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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)

(OJ L 228, 31.8.2010, p. 15)

Amended by:

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							No	page	date
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COMMISSION DECISION

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(notified under document C(2010) 5779)

(Text with EEA relevance)

(2010/470/EU)

THE EUROPEAN COMMISSION,

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

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

Whereas:

- (1) Directive 92/65/EEC lays down the animal health requirements governing trade within the Union of animals, semen, ova and embryos not subject to the animal health requirements laid down in specific Union acts. It includes requirements for trade in semen, ova and embryos of animals of the equine, ovine and caprine species and in ova and embryos of animals of the porcine species ('the commodities'). In addition, it provides for health certificates to be established for trade in the commodities within the Union.
- (2) Annex D to Directive 92/65/EEC, as amended by Commission Regulation (EU) No 176/2010 (²), sets out certain new requirements for the commodities which are to apply from 1 September 2010.
- (3) Annex D to Directive 92/65/EEC, as thus amended by Regulation (EU) No 176/2010, introduces rules concerning semen storage centres and detailed conditions for their approval and supervision. It also sets out detailed conditions for the approval and supervision of embryo collection and production teams, for the collection and processing of *in vivo* derived embryos and the production and processing of *in vitro* fertilised embryos and micromanipulated embryos. Annex D, as thus amended, also amended the conditions to be applied to the donor animals of semen, ova and embryos of animals of the equine, ovine and caprine species and of ova and embryos of porcine species.

^{(&}lt;sup>1</sup>) OJ L 268, 14.9.1992, p. 54.

⁽²⁾ OJ L 52, 3.3.2010, p. 14.

- It is necessary to establish new model health certificates for trade (4) 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.
- In the interests of consistency and simplification of Union legis-(7) lation, 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 $(^1)$.
- In order to ensure full traceability of the commodities, model (8) 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 acts (9) 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.

⁽¹⁾ OJ L 94, 31.3.2004, p. 44.

^{(&}lt;sup>2</sup>) OJ L 182, 2.8.1995, p. 27.
(³) OJ L 185, 4.8.1995, p. 58.

^{(&}lt;sup>4</sup>) OJ L 234, 3.10.1995, p. 30.

⁽⁵⁾ OJ L 275, 18.11.1995, p. 30.

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

HAS ADOPTED THIS DECISION:

Article 1

Subject matter

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

- (a) semen of animals of the equine species;
- (b) ova and embryos of animals of the equine species;
- (c) semen of animals of the ovine and caprine species;
- (d) ova and embryos of animals of the ovine and caprine species;
- (e) ova and embryos of animals of the porcine species.

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.

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

ROP	ΡEA	N UNION		Intra trade certificat		
1.	.1.	Consignor	I.2. Certificate reference No	I.2.a. Local reference No		
		Name Address	I.3. Central competent authority			
		Postal code	I.4. Local competent authority			
5 I.	.5.	Consignee	1.6.			
5		Name				
		Address	1.7.			
		Postal code				
	.8.	Country of origin ISO I.9. Region of origin Code code	I.10. Country of ISO destination code	I.11. Region of Code destination		
I.	.12.	Place of origin	I.13. Place of destination			
		Semen centre	Semen centre	Holding		
		—	_	• —		
•		Name Approval number Address	Name	Approval number		
		Postal code	Address			
		Postal code	Postal code			
1.	.14.		1.15.			
1.	.16.	Means of transport	I.17.			
		Aeroplane 🗌 Ship 🗌 Railway wagon 🗌				
		Road vehicle Other				
		Identification				
-	10	Description of commodity				
1.	.10.	Description of commonly	I.19. Commodity coo	de (HS code) 5 11 99 85		
			1.20.	Quantity		
1.	.21.	Temperature of product	I.22. Number of packages			
		Ambient Chilled	Frozen			
1.	.23.	Seal/container No	1.24.	Type of packaging		
t.	.25.	Commodities certified for: Artificial reproduction				
+			I.27. Transit through Member States			
1.	.26.	Transit through a third country	Member State			
		Third country ISO code	Member State	ISO code		
		Exit point Code	Member State	ISO code		
		Entry point BIP No	Member State	ISO code		
1.	.28.	Export 🗌	1.29.			
		Third country ISO code				
		Exit point Code				
I.	.30.					
1.	.31.	Identification of the commodities				
		Species Breed Donor identity D	Date of collection Approval nu	mber Quantity		
		(scientific name)	of the tea			

	EUROPE	AN UNION			Equine semen — Part A						
	П.	Health inforr	nation		II.a. Certificate reference No	II.b.					
	I, the un	idersigned of	ficial veter	inarian, hereby certify that:							
	11.1.	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(I)(1) of Annex D to Directive 92/65/EEC;									
_	II.1.1.	during the period commencing 30 days prior to the date of first collection of the semen described above until the date the fresh or chilled semen was dispatched or until the 30 days storage period for frozen semen elapsed, the semen collection centre;									
Part II: Certification	II.1.1.1.	was situated on the territory or in the case of regionalisation in a part of the territory (1) of a Member State which was not considered to be infected with African horse sickness in accordance with Article 5(2)(a) and (b) of Directive 2009/156/EC (3);									
art II: C	II.1.1.2.	fulfilled the	conditions	for a holding laid down in Article 4(5) of	Directive 2009/156/EC;						
å	II.1.1.3.	contained c	only equida	e which were free of clinical signs of equ	uine viral arteritis and contagious equi	ne metritis.					
	11.2.	Only equidation the centre.	ae satisfyin	g the conditions laid down in Articles 4 an	nd 5 or Articles 12 to 16 of Directive 20	09/156/EC have been admitted into					
	II.3.	The semen	described	above was collected from donor stallion	s, which:						
	II.3.1.	have not sh was collect		linical sign of an infectious or contagious o	disease at the time of admission into th	ne centre and on the day the semen					
	II.3.2.) days prior to the date of semen collectio equine metritis during that period;	on in holdings where no equine has sh	own any clinical sign of equine viral					
	II.3.3.			or natural mating during at least 30 days points II.3.5.1, II.3.5.2 or II.3.5.3 until the	is prior to the date of first semen collection and from the dates of the first is end of the collection period;						
	II.3.4.	Vaccines fo	r Terrestri		requirements of the relevant chapter of the Manual of Diagnostic Tests and amples taken in accordance with one of the programmes specified in point						
		(¹) either	[11.3.4.1.	an agar-gel immuno-diffusion test (Cogg	ins test) for equine infectious anaemia	a (EIA) with negative result;]					
		(¹) or	[11.3.4.1.	an ELISA for equine infectious anaemia	(EIA) with negative result;]						
	and	(¹) either	[.3.4.2.	a serum neutralisation test for equine vi	ral arteritis (EVA) with negative result	at a serum dilution of one in four;]					
		(¹) or	[11.3.4.2.	a virus isolation test for equine viral arteri of the donor stallion;]	itis (EVA) carried out with negative res	ult on an aliquot of the entire semen					
	and		[11.3.4.3.	an agent identification test for contagiou with an interval of seven days by isolati ejaculatory fluid or a semen sample an urethral fossa with negative result in eac	on of <i>Taylorella equigenitalis</i> after a c d from genital swabs taken at least fi	ultivation of 7 to 14 days from pre-					
	II.3.5.	have been II.3.5.2 and		with the results specified in II.3.4 in each follows:	case to at least one of the test progra	ammes (⁴) detailed in points II.3.5.1,					
	II.3.5.1.	during the p	eriod of co	continuously resident on the semen colie ollection of the semen described above an th status than the donor stallion;							
				n point II.3.4 have been carried out on si the commencement of the residence peri-		en collection and at least 14 days					

	AN UN	IUN					Equin	e semen — Part A
П.	Health	information			II.a. Certific	ate reference No	II.b.	
II.3.5.2.	period contini	of collection of the	semen descri	bed above, but has	s left the cent	0 days prior to the d re under the respor ction centre came ir	nsibility of the centre	e veterinarian for a
	collect		r the semen de) prior to the first sen at least 14 days follo		
and		t described in point I the semen describe			a was last carr	ied out on a sample	of blood taken (⁵) no l	more than 90 days
and	(¹) eith			point II.3.4.2 for equ ribed above was co		s was last carried ou	t on a sample taken (⁽⁵) not more than 30
	(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;]							
and	the test described in point II.3.4.3 for contagious equine metritis was last carried out on samples taken ⁽⁵⁾ not more than 60 days before the semen described above was collected.							
II.3.5.3.		sts described in poin ion period in the yea) prior to the first ser	men collection of the	breeding season or
and		its described in point ion of the semen de			nples taken (⁵)	not less than 14 da	iys and not more tha	n 90 days after the
II.3.6.	Have	undergone the testing	g provided for	in point II .3.5 on sa	mples taken o	n the following dates	:	
		Start date	(⁵)		Date	of sampling for hea	ulth tests (5)	
catior men	amme	Donor residence	Semen collection	EIA II.3.4.1	EVA .3.4.		CEI II.3.4	
Identification of semen	Test programme	brogr			lood Imple	Semen sample	1. sample	2. sample
(¹) eithe	er [11.4	No antibiotics we	re added to the	e semen;]			I	
(1) or	[1].4					produce a concentrat		d semen of not less
								;]
II.5.	The	semen described al	oove was:					
ll.5.1.		ected, processed, sto Directive 92/65/EE0		orted under conditio	ns which comp	ly with the requireme	nts of Chapters II(I)(1) and III(I) of Annex
II.5.2.		to the place of load ring the number indic			ance with poin	t 1.4 of Chapter III(I)	of Annex D to Direc	tive 92/65/EEC and

	IN UNION	II.a. Certificate reference No	Equine semen — Part /
Notes			
Part I:			
Box 1.12:	place of origin shall correspond to the semen collection centr	e of origin of the semen.	
Box I.13:	place of destination shall correspond to the semen collection	or storage centre or to the holding of	semen destination.
Box 1.23:	identification of container and seal number shall be indicated.		
Box I.31:	donor identity shall correspond to the official identification of t	the animal.	
	date of collection shall be indicated in the following format: de	d/mm/yyyy.	
	approval number of the centre shall correspond to the approva collected.	al number of the semen centre indicate	d in Box I.12 where the semen was
Part II:			
Guidance	for the completion of Table in II.3.6:		
Abbrevial	ions:		
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 occ	casion	
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	er CEM-11	
CEM-21	CEM testing second occasion first sample		
CEM-22	CEM testing second occasion second sample taken 7 days	after CEM-21	
Instructio	ns:		
	semen identification in column A in the example below, the tes olumns C and D must be completed with the dates required.	st programme (II.3.5.1, II.3.5.2 and/or II.	3.5.3) must be described in colum
and II.3.	is when samples where taken for laboratory testing prior to the f 5.3, are entered in the upper line of columns 5 to 9 of the table, <i>A</i> -12 in the example below.		
	is when samples were taken for repeat laboratory testing as rec 5 to 9 in table, this being the boxes EIA-2, EVA-B2 or EVA-S		

II.	Health i	information			II.a. C	ertificate reference No	II.b.			
-		Start date	ə (⁵)			Date of sampling for health tests (5)				
Identification of semen	Test programme	Donor residence	Semen collection	EIA II.3.4.1	I	EVA 1.3.4.2		CEM II.3.4.3		
ofs	pro <u>ç</u>				Blood sample	Semen sample	1. sample	2. sample		
•	в	0		EIA-1	EVA-B1	EVA-S1	CEM-11	CEM-12		
A	В	С	D	EIA-2	EVA-B2	EVA-S2	CEM-21	CEM-22		
(⁴) Cro	oss out the	3.7.2010, p. 1. e programme(s) tha n table in point II.3.6			•					
(⁴) Cro (⁵) Ins (⁶) Ins	oss out the ert date in ert names I veterinar	e programme(s) tha n table in point II.3.6 s and concentrations ian (*)	6 (follow guidar		•					
(⁴) Cro (⁵) Ins (⁶) Ins	ert date in ert date ir ert names I veterinar Name	e programme(s) than a table in point II.3.6 and concentrations ian (*) (in capital letters):	6 (follow guidar		•	Qualification and title	:			
(⁴) Cro (⁵) Ins (⁶) Ins	ert date in ert names I veterinar Name Local v	e programme(s) tha n table in point II.3.6 s and concentrations ian (*)	6 (follow guidar		•	LVU No:				
(⁴) Cro (⁵) Ins (⁶) Ins	ert date ir ert names I veterinar Name Local v Date:	e programme(s) than a table in point II.3.6 a and concentrations ian (*) (in capital letters): zeterinary unit:	6 (follow guidar		•		:			
(⁴) Cro (⁵) Ins (⁶) Ins	ert date in ert names I veterinar Name Local v	e programme(s) than a table in point II.3.6 a and concentrations ian (*) (in capital letters): zeterinary unit:	6 (follow guidar		•	LVU No:				

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

EUR	OPEA	N UNION	Intra trade certificate				
	l.1.	Consignor	I.2. Certificate reference No I.2.a. Local reference No				
		Name	I.3. Central competent authority				
		Address					
ted		Postal code	I.4. Local competent authority				
presented	1.5.	Consignee	1.6.				
t pr		Name					
nen		Address	1.7.				
ign		Postal code					
consignment	1.8.	Country of origin ISO I.9. Region of origin Code code	I.10. Country of ISO I.11. Region of Code destination code destination				
° of							
Part I: Details	1.12.	Place of origin	I.13. Place of destination				
ŏ		Semen centre	Semen centre				
art		Name Approval number	Name Approval number				
ä		Address	Address				
		Postal code	Postal code				
	I.14.		I.15.				
	I.16.	Means of transport	I.17.				
		Aeroplane 🗌 Ship 🗌 Railway wagon 🗌					
		Road vehicle Other Other					
		Identification					
	1 18	Description of commodity	I.19. Commodity code (HS code)				
			05 11 99 85				
			I.20. Quantity				
	1.01	Tomporative of product					
	1.21.	Temperature of product	I.22. Number of packages				
		Ambient Chilled	Frozen				
	1.23.	Seal/container No	I.24. Type of packaging				
	1.25.	Commodities certified for:	I				
		Artificial reproduction					
	1.26.	Transit through a third country	I.27. Transit through Member States				
		Third country ISO code	Member State ISO code				
		Exit point Code	Member State ISO code				
		Entry point BIP No	Member State ISO code				
	1.28.	Export	1.29.				
		Third country ISO code					
		Exit point Code					
	1.30.						
	1,31	Identification of the commodities					
			te of collection Approval number Quantity of the team				

	EUROPE	AN UNION		Equine semen — Part B					
	П.	Health information		II.a. Certificate reference No	II.b.				
	I, the ur	ndersigned official vete	rinarian, hereby certify that:						
	II.1.	The semen collection	centre (2), in which the semen described	above was collected, processed and	stored for trade:				
	II.1.1.	is approved and sup	ervised by the competent authority accor	rding to the conditions of Chapter I o	f Annex D to Directive 92/65/EEC;				
Part 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);							
O ⇒ II.1.3. fulfilled during the period commencing 30 days prior to the date of semen collection until the date the semen was dispatched to chilled (1) semen or until the 30 days mandatory storage period for frozen semen elapsed (1), the conditions of Article 4 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;								
	11.2.	All equidae have bee	n admitted into the centre under the prov	visions of Article 4 and 5 of Directive 2	2009/156/EC.				
	II.3.	The semen described	above was collected from donor stallion	is, which:					
	II.3.1. on the day the semen was collected have not shown clinical signs of an infectious or contagious disease;								
	11.3.2.	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 day arteritis;	s prior to collection of the semen have bee	en kept on holdings where no equidae	showed clinical signs of equine viral				
	II <i>.</i> 3.4.	during the last 60 day equine metritis;	rs prior to collection of the semen have be	en kept on holdings where no equidae	showed clinical signs of contagious				
	II.3.5.		lowledge and as far as I could ascertain luring 15 days immediately preceding coll		dae suffering from an infectious or				
	II.3.6 <i>.</i>		ollowing animal health tests, carried out in pecified in point II.3.7;	a laboratory recognised by the compe	etent authority, in accordance with a				
		[11.3.6.7	. an agar gel immunodiffusion test (Cog	ggins test) for equine infectious anaem	ia with negative result;]				
	and	(¹) either [II.3.6.2	2. a serum neutralisation test for equine	uine viral arteritis with negative result at a serum dilution of one in for					
		(¹) or [II.3.6.2	 a virus isolation test for equine viral ar the donor stallion;] 	teritis carried out with negative result o	on an aliquot of the entire semen of				
	and	II.3.6.3	 an agent identification test for contagio the donor stallion with an interval of sev a semen sample and from genital swa negative result in each case; 	ven days by isolation of Taylorella equi	genitalis from pre-ejaculatory fluid or				
	II.3.7.	have been subject to	the one of the following test programme:	s (⁴):					
	II.3.7.1.		s continuously resident in the collection of no equidae in the collection centre came						
		the tests described in point II.3.6 have been carried out on samples taken on							

II. Health information II.a. Certificate reference No II.b. II.3.7.2. the dorse relation was not continuously resident in the collection centre or other equidae in the collection centre came into contact with equidae of lower health tables than the donard station: (*) being within the 14 days particle before a same on a sample taken on	EUROPEAN UNION Equine semen —						
equidae of lower health status than the donor statilon: the tests described in point II.36 have been carried out on samples taken on	П.	Health inform	ation	II.a. Certificate reference No	ll.b.		
contragious equipmentation and at least at the beginning of the breeding seasor: (b) being within the 14 days period before the first semen collected in point II 3.6.1 for equips infectious ansemia was last carried out on a sample of blood taken on	11.3.7.2.			on centre or other equidae in the colle	ection centre came into contact with		
and (*) etiliter [One of the test described in point II.3.6.2 for equine viral arteritis was list carried out on a sample collected on		contagious	equine metritis on a second sample taken on				
(*) being not more than 30 days before the semen described above was collected.) (*) or [(he non-shedder tata of the semen some of the done salion collected on	and						
acried out on an aliquot of the entrie semen described above was collected.] (?), being not more than one year before the semen described above was collected.] II.3.7.3. The tests described in point II.3.6 have been carried out during the 30 days mandatory storage period of frozen semen and not less than 14 days after the collection of the semen on samples taken on	and	(¹) either					
14 days after the collection of the semen on samples taken on		(¹) or	carried out on an aliquot of the entire semen of	the donor stallion collected on			
Chapters II and III of Annex D to Directive 92/65/EEC. Notes Part I: Box 1.12: place of origin shall correspond to the semen collection centre of origin of the semen. Box 1.13: place of destination shall correspond to the semen collection or storage centre or to the holding of semen destination. Box 1.23: identification of container and seal number shall be indicated. Box 1.31: 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 1.12 where the semen was collected. Part II: (*) Delete as appropriate. (*) Only approved semen collection centres listed in accordance with Artole 11(4) of Council Directive 92/65/EEC on the Commission website: http://ec.europa.eu/food/anima/approved_establishments_vet_field_en.htm (*) Otal 192, 23.7.2010, p. 1. (*) Cross out the programme(s) that do(es) not apply to the consignment. (*) Insert date. Official veterinarian or official inspector (*) Name (in capital letters): Local veterinary unit: Local veterinary unit: Date: Stamp:	11.3.7.3.	14 days aft	er the collection of the semen on samples taken o	on (⁵) and in the			
Part I: Box 1.12: place of origin shall correspond to the semen collection centre of origin of the semen. Box 1.13: place of destination shall correspond to the semen collection or storage centre or to the holding of semen destination. Box 1.23: identification of container and seal number shall be indicated. Box 1.31: 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 1.12 where the semen was collected. Part II: (1) Delete as appropriate. (2) Only approved semen collection centres listed in accordance with Article 11(4) of Council Directive 92/65/EEC on the Commission website: http://ec.europa.eu/food/anima/approved_establishments/establishments_vet_field_en.htm (2) OL 1 192, 23.7.2010, p. 1. (3) Cross out the programme(s) that do(es) not apply to the consignment. (4) Insert date. Official veterinarian or official inspector (*) Name (in capital letters): Qualification and title: Local veterinary unit: LVU No: Date: Signature: Stamp: Signature:	11.4.			and transported under conditions wh	ich comply with the requirements of		
Box 1.12: place of origin shall correspond to the semen collection centre of origin of the semen. Box 1.13: place of destination shall correspond to the semen collection or storage centre or to the holding of semen destination. Box 1.23: identification of container and seal number shall be indicated. Box 1.31: 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 1.12 where the semen was collected. Part II: 1) Delete as appropriate. (?) 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/anima/approved_establishments/establishments_vet_field_en.htm (?) OL 192, 23.7.2010, p. 1. (d) Cross out the programme(s) that do(es) not apply to the consignment. (?) Insert date. Official veterinarian or official inspector (*) Name (in capital leiters): Local veterinary unit: Local	Notes						
Box 1.13: place of destination shall correspond to the semen collection or storage centre or to the holding of semen destination. Box 1.23: identification of container and seal number shall be indicated. Box 1.31: 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 1.12 where the semen was collected. Part II: (1) Delete as appropriate. (2) Only approved semen collection centres listed in accordance with Article 11(4) of Council Directive 92/65/EEC on the Commission website: http://ec.europa.eu/food/animal/approved_establishments/vet_field_en.htm (3) OJ L 192, 23.7.2010, p. 1. (4) Cross out the programme(s) that do(es) not apply to the consignment. (5) Insert date. (7) Insert date. (7) Insert date. (7) Name (in capital letters): Local veterinary unit: LVU No: Date: Stamp:	Part I:						
Box 1.23: identification of container and seal number shall be indicated. Box 1.23: identification of container and seal number shall be indicated. Box 1.31: 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 1.12 where the semen was collected. Part II: 1) Delete as appropriate. (?) 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/tood/animal/approved_establishments/establishments_vet_field_en.htm (?) OL L 192, 23.7.2010, p. 1. (*) Cross out the programme(s) that do(es) not apply to the consignment. (*) Insert date. Official veterinarian or official inspector (*) Name (in capital letters): Local veterinary unit: Local veterinary unit: Local veterinary unit: Local veterinary unit: Stamp:	Box 1.12	2: place of ori	gin shall correspond to the semen collection centr	e of origin of the semen.			
Box I.31: donor identity shall correspond to the official identification of the animal. date of collection shall be indicated in the following format: dd/mm/yyyy. approval number of the centre shall correspond to the approval number of the semen centre indicated in Box I.12 where the semen was collected. Part II: (¹) Delete as appropriate. (²) Only approved semen collection centres listed in accordance with Article 11(4) of Council Directive 92/65/EEC on the Commission website: http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm (²) OL L 192, 23.7.2010, p. 1. (²) Cross out the programme(s) that do(es) not apply to the consignment. (?) Insert date. Official veterinarian or official inspector (*) Name (in capital letters): Qualification and title: Local veterinary unit: LVU No: Date: Signature: Stamp: Stamp:	Box 1.13	3: place of de	stination shall correspond to the semen collection	or storage centre or to the holding of	semen destination.		
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 1.12 where the semen was collected. Part II: (1) Delete as appropriate. (2) Only approved semen collection centres listed in accordance with Article 11(4) of Council Directive 92/65/EEC on the Commission website: http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm (2) OL L 192, 23.7.2010, p. 1. (4) Cross out the programme(s) that do(es) not apply to the consignment. (5) Insert date. Official veterinarian or official inspector (*) Name (in capital letters): Qualification and title: Local veterinary unit: LVU No: Date: Signature: Stamp:	Box 1.23	3: identification	n of container and seal number shall be indicated.				
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: (*) Delete as appropriate. (*) 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_estabilishments/estabilishments_vet_field_en.htm (*) OJ L 192, 23.7.2010, p. 1. (*) Cross out the programme(s) that do(es) not apply to the consignment. (*) Insert date. Official veterinarian or official inspector (*) Name (in capital letters): Local veterinary unit: Date: Stamp:	Box I.31	1: donor ident	ity shall correspond to the official identification of t	he animal.			
collected. Part II: (1) Delete as appropriate. (2) Only approved semen collection centres listed in accordance with Article 11(4) of Council Directive 92/65/EEC on the Commission website: http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm (3) 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. Official veterinarian or official inspector (*) Name (in capital letters): Local veterinary unit: Local veterinary unit: Date: Stamp:		date of colle	ection shall be indicated in the following format: do	d/mm/yyyy.			
 (1) Delete as appropriate. (2) Only approved semen collection centres listed in accordance with Article 11(4) of Council Directive 92/65/EEC on the Commission website: http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm (3) OJ L 192, 23.7.2010, p. 1. (4) Cross out the programme(s) that do(es) not apply to the consignment. (5) Insert date. Official veterinarian or official inspector (*) Name (in capital letters): Qualification and title: Local veterinary unit: LVU No: Date: Signature: 			mber of the centre shall correspond to the approva	I number of the semen centre indicate	d in Box I.12 where the semen was		
(*) 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 (*) OJ L 192, 23.7.2010, p. 1. (*) Cross out the programme(s) that do(es) not apply to the consignment. (*) Insert date. Official veterinarian or official inspector (*) Name (in capital letters): Local veterinary unit: Date: Stamp:	Part II:						
http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm (*) OJ L 192, 23.7.2010, p. 1. (*) Cross out the programme(s) that do(es) not apply to the consignment. (*) Insert date. Official veterinarian or official inspector (*) Name (in capital letters): Local veterinary unit: Date: Stamp:	(¹) Dele	te as appropri	ate.				
(*) Cross out the programme(s) that do(es) not apply to the consignment. (5) Insert date. Official veterinarian or official inspector (*) Name (in capital letters): Qualification and title: Local veterinary unit: LVU No: Date: Signature: Stamp:					5/EEC on the Commission website:		
(5) Insert date. Official veterinarian or official inspector (*) Name (in capital letters): Qualification and title: Local veterinary unit: LVU No: Date: Signature: Stamp:							
Official veterinarian or official inspector (*) Name (in capital letters): Qualification and title: Local veterinary unit: LVU No: Date: Signature: Stamp:			rramme(s) that do(es) not apply to the consignmer	nt.			
Name (in capital letters): Qualification and title: Local veterinary unit: LVU No: Date: Signature:			official inspector (*)				
Local veterinary unit: LVU No: Date: Signature: Stamp:				Qualification	and title:		
Date: Signature: Signature:							
Stamp:							
				Ŭ			
(*) The colour of the stamp and signature must be different from that of the other particulars in the certificate.		I					
(*) The colour of the stamp and signature must be different from that of the other particulars in the certificate.							
	(*) The c	olour of the stan	np and signature must be different from that of the other p	articulars 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

RO	PEA	N UNION										Intra trac	le certificate
٦	1.1.	Consignor					1.2.	Certificate	reference	No	1.:	2.a. Local referen	nce No
		Name											
		Address					I.3. Central competent authority						
		Postal code					1.4.	Local com	petent aut	hority			
	1.5.	Consignee							elated origi	nal	No(s)	No(s) of accompanying	
.		Name					0	certificates			docur	ments	
		Address					1.7.						
		Postal code					1.7.						
┢	Iß	Country of origin	ISO		ion of origin	Code	110	Country	of	ISO	1.1.1	Region of	Code
	1.0.	obunity of origin	code	no. nog	ion of origin	obac	1.10.	destinatio	on	code		destination	0000
F	1.12.	Place of origin					1.13.	Place of	destination				I
			Semen cen	tre 🗖				Se	emen centr	• □		Holding	
		Name			al number							_	
		Address		Abbiova	al number			Name				Approval numbe	er
								Address					
		Postal code						Postal co	de				
	1.14.						I.15.						
┢	1.16.	6. Means of transport					1.17.						
		Aeroplane 🗌 Ship 🗌 Railway wagon 🗌											
		Road vehicle D Other											
		Identification											
	1.18.	Description of com	modity						I.19. Com	nmodity co	de (HS	3 code)	
										0	5 11 9	9 85	
										1.20.	Quant	tity	
ŀ	1.21.	Temperature of pro	duct										
		Ambient 🗖		Cł	nilled 🔲		Frozen I.22. Number of packages						
F	1.23.	Seal/container No								1.24.	Type	of packaging	
┝	1.25	Commodities certifi	ad for:								.,,		
	1.20.	Artificial reproduction											
$\left \right $	1.26.	Transit through a th	nird country				1.27.	Transit th	rough Men	nber State	s	[
		Third country	ISO co		—			Member	-			- O code	
		Exit point	Code					Member				O code	
		Entry point	BIP N	c				Member				O code	
ŀ	1.28	Export					1.29.						
		Third country	ISO co	-									
		Exit point	Code										
╞	1.30.	-											
	1.31.	Identification of the	commoditie	es									
		Species	I	Breed	Donor ider	ntity	Date	of collect	tion		oval nu		Quantity
		(scientific name)								of	the tea	am	

ļ	EUROPEA	N UNION			Equine semen — Part C
	п. ⊢	lealth inforr	nation	II.a. Certificate reference No	ll.b.
	I, the und	lersigned of	fficial veterinarian, hereby certify that the semen des	scribed above	
	(¹) either	(II.1.	was collected, processed and stored for a minimum collection centre (²) situated in the Member State c Chapter I(I)(1) and Chapter I(II)(1) of Annex D to Di storage centre detailed in Box I.12 situated in the veterinary certification conditions at least as strict a	of origin of the semen and operated a rective 92/65/EEC, and from where the same Member State of origin of the	and supervised in accordance with e semen was moved to the semen
		(¹) or	[Part B of Annex I to Decision 2010/470/EU;]		
		(¹) or	[Decision 95/307/EC;]		
	(¹) or	[1].1.	was collected, processed and stored for a minimum collection centre $\binom{2}{2}$ situated in the European Unior Chapter I(II)(1) of Annex D to Directive 92/65/EEC accordance with:	n and operated and supervised in ac	cordance with Chapter I(I)(1) and
		(¹) either	[Part A of Annex I to Decision 2010/470/EU;]		
_		(¹) or	[Part B of Annex I to Decision 2010/470/EU;]		
		(¹) or	[Part C of Annex I to Decision 2010/470/EU;]		
		(¹) or	[Decision 95/307/EC;]		
	(¹) or	[II.1.	was collected, processed and stored in an approve listed in columns 2 and 4 of Annex I to Commission with Chapter I(I)(1) and Chapter I(II)(1) of Annex D t the conditions of Article 4 of Decision 2004/211/EC	Decision 2004/211/EC which is operate to Directive 92/65/EEC, and was import	ated and supervised in accordance
		(¹) either	[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 (with Chapter I(I)(2) and Chapter I(II)(2) of Annex D		ated and supervised in accordance
		II.3.	was sent to the place of loading in a sealed contail 92/65/EEC and bearing the number indicated in Bo		napter III(I) of Annex D to Directive
	Notes				
	Part I:				
		panied the	spond to the serial number of the individual official semen described above from the approved semen c l(s) of this/these document(s) or certificate(s) or the c	ollection centre of its origin to the desc	ribed above semen storage centre.
	Box 1.12:	place of o	rigin shall correspond to the semen storage centre of	of dispatch of the semen.	
l					

UROPEAN 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 of	or storage centre or to the holding of	destination of the semen.				
Box I.23: identification of container and seal number shall be indicated.							
Box I.31: donor identity shall o	correspond to the official identification of th	ne animal.					
date of collection sh	all be indicated in the following format: dd	l/mm/yyyy.					
approval number of	the centre shall correspond to the approva	al number of the semen collection cer	ntre of origin of the semen.				
Part II:							
(¹) Delete as appropriate.							
(²) Only approved semen colle the Commission websites:	ction or storage centres listed in accordance	ce with Article 11(4) or Article 17(3)(b)	of Council Directive 92/65/EEC on				
	mal/approved_establishments/establishmen mal/semen_ova/equine/index_en.htm	its_vet_field_en.htm					
Official veterinarian or official i	nspector (*)						
Name (in capital lette	rs):	Qualification and title:					
Local veterinary unit:		LVU No:					
Date:		Signature:					
Stamp:							
(*) The colour of the stamp and sig	nature must be different from that of the other pa	articulars in the certificate.					

ANNEX II

Model health certificates for trade within the Union in consigments 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

OPEA	IN UNION	Intra trade certifi			
l.1.	Name	1.2. Certificate reference No 1.2.a. Local reference No			
	Address	I.3. Central competent authority			
	Postal code	1.4. Local competent authority			
1.5.	Consignee	1.6.			
	Name				
	Address	1.7.			
	Postal code				
1.8.	Country of origin ISO code I.9. Region of origin Code	I.10. Country of ISO I.11. Region of Code destination code destination			
1.12	. Place of origin	I.13. Place of destination			
	Embryo team	Holding Embryo team			
	Name Approval number				
	Name Approval number Address	Name Approval number			
	Postal code	Address Postal code			
I. 1 4		I.15.			
110	Manua of house of	1.17.			
1.10	. Means of transport Aeroplane 🗌 Ship 🗌 Railway wagon 🗌	1.17.			
	Road vehicle Other				
	Identification				
1.18	. Description of commodity	I.19. Commodity code (HS code)			
		05 11 99 85			
		I.20. Quantity			
121	. Temperature of products	I.22. Number of packages			
	Ambient	Frozen			
1.23	. Seal/Container No	I.24. Type of packaging			
1.25	Commodities certified for: Artificial reproduction				
1.26	Transit through third country	I.27. Transit through Member States			
	Third country ISO code	Member State ISO code			
	Exit point Code	Member State ISO code			
	Entry point BIP No	Member State ISO code			
1.28	Export	1.29.			
	Third country ISO code				
	Exit point Code				
1.30					
1					
1.31	I.31. Identification of the commodities				
1.31	Identification of the commodities Species Breed Category Donor identi	ty Date of collection Approval number Quantit			

	EUROPEAN UNION			Equine ova and embryos — Part A					
	II. He	alth info	rmation	II.a. Certificate reference No	II.b.				
	I, the unde	rsigned (official veterinarian, hereby certify that:						
	(¹) either	(¹) either [II.1. the in vivo derived embryos/in vivo derived ova (¹) described above were collected, processed and stored by an embryor collection team (²) approved and supervised in accordance with Chapter I(III)(1) of Annex D to Directive 92/65/EEC;]							
E	(†) or	[.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;]					
Part II: Certification	(¹) either	[11.2.	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;]						
	(¹) or	[11.2.	the <i>in vivo</i> derived ova described above meet the requirements of Chapter III(II)(2) of Annex D to Directive 92/65/EEC;]						
Ра	(¹) or	[11.2.	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	[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 come from do	onor mares which:					
		II.3.1.	coming from holdings fulfilling the conditions laid di satisfying the conditions laid down in Articles 4 an						
		II.3.2.	meet the additional requirements of Chapter IV(4) of	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.3						
		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	e 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 for cultivation of 7 to 14 days carried out with negative r date of the first collection of ova or embryos fro consecutives cestrus 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) either	[11.4.	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	[11.4.	the embryos described above were conceived as a of Chapter III(II) of Annex D to Directive 92/65/EEC v conditions which comply with the requirements of C	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 the	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 at						
	Notes								
	Part I:								
	Box 1.12: F	Place of	origin shall correspond to the embryo collection team	a or embryo production team of ova/e	mbryos collection/production.				
		Place of lestinatio	destination shall correspond to the embryo collect on.	tion team, embryo production team	or to the holding of ova/embryos				
	Box I.23: Identification of container and seal number shall be indicated.								

EUBOPEAN UNION

C	Category: specify if: <i>in vivo</i> derived embryos, <i>in vivo</i> derived of Donor identity shall correspond to the official identification of t		nicromanipulated embryos.
	Donor identity shall correspond to the official identification of t	he animal.	
0			
	Date of collection shall be indicated in the following format: do	l/mm/yyyy.	
	Approval number of the team shall correspond to the e collection/production.	mbryo collection team or embryo	production team of ova/embryos
art II:			
) Delete	as appropriate.		
2) Only ap website	pproved embryo collection or production teams listed in accorda e:	ance with Article 11(4) of Council Dir	ective 92/65/EEC on the Commission
http://ec	c.europa.eu/food/animal/approved_establishments/establishmer	nts_vet_field_en.htm	
³) Insert d	late.		
⁴) OJ L 1!	92, 23.7.2010, p. 1.		
fficial vet	erinarian or official inspector (*)		
Name ((in capital letters):		Qualification and title:
Local v	reterinary unit:		LVU No:
Date:			Signature:
Stamp:			

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

OPEA	N UNION	Intra trade certific			
l.1.	Consignor Name	I.2. Certificate reference No I.2.a. Local reference No			
	Address	I.3. Central competent authority			
	Postal code	I.4. Local competent authority			
1.5.	Consignee	1.6.			
	Name Address				
		1.7.			
	Postal code				
I.5. I.8. I.12.	Country of origin ISO code I.9. Region of origin Code	e I.10. Country of ISO I.11. Region of Code destination code destination			
1.12.	Place of origin	I.13. Place of destination			
	Embryo team 🗖	Holding Embryo team			
	Name Approval number	Name Approval number			
	Address	Address			
	Postal code	Postal code			
I.14.		L.15.			
I.16.	Means of transport	l.17.			
	Aeroplane 🗌 Ship 🗌 Railway wagon 🗌				
	Road vehicle 🗌 Other 🗌				
	Identification				
I.18.	Description of commodity	I.19. Commodity code (HS code)			
		05 11 99 85 1.20. 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			
	Third country ISO code	Member State ISO code			
	Exit point Code	Member State ISO code			
	Entry point BIP No	Member State ISO code			
1.28.	Export	1.29.			
	Third country ISO code				
	Exit point Code				
1.30.					
1.31	Identification of the commodities				
	Species Breed Category Donor ide (Scientific name)	entity Date of collection Approval number Quanti of the team			

▼ <u>B</u>	
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l	EUROPEA	N UNION	Equine ova and embryos — Part B				
	ΙΙ. Η	ealth information	II.a. Certificate reference No	II.b.			
	I, the unde	ersigned official veterinarian, hereby certify that:					
	II.1. Ova/embryos (1) described above were collected by a collection team (2) approved by the competent authority and processed in appropriate laboratory; II.2. Ova/embryos (1) were collected from donor mares which:						
fication	II.2 .1 <i>.</i>	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 which on the day of collection fulfilled the conditions of Article 4 of Directive 2009/156/EC;					
Pa	II.2.3.	have been kept prior to the collection in holdings free from	clinical signs of contagious equine me	etritis for 60 days;			
	II.2.4 <i>.</i>	have not been used for natural breeding during the period of	of 30 days prior to the collection of ov	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 an	infectious or contagious disease;				
II.3. Ova/embryos (¹) were collected, processed, stored and transported under conditions which comply with the requirem of Directive 92/65/EEC;							
	II.4 .	The semen used for the artificial insemination of 92/65/EEC $(^{4})$ $(^{1});$	the donor mares complies with	the requirements of Directive			
	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 collect 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 t	the animal.				
		Date of collection shall be indicated in the following format: de	d/mm/yyyy.				
		Approval number of the team shall correspond to the embryo	collection team of ova/embryos collection	ation.			
	Part II:						
	(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/establishmer	nts_vet_field_en.htm				
	(³) OJ L 1	92, 23.7.2010, p. 1.					
	(⁴) Does not apply to ova.						

EURO	PEAN UNION		Equine ova and embryos — Part B
II.	Health information	II.a. Certificate reference No	II.b.
Officia	al veterinarian or official inspector (*)		
N	lame (in capital letters):		Qualification and title:
L	ocal veterinary unit:		LVU No:
0	ate:		Signature:
ຣ	tamp:		
(*) The	colour of the stamp and signature must be different from that of the other pa	articulars in the certificate.	

ANNEX III

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

▼<u>M1</u>

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

EUF	ROPEAN UNION Intra trade certifi					
	l.1.	Consignor Name	I.2. Certificate reference No I.2.a. Local reference No			
		Address	I.3. Central competent authority			
nted		Postal code	I.4. Local competent authority			
rese	1.5.	Consignee	1.6.			
t		Name Address				
nme		Postal code	1.7.			
nsig	1.8.	Country of origin ISO code I.9. Region of origin Code	I.10. Country of ISO code I.11. Region of Code			
of consignment presented	1.0.		destination destination			
ails	1.12.	Place of origin	I.13. Place of destination			
Det		Semen centre 🔲	Semen centre 🔲 Holding 🗌			
Part I: Details		Name Approval number Address	Name Approval number Address			
_		Postal code	Postal code			
	l.14.		l.15.			
	I.16.	Means of transport	1.17.			
		Aeroplane Ship Railway wagon				
		Road vehicle Other				
		Identification				
	l.18.	Description of commodity	I.19. Commodity code (CN code)			
			05 11 99 85			
	1.04					
	1.21.	Temperature of products Ambient Chilled Froz	zen			
		Seal/Container No	I.24. Type of packaging			
	1.25.	Commodities certified for: Artificial reproduction				
	1.26.	Transit through third country	I.27. Transit through Member States			
		Third country ISO code	Member State ISO code			
		Exit point Code	Member State ISO code			
		Entry point BIP No	Member State ISO code			
	1.28.	Export 🛛	1.29.			
		Third country ISO code Exit point Code				
		Exit point Code				
	1.30.					
	1.31.	Identification of the commodities				
		Species Breed Donor identity [(Sclentific name)	Date of collection Approval number of the Quantity centre			

▼<u>M1</u>

ا	EUROPEAN UNION			Ovine and caprine semen — Part A				
	II. Healt	h informatio	n	II.a. Certificate reference No	II.b.			
	I, the under	signed offici	al veterinarian, hereby certify that:					
		II.1.	The semen described above:					
		II.1.1. was collected, processed and stored in a semen collection centre (²) approved and supervised by the competent authority in accordance with Chapter I(I)(1) and Chapter I(II)(1) of Annex D to Directive 92/65/EEC;						
lion		II.1.2.	comes from donor animals which meet the requirements of Chapter II(II) of Annex D to Directive 92/65/EEC;					
Certificat		II.1.3.	was collected, processed, stored and transported under conditions which comply with the requirements of Chapters II(II) and III(I) of Annex D to Directive 92/65/EEC;					
Part II: Certification	(¹) either	[.1.4.	was collected from animals which have been kept continuously since birth on a holding or holdings recognised as having a negligible or controlled risk of classical scraple according to point 1 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001.]					
(¹) or [II.1.4. was collected from animals which have been kept continuously for the last three years before the collection on a holdings which has/have complied for the last three years before the collection with the requirements laid down 1.3(a) to (f) of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001.]								
(¹) or [II.1.4. was collected from animals which have been kept continuously since birth in a Member State or zone of a Me with a negligible risk status for classical scrapie approved in accordance with point 2.2 of Section A of Chapter . VIII to Regulation (EC) No 999/2001.]								
	(¹) or	[11.1.4.	was collected from ovine animals of the ARR/A	RR prion protein genotype;]				
	II.1.5. was sent to the place of loading in a sealed container in accordance with point 1.4 of Chapter III(I) of Annex D to I 92/65/EEC and bearing the number detailed in Box I.23.							
	(1) either	[.2.	No antibiotics or no mixture of antibiotics were	added to the semen.]				
	(¹) or [II.2. The following antibiotic or combination of antibiotics was added to produce a concentration in the final diluted a not less than (³):							
	Notes							
	Part I:							
	Box 1.12.: /	Place of orig	yin shall correspond to the semen collection cent	re of origin of the semen.				
	Box I.13.: /	Place of des	tination shall correspond to the semen collection	or storage centre or to the holding	of semen destination.			
	Box 1.23.: 1	dentification	of container and seal number shall be indicated	l.				
	Box 1.31.: 1	Donor identi	ty shall correspond to the official identification of	the animal.				
			ection shall be indicated in the following format: o					
		Approval nur collected.	mber of the centre shall correspond to the approve	al number of the semen centre indica	ted in Box I.12 where the semen was			
	Part II:							
	(1) Delete a	is appropria	te.					
			en collection centres listed in accordance with A pod/animal/approved_establishments/establishmer		n the Commission website:			
	(³) Insert na	ames and co	oncentrations.					
	— The colour of the stamp and signature must be different from that of the other particulars in the certificate.							

▼<u>M1</u>

EUROPEAN UNION	Ovine and caprine semen — Part A
II. Health information	II.a. Certificate reference No II.b.
Official veterinarian or official inspector	
Name (in capital letters):	Qualification and title:
Local veterinary unit:	LVU No:
Date:	Signature:
Stamp:	

PART B

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

EURO	OPEA	N UNION		Intra trade certificate		
	l.1.		I.2. Certificate reference No	I.2.a. Local reference No		
		Name Address Postal code	I.3. Central competent authority			
nted			I.4. Local competent authority			
t presei	1.5.	Consignee Name	1.6.			
ignmen		Address Postal code	1.7.			
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	I.11. Region of Code destination		
etails	1.10					
۵ ۳	1.12.	Place of origin Semen centre	I.13. Place of destination Semen centre	Holding		
Part		_	_			
		Name Approval number Address	Name Address	Approval number		
		Postal code	Postal code			
	1.14.		1.15.			
	1.16.	Means of transport	1.17.			
		Aeroplane 🗌 Ship 🗌 Railway wagon 🗌				
		Road vehicle 🗌 Other 🗌				
		Identification				
	1.18.	Description of commodity	I.19. Commodity co 05	de (HS code) 5 11 99 85		
			1.20.	Quantity		
	1.21.	Temperature of products Amblent Chilled	Frozen	Number of packages		
	1.23.	Seal/Container No		Type of packaging		
	1.25.	Commodities certified for:				
		Artificial reproduction				
	1.26.	Transit through third country	I.27. Transit through Member States	s 🗌		
		Third country ISO code	Member State	ISO code		
		Exit point Code	Member State Member State	ISO code ISO code		
		Entry point BIP No	1.29.			
	1.28.	Export Third country ISO code	1.29.			
		Exit point Code				
	1.30.					
	1.31.	Identification of the commodities				
		Species Breed Donor identity (Scientific name)	Date of collection Approval of the c			

	EUROPE	N UNION		0	Ovine and caprine semen — Part B		
	II. H	lealth info	rmation	II.a. Certificate reference No	II.b.		
	I, the und	lersigned (official veterinarian, hereby certify that the semen de	scribed 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;					
ų		II.2. comes from the donor animals which meet the requirements of Chapter II(II) of Annex D to Directive 92/65/EEC;					
Part II: Certification	II.3. was collected, processed, stored and transported under conditions which comply with the requirements of Chapters II(II) and III of Annex D to Directive 92/65/EEC;						
rt II: C	(¹) either	[11.4.	meets the requirements of Chapter A(I) of Annex \boldsymbol{V}	/III to Regulation (EC) No 999/2001;]			
(1) or [II.4. meets the requirements of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and is destined for a M which benefits, for all or part of its territory, from the provisions laid down in point (b) or (c) of Chapter A(I) of Regulation (EC) No 999/2001 and the donor animals comply regarding scrapie with the guarantees provide programmes referred to in that point and with the guarantees (³) requested by the Member State of destination				r (c) of Chapter A(I) of Annex VIII to the guarantees provided for by the			
	Notes						
	Part I:						
	Box 1.12:	Place of	origin shall correspond to the semen collection centr	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 o	f semen destination.		
	Box 1.23:	Identificat	ion of container and seal number shall be indicated.				
	Box 1.31:	Donor ide	entity shall correspond to the official identification of t	the animal.			
		Date of c	collection shall be indicated in the following format: d	d/mm/yyyy.			
		Approval collected.	number of the centre shall correspond to the approva	al number of the semen centre indicate	ed in Box I.12 where the semen was		
	Part II:						
	(1) Delete	e as appro	priate.				
			semen collection centres listed in accordance with A eu/food/animal/approved_establishments/establishme		5/EEC on the Commission website:		
	(³) Additi	onal guara	ntees as laid down in Article 2 of Regulation (EC) N	lo 546/2006 (OJ L 94, 1.4.2006, p. 2	8).		
	Official v	eterinarian	or official inspector (*)				
	Name	(in capita	l letters):	G	Qualification and title:		
	Local	veterinary	unit:	L	VU No:		
	Date:			S	ignature:		
	Stamp):					
	(*) The c	olour of the	stamp and signature must be different from that of the other	particulars in the certificate.			

PART C

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

EURC	ROPEAN UNION Intra trade certificate							
	l.1.	Consignor	I.2. Certificate reference No I.2.a. Local reference No					
		Name Address	I.3. Central competent authority					
ted		Postal code	I.4. Local competent authority					
preser	1.5.	Consignee Name	I.6. No(s) of related original certificates No(s) of accompanying documents					
ent		Address	1.7.					
muß		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 I.11. Region of Code destination destination					
ails	112	Place of origin	I.13. Place of destination					
l: Det		Semen centre	Semen centre Holding					
Part		Name Approval number Address	Name Approval number Address					
		Postal code	Postal code					
	1.14.		1.15.					
	l.16.	Means of transport	1.17.					
		Aeroplane 🗌 Ship 🗌 Railway wagon 🗌						
		Road vehicle 🔲 Other 🔲						
		Identification						
	l.18.	Description of commodity	1.19. 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	1.24. Type of packaging					
	1.25.	Commodities certified for:						
		Artificial reproduction 🔲						
	1.26.	Transit through third country	1.27. Transit through Member States					
		Third country ISO code	Member State ISO code					
		Exit point Code	Member State ISO code Member State ISO code					
	1.00	Entry point BIP No	1.29.					
	1.20.	Export Third country ISO code	1.29.					
		Exit point Code						
	1.30.							
	1.31.	Identification of the commodities						
		Species Breed Donor identity (Scientific name)	Date of collection Approval number Quantity of the centre					

	EUROPEA	N UNION		Ovine and caprine semen — Part C			
	II. F	lealth informa	tion	II.a. Certificate reference No	II.b.		
	I, the und	lersigned offic	ial veterinarian, hereby certify that the semen des	cribed above:			
	(¹) either	[11.1.	was collected, processed and stored for a minir semen collection centre ${}^{(2)}$ situated in the Me accordance with Chapter ${l}(l)(1)$ and Chapter ${l}(l)$ moved to the semen storage centre detailed in P animal health and veterinary certification at least	mber State of origin of the semer (1) of Annex D to Directive 92/65/EE art I.12 situated in the same Member	and operated and supervised in C, and from where the semen was		
cation		(¹) either	[Part A of Annex III to Decision 2010/470/EU;]				
Certific		(1) or	[Part B of Annex III to Decision 2010/470/EU;]				
Part II: Certification		(¹) or	[Decision 95/388/EC;]]				
e	(¹) or	[.1.	was collected, processed and stored for a minir semen collection centre (²) situated in the Europ and Chapter I(II) of Annex D to Directive 92/65/E accordance with:	ean Union and operated and supervi	sed in accordance with Chapter I(I)		
		(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	[.1.	was collected, processed and stored for a minir semen collection centre (²) situated in a third cou operated and supervised in accordance with Chap imported into the European Union under the con	intry or part(s) thereof listed in Annex oter I(I)(1) and Chapter I(II)(1) of Anne:	I to Decision 2010/472/EU which is C D to Directive 92/65/EEC, and was		
		(¹) either	[Section A of Part 2 of Annex II to Decision 2010	D/472/EU;]			
		(¹) or	[Section B of Part 2 of Annex II to Decision 2010	0/472/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		rated and supervised in accordance		
		II.3.	was sent to the place of loading in a sealed conta 92/65/EEC and bearing the number indicated in		Chapter III(I) of Annex D to Directive		
	Notes						
	Part I:						
	Box I.6:	panied the se	bond to the serial number of the individual official amen described above from the approved semen c of this/these document(s) or certificate(s), or the o	ollection centre of its origin to the des	cribed above semen storage centre.		
	Box 1.12:	Place of orig	in shall correspond to the semen storage centre of	of dispatch of the semen.			
	Box I.13:	Place of des	tination shall correspond to the semen collection	or storage centre or to the holding of	destination of the semen.		
	Box 1.23:	Identification	of container and seal number shall be indicated.				
	Box I.31:	Date of colle	y shall correspond to the official identification of t ction shall be indicated in the following format: do nber of the centre shall correspond to the approv	l/mm/yyyy.	entre of the semen origin.		

EUROPEAN UNION	Equine ova and embryos — Part					
II. Health information	II.a. Certificate reference No	II.b.				
Part II:		14 ²				
(¹) Delete as appropriate.						
(²) Only approved semen collection or storage centres listed in accordar the Commission websites:	nce with Article 11(4) or Article 17(3)(b) o	of Council Directive 92/65/EEC on				
http://ec.europa.eu/food/animal/approved_establishments/establishme http://ec.europa.eu/food/animal/semen_ova/ovine/index_en.htm	nts_vet_field_en.htm					
Official veterinarian or official inspector (*)						
Name (in capital letters):	Qua	lification and title:				
Local veterinary unit:	LVU	LVU No:				
Date:	Sigr	nature:				
Stamp:						
(*) The colour of the stamp and signature must be different from that of the other particular that the other particular th	articulars in the certificate.					

ANNEX IV

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

▼<u>M1</u>

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

EUR	OPE/	AN 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			
Ited		Postal code	I.4. Local competent authority			
sen	I.5.	Consignee	1.6.			
pre		Name				
hent		Address	1.7.			
gnn		Postal code				
Part I: Details of consignment presented	I.8.	Country of origin ISO code I.9. Region of origin Code	I.10. Country of ISO code I.11. Region of Code destination			
s o	140	Direct of exists				
etail	1.12.	Place of origin	I.13. Place of destination			
<u>0</u>		Embryo team	Holding 🗌 Embryo team 🗌			
Part		Name Approval number Address	Name Approval number Address			
		Postal code	Postal code			
	I.14.		1.15.			
	I.16.	Means of transport	1.17.			
		Aeroplane Ship Railway wagon				
		Road vehicle Other				
		Identification				
	1.40		I.19. Commodity code (CN code)			
	1.10.	Description of commodity	05 11 99 85			
			I.20. Quantity			
	I.21.	Temperature of products	I.22. Number of packages			
		Ambient Chilled Froz				
	1.23.	Seal/Container No	I.24. Type of packaging			
	1.25.	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			
	1.28.	Export 🗌	1.29.			
		Third country ISO code				
		Exit point Code				
	I.30.					
	1.31.	Identification of the commodities				
		Species Breed Category Donor identity (Scientific name)	v Date of collection Approval number of Quantity the team			

▼<u>M1</u>

	EUROPEAN	JNION		Ovine and caprine ova/embryos — Part A				
	II. Hea	th informatio	on	II.a. Certificate reference No	II.b.			
	I, the unders	igned officia	l veterinarian, hereby certify that:					
	(¹) either	[11.1.	the <i>in vivo</i> derived embryos (¹)/ <i>in vivo</i> derived embryo <i>collection</i> team (²) approved and su 92/65/EEC;]					
tion	(¹) or	[.1.	the <i>in vitro</i> produced embryos (¹)/micromanipul by an embryo production team (²) approved an Directive 92/65/EEC;]					
Part II: Certification	(¹) either	[11.2.	the <i>in vivo</i> derived embryos described abov 92/65/EEC;]	e meet the requirements of Chapter	III(II)(1) of Annex D to Directive			
Part II:	(†) or	[11.2.	the in vivo derived ova described above meet	the requirements of Chapter III(II)(2) o	f Annex D to Directive 92/65/EEC;]			
	(¹) or	[.2.	the <i>in vitro</i> produced embryos described abc 92/65/EEC;]	ove meet the requirements of Chapte	r III(II)(3) of Annex D to Directive			
	(1) or	[.2.	the micromanipulated embryos described abc 92/65/EEC:]	we meet the requirements of Chapte	r III(II)(4) of Annex D to Directive			
		<i>(¹)</i> [II.3.	the consignment consists of embryos of the ov	vine or caprine species which:				
		(¹) either	[were collected from animals which have bee having a negligible or controlled risk of classica Regulation (EC) No 999/2001;]]					
		(¹) or	[were collected from animals which have been I or holdings which have complied for the last thi to (f) of Section A of Chapter A of Annex VIII	ree years before collection with the rec				
		(¹) or	[were collected from animals which have beer State with a negligible risk status for classical s Annex VIII to Regulation (EC) No 999/2001;]]					
		(¹) or	[were collected from ovine animals of the ARF	VARR prion protein genotype;]]				
		II .4.	the ova or embryos described above come trequirements of Chapter IV(3) of Annex D to D		aprine species (¹) which meet the			
	(¹) either	[11.5.	the embryos described above were conceived a was collected, processed, stored and transport II(I) and III(I) of Annex D to Directive 92/65/EE	ted under conditions which comply wit				
	(¹) or	[11.5.	the embryos described above were conceived Chapter III(II)(2) of Annex D to Directive 92/65/ under conditions which comply with the require	EEC with semen which was collected,	processed, stored and transported			
	(¹) or	[11.5.	the ova have not been in contact with semen	of the ovine and caprine species;]				
		II.6.	the ova or embryos described above were sent Chapter III(II) of Annex D to Directive 92/65/EE					
	Notes							
	Part I:							
	Box 1.12.: P	lace of origi	n shall correspond to the embryo collection tear	m or embryo production team of embr	yos collection/production.			
		<i>lace of des</i> estination.	<i>tination</i> shall correspond to the embryo collec	stion team, embryo production team	or to the holding of ova/embryos			
	Box 1.23.: 1d	entification of	of container and seal number shall be indicated	Ι.				

▼<u>M1</u>

EUROPI	EAN UNION	Ovine	Ovine and caprine ova/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 v	vivo derived ova, in vitro produced embryos or	micromanipulated embryos.					
	Donor identity shall correspond to the official iden	ntification of the animal.						
	Date of collection shall be indicated in the following format: dd/mm/yyyy.							
	Approval number of the team shall correspond to the embryo collection team or embryo production team of ova/embryos collection/ production.							
Part II:								
(¹) Dele	ete as appropriate.							
	y approved embryo collection or production teams liste ://ec.europa.eu/food/animal/approved_establishments/e		2/65/EEC on the Commission website:					
— The	colour of the stamp and signature must be different	from that of the other particulars in the certifica	ate.					
Official	veterinarian or official inspector							
Nan	ne (in capital letters):	Qualif	cation and title:					
Loc	al veterinary unit:	LVU N	lo:					
Date	e:	Signat	ure:					
Star	Stamp:							

PART B

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

EUR	OPE	AN UNION							h	ntra trade	certificate
	1.1.	Consignor				1.2. (Certificate reference No		I.2.a. Local	reference I	No
		Name				-					
		Address				1.3. (Central competent author	ity			
Ited		Postal code				1.4. 1	_ocal competent authority	'			
sen	1.5.	Consignee									
bre		Name									
Jent		Address				1.7.	-				
ignn		Postal code									
Part I: Details of consignment presented		Country of origin	ISO code	I.9. Region of origin	Code	1.10.	Country of destination	ISO code	I.11. Reç des	gion of tination	Code
etaile	1.12	. Place of origin				1.13.	Place of destination				
ŏ		i i i i i i i i i i i i i i i i i i i	Embryo tear	m 🗖			Holdir	ng 🗖	Em	bryo team [
art		Name		Approval number			Name		App	proval numb	ber
Ċ.		Address					Address				
		Postal code					Postal code				
	1.14	l,				1.15.					
	1.16	. Means of transport				1.17.					
		Aeroplane 🗖	Ship 🔲	Railway wagon							
		Road vehicle 🗌	Other 🗌								
		Identification				_					
	1.18	. Description of commo	dity				I.19. Commodi	ty code	(HS code)		
									05 11 99 8	35	
									.20. Quantity		
	1.21	. Temperature of produc	ots	_			_	1	.22. Number o	of packages	i
		Ambient		Chilled 🗌		Fro	zen 🗌				
	1.23	. Seal/Container No						1	.24. Type of p	backaging	
	1.25	. Commodities certified	for:								
		Artificial reproduction									
	1.26	. Transit through third c	ountry			1.27.	Transit through Member	States			
		Third country		ISO code			Member State		I	ISO code	
		Exit point		Code			Member State			ISO code	
		Entry point		BIP No			Member State			ISO code	
	1.28	. Export [1.29.					
		Third country		ISO code							
		Exit point		Code							
	1.30										
	1.31	. Identification of the co	mmodities								
		Species	Breed	Category	Donor i	dentity	Date of		oval number	Q	uantity
		(Scientific name)					collection	of	the team		

EUR	OPEA	N UNION		Ovine and caprine ova/embryos — Part B					
Ш.		Health int	formation	II.a. Certificate reference No	II.b.				
I, th	ne und	ersigned (official veterinarian, hereby certify that the ova/embry	os (1) described above:					
	II.1. were collected, processed and stored under conditions which meet the requirements of Directive 92/65/EEC;								
	II.2. come from female donors of the ovine/caprine species (1) which meet the requirements of Chapter IV of Annex D to Directiv 92/65/EEC;								
	i) either [II.3. meet the requirements of Chapter III of Annex D to Directive 92/65/EEC and of Chapter A(I) of Annex VIII to Regulation (EC No 999/2001.]								
	(1) or [II.3. meet the requirements of Chapter III of Annex D to Directive 92/65/EEC and of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and are destined for a Member State which benefits, for all or part of its territory, from the provisions laid down in point (b) or (c) of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and the donor animals comply regarding scrapie with the guarantees provided for by the programmes referred to in that point and with the guarantees (²) requested by the Member State of destination.]								
(1)	either	[11.4.	in the case of embryos, the semen used for fertilisat of Chapter A(I) of Annex VIII to Regulation (EC) No		tive 92/65/EEC and the requirements				
(¹) or [II.4. in the case of embryos, the semen used for fertilisation meets the requirements of Directive 92/65/EEC and the recommendation of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and is destined for a Member State which benefits, for 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 its destined for a Member State which benefits, for and the donor animals comply regarding scraple with the guarantees provided for by the programmes referred to in and with the guarantees (²) requested by the Member State of destination.]									
Not	tes								
Par	t I:								
Вох	< 1.12:	Place of	origin shall correspond to the embryo collection tean	n of ova/embryos collection.					
Box	< l.13:	Place of destination	destination shall correspond to the embryo collec	tion team, embryo production tean	n or to the holding of ova/embryos				
Box	< I.23:	Identifica	tion of container and seal number shall be indicated.						
Box	< l.31:	Donor id Date of o	r: specify if: in vivo derived embryos, in vivo derived entity shall correspond to the official identification of collection shall be indicated in the following format: d number of the team shall correspond to the embryo	the animal. d/mm/yyyy.					
Par	t II:								
		as appro nal guara	priate. ntees as laid down in Article 2 of Regulation (EC) N	o 546/2006 (OJ L 94, 1.4.2006, p. 2	28).				
Offi	icial ve	terinarian	or official inspector (*)						
	Nam	ie (in capi	ital letters):	Q	ualification and title:				
	Local veterinary unit: LVU No:								
	Date Stan			S	ignature:				
	5.01	- Province of the second s							
(*) 1	The col	our 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

EUR	OPE	AN UNION						Intra trade	certificate
	1.1.	Consignor		I.2. Certificate	e reference No		I.2.a. Loo	cal reference l	No
		Name							
	Address				ompetent author	rity			
ented		Postal code		I.4. Local cor	npetent authorit	У			
rese	1.5.	Consignee		I.6.					
nt p		Name							
mer		Address		1.7.					
ign		Postal code							
Part I: Details of consignment presented	1.8.	Country of ISO code origin	I.9. Region of origin Code	I.10. Country destinati		ISO code		Region of lestination	Code
ls o		- -							
etai	l.12	. Place of origin		I.13. Place of					
0 ;;		Embryo team			Hold	ling 🔲		Embryo team	
art			Approval number	Name				Approval num	ber
۹.		Address		Address					
		Postal code		Postal c	ode				
	1.14	4		I.15.					
	116	. Means of transport		1.17.					
		Aeroplane 🗌 Ship 🗌	Railway wagon 🗖						
		Road vehicle Other							
		Identification							
	I.18	. Description of commodity		I.19. Commodity code (HS code)					
				05 11 99 85 I.20. Quantity					
						1.20. Qua	ntity		
	1.21	. Temperature of products		I.22. Number of packages					
		Ambient Chil	lled 🗌 Fro:	zen 🗖					
	1.23	. Seal/Container No				1.24. Туре	e of pack	aging	
	1.25	. Commodities certified for:							
		Artificial reproduction							
	1.26	. Transit through third country]	1.27. Transit th	rough Member	States			
		Third country	SO code	Member	State			ISO code	
		Exit point C	Code	Member	-			ISO code	
		Entry point E	3IP No	Member	State			ISO code	
	1.28	. Export		1.29.					
			SO code Code						
	1.30	•							
	1.31	Identification of the commodities Species Breed (Scientific name)	Category Donor iden	tity Date colle		Approval r of the t		Qı	antity

	11	t						
11.	Health int	formation	II.a. Certificate reference No	II.b.				
I, the ur	ndersigned (official veterinarian, hereby certify that the ova/embry	ros (1) described above:					
	II.1.	were produced/collected (1), processed and stored b accordance with Chapter I(III) of Annex D to Directi) team (²) approved and supervised				
	II.2.	meet the requirements of Chapter III(II) of Annex D	to Directive 92/65/EEC;					
	II.3.	come from donor females of the porcine species 92/65/EEC;	which meet the requirements of Cl	napter IV(2) of Annex D to Direct				
(¹) eithe	r [.4.	are in vivo derived embryos which:						
	II.4. 1 .	were conceived as a result of artificial insemination with semen meeting the requirements of Directive 90/429/EEC,						
	II.4.2.	originate from a Member State or region thereof:						
	(¹) either	[listed in Annex I to Decision 2008/185/EC and are d 2008/185/EC;]	destined for a Member State or regio	n thereof listed in Annex I to Decis				
	(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	on thereof not listed in Annex I or II				
	(¹) or	[listed in Annex II to Decision 2008/185/EC and are c 2008/185/EC and have been washed with trypsin;]	destined for a Member State or regio	n thereof listed in Annex I to Decis				
	(¹) or	[listed in Annex II to Decision 2008/185/EC and are c 2008/185/EC;]	destined for a Member State or regio	n thereof listed in Annex II to Decis				
	(¹) or	[not listed in Annex I or II to Decision 2008/185/EC a to Decision 2008/185/EC and have been washed w		or region thereof listed in Annex I o				
	(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 Anne				
(¹) or	[.4.	are in vitro produced/micromanipulated (1) embryos	which:					
	II.4.1.	were conceived as a result of in vitro fertilisation wi	ith semen meeting the requirements	of Directive 90/429/EEC,				
	II.4.2.	originate from a Member State or region thereof:						
	(¹) either	[listed in Annex I to Decision 2008/185/EC and are d 2008/185/EC;]	destined for a Member State or regio	n thereof listed in Annex I to Decis				
	(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	on thereof not listed in Annex I or I				
	(¹) or	[listed in Annex II to Decision 2008/185/EC and are c 2008/185/EC and the donor females of the ova use 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 region	n thereof listed in Annex II to Decis				
	(¹) or	[not listed in Annex I or II to Decision 2008/185/EC a to Decision 2008/185/EC and the donor females of t Decision 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 Anne				

II.	Health info	mation	II.a. Certificate reference No	II.b.						
(¹) or	[II.4. are in vivo derived ova which originate from a Member State or region thereof:									
	(¹) either	(¹) either [listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC;]								
	(¹) or	or [listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;]								
	(¹) or		/EC and are destined for a Member State or nor females complying with the conditions o							
	(¹) or	[listed in Annex II to Decision 2008/185, 2008/185/EC;]	/EC and are destined for a Member State or	region thereof listed in Annex II to Decisio						
	(¹) or		008/185/EC and are destined for a Member S come from donor females complying with							
	(¹) or	[not listed in Annex I or II to Decision 2 or II to Decision 2008/185/EC;]]	008/185/EC and are destined for a Member a	State or region thereof not listed in Annex						
	II.5.	were sent to the place of loading in a s Directive 92/65/EEC and bearing the n	sealed container under conditions complying umber detailed in Box I.23.	with point 6 of Chapter III(II) of Annex D t						
Notes										
Part I:										
3ox 1.12	: place of	origin shall correspond to the embryo c	ollection team or embryo production team of	ova/embryos collection/production.						
Box 1.13	B: place of (destination shall correspond to the embry	yo collection team, embryo production team o	r to the holding of ova/embryos destination						
Box 1.23	3: identifica	tion of container and seal number shall	be indicated.							
Box I.31	: category: specify if: in vivo derived embryos, in vivo derived ova, in vitro produced embryos or micromanipulated embryos.									
	donor ide	donor identity shall correspond to the official identification of the animal.								
	date of c	collection shall be indicated in the follow	ing format: dd/mm/yyyy.							
		number of the team shall correspond n indicated in Box I.12.	to the embryo collection team or embryo p	roduction team of ova/embryos collection						
Part II:										
• •	te as appro									
(²) Only web:		mbryo collection or production teams lis	sted in accordance with Article 11(4) of Counc	cil Directive 92/65/EEC on the Commissio						
http:	//ec.europa.	eu/food/animal/approved_establishments	/establishments_vet_field_en.htm							
Official	veterinarian	or official inspector (*)								
Na	ame (in capi	tal letters):	Qualification and title:							
Local veterinary unit: LVU No:										
Da	Date: Signature:									
St	amp:									

PART B

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

EUR	OPE/	AN UNION									Intra trade o	certificate
		Consignor				1.2.	Certificate	e reference No		I.2.a. Loca	al reference N	No
	Name Address					I.3. Central competent authority						
Part I: Details of consignment presented	-											
	Postal code					I.4. Local Competent Authority						
		Consignee				1.6.						
t		Name										
Isignmei		Address Postal code						1.7.				
s of consi		Country of origin	ISO code	I.9. Region of origin	Code	l.10.	Country destinati		ISO code	I.11. Re de	egion of stination	Code
Detail	1.12.	2. Place of origin Embryo team □			l.13.	Place of	destination	ng 🔲		nbryo team [۰ــــــــــــــــــــــــــــــــــــ	
Ë		Name	Linbiyo teal	Approval number			Name	TIOU	ng 🗀		proval numb	
Ъа		Address		Approvar number			Address			~F	provar numb	01
		Postal code					Postal c	ode				
	1.14					1.15.						
	1.16.	Means of transport				1.17.						
		Aeroplane 🔲	Ship 🗖	Railway wagon								
		Road vehicle 🗌	Other 🗌									
		Identification										
	1.18.	Description of commo	dity					I.19. Commod	ity code			
									1.20. Qu	05 11 99 antity	85	
										•		
	1.21.	Temperature of produce		illed 🗌	Fro	zen []		1.22. Nur	mber of pac	kages	
	1.23.	Seal/Container No							I.24. Typ	be of packa	ging	
	1.25.	Commodities certified	for:									
		Artificial reproduction										
	1.26.	Transit through third c	ountry			1.27.	Transit th	nrough Member	States			
		Third country		ISO code			Member	State			ISO code	
		Exit point		Code			Member	State			ISO code	
		Entry point		BIP No			Member	State			ISO code	
	1.28.	Export				1.29.						
		Third country		ISO code								
		Exit point		Code								
	1.30.											
	1.31.	Identification of the col	mmodities									
		Species (Scientific name)	Breed	Category D	onor ide	ntity		ate of lection		val number he team	Q	uantity

	EUROPEA		4	Porcine ova/embryos — Part E							
	11.	Health in	formation	II.a. Certificate reference No	II.b.						
	I, the und	undersigned official veterinarian, hereby certify that:									
		II. 1 .	The ova/embryos (1) described above:								
		II.1.1.	were collected, processed and stored under condition	ons which meet the requirements of Directive 92/65/EEC;							
		ll.1.2.	2. come from donor female swine which meet the requirements of Chapter IV of Annex D to Directive 92/65/EEC;								
Part II: Certification		II.1.3.	meet the requirements of Chapter III of Annex D to Directive 92/65/EEC.								
	(¹) either	[.2.	In the case of embryos,								
		II.2.1.	the semen used for fertilisation meets the requirements of Directive 90/429/EEC;								
		II.2.2.	.2.2. the embryos have been washed with trypsin (²).]								
	(¹) or	or [II.2. In the case of ova, the ova comes from a donor female swine which meets the conditions of Article 1 of I 2008/185/EC (²).]									
	Notes										
	Part I:										
	Box I.12: place of origin shall correspond to the embryo collection team of ova/embryos collection.										
	Box I.13:	 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 I.23: identification of container and seal number shall be indicated. 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. 									
	Box 1.23:										
	Box 1.31:										
		approval	ction indicated in Box I.12.								
	(1) Delete	as appro	opriate.								
	2008/ Memb	This condition applies only to ova and embryos which originate in the Member States or regions thereof not listed in Annexes I and II to Decision 2008/185/EC (OJ L 59, 4.3.2008, p. 19) and destined to the Member States or regions thereof so listed. It shall also apply to movements from Member States or regions thereof listed in Annex I of Decision 2008/185/EC to Member States or regions thereof listed in Annex I of Decision 2008/185/EC.									
Official veterinarian or official inspector (*)											
	Nan	ne (in cap	ital letters):	Qualification and title:							
	Loc	al veterina	ary unit:	LVU No:							
	Date	э:		Signature:							
	Star										
	(*) The co	our of the s	stamp and signature must be different from that of the other \mathbf{p}	articulars in the certificate.							