 🔲	Chilled		Frozen		6		
1.23	. Seal/container No					1.24.	Type of packaging	
1.25	. Commodities certified for:							
	Artificial reproduction 🔲							
1.26	. Transit through a third count	ry 🗆		I.27. Transit th	nrough Membe	r States	;	
	Third country	ISO code		Member	State		ISO code	
	Exit point	Code		Member			ISO code	
	Entry point	BIP No		Member			ISO code	
1.29	. Export			1.29.				
1.20	Third country			1.23.				
	Exit point	Code					_	
1.30		0006						
31	. Identification of the commodi	ties						
			. .					Quantit
	Species Bre (scientific name)	ed Donor identity	Dat	e of collection		proval ni of the te		Quantity

	EUROPE	AN UNION			Equine semen — Part B				
	П.	Health information		II.a. Certificate reference No	ll.b.				
	I, the ur	ndersigned official veteri	narian, hereby certify that:						
	II.1.	The semen collection centre (2), in which the semen described above was collected, processed and stored for trade:							
	∥.1.1.	is approved and supervised by the competent authority according to the conditions of Chapter I of Annex D to Directive 92/65/EEC;							
II: Certification	II.1.2.	is situated on the territory or in the case of regionalisation in a part of the territory (¹) of a Member State which was on the day semen was collected until the date the semen was dispatched as fresh/chilled (¹) semen or until the 30 days mandatory storage period for frozen semen elapsed (¹) not considered to be infected with African horse sickness in accordance with Article 5(2)(a) and (b) of Directive 2009/156/EC (³);							
Part II: C	II.1.3.	fulfilled during the period commencing 30 days prior to the date of semen collection until the date the semen was dispatched as fresh/ chilled (¹) semen or until the 30 days mandatory storage period for frozen semen elapsed (¹), the conditions of Article 4 of Directive 2009/156/EC;							
	II.1.4.	contained during the period commencing 30 days prior to the date of semen collection until the date the semen was dispatched as fresh/ chilled (¹) semen or until the 30 days mandatory storage period for frozen semen elapsed (¹) only equidae which were free of clinical signs of equine viral arteritis and contagious equine metritis;							
	∥.2.	All equidae have been	admitted into the centre under the prov	isions of Article 4 and 5 of Directive 2	2009/156/EC.				
	II.3.	The semen described	above was collected from donor stallions	s, which:					
	II.3.1.	on the day the semen	was collected have not shown clinical s	igns of an infectious or contagious dis	sease;				
	II.3.2.	during at least 30 days prior to collection of the semen have not been used for natural service;							
	II.3.3.	during the last 30 days arteritis;	prior to collection of the semen have bee	en kept on holdings where no equidae s	showed clinical signs of equine viral				
	II.3.4.	during the last 60 days equine metritis;	prior to collection of the semen have been	en kept on holdings where no equidae	showed clinical signs of contagious				
	II.3.5.		owledge and as far as I could ascertain ring 15 days immediately preceding colle		dae suffering from an infectious or				
	II.3.6.	have undergone the fo test programme as spe	Ilowing animal health tests, carried out in ecified in point II.3.7;	a laboratory recognised by the compe	tent authority, in accordance with a				
		[II.3.6.1.	an agar gel immunodiffusion test (Cog	gins test) for equine infectious anaem	ia with negative result;]				
	and	(¹) either [II.3.6.2.	a serum neutralisation test for equine	viral arteritis with negative result at a	serum dilution of one in four; and]				
		(¹) or [II.3.6.2.	a virus isolation test for equine viral art the donor stallion;]	teritis carried out with negative result c	on an aliquot of the entire semen of				
	and	II.3.6.3.	an agent identification test for contagiou the donor stallion with an interval of sev a semen sample and from genital swa negative result in each case;	ren days by isolation of Taylorella equig	<i>penitalis</i> from pre-ejaculatory fluid or				
	II.3.7.	have been subject to t	the one of the following test programmes	\$ (⁴):					
	II.3.7.1.		continuously resident in the collection of no equidae in the collection centre came of						
		contagious equine met	point II.3.6 have been carried out on sar tritis on a second sample taken on above residence period and at least at		at least 14 days after the				

EUROPEAN UNION

EUROPEAN UNION Equine semen — Par				
II. –	Health informa	ation	II.a. Certificate reference No	ll.b.
II.3.7.2.		allion was not continuously resident in the collectic ower health status than the donor stallion;	on centre or other equidae in the colle	ction centre came into contact with
	contagious e	scribed in point II.3.6 have been carried out on sa equine metritis on a second sample taken on en collection and at least at the beginning of the		(⁵) and in the case of within the 14 days period before
and		the test described in point II.3.6.1 for equine in		
and	(¹) either	[one of the tests described in point II.3.6.2 for		
	(¹) or	[the non-shedder state of the seropositive stallion carried out on an aliquot of the entire semen of t than one year before the semen described above	the donor stallion collected on	
II.3.7.3.	14 days afte	escribed in point II.3.6 have been carried out during ar the collection of the semen on samples taken o d sample taken on	n (⁵) and in the	
II.4.		described above was collected, processed, stored and III of Annex D to Directive 92/65/EEC.	and transported under conditions whi	ch comply with the requirements of
Notes				
Part I:				
Box I.12	place of orig	in shall correspond to the semen collection centre	e of origin of the semen.	
Box I.13	place of des	stination shall correspond to the semen collection	or storage centre or to the holding of	semen destination.
Box 1.23	dentification	of container and seal number shall be indicated.		
Box I.31	donor identi	ty shall correspond to the official identification of th	ne animal.	
	date of colle	ection shall be indicated in the following format: do	l/mm/yyyy.	
	approval nui collected.	mber of the centre shall correspond to the approval	number of the semen centre indicated	d in Box I.12 where the semen was
Part II:				
(1) Delet	e as appropria	ate.		
		nen collection centres listed in accordance with A food/animal/approved_establishments/establishmen		/EEC on the Commission website:
(³) OJ L	192, 23.7.201	0, p. 1.		
		ramme(s) that do(es) not apply to the consignmen	t.	
(⁵) Insert				
Official v	eterinarian or	official inspector (*)		
Nan	ne (in capital	letters):	Qualification	and title:
Loci	al veterinary ι	nit:	LVU No:	
Date	э:		Signature:	
Star	mp:			
(*) The co	lour of the star	n and aignature must be different from that of the other an	articulars in the certificate	
() ine co	our or the stam	p and signature must be different from that of the other pa	anuculars in the certificate.	

PART C

Model health certificate IC for trade within the Union in consignments of semen of animals of the equine species collected, processed and stored in accordance with Council Directive 92/65/EEC after 31 August 2010 and in consignments of stocks of semen of animals of the equine 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	OPEA	N UNION							Intra	trade	certificate
	1.1.	Consignor				1.2.	Certificate reference	No	I.2.a. Local re	ference	e No
		Name					-				
		Address				I.3. Central competent authority					
eq		Postal code			1.4.	Local competent aut	hority				
sent	1.5.	5. Consignee			I.6. I	No(s) of related origi	nal	No(s) of accompa	nying		
pres		Name					certificates		documents		
ent		Address				1.7.					
consignment presented		Postal code			1.7.						
nsig	18	Country of origin	ISO	I.9. Region of origin	Code	110	Country of	ISO	I.11. Region of	(Code
S		country of origin	code			1.10.	destination	code	destination		5040
s of											
Part I: Details	1.12.	. Place of origin				I.13.	Place of destination				
ŏ			Semen cen	tre 🔲			Semen centr	e 🗌	Holding 🔲		
art		Name		Approval number			Name		Approval nu	mber	
۵.		Address					Address				
		Postal code					Postal code				
	1.14.					I.15.					
	I.16.	. Means of transpor	t			1.17.					
		Aeroplane 🗌	Ship [Railway wag	on 🗖						
		Road vehicle 🗌	Other								
		Identification									
	1.18.	. Description of corr	nmodity				I.19. Con	nmodity co	de (HS code)		
								0	5 11 99 85		
								1.20.	Quantity		
	1.21.	. Temperature of pro	oduct								
		Ambient 🔲		Chilled 🔲			Frozen 🗌	1.22.	Number of packag	es	
	1.23.	. Seal/container No						1.24.	Type of packaging		
	125	. Commodities certif	ied for:								
		Artificial reproducti									
	1.26.	. Transit through a t	hird country			1.27.	Transit through Mer	nber State	s		
		Third country	ISO c	ode			Member State		ISO code		
		Exit point	Code				Member State		ISO code		
		Entry point	BIP N	0			Member State		ISO code		
	1.28.	. Export				1.29.					
		Third country	ISO co	ode			_				
		Exit point	Code								
	1.30.										
	1.31.	. Identification of the	commoditie	es							
		Species		Breed Donor iden	tity	Date	of collection	Appro	oval number	Qı	uantity
		(scientific name)						of	the team		

	EUROPEA	AN UNION	l		Equine semen — Part C
	II. F	Health infor	rmation II.a. Certificat	e reference No	II.b.
	I, the und	dersigned o	official veterinarian, hereby certify that the semen described above		
	(¹) either	[II.1.	was collected, processed and stored for a minimum period of 30 d collection centre $(^2)$ situated in the Member State of origin of the Chapter I(I)(1) and Chapter I(II)(1) of Annex D to Directive 92/65/E storage centre detailed in Box I.12 situated in the same Member veterinary certification conditions at least as strict as those provided the storage centre detailed of the storage centre detailed in Box I.12 situated in the same Member veterinary certification conditions at least as strict as the storage provided the storage centre detailed in Box I.12 situated in the same Member veterinary certification conditions at least as strict as the storage provided to the storage provided to the storage centre detailed to the storage provided to the storage centre detailed to the storage provided to the storage centre detailed to t	semen and operated a EC, and from where the er State of origin of the	and supervised in accordance with e semen was moved to the semen
ion		(¹) either	r [Part A of Annex I to Decision 2010/470/EU;]		
ertificat		(¹) or	[Part B of Annex I to Decision 2010/470/EU;]		
Part II: Certification		(¹) or	[Decision 95/307/EC;]		
Ра	(¹) or	[.1.	was collected, processed and stored for a minimum period of 30 d collection centre (²) situated in the European Union and operate Chapter I(II)(1) of Annex D to Directive 92/65/EEC, and was mo accordance with:	d and supervised in ac	cordance with Chapter I(I)(1) and
		(¹) either	r [Part A of Annex I to Decision 2010/470/EU;]		
		(¹) or	[Part B of Annex I to Decision 2010/470/EU;]		
		(¹) or	[Part C of Annex I to Decision 2010/470/EU;]		
		(¹) or	[Decision 95/307/EC;]		
	(¹) or	[.1.	was collected, processed and stored in an approved semen collection in columns 2 and 4 of Annex I to Commission Decision 2000 with Chapter I(I)(1) and Chapter I(II)(1) of Annex D to Directive 92/ the conditions of Article 4 of Decision 2004/211/EC in accordance	4/211/EC which is opera 65/EEC, and was impor	ated and supervised in accordance
		(¹) either	r [Part A of Annex I to Decision 2010/471/EU;]		
		(¹) or	[Part B of Annex I to Decision 2010/471/EU;]		
		(¹) or	[Part C of Annex I to Decision 2010/471/EU;]		
		(¹) or	[Decision 96/539/EC;]]		
		II.2.	was stored in the approved semen storage centre $(^2)$ indicated in with Chapter I(I)(2) and Chapter I(I)(2) of Annex D to Directive 92	Box I.12, which is opera 2/65/EEC;	ated and supervised in accordance
		II.3.	was sent to the place of loading in a sealed container in accorda 92/65/EEC and bearing the number indicated in Box I.23.	nce with point 1.4 of Ch	napter III(I) of Annex D to Directive
	Notes				
	Part I:				
	Box I.6:	panied the	espond to the serial number of the individual official document(s) o e semen described above from the approved semen collection centre nal(s) of this/these document(s) or certificate(s) or the officially endors	e of its origin to the desc	ribed above semen storage centre.
	Box I.12:	place of o	origin shall correspond to the semen storage centre of dispatch of t	the semen.	

EUROPEAN UNION

Equine semen — Part C

II. Health information	II.a. Certificate reference No	II.b.				
Box I.13: place of destination shall correspond to the semen collection or storage centre or to the holding of destination of the semen.						
Box I.23: identification of container and seal number shall be indicated.	Box I.23: identification of container and seal number shall be indicated.					
Box I.31: donor identity shall correspond to the official identification of t	he animal.					
date of collection shall be indicated in the following format: do	d/mm/yyyy.					
approval number of the centre shall correspond to the approv	al number of the semen collection ce	ntre of origin of the semen.				
Part II:						
(¹) Delete as appropriate.						
(²) Only approved semen collection or storage centres listed in accordan the Commission websites:	ce with Article 11(4) or Article 17(3)(b)	of Council Directive 92/65/EEC on				
http://ec.europa.eu/food/animal/approved_establishments/establishmen http://ec.europa.eu/food/animal/semen_ova/equine/index_en.htm	nts_vet_field_en.htm					
Official veterinarian or official inspector (*)						
Name (in capital letters):	Qualification and title:					
Local veterinary unit:	LVU No:					
Date:	Signature:					
Stamp:						
) The colour of the stamp and signature must be different from that of the other particulars in the certificate.						

EUROPEAN UNION

EN

ANNEX II

Model health certificates for trade within the Union in consignents of ova and embryos of animals of the equine species

PART A

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

I.1. Consignor I.2. Certificate reference No I.2.a. Local reference No Name I.3. Central competent authority Address I.4. Local competent authority Postal code of consignment presented l.5. Consignee I.6. Name Address 1.7. Postal code I.8. Country of origin ISO code I.9. Region of origin Code I.10. Country of ISO I.11. Region of destination code destination Part I: Details I.12. Place of origin I.13. Place of destination Embryo team 🗌 Holding 🔲 Embryo team 🗌 Approval number Name Name Approval number Address Address Postal code Postal code 1.14. I.15. l.17. I.16. Means of transport Railway wagon 🗌 Aeroplane Ship 🔲 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 I.22. Number of packages Ambient 🗌 Chilled 🗌 Frozen 🗌 I.23. Seal/Container No I.24. Type of packaging I.25. Commodities certified for: Artificial reproduction I.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 I.28. Export 1.29. Third country ISO code Exit point Code 1.30.

I.31. Identification of the commodities

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

Intra trade certificate

Code

	EUROPEAN UNION		N	Equine ova and embryos — Part							
	П.	Health inf	ormation	II.a. Certificate reference No	II.b.						
	I, the u	Indersigned	l official veterinarian, hereby certify that:								
	(¹) eith	<i>er</i> [II.1.	the <i>in vivo</i> derived embryos/ <i>in vivo</i> derived ova (¹ , collection team (²) approved and supervised 92/65/EEC;]								
Part II: Certification	(¹) or	[.1.		he <i>in vitro</i> produced embryos/micromanipulated embryos (¹) described above were produced, processed and stored by ar embryo production team (²), approved and supervised in accordance with Chapter I(III)(1) and (2) of Annex D to Directive 32/65/EEC;]							
	(¹) eith	<i>er</i> [II.2.	the <i>in vivo</i> derived embryos described above meet the requirements of Chapter III(II)(1) of Annex D to Directive 92/65/EEC;]								
Int II: Ce	(¹) or	[II.2.	the <i>in vivo</i> derived ova described above me 92/65/EEC;]	et the requirements of Chapter I	I(II)(2) of Annex D to Directive						
Ра	(¹) or	[II.2.	the <i>in vitro</i> produced embryos described above 92/65/EEC;]	meet the requirements of Chapter	III(II)(3) of Annex D to Directive						
	(¹) or	[II.2.	the micromanipulated embryos described above 92/65/EEC;]	meet the requirements of Chapter	III(II)(4) of Annex D to Directive						
		II.3.	the ova or embryos described above come from d	onor mares which:							
		II.3.1.	coming from holdings fulfilling the conditions laid d satisfying the conditions laid down in Articles 4 ar								
		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 le the date of the first sample referred to in points II.								
		II.3.4.	have been subjected with negative result to an agar anaemia carried out on a blood samples taken on first collection of ova or embryos and th on	(³), being during the p ne last test was carried out o	ast 30 days prior to the date of the on a sample of blood taken						
		II.3.5.	have been subjected to an agent identification test f cultivation of 7 to 14 days carried out with negative date of the first collection of ova or embryos fro consecutives oestrus periods on	results in each case on samples taken om mucosal surfaces of the clitoral and on	during the past 30 days prior to the fossa and clitoral sinuses on two						
	(¹) eith	<i>er</i> [II.4.	the embryos described above were conceived as a collected, processed, stored and transported under III(I) of Annex D to Directive 92/65/EEC;]								
	(¹) or	[1].4.	the embryos described above were conceived as a of Chapter III(II) of Annex D to Directive 92/65/EEC conditions which comply with the requirements of 0	with semen which was collected, proce	essed, stored and transported under						
	(¹) or	[11.4.	the ova have not been in contact with semen of th	e equine species;]							
		II.5.	the ova or embryos described above were sent to Chapter III(II) of Annex D to Directive 92/65/EEC a								
	Notes										
	Part I:										
	Box I.1	2: Place of	f origin shall correspond to the embryo collection tean	n or embryo production team of ova/e	mbryos collection/production.						
	Box I.1	3: Place o destinat	of destination shall correspond to the embryo collec ion.	tion team, embryo production team	or to the holding of ova/embryos						
	Box 1.2	3: Identific	ation of container and seal number shall be indicated.								

EUROPEAN UNION

Equine ova and embryos - Part A

II.	Health information	II.a. Certificate reference No	II.b.		
Box I.3	1: Category: specify if: in vivo derived embryos, in vivo derived o	ova, <i>in vitro</i> produced embryos or mic	romanipulated embryos.		
	Donor identity shall correspond to the official identification of t	he animal.			
	Date of collection shall be indicated in the following format: do	d/mm/yyyy.			
	Approval number of the team shall correspond to the e collection/production.	mbryo collection team or embryo	production team of ova/embryos		
Part II:					
(¹) Dele	ete as appropriate.				
(²) Only web	v approved embryo collection or production teams listed in accordance site:	ance with Article 11(4) of Council Direct	tive 92/65/EEC on the Commission		
http	//ec.europa.eu/food/animal/approved_establishments/establishmer	nts_vet_field_en.htm			
(³) Inse	rt date.				
(⁴) OJ	L 192, 23.7.2010, p. 1.				
Official	veterinarian or official inspector (*)				
Nan	ne (in capital letters):	Q	ualification and title:		
Loca	al veterinary unit:	L۱	/U No:		
Date	9:	Si	gnature:		
Star	np:				
(*) 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 IIB for trade within the Union in consignments of stocks of ova and embryos of animals of the equine 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

EURO	OPEA	N UNION	Intra trade certificat			
	l.1.	Consignor	I.2. Certificate reference No I.2.a. Local reference No			
		Name				
		Address	I.3. Central competent authority			
		Destal sode	I.4. Local competent authority			
fed		Postal code				
consignment presented	1.5.		1.6.			
pre		Name				
ent		Address	1.7.			
Ĕ		Postal code				
Isig	1.8.	Country of origin ISO code I.9. Region of origin Code	I.10. Country of ISO I.11. Region of Code			
0 S			destination code destination			
s of						
Part I: Details	1.12.	Place of origin	I.13. Place of destination			
Ď		Embryo team 🔲	Holding 🗌 🛛 Embryo team 🗖			
L L						
۳ ۳		Name Approval number	Name Approval number			
		Address	Address			
		Postal code	Postal code			
	1.14.		l.15.			
	116	Means of transport	1.17.			
		Aeroplane 🗌 Ship 🗌 Railway wagon 🗌				
		Road vehicle Other				
	1.18	Identification Description of commodity	I.19. Commodity code (HS code)			
			05 11 99 85			
			I.20. Quantity			
	1.21.	Temperature of products	I.22. Number of packages			
		Ambient Chilled	Frozen			
	1.23.	Seal/Container No	I.24. Type of packaging			
	125	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 Third country ISO code	1.29.			
		Exit point Code				
	1.30.					
	1.30.					
	1.31.	Identification of the commodities				
		Species Breed Category Donor ident (Scientific name)	ity Date of collection Approval number Quantity of the team			
			or the team			

	EUROPEA	N UNION	Equine ova and embryos — Par						
	II. Н	ealth information	II.a. Certificate reference No	II.b.					
	I, the unde	ersigned official veterinarian, hereby certify that:							
	II.1 <i>.</i>	Ova/embryos (¹) described above were collected by a colle appropriate laboratory;	ction team $(^2)$ approved by the compe	tent authority and processed in an					
	II.2.	Ova/embryos (1) were collected from donor mares which:							
fication	II.2.1.	on the day of collection have been located in premises situated on the territory or in the case of regionalisation in a part of the territory of a Member State which is not considered to be infected with African horse sickness in accordance with Article 5(2)(a) and (b) of Directive 2009/156/EC (³);							
Part II: Certification	II.2.2.	have been located in holdings under veterinary supervision w 2009/156/EC;	hich on the day of collection fulfilled th	e conditions of Article 4 of Directive					
Part	II.2.3.	have been kept prior to the collection in holdings free from	clinical signs of contagious equine m	etritis for 60 days;					
	II.2.4.	have not been used for natural breeding during the period	of 30 days prior to the collection of o	va/embryos (¹);					
	II.2.5.	to the best of my knowledge and as fare as I could ascerta contagious disease during the 15 days immediately precedi		idae suffering from an infectious or					
	II.2.6.	have on the day of collection not shown clinical signs of ar	n infectious or contagious disease;						
	II.3.	Ova/embryos (¹) were collected, processed, stored and trans of Directive 92/65/EEC;	sported under conditions which comply	v with the requirements of Annex D					
II.4. The semen used for the artificial insemination of the donor mares complies with the requirements of I $92/65/EEC$ (⁴) (¹);									
	II.5.	The ova used for the in vivo production of embryos comply	with the requirements of Directive 92	/65/EEC (¹).					
	Notes								
	Part I:								
	Box 1.12:	Place of origin shall correspond to the embryo collection team	n of ova/embryos collection.						
	Box I.13:	Place of destination shall correspond to the embryo collec destination.	tion team, embryo production team	or to the holding of ova/embryos					
	Box 1.23:	Identification of container and seal number shall be indicated.							
	Box 1.31:	Category: specify if: in vivo derived embryos, in vivo derived	ova, <i>in vitro</i> produced embryos or mic	romanipulated embryos.					
		Donor identity shall correspond to the official identification of	the animal.						
		Date of collection shall be indicated in the following format: d	d/mm/yyyy.						
Approval number of the team shall correspond to the embryo collection team of ova/embryos collection.									
									(¹) Delete as appropriate.
	(²) Only a	pproved embryo collection teams listed in accordance with A	rticle 11(4) of Council Directive 92/65	/EEC on the Commission website:					
	http://e	c.europa.eu/food/animal/approved_establishments/establishme	nts_vet_field_en.htm						
	(³) OJ L 1	192, 23.7.2010, p. 1.							
	(4) Does r	not apply to ova.							
	1								

EUROPEAN UNION

Equine ova and embryos - Part B

II.a. Certificate reference No	II.b.					
1						
	Qualification and title:					
	LVU No:					
	Signature:					
(*) The colour of the stamp and signature must be different from that of the other particulars in the certificate.						

Intra trade certificate

ANNEX III

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

PART A

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

EUROPEAN UNION

	l.1.	Consignor	I.2. Certificate reference No I.2.a. Local reference No				
		Name Address	I.3. Central competent authority				
pe		Postal code	I.4. Local competent authority				
sente	l.5.	Consignee	1.6.				
pre		Name Address	1.7				
nent			1.7.				
ignr		Postal code					
Part I: Details of consignment presented	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			
ails	l.12.	Place of origin	I.13. Place of destination				
Det		Semen centre 🔲	Semen centre 🗖	Holding			
art I:		Name Approval number	Name	Approval number			
å		Address	Address				
		Postal code	Postal code				
	l.14.		l.15.				
	l.16.	Means of transport	l.17.				
		Aeroplane 🗌 Ship 🗌 Railway wagon 🗌					
		Road vehicle Other					
		Identification					
	l.18.	Description of commodity	I.19. Commodity code (HS code) 05 11 99 85				
				I.20. Quantity			
	1.01	Temperature of products		· · · · · · · · · · · · · · · · · · ·			
	1.21.		Frozen	.22. Number of packages			
	1.23.	Seal/Container No		.24. Type of packaging			
	125	Commodities certified for:					
	1.20.	Artificial reproduction					
	1.26.	Transit through third country	I.27. Transit through Member S	States			
		Third country ISO code	Member State	ISO code			
		Exit point Code	Member State	ISO code			
		Entry point BIP No	Member State	ISO code			
	I.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)		proval number Quantity f the centre			

	EUROPEAN UNION			Ovine and caprine semen — Part /						
	П.	Health info	ormation	II.a. Certificate reference No	II.b.					
	I, the un	dersigned	official veterinarian, hereby certify that:							
		II.1.	the semen described above:							
		II.1.1.	was collected, processed and stored in a semen cracordance with Chapter I(I)(1) and Chapter I(II)(1) and Chapter		rvised by the competent authority in					
tion		II.1.2.	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.1.3.	was collected, processed, stored and transported u III(I) of Annex D to Directive 92/65/EEC;	under conditions which comply with the	e requirements of Chapters II(II) and					
	(¹) either	· [II.1.4.	meets the requirements of Chapter A(I) of Annex $\ensuremath{^{\circ}}$	/III to Regulation (EC) No 999/2001;]						
Par	(¹) or	[11.1.4.	meets the requirements of Chapter A(I) of Annex which benefits, for all or part of its territory, from th Regulation (EC) No 999/2001 and the donor anii programmes referred to in that point and with the	ne provisions laid down in point (b) or mals comply regarding scrapie with t	(c) of Chapter A(I) of Annex VIII to he guarantees provided for by the					
	_	ll.1.5.	was sent to the place of loading in a sealed conta 92/65/EEC and bearing the number detailed in Bo		Chapter III(I) of Annex D to Directive					
	(¹) either	[11.2.	no antibiotics or no mixture of antibiotics were add	led to the semen;]						
	(¹) or	[II.2.	less than (⁴):							
	Notes	Notes								
	Part I:									
	Box I.12:	Place of	origin shall correspond to the semen collection cent	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	f 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							
		Date of c	collection shall be indicated in the following format: d							
		Approval collected		val number of the semen centre indicated in Box I.12 where the semen was						
	Part II:									
	(1) Delet	e as appro	ppriate.							
	5/EEC on the Commission website:									
	(³) Addit	onal guara	antees as laid down in Article 2 of Regulation (EC) N	lo 546/2006 (OJ L 94, 1.4.2006, p. 28	3).					
(⁴) Insert names and concentrations.										
	Official veterinarian or official inspector (*)									
	Name	e (in capita	al letters):	Q	ualification and title:					
	Local	veterinary	r unit:	Ľ	VU No:					
	Date:	-		S	ignature:					
	Stam	o:								

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

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

EURO	OPEA	N UNION	Intra trade certificate							
	l.1.	Consignor Name Address			1.2. C	ertificate refer	rence No	1.2.	a. Local ref	erence No
		Postal code			1.3. C	entral compet	tent autho	rity		
nted					1.4. Lo	ocal compete	nt authorit	y		
t prese	1.5.	Consignee Name			1.6.					
consignment presented		Address Postal code			1.7.					
	1.8.	Country ISO code of origin	I.9. Region of origin	Code		Country of destination	ISO co	de I.11.	Region of destination	Code
Part I: Details of	I.12.	Place of origin			I.13. Place of destination					
art		Semen centre [Semen o	entre 🗌		Holding	
		Name Address	Approval number		1	Name Address			Approval r	number
		Postal code			F	Postal code				
	1.14.				l.15.					
	I.16. Means of transport Aeroplane Ship Railway wagon				l.17.					
		Road vehicle Othe	er 🗖							
	Identification									
	l.18.	Description of commodity			I.19. Commodity code (HS code) 05 11 99 85					
		-						I.20. Quantit	y	
	1.21.	Temperature of products Ambient	Chilled		Frozen			I.22. Numbe	r of packag	es
	1.23.	Seal/Container No						I.24. Type o	f packaging	
	1.25.	Commodities certified for:								
		Artificial reproduction								
	I.26. Transit through third country				I.27. T	ransit through	n Member	States		
	Third country ISO code				Member State			O code		
	Exit point Code			Vember State Vember State			O code O code			
	Entry point BIP No			1.29.	vieniber State	,	10	O COUE		
	I.28. Export ISO code Third country ISO code Exit point Code		1.29.							
	1.30.									
	1.31.	Identification of the commod	ities							
			reed Donor identity		Date of	collection	Apj o	proval number f the centre		Quantity

	EUROPEAN UNION Ovine and caprine semen —									
	Π.	Н	ealth infor	mation	II.a. Certificate reference No	ll.b.				
I, the undersigned official veterinarian, hereby certify that the semen described above:										
		II.1. was collected, processed and stored in a semen collection centre (²) approved and supervised by the competent authority in accordance with Chapter I(I) and Chapter I(II) of Annex D to Directive 92/65/EEC;								
on			II.2.	comes from the donor animals which meet the requ	uirements of Chapter II(II) of Annex D	to Directive 92/65/EEC;				
Part II: Certification			II.3.	was collected, processed, stored and transported un III of Annex D to Directive 92/65/EEC;	nder conditions which comply with the	requirements of Chapters II(II) and				
IT II: C	(¹)	either	[11.4.	meets the requirements of Chapter A(I) of Annex V	III to Regulation (EC) No 999/2001;]					
Ра	(')	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 (³) requested by the Member State of destination.]						
	No	tes								
	Pa	rt I:								
	Bo	x I.12:	Place of c	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 I.23: Identification of container and seal number shall be indicated.									
	Box	x I.31:	Donor ide	ntity shall correspond to the official identification of t	he animal.					
				ollection shall be indicated in the following format: do						
			Approval collected.	number of the centre shall correspond to the approval	I number of the semen centre indicated	d in Box I.12 where the semen was				
	Pa	rt II:								
	(1)	Delete	as approj	oriate.						
	(²) Only approved semen collection centres listed in accordance with Article 11(4) of Council Directive 92/65/EEC on the Commission website: http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm									
	(³) Additional guarantees as laid down in Article 2 of Regulation (EC) No 546/2006 (OJ L 94, 1.4.2006, p. 28).									
	Official veterinarian or official inspector (*)									
	Name (in capital letters):Qualification and title:Local veterinary unit:LVU No:									
		Date:			Siç	gnature:				
		Stamp								
	(*) The colour of the stamp and signature must be different from that of the other particulars in the certificate.									

PART C

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

Name Address I.3. Central competent authority Postal code I.4. Local competent authority I.5. Consignee Name Address I.6. No(s) of related original centificates No(s) docum Postal code I.7. Postal code I.7. I.8. Country ISO code I.10. Country of destination ISO code I.11. I.12. Place of origin Semen centre Name Address Name Address Name Address Postal code I.13. Place of destination Semen centre Name Address Name Address Name Address Postal code I.14. I.15. I.15. I.16. Means of transport I.14. I.15. I.17. Aeroplane Ship Railway wagon Name Address Road vehicle Other I.17. I.19. Commodity code (HS 05 11 99 I.19. Commodity code (HS 05 11 99 I.12. Temperature of products Ambient Chilled Frozen I.22. Numbe I.23. Seal/Container No I.24. Type o I.24. Type o I.26. Transit through third country I.27. Transit through Member States I.27. Transit through Member State Third country ISO code Member State I.57<	Intra trade certificate				
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I.28. Export I.29. Third country ISO code Exit point Code	SO code				
Third country ISO code Exit point Code	SO code				
1.30.					
I.31. Identification of the commodities					
Species Breed Donor identity Date of collection Approval number (Scientific name) of the centre	er Quantity				

EUROPEA	N UNION		O	vine and caprine semen — Part			
II. F	lealth inform	nation I	I.a. Certificate reference No	II.b.			
I, the und	lersigned off	icial veterinarian, hereby certify that the semen desc	ribed above:				
(¹) either	[11.1.	was collected, processed and stored for a minimus semen collection centre (²) situated in the Mem accordance with Chapter I(I)(1) and Chapter I(II)(1 moved to the semen storage centre detailed in Par animal health and veterinary certification at least a	ber State of origin of the semer) of Annex D to Directive 92/65/EE rt I.12 situated in the same Membe	n 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 minimum period of 30 days immediately following collection in an appr semen collection centre (2) situated in the European Union and operated and supervised in accordance with Chapter and Chapter I(II) of Annex D to Directive 92/65/EEC and was moved to the semen storage centre detailed in Box I.1 accordance with:							
	(¹) either	[Part A of Annex III to Decision 2010/470/EU;]					
	(¹) or	[Part B of Annex III to Decision 2010/470/EU;]					
	(¹) or	[Decision 95/388/EC;]]					
(¹) or [II.1. was collected, processed and stored for semen collection centre (²) situated in a th operated and supervised in accordance wit imported into the European Union under the function of the function of the function of the function.			try or part(s) thereof listed in Annex er I(I)(1) and Chapter I(II)(1) of Anne	I to Decision 2010/472/EU which i x D to Directive 92/65/EEC, and wa			
	(¹) either [Section A of Part 2 of Annex II to Decision 2010/472/EU;]						
	(¹) or	[Section B of Part 2 of Annex II to Decision 2010/	472/EU;]				
	(¹) or	[Annex II to Decision 2008/635/EC;]					
	II.2.	was stored in the approved semen storage centre (with Chapter I(I)(2) and Chapter I(II)(2) of Annex D		arated and supervised in accordanc			
	II.3.	was sent to the place of loading in a sealed contain 92/65/EEC and bearing the number indicated in B		Chapter III(I) of Annex D to Directiv			
Notes							
Part I:							
Box I.6:	Box I.6: Shall correspond to the serial number of the individual official document(s) or health certificate(s) (either INTRA or CVED) that accor panied the semen described above from the approved semen collection centre of its origin to the described above semen storage centr The original of this/these document(s) or certificate(s), or the officially endorsed copy/copies thereof must be attached to this certificate						
Box I.12:	Box I.12: Place of origin 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 o	f destination of the semen.			
Box 1.23:	Identification	n of container and seal number shall be indicated.					
Box I.31:	Box I.31: Donor identity shall correspond to the official identification of the animal. Date of collection shall be indicated in the following format: dd/mm/yyyy. Approval number of the centre shall correspond to the approval number of the semen collection centre of the semen origin.						

EUROPEAN UNION Equine ova and embryos - Part C ١١. Health information II.a. Certificate reference No II.b. Part II: (1) Delete as appropriate. (2) Only approved semen collection or storage centres listed in accordance with Article 11(4) or Article 17(3)(b) of Council Directive 92/65/EEC on the Commission websites: http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm http://ec.europa.eu/food/animal/semen_ova/ovine/index_en.htm Official veterinarian or official inspector (*) Name (in capital letters): Qualification and title: LVU No: Local veterinary unit: Date: Signature: Stamp: (*) The colour of the stamp and signature must be different from that of the other particulars in the certificate.

ANNEX IV

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

PART A

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

EUROPEAN UNION

Intra trade certificate

	I.1. Consignor Name					I.2. Certificate reference No I.2.a. Local reference No				No	
		Address				I.3. Central competent authority					
pe	Postal code				I.4. Local co	mpetent authoril	у				
sent	l.5.	Consignee				I.6.					
pre		Name									
ent		Address Postal code			1.7.						
Ē											
Part I: Details of consignment presented		Country of origin	ISO code	I.9. Region of origin	Code	I.10. Country destina		ISO code	I.11. Reg des	ion of tination	Code
etails	112	Place of origin				I.13. Place c	f destination				
ă 			Embryo tear	n 🗖		1.15. Flace C		ng 🗖	Em	bryo team	
art		Name		Approval number		Name		• —		oroval numb	
₽		Address				Addres	S				
		Postal code				Postal	code				
	1.14					l.15.					
	I.16.	Means of transport				l.17.				_	
		Aeroplane 🔲	Ship 🔲	Railway wagon							
	Road vehicle Other Other										
		Identification									
	I.18	Description of commodi	ty				I.19. Commod	lity code	(HS code)		
						05 11 99 85					
								1.2	20. Quantity		
	I.21.	Temperature of product	s					1.2	2. Number o	of packages	i
		Ambient 🔲		Chilled 🗌		Frozen 🗌					
	1.23.	Seal/Container No						1.2	24. Type of p	ackaging	
	1.25.	Commodities certified for	or:								
		Artificial reproduction]								
	1.26	Transit through third co	untry			I.27. Transit f	hrough Member	States			
		Third country		ISO code		Member	State		I	SO code	
	Exit point Code Entry point BIP No			Member	State		I	SO code			
				Member	State		I	SO code			
	Third country ISO code				1.29.						
	Exit point Code										
	1.30										
	1.31.	Identification of the com	modities								
		Species (Scientific name)	Breed	Category	Donor ic		Date of collection		val number the team	Qu	antity

EUROPEAN UNION

Ovine and caprine ova/embryos - Part A

	Ш.	Health inf	th information II.a. Certificate reference No II.b.							
	I, the unc	ersigned o	official veterinarian, hereby certify that:							
	(¹) either	[.1.	the <i>in vivo</i> derived embryos/ <i>in vivo</i> derived ova $(^1)$ collection team $(^2)$ approved and supervised in acco							
ation	(¹) or	[II.1. the <i>in vitro</i> produced embryos/micromanipulated embryos (¹) described above were produced, processed and stored by an embryo production team (²) approved and supervised in accordance with Chapter I(III)(1) and (2) of Annex D to Directive 92/65/EEC;]								
Certifica	(¹) either	[II.2.	the in vivo derived embryos described above meet the requirements of Chapter III(II)(1) of Annex D to Directive 92/65/EEC;]							
Part II: Certification	(¹) or	[11.2.	the in vivo derived ova described above meet the requirements of Chapter III(II)(2) of Annex D to Directive 92/65/EEC;]							
ď	(¹) or	[11.2.	2. the <i>in vitro</i> produced embryos described above meet the requirements of Chapter III(II)(3) of Annex D to Directive 92/65/EEC;]							
	(¹) or	[11.2.	the micromanipulated embryos described above meet the requirements of Chapter III(II)(4) of Annex D to Directive 92/65/EEC;]							
		II.3. the ova or embryos described above:								
	(¹) either	[II.3.1.	meet the requirements of Chapter A(I) of Annex VIII	to Regulation (EC) No 999/2001;]						
	(¹) or	[II.3.1.	meet the requirements of Chapter A(I) of Annex VIII which benefits, for all or part of its territory, from the Regulation (EC) No 999/2001 and the donor anim programmes referred to in that point and with the g	e provisions laid down in point (b) or als comply regarding scrapie with t	(c) of Chapter A(I) of Annex VIII to he guarantees provided for by the					
		II.3.2.	come from female donors of the ovine/caprine specific principle (2016) Directive 92/65/EEC;	ecies (¹) which meet the requiremen	ts of Chapter IV(3) of Annex D to					
	(¹) either	[11.4.		a result of artificial insemination of the donor females with semen which was conditions which comply with the requirements of Chapters I(I), II(I) and III(I)						
	(¹) or	[11.4.	III(II)(2) of Annex D to Directive 92/65/EEC with s	a result of <i>in vitro</i> fertilisation of ova complying with the conditions in Chapter ith semen which was collected, processed, stored and transported under of Chapters I(I), II(I) and III(I) of Annex D to Directive 92/65/EEC;]						
	(¹) or	[11.4.	the ova have not been in contact with semen of the	ovine and caprine species;]						
		II.5.	the ova or embryos described above were sent to the Chapter III(II) of Annex D to Directive 92/65/EEC and							
	Notes									
	Part I:									
	Box I.12:	Box I.12: Place of origin shall correspond to the embryo collection team or embryo production team of embryos collection/production.								
	Box I.13:	I.13: Place of destination shall correspond to the embryo collection team, embryo production team or to the holding of ova/embryos des nation.								
	Box 1.23:	.23: Identification of container and seal number shall be indicated.								
	Box 1.31	Category.	specify if: in vivo derived embryos, in vivo derived o	ova, <i>in vitro</i> produced embrvos or mi	cromanipulated embrvos.					
	20/ 101	• •	intity shall correspond to the official identification of t							
			ollection shall be indicated in the following format: do							
			number of the team shall correspond to the embryo		on team of ova/embryos collection/					

EUROPEAN UNION

Ovine and caprine ova/embryos - Part A

II. Health information	II.a. Certificate reference No	II.b.							
Part II:									
(¹) Delete as appropriate.									
(²) Only approved embryo collection or production teams listed in accordance with Article 11(4) of Council Directive 92/65/EEC on the Commission website: http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm									
(³) Additional guarantees as laid down in Article 2 of Regulation (EC) N	o 546/2006 (OJ L 94, 1.4.2006, p. 2	8).							
Official veterinarian or official inspector (*)									
Name (in capital letters):		Qualification and title:							
Local veterinary unit:		LVU No:							
Date:		Signature:							
Stamp:	Stamp:								
(*) The colour of the stamp and signature must be different from that of the other particulars in the certificate.									

PART B

Model health certificate 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

ROPE	AN UNION			Intra trade certificate
1.1.	Consignor		I.2. Certificate reference No	I.2.a. Local reference No
	Name		I.3. Central competent authority	
	Address			
	Postal code		I.4. Local competent authority	
1.5.	Consignee		1.6.	
	Name			
	Address		1.7.	
	Postal code			
I.5.	Country of ISO coo origin	le I.9. Region of origin Code	I.10. Country of ISO destination code	J
1.12	. Place of origin		I.13. Place of destination	
	Embryo t	eam 🗖	Holding	Embryo team 🔲
	Name	Approval number	Name	Approval number
	Address		Address	
	Postal code		Postal code	
1.1	4.		l.15.	
1.10	. Means of transport		l.17.	
	Aeroplane 🗌 Ship 🗌			
	Road vehicle]		
	Identification			
1.18	. Description of commodity		I.19. Commodity code	e (HS code) 05 11 99 85
				I.20. Quantity
1.2	. Temperature of products			I.22. Number of packages
	Ambient	Chilled 🔲	Frozen 🔲	,
1.23	. Seal/Container No			I.24. Type of packaging
1.28	. 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 Exit point	ISO code Code		
1.30		Code		
1.30	·			
1.3	. Identification of the commodities			
	Species Breed	Category Donor i		roval number Quantity
	(Scientific name)		collection o	f the team

EUROPEAN UNION

Ovine and caprine ova/embryos — Part B

	II.		Health in	formation	II.a. Certificate reference No	II.b.				
	I, the undersigned official veterinarian, hereby certify that the ova/embryos (1) described above:									
			II.1. were collected, processed and stored under conditions which meet the requirements of Directive 92/65/EEC;							
			II.2. come from female donors of the ovine/caprine species (¹) which meet the requirements of Chapter IV of Annex D to Direct 92/65/EEC;							
tification	(')	either	[11.3.	 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.] 						
Part II: Certification	(')	or	[11.3.	.3. meet the requirements of Chapter III of Annex D to Directive 92/65/EEC and of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and are destined for a Member State which benefits, for all or part of its territory, from the provisions laid down in point (b) or (c) of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and the donor animals comply regarding scrapie with the guarantees provided for by the programmes referred to in that point and with the guarantees (²) requested by the Member State of destination.]						
	(')	either	[II.4. in the case of embryos, the semen used for fertilisation meets the requirements of Directive 92/65/EEC and the requirements of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001.]							
	(¹) or [II.4. in the case of embryos, the semen used for fertilisation meets the requirements of Directive 92/65/EEC and the requirement of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and is destined for a Member State which benefits, for all or part of its territory, from the provisions laid down in point (b) or (c) of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and the donor animals comply regarding scrapie with the guarantees provided for by the programmes referred to in that point and with the guarantees (²) requested by the Member State of destination.]									
	No	tes								
	Pa	rt I:								
	Bo	k I.12:	Place of	origin shall correspond to the embryo collection tean	n of ova/embryos collection.					
	Bo	k I.13:	Place of destination	destination shall correspond to the embryo collec	tion team, embryo production tea	m or to the holding of ova/embryos				
	Bo	k I.23:	Identifica	tion of container and seal number shall be indicated.						
	Bo	 Box I.31: Category: specify if: <i>in vivo</i> derived embryos, <i>in vivo</i> derived ova, <i>in vitro</i> produced embryos or micromanipulated embryos. Donor identity shall correspond to the official identification of the animal. Date of collection shall be indicated in the following format: dd/mm/yyyy. Approval number of the team shall correspond to the embryo collection team of ova/embryos collection indicated in Box I.12. 								
	Part II:									
	 (¹) Delete as appropriate. (²) Additional guarantees as laid down in Article 2 of Regulation (EC) No 546/2006 (OJ L 94, 1.4.2006, p. 28). 									
	Official veterinarian or official inspector (*)									
	Name (in capital letters): Qualification and title:									
Local veterinary unit:						LVU No:				
	Date: Signature:									
	Stamp:									
	(*)	The col	lour of the s	stamp and signature must be different from that of the other p	articulars in the certificate.					

ANNEX V

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

PART A

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

EUROPEAN UNION

Intra trade certificate

		Consignor			1.2.	I.2. Certificate reference No I.2.a. Local reference No					
		Name Address	1.3.	I.3. Central competent authority							
ð		Postal code . Consignee									
Part I: Details of consignment presented						I.4. Local competent authority					
pres											
ent		Name			1.7.						
ũuấ		Address									
onsiç		Postal code Country of ISO code I.9. Region of origin Code				I.10. Country of ISO I.11. Region of Code					
of CC		Country of origin	ISO code	I.9. Region of origin Code	1.10	destinal		ISO code		estination	Code
ils o					_						
Deta	1.12	. Place of origin	Embryo tear	~ □	1.10	3. Place o	f destination	ding 🗖	-	mbryo team	
÷		Name	Embryo tear	Approval number		Name				ambryo team Approval num	
Par		Address				Address	-		~	ippioval num	
		Postal code			_	Postal o	code				
	1.14	4.			1.15	5.					
	1.16	. Means of transport			1.17	7.					
		Aeroplane 🔲	Ship 🔲	Railway wagon 🔲							
		Road vehicle 🗌	Other 🗌								
	Identification										
	I.18. Description of commodity						I.19. Commo	dity code (
								1.20. Qua	05 11 99 antity	9 85	
									-		
	1.21	I.21. Temperature of products						1.22. Nur	nber of pa	ickages	
	Ambient Chilled Froz										
	1.23	. Seal/Container No						1.24. Тур	e of pack	aging	
	1.25	. Commodities certified				•					
	Artificial reproduction 🔲										
	1.26	. Transit through third c	ountry		1.27	. Transit t	hrough Membe	r States			
		Third country		ISO code		Member	State			ISO code	
		Exit point		Code		Member	State			ISO code	
		Entry point		BIP No		Member	State			ISO code	
	I.28. Export				1.29						
		Third country ISO code									
		Exit point		Code							
	1.30										
	1.31	. Identification of the co	mmodities								
		Species	Breed	Category Donor id	entity		te of ection	Approval of the		Q	uantity
		(Scientific name)				COILE	GUUN	or the	leann		

EUROPEAN UNION

	EUROPEA			Porcine ova/embryos — Part A						
	II.	Health inf	formation	II.a. Certificate reference No	II.b.					
	I, the und	lersigned o	official veterinarian, hereby certify that the ova/embryc	os (1) described above:						
		II.1.	were produced/collected (¹), processed and stored by accordance with Chapter I(III) of Annex D to Directiv		eam (²) approved and supervised in					
		II.2.	meet the requirements of Chapter III(II) of Annex D to Directive 92/65/EEC;							
Part II: Certification		II.3.	come from donor females of the porcine species which meet the requirements of Chapter IV(2) of Annex D to Directive 92/65/EEC;							
ll: Cert	(¹) either	[11.4.	are in vivo derived embryos which:							
Part		II.4.1.	were conceived as a result of artificial insemination	with semen meeting the requirement	s of Directive 90/429/EEC,					
		II.4.2.	originate from a Member State or region thereof:							
		(¹) either	[listed in Annex I to Decision 2008/185/EC and are de 2008/185/EC;]	estined for a Member State or region	thereof listed in Annex I to Decision					
		(1) 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	[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					
		(¹) or	[listed in Annex II to Decision 2008/185/EC and are de 2008/185/EC;]	estined for a Member State or region	thereof listed in Annex II to Decision					
		(¹) or	[not listed in Annex I or II to Decision 2008/185/EC ar to Decision 2008/185/EC and have been washed wi		region thereof listed in Annex I or II					
		(¹) or	[not listed in Annex I or II to Decision 2008/185/EC ar or II to Decision 2008/185/EC;]]	nd are destined for a Member State o	r region thereof not listed in Annex I					
	(¹) or	[11.4.	are in vitro produced/micromanipulated (1) embryos v	which:						
		II.4.1.	were conceived as a result of in vitro fertilisation wit	h semen meeting the requirements o	of Directive 90/429/EEC,					
		II.4.2.	originate from a Member State or region thereof:							
		(¹) either	[listed in Annex I to Decision 2008/185/EC and are de 2008/185/EC;]	estined for a Member State or region	thereof listed in Annex I to Decision					
		(¹) 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	[listed in Annex II to Decision 2008/185/EC and are d 2008/185/EC and the donor females of the ova used 2008/185/EC;]							
		(¹) or	[listed in Annex II to Decision 2008/185/EC and are de 2008/185/EC;]	estined for a Member State or region	thereof listed in Annex II to Decision					
		(¹) or	[not listed in Annex I or II to Decision 2008/185/EC ar to Decision 2008/185/EC and the donor females of th Decision 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;]]	nd are destined for a Member State o	r region thereof not listed in Annex I					

EUROPEAN UNION

Porcine ova/embryos — Part A

П.	Health info	rmation	II.a. Certificate reference No	II.b.			
(¹) or	[1].4.	are in vivo derived ova which originate from a Mem	ber State or region thereof:				
	(¹) either	[listed in Annex I to Decision 2008/185/EC and are c 2008/185/EC;]	lestined for a Member State or regior	n thereof listed in Annex I to Decision			
	(¹) or	[listed in Annex I to Decision 2008/185/EC and are o Decision 2008/185/EC;]	destined for a Member State or regio	n thereof not listed in Annex I or II to			
	(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 Deci 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 c 2008/185/EC;]	destined for a Member State or regior	n thereof listed in Annex II to Decision			
	(¹) or	[not listed in Annex I or II to Decision 2008/185/EC a to 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 contair Directive 92/65/EEC and bearing the number detaile		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 to	eam, embryo production team or to th	he holding of ova/embryos destination.			
Box 1.23	3: identifica	tion of container and seal number shall be indicated.					
Box I.31	I: category	: specify if: <i>in vivo</i> derived embryos, <i>in vivo</i> derived o	ova, <i>in vitro</i> produced embryos or mi	icromanipulated embryos.			
	donor ide	entity shall correspond to the official identification of t	he animal.				
		collection shall be indicated in the following format: do					
		number of the team shall correspond to the embry on indicated in Box I.12.	o collection team or embryo produc	tion team of ova/embryos collection/			
Part II:							
	te as appro	•		offer op/or/FEO on the Commission			
webs	site:	embryo collection or production teams listed in accord		ective 92/65/EEC on the Commission			
http:/	//ec.europa.	eu/food/animal/approved_establishments/establishmer	its_vet_field_en.htm				
Official veterinarian or official inspector (*)							
Na	ame (in capi	ital letters):	Qualification and title:				
Local veterinary unit:			LVU No:				
Date:			Signature:				
Stamp:							
(*) The c		stamp and signature must be different from that of the other p	articulars in the certificate				
1, 1100	*) 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

In Consignor I.2. Cartificate reference No I.2.a. Local reference Address I.3. Certral competent authority I.3. Certral competent Authority Postal code I.4. Local Competent Authority I.4. Local Competent Authority Address I.3. Certral competent Authority I.4. Local Competent Authority Address I.7. I.6. Address I.7. Postal code I.7. I.8. Country of argin ISO code I.12. Place of origin I.13. Place of destination I.14. Place of origin I.13. Place of destination I.14. Place of origin I.14. Place of destination I.15. Commodity of argin I.16. Commodity code (HS code) I.16. Means of transport I.17. Address Postal code I.16. Means of transport I.17. Address Other Identification I.17. I.18. Description of commodity I.19. Commodity code (HS code) I.20. Outanity I.24. Type of packages Ambient Chilled Frozen I.23. Seal/Container No I.24. Type of packages I.24. Type of packages <	de certificate				
Address I.3. Central competent authority Postal code I.4. Local Competent Authority I.5. Consignee I.6. Name I.7. Address I.7. Postal code I.7. Postal code I.1. Country of origin I.1.2. Place of origin ISO code I.9. Region of origin I.1.2. Place of origin I.1.1. Region of destination Name Approval number Address Postal code I.1.2. Place of origin I.1.3. Place of destination Name Approval number Address Postal code I.1.4. I.1.5. I.1.4. I.1.5. I.1.4. I.1.5. I.1.5. I.1.7. Address Postal code I.1.4. I.1.5. I.1.5. I.1.7. Acdress Postal code I.1.6. Means of transport I.1.7. Acdress I.1.7. I.1.8. Description of commodity I.1.7. I.1.8. Description of commodity I.1.1.9. Commodity code (HS code) I.2.0. Quantly I.2.0. Quantly </th <th>ce No</th>	ce No				
Postal code 14. Local Competent Authority 15. Consignee 16. Name 17. Address 17. Postal code 17. Its. Country of origin ISO code 19. Region of origin Code 112. Place of origin ISO code 19. Region of origin Code 112. Place of origin Embryo team 113. Place of destination Name Approval number Address Postal code Postal code Postal code 114. I15. I16. 114. I16. Embryo team Name Approval number Name Address Postal code Postal code 114. I15. I17. Aeropiane Ship Railway wagon I17. Road vehicle Other I18. Description of commodity I17. I18. Description of commodity I118. Commodity code (HS code) 05 11 99 85 I20. Quantity I210. Quantity I22. Number of packages Ambient Chilled Frozen I24. Type of packages I28. Transit through hird country ISO code <th></th>					
15. Consignee 16. Name Address Potal code 17. 18. Country of origin ISO code 19. Region of origin Code 11. Potal code 11. Country of origin ISO code 11. Protal code 11. Country of destination ISO code 11. Protation Embryo team 11. Protocol Name Approval number 11. Protocol Holding Name Approval number Address Postal code Postal code Postal code 11. Protocol Embryo team Name Approval number Address Postal code 11. Protocol 1.14. 1.15. 11. Protocol 11. Protocol 11. Protocol 1.14. 1.15. 11. Protocol 11. Protocol 11. Protocol 1.14. 1.15. 11. Protocol 12. Protocol 12. Protocol 1.18. Description of commodity 1.17. 12. Protocol 12. Protocol 12. Protocol 1.13. Sea/Container No 1.20. Quantity 12. Protocol 12. Protocol 12. Protocol 12. Protocol 12. Protocol 12. Protocol 12. Protocol </th <th></th>					
Address I.7. Postal code I.7. I.8. Country of origin ISO code I.9. Region of origin Code I.10. Country of destination ISO code I.11. Region of destination I.1.2. Place of origin I.1.2. Place of origin I.1.3. Place of destination I.1.4. I.1.5. Place of destination I.1.7. Name Approval number Address Postal code Postal code I.1.4. I.1.4. I.1.5. I.1.6. I.1.6. I.1.7. Address Postal code Special code Postal code I.1.7. Address Postal code Other C I.1.6. I.1.7. I.1.7. I.1.4. I.1.5. I.1.7. I.1.7. I.1.7. Read vehicle					
Address I.7. Postal code I.7. I.8. Country of origin ISO code I.9. Region of origin Code I.12. Place of origin I.11. Region of origin I.13. Place of destination I.14. I.14. I.15. Holding Embryo team Holding Embryo team Name Approval number Address Postal code Postal code I.14. I.14. I.15. I.16. Means of transport Address Postal code I.17. Road vehicle Other I.16. Means of transport I.17. I.17. Road vehicle Other I.19. Commodity code (HS code) 05 11 99 85 I.20. Ouantity I.20. Ouantity I.20. Ouantity I.20. Ouantity I.21. Temperature of products I.22. Number of packages Ambient I.20. Ouantity I.23. Seal/Container No I.24. Type of packaging I.26. Transit through third country I.27. Transit through Member State ISO code Entry point BiP No Member State ISO code Member State ISO code Exit point Code Member State ISO code ISO code <th colspan="5">1.6.</th>	1.6.				
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Secondation Octown Decondation Octown Decondation 1.12. Place of origin I.13. Place of destination Holding I Embryo team Name Approval number Name Approval number Address Postal code Postal code Intra- 1.14. I.15. Intra- Intra- Intra- I.16. Means of transport Railway wagon I Intra- Intra- Road vehicle Other I Other I Intra- Intra- I.18. Description of commodity Intra- Intra- Intra- I.19. Commodity code (HS code) 05 11 99 85 Intra- Intra- I.20. Quantity Intra- Intra- Intra- Intra- I.21. Temperature of products Intra- Intra- Intra- Intra- I.23. Seal/Container No Intra- Intra- Intra- Intra- Intra- I.26. Transit through third country Intra- Intra- Intra- Intra- Intra- I.28. Export Intra- Intra- Intra- Intra- Intra- Intra- I.28. Export Intra- Intra-	1.7.				
Totagent Contractor:					
Address Address Postal code Postal code 1.14. 1.15. 1.14. 1.15. I.16. Means of transport 1.17. Aeroplane Ship Railway wagon Road vehicle Other Identification I.17. I.18. Description of commodity I.19. Commodity code (HS code) 05 11 99 85 I.20. Quantity I.21. Temperature of products I.22. Number of packages Ambient Chilled Frozen I.23. Seal/Container No I.24. Type of packaging I.25. Commodities certified for: Artificial reproduction I.26. Transit through third country I.27. Transit through Member States ISO code Exit point Code Member State ISO code Exit point Code Member State ISO code I.30. I.29. I.29. I.29. I.31. Identification of the commodities Species Breed Category Donor identity Date of Approval number	Code				
Address Address Postal code Postal code 1.14. 1.15. 1.14. 1.15. I.16. Means of transport 1.17. Aeroplane Ship Railway wagon Road vehicle Other Identification I.17. I.18. Description of commodity I.19. Commodity code (HS code) 05 11 99 85 I.20. Quantity I.21. Temperature of products I.22. Number of packages Ambient Chilled Frozen I.23. Seal/Container No I.24. Type of packaging I.25. Commodities certified for: Artificial reproduction I.26. Transit through third country I.27. Transit through Member States ISO code Exit point Code Member State ISO code Exit point Code Member State ISO code I.30. I.29. I.29. I.29. I.31. Identification of the commodities Species Breed Category Donor identity Date of Approval number	 m				
Address Address Postal code Postal code 1.14. 1.15. 1.14. 1.15. I.16. Means of transport 1.17. Aeroplane Ship Railway wagon Road vehicle Other Identification I.17. I.18. Description of commodity I.19. Commodity code (HS code) 05 11 99 85 I.20. Quantity I.21. Temperature of products I.22. Number of packages Ambient Chilled Frozen I.23. Seal/Container No I.24. Type of packaging I.25. Commodities certified for: Artificial reproduction I.26. Transit through third country I.27. Transit through Member States ISO code Exit point Code Member State ISO code Exit point Code Member State ISO code I.30. I.29. I.29. I.29. I.31. Identification of the commodities Species Breed Category Donor identity Date of Approval number					
Postal code Postal code 1.14. 1.15. 1.16. Means of transport Aeroplane Aeroplane Ship Road vehicle Other Identification 1.17. 1.18. Description of commodity 1.17. 1.18. Description of commodity 1.19. Commodity code (HS code) 05 11 99 85 1.20. Quantity 1.21. Temperature of products 1.22. Number of packages Ambient Chilled Frozen 1.23. Seal/Container No 1.24. 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 Exit point Code Member State ISO code Exit point Code I.30. 1.29. 1.31. Identification of the commodities Species Breed Category Donor identity Date of 1.30. I.31. Identification of the commodities Species Approval number	mbol				
1.14. I.15. 1.16. Means of transport I.17. Aeroplane Ship Railway wagon Road vehicle Other Identification I.17. 1.18. Description of commodity I.19. Commodity code (HS code) 05 11 99 85 I.20. Quantity 1.21. 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 I.90 code Exit point Code Member State ISO code Exit point Code Member State ISO code I.28. Export I.29. Third country ISO code I.29. Third country ISO code I.29. Third country ISO code I.29. Third country ISO code I.29. Third country ISO code I.29. Third country ISO code I.29. Third country ISO code I.29.					
1.16. Means of transport 1.17. Aeroplane Ship Railway wagon Road vehicle Other 1.17. Identification 1.18. Description of commodity 1.19. Commodity code (HS code) 05 11 99 85 1.20. Quantity 1.21. Temperature of products 1.22. Number of packages Ambient Chilled Frozen 1.23. Seal/Container No 1.24. Type of packaging 1.25. Commodities certified for: Artificial reproduction Artificial reproduction 1.27. Transit through Member States I Third country ISO code Member State ISO code Exit point Code Member State ISO code I.28. Export I.29. Third country ISO code I.30. I.31. Identification of the commodities Species Breed Category Donor identity Date of Approval number					
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	EUROPEA	N UNION	I	Porcine ova/embryos — Part B					
	II.	Health int	formation	II.a. Certificate reference No	II.b.				
	I, the und	ersigned	official veterinarian, hereby certify that:						
		ll.1.	The ova/embryos (1) described above:						
II.1.1. were collected, processed and stored under conditions which meet the requirements of Directive 92/65/E									
		ll.1.2.	come from donor female swine which meet the requ	meet the requirements of Chapter IV of Annex D to Directive 92/65/EEC;					
tion		ll.1.3.	meet the requirements of Chapter III of Annex D to	Directive 92/65/EEC.					
Part II: Certification	(¹) either	[II.2.	n the case of embryos,						
ö ≓		ll.2.1.	the semen used for fertilisation meets the requirement	ents of Directive 90/429/EEC;					
Ра		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 donc 2008/185/EC (²).]	or female swine which meets the c	conditions of Article 1 of Decision				
	Notes								
	Part I:								
	Box 1.12:	place of	origin shall correspond to the embryo collection team	n of ova/embryos collection.					
	Box I.13: place of destination shall correspond to the embryo collection team, embryo production team or to the holding of ova/embryo destination.								
	Box 1.23:	identifica	tion of container and seal number shall be indicated.						
	Box I.31:		: specify if: <i>in vivo</i> derived embryos, <i>in vivo</i> derived o entity shall correspond to the official identification of t		cromanipulated embryos.				
			collection shall be indicated in the following format: do						
		approval	number of the team shall correspond to the embryo	collection team of ova/embryos colle	ction indicated in Box I.12.				
	Part II:								
	(¹) Delete	as appro	priate.						
	(²) This condition applies only to ova and embryos which originate in the Member States or regions thereof not listed in Annexes I and II to Decis 2008/185/EC (OJ L 59, 4.3.2008, p. 19) and destined to the Member States or regions thereof so listed. It shall also apply to movements fr Member States or regions thereof listed in Annex II of Decision 2008/185/EC to Member States or regions thereof listed in Annex I of Decision 2008/185/EC.								
	Official ve	eterinarian	or official inspector (*)						
	Nan	ne (in capi	ital letters):	Qualification and title:					
	Loca	al veterina	ry unit:	LVU No:					
	Date	9:		Signature:					
	(*) The col	our of the s	stamp and signature must be different from that of the other ${\mathfrak p}_i$	articulars in the certificate.					