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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:

Official Journal

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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)

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

OPEA	IN UNION								Intra t	rade certificat	
l.1.	Consignor				1.2.	Certificate	reference N	10	I.2.a. Local refer	rence No	
	Name					1.2 Central competent authority					
	Address					I.3. Central competent authority					
	Postal code	1.4.	I.4. Local competent authority								
1.5.	Consignee	I.6.									
	Name										
	Address					1.7.					
	Postal code										
I.5. I.8.	Country of origin ISO code	I.9. Region	of origin	Code	l.10.	Country o destinatio		ISO code	I.11. Region of destination	Code	
1.12	. Place of origin				113	Place of r	destination				
	Semen ce	untro 🗖			1.10.				Halatian 🗖		
	Semen de					Se	men centre		Holding 🔲		
	Name	Approval n	umber			Name			Approval num	nber	
	Address					Address					
	Postal code					Postal co	de				
1.14					l.15.						
1.16	. Means of transport				1.17.						
	Aeroplane 🗌 Ship	_	Railway wag	~~ ¬							
		—	naliway way								
	Road vehicle Othe	r 🛄									
	Identification						-				
1.18	. Description of commodity						I.19. Comr	nodity co	de (HS code)		
						l		0	5 11 99 85		
								1.20.	Quantity		
1.21	. Temperature of product							1.22	Number of package	c	
	Ambient	Chille	ed 🔲			Frozen]	1.22.	Number of package	5	
1.23	. Seal/container No							1.24.	Type of packaging		
1.25	. Commodities certified for: Artificial reproduction 🗌							- 1			
1.26	. Transit through a third countr	v D			1.27.	Transit thr	ough Memb	per State	s		
1.20	-		4			Member	-		ISO code		
	Third country	ISO code Code				Member			ISO code		
	Exit point					Member			ISO code		
	Entry point	BIP No			_	Member	Olalo		100 0006		
1.28	. Export				1.29.						
	Third country	ISO code									
	Exit point	Code									
1.30											
1.31	. Identification of the commodi	ties									
	Species	Breed	Donor ider	ntity I	Date of	collection	Ap	proval nu	imber	Quantity	
	(scientific name)							of the tea		-	

	AN UNION			Equine semen — Part
11.	Health information	I	II.a. Certificate reference No	II.b.
I, the un	ndersigned official v	veterinarian, hereby certify that:		
ll.1.			scribed above was collected, processed a th Chapter I(I)(1) and Chapter I(I)(1) of An	
II.1.1.			of first collection of the semen described ab od for frozen semen elapsed, the semen c	
II.1.1.1.			on in a part of the territory (¹) of a Member § Article 5(2)(a) and (b) of Directive 2009/15	
II.1.1.2.	fulfilled the condit	tions for a holding laid down in Article	4(5) of Directive 2009/156/EC;	
II.1.1.3.	contained only ec	quidae which were free of clinical signs	of equine viral arteritis and contagious equ	uine metritis.
11.2.	Only equidae sati the centre.	isfying the conditions laid down in Article	es 4 and 5 or Articles 12 to 16 of Directive 2	009/156/EC have been admitted int
II.3.	The semen descr	ribed above was collected from donor a	stallions, which:	
II.3.1.	have not shown a was collected;	any clinical sign of an infectious or conta	gious disease at the time of admission into	the centre and on the day the seme
II.3.2.		or 30 days prior to the date of semen o gious equine metritis during that period;	ollection in holdings where no equine has s	hown any clinical sign of equine vira
II.3.3.		sed for natural mating during at least 30 to in points II.3.5.1, II.3.5.2 or II.3.5.3 ur	0 days prior to the date of first semen colle ntil the end of the collection period;	ection and from the dates of the firs
II.3.4.	Vaccines for Terr		the requirements of the relevant chapter of on samples taken in accordance with one or rity:	
	(¹) either [II.3.4	4.1. an agar-gel immuno-diffusion test	(Coggins test) for equine infectious anaem	ia (EIA) with negative result;]
	(1) or [11.3.4	4.1. an ELISA for equine infectious an	naemia (EIA) with negative result;]	
and	(¹) either [.3.4	4.2. a serum neutralisation test for eq	uine viral arteritis (EVA) with negative resul	It at a serum dilution of one in four
	(¹) or [.3.4	4.2. a virus isolation test for equine vira of the donor stallion;]	al arteritis (EVA) carried out with negative re	sult on an aliquot of the entire seme
and	[.3.4	with an interval of seven days by	ntagious equine metritis (CEM) carried out isolation of <i>Taylorella equigenitalis</i> after a ple and from genital swabs taken at least i in each case;	cultivation of 7 to 14 days from pre
II.3.5.	have been subjec II.3.5.2 and II.3.5.		n each case to at least one of the test prog	rammes (⁴) detailed in points II.3.5.1
II.3.5.1.	during the period		n collection centre for at least 30 days prior ove and no equidae on the semen collection	
		ed in point II.3.4 have been carried ou e of the commencement of the residenc	t on samples taken (⁵) prior to the first ser se period of at least 30 days;	men collection and at least 14 day

▼	B
	-

UROPE	AN UN	ION					Equin	e semen — Part			
ΙΙ.	Health	information			II.a. Cer	tificate reference No	II.b.				
1.3.5.2.	period contini	the donor stallion was resident on the semen collection centre for at least 30 days prior to the date of the first collection and during the period of collection of the semen described above, but has left the centre under the responsibility of the centre veterinarian for a continuous period of less than 14 days, and/or other equidae on the collection centre came into direct contact with equidae of lower health status;									
	collect	ion period in the ye	ibed in point II.3.4 have been carried out on samples taken (⁵) prior to the first semen collection of the breeding season or d in the year the semen described above was collected and at least 14 days following the date of the commencement of period of at least 30 days;								
and	the test described in point II.3.4.1 for equine infectious anaemia was last c before the semen described above was collected;					carried out on a sample	of blood taken (⁵) not	more than 90 days			
and	(¹) eith		sts described in he semen desc			eritis was last carried ou	ut on a sample taken (⁵) not more than 30			
	(¹) or	donor stallion	taken (5) not me	ore than six m	onths before the s	ut with negative result semen described above on test for equine viral	was collected and a	blood sample taker			
and			ribed in point II. he semen desc			tritis was last carried ou	ut on samples taken ^{(;}	⁵⁾ not more than 6(
1.3.5.3.		sts described in poi ion period in the ye				n (⁵) prior to the first se	men collection of the	breeding season o			
and		sts described in poir ion of the semen de			on samples taker	n (⁵) not less than 14 da	ays and not more tha	n 90 days after the			
1.3.6.	Have	undergone the testir	ng provided for	in point II.3.5	on samples take	n on the following dates	s:				
_		Start dat	e (⁵)		C	ate of sampling for he	alth tests (⁵)				
men	Test programme	Donor Semen residence collection	Semen collection	EIA II.3.4.1		EVA 3.4.2	CEM II.3.4.3				
Identification of semen	progr				Blood sample	Semen sample	1. sample	2. sample			
				_							
				-							
				_							
^{,1}) eithe	er [1].4	No antibiotics we	ere added to the	e semen:1							
) 010.0	. [
1) or	[.4	than (⁶):				to produce a concentra					
.5.	The	semen described a									
1.5.1.		ected, processed, st Directive 92/65/EE		orted under c	onditions which co	mply with the requireme	ents of Chapters II(I)(1) and III(I) of Anne:			
.5.2.		t to the place of loar ring the number ind			accordance with p	ooint 1.4 of Chapter III(I)) of Annex D to Direc	tive 92/65/EEC and			

I. I	Health information	II.a. Certificate reference No	ll.b.
Votes			
Part I:			
Box 1.12:	place of origin shall correspond to the semen	n collection centre of origin of the semen.	
3ox 1.13:	place of destination shall correspond to the s	emen collection or storage centre or to the holding	g of semen destination.
Box 1.23:	identification of container and seal number sh	nall be indicated.	
Box 1.31:	donor identity shall correspond to the official	identification of the animal.	
	date of collection shall be indicated in the fol	lowing format: dd/mm/yyyy.	
	approval number of the centre shall correspon collected.	nd to the approval number of the semen centre indi	cated in Box I.12 where the semen was
Part II:			
Guidance	e for the completion of Table in II.3.6:		
Abbrevia	tions:		
EIA-1	Equine infectious anaemia (EIA) testing first	t occasion	
EIA-2	EIA testing second occasion		
EVA-B1	Equine viral arteritis (EVA) testing on blood	sample first occasion	
EVA-B2	2 EVA testing on blood sample second occas	sion	
EVA-S1	EVA testing on semen sample first occasion	n	
EVA-S2	2 EVA testing on semen sample second occa	asion	
CEM-11	1 Contagious equine metritis (CEM) testing fir	rst occasion first sample	
CEM-12	2 CEM testing first occasion second sample t	taken 7 days after CEM-11	
CEM-21	1 CEM testing second occasion first sample		
CEM-22	2 CEM testing second occasion second samp	ole taken 7 days after CEM-21	
Instructio	ns:		
	h semen identification in column A in the examp olumns C and D must be completed with the o	ble below, the test programme (II.3.5.1, II.3.5.2 and/ dates required.	or II.3.5.3) must be described in columr
and II.3.		sting prior to the first collection of the semen describ to 9 of the table, this being the boxes marked with	
The date	es when samples were taken for repeat laborate	ory testing as required in accordance with II.3.5.2 o	r II.3.5.3 are entered in the lower line σ

II.	Health i	nformation			1	ll.a. Cert	ificate reference No	II.b.			
-		Start date (5)			I	Date of sampling for health tests (5)					
Identification of semen	Test	Donor residence				EVA .3.4.2		CEM II.3.4.3			
Ident of s	brog				Bloo samp		Semen sample	1. samp	le	2. sample	
٨	в	С	D	EIA-1	EVA-E	B1	EVA-S1	CEM-1	1	CEM-12	
A	Б	C		EIA-2	EVA-E	B2	EVA-S2	CEM-2	1	CEM-22	
(⁴) Cro (⁵) Ins	oss out the ert date in	3.7.2010, p. 1. e programme(s) tha n table in point II.3.6 , and concentrations	3 (follow guidan		•						
(⁴) Cro (⁵) Ins (⁶) Ins	ert date ir ert date ir ert names	e programme(s) than a table in point II.3.0 and concentrations ian (*)	3 (follow guidan		•		Qualification and this				
(⁴) Cro (⁵) Ins (⁶) Ins	oss out the ert date ir ert names I veterinar Name	e programme(s) that a table in point II.3. and concentrations ian (*) (in capital letters):	3 (follow guidan		•		Qualification and title				
(⁴) Cro (⁵) Ins (⁶) Ins	oss out the ert date ir ert names I veterinar Name	e programme(s) than a table in point II.3.0 and concentrations ian (*)	3 (follow guidan		•						
(⁴) Cro (⁵) Ins (⁶) Ins	oss out the ert date ir ert names I veterinar Name Local v	e programme(s) than a table in point II.3. and concentrations ian (*) (in capital letters): veterinary unit:	3 (follow guidan		•		LVU No:				
(⁴) Cro (⁵) Ins (⁶) Ins	oss out the ert date ir ert names I veterinar Name Local v Date:	e programme(s) than a table in point II.3. and concentrations ian (*) (in capital letters): veterinary unit:	3 (follow guidan		•		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

EURO	OPEA	N UNION	Intra trade certificate					
	1.1.	Consignor	I.2. Certificate reference No I.2.a. Local reference No					
		Name	1.2. Control compotent authority					
		Address	I.3. Central competent authority					
ted		Postal code	I.4. Local competent authority					
presented	1.5.	Consignee	1.6.					
bre		Name						
nent		Address	1.7.					
ignn		Postal code						
consignment	1.8.	Country of origin ISO I.9. Region of origin Code	I.10. Country of ISO I.11. Region of Code					
oľo		code	destination code destination					
ils								
Part I: Details	1.12.	Place of origin	I.13. Place of destination					
		Semen centre	Semen centre 🔲 Holding 🗌					
Part		Name Approval number	Name Approval number					
		Address	Address					
		Postal code	Postal code					
	1.14.		I.15.					
	1.16.	Means of transport	1.17.					
		Aeroplane 🗌 Ship 🗌 Railway wagon 🗌						
		Road vehicle Other Other						
	1 1 0	Identification Description of commodity						
	1.10.	Description of commonly	I.19. Commodity code (HS code) 05 11 99 85					
			I.20. Quantity					
	1.21.	Temperature of product						
		Ambient Chilled	Frozen					
	1.23.	Seal/container No						
			I.24. Type of packaging					
	1.25.	Commodities certified for:	·					
		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						
		Species Breed Donor identity Da (scientific name)	te of collection Approval number Quantity of the team					
		(

	EUROPE	AN UNION				Equine semen — Part B					
	11.	Health informa	ation		II.a. Certificate reference No	II.b.					
	I, the un	dersigned offic	cial veterin	arian, hereby certify that:							
	II.1.	The semen c	ollection c	entre (²), in which the semen described	above was collected, processed and	stored for trade:					
	IL1.1.	is approved and supervised by the competent authority according to the conditions of Chapter I of 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 semi- collected until the date the semen was dispatched as fresh/chilled (¹) semen or until the 30 days mandatory storage period for semen elapsed (¹) not considered to be infected with African horse sickness in accordance with Article 5(2)(a) and (b) of Di 2009/156/EC (³);										
Part II: C	II.1.3.	fulfilled during the period commencing 30 days prior to the date of semen collection until the date the semen was dispatched as fresh/ chilled (1) semen or until the 30 days mandatory storage period for frozen semen elapsed (1), the conditions of Article 4 of Directive 2009/156/EC;									
	II.1.4.	chilled (1) sen	nen or unti	riod commencing 30 days prior to the da I the 30 days mandatory storage period t and contagious equine metritis;							
	11.2.	All equidae h	ave been	admitted into the centre under the prov	isions of Article 4 and 5 of Directive 2	009/156/EC.					
	II.3.	The semen d	lescribed a	above was collected from donor stallion	s, which:						
	II.3.1.	on the day th	ne semen i	was collected have not shown clinical s	signs of an infectious or contagious disease;						
	11.3.2.	during at leas	st 30 days	prior to collection of the semen have r	not been used for natural service;						
	II.3.3.	during the last 30 days prior to collection of the semen have been kept on holdings where no equidae showed clinical signs of equine v arteritis;									
	II.3.4.	during the las equine metrit		prior to collection of the semen have bee	een kept on holdings where no equidae showed clinical signs of contagious n, have not been in contact with equidae suffering from an infectious or ilection of the semen;						
	II.3.5.			vledge and as far as I could ascertain ing 15 days immediately preceding colle							
	II.3.6.			owing animal health tests, carried out in cified in point II.3.7;	a laboratory recognised by the compe	tent authority, in accordance with a					
			[11.3.6.1.	an agar gel immunodiffusion test (Cog	gins test) for equine infectious anaemi	a with negative result;]					
	and	(1) either	[11.3.6.2.	a serum neutralisation test for equine	viral arteritis with negative result at a	serum dílution of one in four; and]					
		(¹) or	[11.3.6.2.	a virus isolation test for equine viral an the donor stallion;]	teritis carried out with negative result o	n an aliquot of the entire semen of					
	and		II.3.6.3.	an agent identification test for contagion the donor stallion with an interval of sev a semen sample and from genital swa negative result in each case;	en days by isolation of Taylorella equig	enitalis from pre-ejaculatory fluid or					
	II.3.7.	have been su	ubject to th	ne one of the following test programmes	\$ (⁴):						
	II.3.7.1.		iod, and ne	continuously resident in the collection o o equidae in the collection centre came o							
		contagious e	quine metr	point II.3.6 have been carried out on sai itis on a second sample taken on above residence period and at least at		at least 14 days after the					

▼	B

UROPE	AN UNION			Equine semen — Part B		
Ш.	Health inforr	nation	II.a. Certificate reference No	ll.b.		
11.3.7.2.		stallion was not continuously resident in the collection of lower health status than the donor stallion;	on centre or other equidae in the coll	ection centre came into contact with		
	contagious	lescribed in point II.3.6 have been carried out on sa e equine metritis on a second sample taken on men collection and at least at the beginning of the				
and	the test described in point II.3.6.1 for equine infectious anaemia was last carried out on a sample of blood					
and	(¹) either	[one of the tests described in point II.3.6.2 fo 				
	(¹) or	[the non-shedder state of the seropositive stallion carried out on an aliquot of the entire semen of than one year before the semen described abov	the donor stallion collected on			
II.3.7.3.	14 days a	described in point II.3.6 have been carried out during fter the collection of the semen on samples taken c nd sample taken on	m (⁵) and in th			
II.4.		n described above was collected, processed, stored I and III of Annex D to Directive 92/65/EEC.	and transported under conditions where	nich comply with the requirements of		
Notes						
Part I:						
Box I.12	: place of o	rigin shall correspond to the semen collection centre	e of origin of the semen.			
Box I.13	place of d	estination shall correspond to the semen collection	or storage centre or to the holding o	f semen destination.		
Box 1.23	identificatio	on of container and seal number shall be indicated.				
Box I.31	: donor ider	tity shall correspond to the official identification of t	he animal.			
	date of co	llection shall be indicated in the following format: do	d/mm/yyyy.			
	approval n collected.	umber of the centre shall correspond to the approva	I number of the semen centre indicate	ed in Box I.12 where the semen was		
Part II:						
(¹) Delet	e as approp	riate.				
http://	/ec.europa.e	emen collection centres listed in accordance with A u/food/animal/approved_establishments/establishmer		5/EEC on the Commission website:		
	192, 23.7.2	010, p. 1. ogramme(s) that do(es) not apply to the consignmer				
(⁵) Inser			n.			
Official v	veterinarian d	or official inspector (*)				
Nar	me (in capita	l letters):	Qualificatio	n and title:		
Loc	al veterinary	unit:	LVU No:			
Dat	e:		Signature:			
Sta	mp:					
*) The co	blour of the sta	mp 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

EURO	PEA	IN UNION		Intra trade certificate				
	1.1.	Consignor	I.2. Certificate reference No I.2.	a. Local reference No				
		Name						
		Address	I.3. Central competent authority					
pa		Postal code	I.4. Local competent authority					
Part I: Details of consignment presented	1.5.	Consignee	I.6. No(s) of related original No(s) of	No(s) of accompanying				
pres		Name	certificates docum	ents				
t		Address	17					
me		Postal code	1.7.					
sigı								
con	1.8.	Country of origin ISO I.9. Region of origin Code code	I.10. Country of ISO I.11. F destination code c	Region of Code destination				
oţ								
ails	112	Place of origin	I.13. Place of destination					
Det		Semen centre						
÷				Holding 🔲				
Pai		Name Approval number		Approval number				
		Address	Address					
		Postal code	Postal code					
	1.14.		l.15.					
	116	Means of transport	I.17.					
	1.10.	·	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. Quantit	у				
	1.21.	Temperature of product						
		Ambient 🗌 Chilled 🗖	Frozen 🗌 1.22. Numbe	r of packages				
	1.23.	Seal/container No	І.24. Туре о	f packaging				
	1.25.	Commodities certified for:						
		Artificial reproduction						
	1.26.	Transit through a third country	I.27. Transit through Member States					
		Third country ISO code	Member State ISC) code				
		Exit point Code	Member State ISC) code				
		Entry point BIP No	Member State ISC) code				
	1.28.	Export	1.29.					
		Third country ISO code						
		Exit point Code						
	1.30.							
	1.31.	Identification of the commodities						
		Species Breed Donor identity	Date of collection Approval num	nber Quantity				
		(scientific name)	of the tear					

	PEAN UNION			Equine semen — Part				
11.	Health infor	mation	II.a. Certificate reference No	II.b.				
I, the	undersigned o	fficial veterinarian, hereby certify that the semen des	cribed above					
(¹) ei	<i>ther</i> [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 as	f origin of the semen and operated a ective 92/65/EEC, and from where th same Member State of origin of th	and supervised in accordance with e semen was moved to the semer				
(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	[Decision 95/307/EC;]						
(1) or [Part B of Annex I to Decision 2010/470/EU;] (1) or [Decision 95/307/EC;] (1) or [II.1. was collected, processed and stored for a minimum period of 30 days immediately following collection in an appro collection centre (²) situated in the European Union and operated and supervised in accordance with Chapter Chapter I(II)(1) of Annex D to Directive 92/65/EEC, and was moved to the semen storage centre detailed in accordance with:				ccordance 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 approved 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 oper Directive 92/65/EEC, and was impo	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 (2 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 contain 92/65/EEC and bearing the number indicated in Bo		hapter III(I) of Annex D to Directive				
Note	5							
Part	1:							
Box	panied the	spond to the serial number of the individual official of semen described above from the approved semen co al(s) of this/these document(s) or certificate(s) or the c	pliection centre of its origin to the desc	cribed above semen storage centre				
Box	.12: place of c	rigin shall correspond to the semen storage centre c	f dispatch of the semen.					

EUROPE	AN UNION		Equine semen — Part C			
П.	Health information	II.a. Certificate reference No	II.b.			
Box I.1	Box I.13: place of destination shall correspond to the semen collection or storage centre or to the holding of destination of the semen.					
Box I.20	Box I.23: identification of container and seal number shall be indicated.					
Box I.3	I: donor identity shall correspond to the official identification of t	he animal.				
	date of collection shall be indicated in the following format: do	i/mm/yyyy.				
	approval number of the centre shall correspond to the approv	al number of the semen collection c	entre of origin of the semen.			
Part II:						
(1) Dele	te as appropriate.					
(²) Only the (approved semen collection or storage centres listed in accordan Commission websites:	ce with Article 11(4) or Article 17(3)(b) of Council Directive 92/65/EEC on			
http:/	//ec.europa.eu/food/animal/approved_establishments/establishmer	nts_vet_field_en.htm				
http:/	//ec.europa.eu/food/animal/semen_ova/equine/index_en.htm					
Official	veterinarian or official inspector (*)					
	Name (in capital letters):	Qualification and title:				
	Local veterinary unit:	LVU No:				
	Date:	Signature:				
	Stamp:					
(*) The c	olcur of the stamp and signature must be different from that of the other p	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

RO	IOPEAN UNION Intra trade certificate					
	l.1.	Consignor	1.2. Certificate reference No 1.2.a. Local reference No			
		Name Address	I.3. Central competent authority			
p		Postal code	I.4. Local competent authority			
sell	1.5.	Consignee	1.6.			
b d		Name				
		Address	1.7.			
		Postal code				
5	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			
3		Embryo team	Holding Embryo team			
		· _				
-		Name Approval number Address	Name Approval number			
		Postal code	Address			
+			Postal code			
	1.14.		1.15.			
	I.16.	Means of transport	1.17.			
		Aeroplane 🗌 Ship 🗌 Railway wagon 🗌				
		Road vehicle Other				
		Identification				
	1.18.	Description of commodity	I.19. Commodity code (HS code)			
			05 11 99 85 1.20. Quantity			
ſ	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				
ſ	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			
ļ		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 iden (Scientific name)	ity Date of collection Approval number Quantity of the team			

	ROPEAN		· · · · · · · · · · · · · · · · · · ·		Equine ova and embryos — Part			
II.	He	alth info	rmation	II.a. Certificate reference No	II.b.			
I,	the unde	rsigned	official veterinarian, hereby certify that:					
Ċ) either	[1].1.	the <i>in vivo</i> derived embryos <i>/in vivo</i> derived ova (¹) collection team (²) approved and supervised i 92/65/EEC;]					
Ċ.) 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:]					
Ċ) 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;]					
Ċ,) or	[11.2.	the <i>in vivo</i> derived ova described above mee 92/65/EEC;]	t the requirements of Chapter	III(II)(2) of Annex D to Directiv			
Ċ,) or	[11.2.	the <i>in vitro</i> produced embryos described above 92/65/EEC;]	meet the requirements of Chapte	er III(II)(3) of Annex D to Directiv			
Ċ,) or	[11.2.	the micromanipulated embryos described above r 92/65/EEC;]	meet the requirements of Chapte	r III(II)(4) of Annex D to Directiv			
		II.3.	the ova or embryos described above come from dor	nor mares which:				
		II.3.1.	coming from holdings fulfilling the conditions laid do satisfying the conditions laid down in Articles 4 and					
II.3.2. meet the additional requirements of Chapter IV(4) of Annex D to Directive 92/65/EEC;								
 II.3.3. have not been used for natural breeding during at least 30 days prior to the date of collection of ova or embryos and be the date of the first sample referred to in points II.3.4 and II.3.5 and the date of the collection of ova and embryos; II.3.4. have been subjected with negative result to an agar-gel immuno-diffusion test (Coggins test) or an ELISA for equine infe anaemia carried out on a blood samples taken on								
				past 30 days prior to the date of th on a sample of blood take				
II.3.5. have been subjected to an agent identification test for contagious equine metritis by isolation of <i>Taylorella</i> erecultivation of 7 to 14 days carried out with negative results in each case on samples taken during the past 3 date of the first collection of ova or embryos from mucosal surfaces of the clitoral fossa and clitoral consecutives cestrus periods on					n during the past 30 days prior to th I fossa and clitoral sinuses on tw			
Ċ,) either	[11.4.	the embryos described above were conceived as a r collected, processed, stored and transported under III(I) of Annex D to Directive 92/65/EEC;]					
Ċ,) or	[11.4.	the embryos described above were conceived as a re of Chapter III(II) of Annex D to Directive 92/65/EEC w conditions which comply with the requirements of Cl	ith semen which was collected, pro-	cessed, stored and transported und			
Ċ) or	[.4.	the ova have not been in contact with semen of the	equine species;]				
		II.5.	the ova or embryos described above were sent to the Chapter III(II) of Annex D to Directive 92/65/EEC and					
N	otes							
Pa	art I:							
B	ox I.12: F	Place of	origin shall correspond to the embryo collection team	or embryo production team of ova/	embryos collection/production.			
B		Place of lestinatio	destination shall correspond to the embryo collection	on team, embryo production team	n or to the holding of ova/embry			
	C	oounatic	лъ.					

EUROPEAN UNION		Equine ova and embryos — Part A		
II. Health informa	lion	II.a. Certificate reference No	II.b.	
Box I.31: Category: sp	ecify if: in vivo derived embryos, in vivo derive	ed ova, in vitro produced embryos or i	micromanipulated embryos.	
Donor identit	Donor identity shall correspond to the official identification of the animal.			
Date of colle	ction shall be indicated in the following format	: dd/mm/yyyy.		
Approval nu collection/pro	mber of the team shall correspond to the duction.	e embryo collection team or embry	vo production team of ova/embryos	
Part II:				
(1) Delete as appropria	te.			
(²) Only approved emb website:	ryo collection or production teams listed in acco	ordance with Article 11(4) of Council D	irective 92/65/EEC on the Commission	
http://ec.europa.eu/f	ood/animal/approved_establishments/establishr	ments_vet_field_en.htm		
(3) Insert date.				
(⁴) OJ L 192, 23.7.201	0, p. 1 .			
Official veterinarian or	official inspector (*)			
Name (in capital let	ters):		Qualification and title:	
Local veterinary uni	ti		LVU No:	
Date:			Signature:	
Stamp:				
(*) The colour of the stamp	and signature must be different from that of the othe	er particulars in the certificate.		

PART B

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

ROPEAN UNION Intra trade certificat					
1	.1.	Consignor Name	I.2. Certificate reference No I.2.a. Local reference No		
		Address	1.3. Central competent authority		
		Postal code	I.4. Local competent authority		
	.5.	Consignee	l.6.		
		Name Address			
		Postal code	1.7.		
	.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		
5					
	.12.	Place of origin	I.13. Place of destination		
		Embryo team 🗖	Holding 🗌 Embryo team 🗌		
		Name Approval number Address	Name Approval number		
		Postal code	Address Postal code		
1	.14.		L15.		
	10	Manage of Incompany	1.17.		
'	.16.	Means of transport Aeroplane Ship Railway wagon	1.17.		
		Road vehicle Other Other			
		Identification			
I	.18.	Description of commodity	I.19. Commodity code (HS code)		
			05 11 99 85		
			1.20. Quanty		
I	.21.	Temperature of products Ambient Chilled	I.22. Number of packages		
1	.23.	Seal/Container No	I.24. Type of packaging		
1	.25.	Commodities certified for: Artificial reproduction			
1	.26.	Transit through third country	I.27. Transit through Member States		
		Third country ISO code	Member State ISO code		
		Exit point Code	Member State ISO code		
		Entry point BIP No	Member State ISO code		
I	.28.	Export	1.29.		
		Third country ISO code Exit point Code			
1	.30.				
+					
	. 3 1.	Identification of the commodities Species Breed Category Donor ident (Scientific name)	ity Date of collection Approval number Quantity of the team		

	EUROPEAN UNION		Equine ova and embryos — Pa			
	ΙΙ. Η	ealth information	II.a. Certificate reference No	II.b.		
	I, the und	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 an 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 situal of a Member State which is not considered to be infected Directive 2009/156/EC (³);				
Part II: Certification	II.2.2.	have been located in holdings under veterinary supervision w 2009/156/EC;	which on the day of collection fulfilled th	e conditions of Article 4 of Directive		
Par	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 30 days prior to the collection of o	/a/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	n infectious or contagious disease;			
	II.3.	Ova/embryos (¹) were collected, processed, stored and trans of Directive 92/65/EEC;	sported under conditions which comply	/ with the requirements of Annex D		
	II.4. The semen used for the artificial insemination of the donor mares complies with the requirements of Directiv 92/65/EEC (4) (1);					
	II.5.	The ova used for the in vivo production of embryos comply	with the requirements of Directive 92	/65/EEC (¹).		
	Notes					
	Part I:					
	Box 1.12:	Place of origin shall correspond to the embryo collection team	n of ova/embryos collection.			
	Box I.13:	Place of destination shall correspond to the embryo 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, in vitro produced embryos or mic	cromanipulated embryos.		
		Donor identity shall correspond to the official identification of	the animal.			
		Date of collection shall be indicated in the following format: d	ld/mm/yyyy.			
		Approval number of the team shall correspond to the embryo	collection team of ova/embryos collection	ction.		
	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/establishme	nts_vet_field_en.htm			
	(3) OJ L ·	192, 23.7.2010, p. 1.				
	(4) Does	not apply to ova.				

EUROPEAN UNION		Equine ova and embryos — Part B		
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:				
(*) The colour of the stamp and signature must be different from that of the other particulars 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

EUR	ROPEAN UNION Intra trade certificate						
	l.1.	Consignor Name Address	1.2. Certificate reference No 1.2.a. Local reference No 1.3. Central competent authority				
nted		Postal code	I.4. Local competent authority				
of consignment presented	Name Address		1.6.				
consigr	1.8.	Postal code Country of origin ISO code I.9. Region of origin Code	I.10. Country of ISO code I.11. Region of Code destination				
ails of	I.12.	Place of origin	I.13. Place of destination				
Part I: Details		Semen centre	Semen centre Holding				
Part		Name Approval number Address	Name Approval number Address				
		Postal code	Postal code				
	1.14.		1.15.				
	I.16.	Means of transport Aeroplane Ship Road vehicle Other Identification	1.17.				
	118	Description of commodity	I.19. Commodity code (CN code)				
	1.10.		05 11 99 85				
	1.01	Temperature of products	I.20. Quantity I.22. Number of packages				
	1.21.		zen				
	I.23.	Seal/Container No	I.24. Type of packaging				
	I.25.	Commodities certified for: Artificial reproduction					
	I.26.	Transit through third country	I.27. Transit through Member States				
		Third country ISO code Exit point Code Entry point BIP No	Member State ISO code Member State ISO code Member State ISO code				
	I.28.	Export 🗌	1.29.				
		Third country ISO code Exit point Code					
	I.30.						
	I.31.	Identification of the commodities					
		Species Breed Donor identity [(Scientific name)	Date of collection Approval number of the Quantity centre				

▼<u>M1</u>

E	UROPEAN	I UNION			Ovine and caprine semen — Part
1	ll. Hea	Ith informatio	on	II.a. Certificate reference No	II.b.
ſ	, the unde	rsigned offic	ial veterinarian, hereby certify that:		
		II.1.	The semen described above:		
		II.1.1.	was collected, processed and stored in a s in accordance with Chapter $I(l)(1)$ and Cha	emen collection centre (²) approved an apter I(II)(1) of Annex D to Directive 92	d supervised by the competent authority 2/65/EEC;
	II.1.2. comes from donor animals which meet the requirements of Chapter II(II) of Annex D to Directive 92/65/EEC;				
		II.1.3.	was collected, processed, stored and trans and III(I) of Annex D to Directive 92/65/EE		/ with the requirements of Chapters II(I
Image: Control of the second secon					
	(¹) or	[11.1.4.	was collected from animals which have bee holdings which has/have complied for the l 1.3(a) to (f) of Section A of Chapter A of A	ast three years before the collection w	ith the requirements laid down in points
- '	(¹) or	[11.1.4.	was collected from animals which have be with a negligible risk status for classical scr VIII to Regulation (EC) No 999/2001.]		
ſ	(¹) or	[11.1.4.	was collected from ovine animals of the A	RR/ARR prion protein genotype;]	
	II.1.5. was sent to the place of loading in a sealed container in accordance with point 1.4 of Chapter III(I) of Annex D to Direc 92/65/EEC and bearing the number detailed in Box 1.23.				t of Chapter III(I) of Annex D to Directive
1	(¹) either	[.2.	No antibiotics or no mixture of antibiotics v	were added to the semen.]	
1	(¹) or	[11.2.	The following antibiotic or combination of a not less than $(^3)$:	antibiotics was added to produce a co	ncentration in the final diluted semen c
	Notos]
	Notes Part I:				
	3ox 1.12.:	Place of ori	gin shall correspond to the semen collection	centre of origin of the semen.	
	3ox I.13.:	Piace of de	stination shall correspond to the semen colle	action or storage centre or to the holdi	ing of semen destination.
1	3ox 1.23.:	Identificatior	n of container and seal number shall be indic	cated.	
	3ox I.31.:	Donor ident	ity shall correspond to the official identification	on of the animal.	
		Date of coll	ection shall be indicated in the following form	nat: 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.				
		collected.			
	Part II:	collected.			
		as appropria	ite.		
	(¹) Delete (²) Only ap	as appropria	ite. ien collection centres listed in accordance w ood/animal/approved_establishments/establis		
	 Delete Only ap http://ed 	as appropria oproved sem c.europa.eu/f	en collection centres listed in accordance w		

▼<u>M1</u>

EUROPEAN UNION	Ov	ine 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

	FLA	N UNION		Intra trade certificate
	l.1.	Consignor Name	I.2. Certificate reference No	I.2.a. Local reference No
		Address Postal code	I.3. Central competent authority	
nted			I.4. Local competent authority	
prese	1.5.	Consignee Name	1.6.	
nent		Address	1.7.	
ignn		Postal code		
Part I: Details of consignment presented	1.8.	Country ISO code I.9. Region of origin Code of origin	I.10. Country of ISO code destination	I.11. Region of Code destination
tails				
De	1.12.	Place of origin	I.13. Place of destination	
art I		Semen centre	Semen centre 🗖	Holding
۵.		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		
	I.18.	Description of commodity	I.19. Commodity code 05 1	(HS code) 1 99 85
			1.20. Q	uantity
	1.21.	Temperature of products	I.22. Ni	umber of packages
		Ambient Chilled	Frozen	
	1.23.	Seal/Container No	1.24. Ту	ype of packaging
	1.25.	Commodities certified for:		
		Artificial reproduction		
	1.26	Transit through third country	I.27. Transit through Member States	
	1.20.	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 Donor identity (Scientific name)	Date of collection Approval nu of the cer	

ſ		ealth info		II.a. Certificate reference No	vine and caprine semen — Part				
	II. D	eann mi	omation	I.a. Certificate reference No	II.D.				
	I, the unde	ersigned	official veterinarian, hereby certify that the semen de	scribed above:					
		II.1.	was collected, processed and stored in a semen or accordance with Chapter I(I) and Chapter I(II) of A		ervised by the competent authority in				
		II.2.	comes from the donor animals which meet the rec	uirements of Chapter II(II) of Annex [D to Directive 92/65/EEC;				
		II.3.	was collected, processed, stored and transported u III of Annex D to Directive 92/65/EEC;	nder conditions which comply with th	e requirements of Chapters II(II) an				
	(¹) either	[11.4.	meets the requirements of Chapter A(I) of Annex V	/III to Regulation (EC) No 999/2001;]					
	(¹) or	[11.4.	which benefits, for all or part of its territory, from the Regulation (EC) No 999/2001 and the donor anim	nex VIII to Regulation (EC) No 999/2001 and is destined for a Member Stat im the provisions laid down in point (b) or (c) of Chapter A(I) of Annex VIII t animals comply regarding scrapie with the guarantees provided for by th the guarantees (³) requested by the Member State of destination.]					
	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:	Identifica	ation of container and seal number shall be indicated.						
	Box I.31:	Donor id	lentity shall correspond to the official identification of	the animal.					
		Date of	collection shall be indicated in the following format: d	d/mm/yyyy.					
		Approva collectec	I number of the centre shall correspond to the approva d.	I number of the semen centre indicate	ed in Box I.12 where the semen wa				
	Part II:								
	(1) Delete	as appr	opriate.						
			semen collection centres listed in accordance with / .eu/food/animal/approved_establishments/establishme		5/EEC on the Commission website				
	(³) Additio	nal guar	antees as laid down in Article 2 of Regulation (EC) N	o 546/2006 (OJ L 94, 1.4.2006, p. 2	8).				
	Official vel	terinariar	n or official inspector (*)						
	Name	(in capita	al letters):	C	Qualification and title:				
	Local v	/eterinary	y unit:	LVU No:					
	Date:			S	lignature:				
	Stamp:								
1	(*) The co	lour of the	e 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

URC	PEA	N UNION	Intra trade certificate					
	l.1.	5	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	1.6. No(s) of related original certificates No(s) of accompanying documents					
ŗ		Address	.7.					
Bunne		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					
ils	1.40	Discuss of a solution						
Deta	1.12.	Place of origin Semen centre	I.13. Place of destination Semen centre Holding					
#								
Par		Name Approval number Address	Name Approval number Address					
		Postal code	Postal code					
	I.14.		1.15.					
	l.16.	Means of transport	1.17.					
		Aeroplane 🗌 Ship 🗌 Railway wagon 🗌						
		Road vehicle 🗌 Other 🗌						
		Identification						
	l.18.	Description of commodity	I.19. Commodity code (HS code) 05 11 99 85					
			I.20. Quantity					
	I.21.	Temperature of products	I.22. Number of packages					
		Ambient Chilled	Frozen					
	1.23.	Seal/Container No	I.24. Type of packaging					
	1.25.	Commodities certified for:	1					
		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						
	1.30.							
	1.31.	Identification of the commodities						
		Species Breed Donor identity (Scientific name)	Date of collection Approval number Quantity of the centre					

▼	B

EUROPEA	N UNION			Ovine and caprine semen — Part
Π. ⊢	lealth inform	ation	II.a. Certificate reference No	II.b.
I. the und	ersianed offi	۔ cial veterinarian, hereby certify that the semen desc	ribed above:	
(¹) either	[11.1.	was collected, processed and stored for a minim semen collection centre $(^2)$ situated in the Mer accordance with Chapter $(0)(1)$ and Chapter $(0)(1)$ moved to the semen storage centre detailed in Pa animal health and veterinary certification at least a	nber State of origin of the seme) of Annex D to Directive 92/65/E rt I.12 situated in the same Memb	en and operated and supervised i EC, and from where the semen wa
	(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].1.	was collected, processed and stored for a minim semen collection centre (²) situated in the Europe and Chapter I(II) of Annex D to Directive 92/65/EE accordance with:	an Union and operated and super	rvised in accordance with Chapter 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 minim semen collection centre (²) situated in a third cour operated and supervised in accordance with Chapt imported into the European Union under the cond	try or part(s) thereof listed in Anne er I(I)(1) and Chapter I(II)(1) of Ann	ex I to Decision 2010/472/EU which i ex D to Directive 92/65/EEC, and wa
	(¹) either	[Section A of Part 2 of Annex II to Decision 2010/	(472/EU;]	
	(1) or	[Section B of Part 2 of Annex II to Decision 2010/	(472/EU;]	
	(1) or	[Annex II to Decision 2008/635/EC;]		
	II.2.	was stored in the approved semen storage centre with Chapter I(I)(2) and Chapter I(I)(2) of Annex I		perated and supervised in accordanc
	II.3.	was sent to the place of loading in a sealed contai 92/65/EEC and bearing the number indicated in B		f Chapter III(I) of Annex D to Directiv
Notes				
Part I:				
Box 1.6:	panied the s	pond to the serial number of the individual official d emen described above from the approved semen cc of this/these document(s) or certificate(s), or the off	llection centre of its origin to the de	escribed above semen storage centr
Box 1.12:	Place of ori	gin shall correspond to the semen storage centre of	f dispatch of the semen.	
Box 1.13:	Place of de	stination shall correspond to the semen collection o	r storage centre or to the holding	of destination of the semen.
Box 1.23:	Identification	of container and seal number shall be indicated.		
Box 1.31:		ty shall correspond to the official identification of th		
		ection shall be indicated in the following format: dd/		and the same of the
	Approvar hu	mber of the centre shall correspond to the approva	i number of the semen collection i	centre of the semen origin.

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

ANNEX IV

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

▼<u>M2</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

l.1.	Consignor					1.2.	Certificate refe	rence No	1.2.	a. Local refe	rence No		
	Name					1.2	Control compo	topt outbority					
	Address					I.3. Central competent authority							
	Postal code				1.4.	Local compete	nt authority						
1.5.	Consignee Name Address Postal code					1.6.							
						1.7.							
1.8.	Country of origin	ISO code	1.9.	Region of origin	Code	l.10.	Country of destination	ISO code	l.11.	Region of destination	Code		
l.12.	Place of origin Embryo team 🗖					1.13.	I.13. Place of destination Holding □ Embryo team □						
	Name Approval number Address						Name Address	م	∖pprova	l number			
	Postal code	9					Postal code						
l.14.					_	1.15.							
l.16.	Means of transport					l.17.							
	Road vehic Identificatio	le 🗖	Other C	_	on 🗖								
l.18.	Description	of comm	odity						modity 1 99 85	code (CN code	:)		
									1.20.	Quantity			
1.21.	Temperatur Ambient 🗖	e of prod		Chilled 🗖		Froze	en 🗆		I.22. Number of packages				
1.23.	Seal/Conta	iner No							1.24.	Type of pack	kaging		
1.25.	Commoditie Artificial rep												
1.26.	Transit thro	ugh third	country			1.	27. Transit thr	ough Member	States				
	Third count	ry	ISO coo	de			Member S		Code C				
	Exit point						Member S		D code				
	Entry point		BIP No			Member State ISO code							
1.28.				to.		1.29.							
	Third country ISO code Exit point Code												
1.30.													
1.31.	Identificatio	n of the c	ommodi	ties									
	Identification of the commodities Species Category Donor identity						of collection			the team Qu			

11.	N UNION Health inf	ormation II.a. Certificate reference number II.b.								
<u>.</u>	ersigned (official veterinarian, hereby certify that:								
(¹) either	[II.1.	the <i>in vivo</i> derived embryos (¹)/ <i>in vivo</i> derived ova (¹) described above were collected, processed stored by an embryo collection team (²) approved and supervised in accordance with Chapter I(III)(Annex D to Directive 92/65/EEC;]								
(¹) or	[II.1.	he <i>in vitro</i> produced embryos (¹)/micromanipulated embryos (¹) described above were produc processed and stored by an embryo production team (²) approved and supervised in accordance Chapter I(III)(1) and (2) of Annex D to Directive 92/65/EEC;]								
(¹) either	[II.2.	the <i>in vivo</i> derived embryos described above meet the requirements of Chapter III(II)(1) of Annex D Directive 92/65/EEC;]								
(¹) or	[II.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;]								
(¹) or	[11.2.	the <i>in vitro</i> produced embryos described above meet the requirements of Chapter III(II)(3) of Annex D Directive 92/65/EEC;]								
(¹) or	[II.2.	the micromanipulated embryos described above meet the requirements of Chapter III(II)(4) of Annex D Directive 92/65/EEC;]								
	(¹) [II.3.	the consignment consists of embryos of the ovine or caprine species which:								
	(¹) either	[were collected from animals which have been kept continuously since birth on a holding or holdings recognised as having a negligible or a controlled risk of classical scrapie in accordance with point 1 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001;]]								
	(¹) or	[were collected from animals which have been kept continuously for the last three years before the collection on a holding or holdings which have complied for the last three years before collection with the requirements laid down in points (a) to (f) of point 1.3. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001;]]								
	(¹) or	[were collected from animals which have been kept continuously since birth in a Member State or zone of Member State with a negligible risk status for classical scrapie approved in accordance with the fir subparagraph of point 2.2. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001;]]								
	(¹) or	[were collected from ovine animals and								
		(¹) either [are of the ARR/ARR prion protein genotype;]]								
		(¹) or [carry at least one ARR allele and were collected after the date of 1 January 2015;]]								
	II.4.	the ova or embryos described above come from female donors of the ovine (1)/caprine species (1) whit meet the requirements of Chapter IV(3) of Annex D to Directive 92/65/EEC;								
(¹) either	[II.5.	the embryos described above were conceived as a result of artificial insemination of the donor females wi semen which was collected, processed, stored and transported under conditions which comply with the requirements of Chapters I(I), II(I) and III(I) of Annex D to Directive 92/65/EEC;]								
(¹) or	[11.5.	the embryos described above were conceived as a result of <i>in vitro</i> fertilisation of ova complying with the conditions in Chapter III(II)(2) of Annex D to Directive 92/65/EEC with semen which was collected processed, stored and transported under conditions which comply with the requirements of Chapters I(II(I) and III(I) of Annex D to Directive 92/65/EEC;]								
(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 to the place of loading in a sealed container in accordance with point 6 of Chapter III(II) of Annex D to Directive 92/65/EEC and bearing the number detailed Box I.23.								
Notes										
Part I:										
Box I.12:		of origin shall correspond to the embryo collection team or embryo production team of embry on/production.								
Box I.13:		f destination shall correspond to the embryo collection team, embryo production team or to the holding bryos destination.								

▼<u>M2</u>

EURO	OPEA	N UNION		Ovine a	and caprine ova/embryos — Part A					
11.		Health information	ll.a.	Certificate reference number	II.b.					
Box	1.23:	Identification of container and seal	number	shall be indicated.						
Box	I.31:	Category: specify if: in vivo derive embryos.	Category: specify if: in vivo derived embryos, in vivo derived ova, in vitro produced embryos or micromanipulated embryos.							
		Donor identity shall correspond to t	he offici	al identification of the animal.						
		Date of collection shall be indicated in the following format: dd/mm/yyyy.								
		Approval number of the team shall correspond to the embryo collection team or embryo production team of ova/embryos collection/production.								
Part	11:									
(1)	Dele	ete as appropriate.								
(²)		 approved embryo collection or proc imission website; 	luction t	eams listed in accordance with Article	11(4) of Directive 92/65/EEC on the					
	http:	//ec.europa.eu/food/animal/approve	d_establ	ishments/establishments_vet_field_en.	htm.					
_	The	colour of the stamp and signature m	ust be d	ifferent from that of the other particulars	s in the certificate.					
Offic	cial ve	eterinarian or official inspector								
		Name (in capital letters):		Qualification and title:						
		Local veterinary unit:		LVU No:						
		Date: Signature:								
		Stamp:								

PART B

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

EUR	OPE	AN UNION							li li	ntra trade	certificate	
	1.1.	Consignor				1.2.	Certificate reference No		I.2.a. Local	reference I	No	
		Name										
		Address					I.3. Central competent authority					
ented		Postal code					Local competent authority	/				
ser	1.5.	Consignee	1.6.									
nt pre		Name										
lent		Address				1.7.						
ignm		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. Reg des	gion of tination	Code	
etail	1.12	. Place of origin				1.13.	Place of destination					
			Embryo tear	n 🗖			Holdi	ng 🗖	Em	bryo team [
art		Name		Approval number			Name		App	proval numb	per	
<u>.</u>		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	.18. Description of commodity					I.19. Commodi	ty code	(HS code)			
						05 11 99 85						
									.20. Quantity			
	1.21	. Temperature of produ	•				_	1	.22. Number o	of packages		
		Ambient 🗌		Chilled 🗌		Fro	ozen 🗌					
	1.23	. Seal/Container No							.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	SO code		
		Exit point		Code			Member State		I	SO code		
		Entry point		BIP No			Member State		l	SO 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		roval number	Q	uantity	
		(Scientific name)					collection	0	f the team			

					Ovine and caprine ova/embryos — F						
"		Health i	information	II.a. Certificate reference No	II.b.						
_ I,	the un	dersigned	I official veterinarian, hereby certify th	t the ova/embryos (¹) described above:							
		II.1. were collected, processed and stored under conditions which meet the requirements of Directive 92/65/EEC;									
		II.2.	come from female donors of the ov 92/65/EEC;	ne/caprine species (1) which meet the requiremen	ts of Chapter IV of Annex D to Directiv						
(¹) eithei	- [II.3.	meet the requirements of Chapter I No 999/2001.]	of Annex D to Directive 92/65/EEC and of Chap	ter A(I) of Annex VIII to Regulation (E						
(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 Regu No 999/2001 and are destined for a Member State which benefits, for all or part of its territory, from the provisions la point (b) or (c) of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and the donor animals comply regard with the guarantees provided for by the programmes referred to in that point and with the guarantees (²) reques Member State of destination.]									
(¹) eithei	⁻ [II.4.	in the case of embryos, the semen of Chapter A(I) of Annex VIII to Re	used for fertilisation meets the requirements of Di gulation (EC) No 999/2001.]	rective 92/65/EEC and the requiremen						
(1) or	[.4.	of Chapter A(I) of Annex VIII to Reg of its territory, from the provisions I and the donor animals comply rega	used for fertilisation meets the requirements of Di llation (EC) No 999/2001 and is destined for a Me id down in point (b) or (c) of Chapter A(l) of Ann ding scrapie with the guarantees provided for by d by the Member State of destination.]	mber State which benefits, for all or pa ex VIII to Regulation (EC) No 999/200						
\ \	lotes										
F	Part I:										
E	Box 1.12	: Place o	of origin shall correspond to the embr	o collection team of ova/embryos collection.							
E	3ox 1.13	: Place (destina		e embryo collection team, embryo production te	eam or to the holding of ova/embryd						
E	3ox 1.23	: Identific	cation of container and seal number s	all be indicated.							
E	3ox I.31	Donor i Date of	identity shall correspond to the officia f collection shall be indicated in the fo								
F	Part II:										
1.1		e as appr ional guar	,	gulation (EC) No 546/2006 (OJ L 94, 1.4.2006,	p. 28).						
c	Official v	eterinaria	n or official inspector (*)								
	Na	me (in ca	pital letters):		Qualification and title:						
	Loc	al veterin	hary unit:		LVU No:						
	Dat				Signature:						
	Sta	mp:									

ANNEX V

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

PART A

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

EUR	OPE.	AN UNION							Intra trade	certificate
	1.1.	Consignor			I.2. Certificat	e reference No	1	I.2.a. Lo	cal reference	No
		Name								
	Address					competent authori	ty			
Part I: Details of consignment presented		Postal code			I.4. Local co	mpetent authority				
rese	I.5. Consignee I.6									
nt p	Name									
ner		Address	1.7.			_				
ign		Postal code								
cons	I.8.		O code	I.9. Region of origin Code	I.10. Country		ISO		Region of	Code
of		origin	1	1	destinat	ion I	code	C	destination	1
tails	112	. Place of origin			I.13. Place o	f destination				
Det	1.12	•	oryo tean	nΠ	1.10. 1 1205 0	Holdir	na 🗖	E	Embryo team	
μ		Name		Approval number	Name		-		Approval num	
Pai		Address			Address	5			11	
		Postal code			Postal	odo				
						5008				
	1.14	l. 			1.15.					
	I.16	. Means of transport			l.17.					
		Aeroplane 🗌 Shi	nip 🗖	Railway wagon 🗖						
		Road vehicle Oth	her 🔲							
		Identification								
	I.18	. Description of commodity			I.19. Commodity code (HS code) 05 11 99 85					
						L[I	.20. Quar			
	1.21	. Temperature of products			1.22			Number of packages		
		Ambient 🔲	Ch	illed 🔲 Fro	zen 🗌					
	1.23	. Seal/Container No			1.24			. Type of packaging		
	1.25	. Commodities certified for:								
	1.20	Artificial reproduction								
	1.26	. Transit through third countr	nı		1.27 Transit t	hrough Member :	States			
	1.20	Third country	·	LISO code	Member	5	Jiaiba		ISO code	
		Exit point		Code	Member				ISO code	
		Entry point		BIP No	Member	-			ISO code	
	1.00				1.00					
	1.28	Export		100	1.29.					
		Third country Exit point		ISO code Code						
	1.30									
	1.31	Identification of the common Species B (Scientific name)	dities Breed	Category Donor iden		e of A ection	Approval r of the ti		Q	uantity

	EUROPEA				Porcine ova/embryos — Part A							
	II.	Health int	formation	II.a. Certificate reference No	II.b.							
	I, the und	lersigned (official veterinarian, hereby certify that the ova/embry	os (¹) described above:								
Part II: Certification	(¹) either	II. 1 .	were produced/collected (¹), processed and stored by an embryo collection/production (¹) team (²) approved and supervised in accordance with Chapter I(III) of Annex D to Directive 92/65/EEC;									
		II.2.	meet the requirements of Chapter III(II) of Annex D to Directive 92/65/EEC;									
		II.3.	come from donor females of the porcine species which meet the requirements of Chapter IV(2) of Annex D to Directive 92/65/EEC;									
		[.4.	are <i>in vivo</i> derived embryos which:									
		II.4. 1 .	were conceived as a result of artificial insemination with semen meeting the requirements of Directive 90/429/EEC,									
		II.4.2.	originate from a Member State or region thereof:									
		(¹) either	[listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC;]									
		(¹) or	[listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Anr Decision 2008/185/EC;]									
		(1) or	[listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC and have been washed with trypsin;]									
		(¹) or	[listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex II to Decision 2008/185/EC:]									
		(¹) or	[not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I or II to Decision 2008/185/EC and have been washed with trypsin;]									
		(1) or	[not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;]]									
	(¹) or	[.4.	are <i>in vitro</i> produced/micromanipulated (¹) embryos which:									
		II.4.1.	were conceived as a result of in vitro fertilisation with semen meeting the requirements of Directive 90/429/EEC,									
		II.4.2.	originate from a Member State or region thereof:									
		(¹) either	r [listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC;]									
		(¹) or	[listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;]									
		(¹) or	[listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC and the donor females of the ova used for their production comply with the conditions of Article 1 of Decision 2008/185/EC;]									
		(1) or	[listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex II to Decisio 2008/185/EC;]									
		(¹) or		008/185/EC and are destined for a Member State or region thereof listed in Annex I or II r females of the ova used for their production comply with the conditions of Article 1 of								
		(¹) or	[not listed in Annex I or II to Decision 2008/185/EC a or II to Decision 2008/185/EC;]]	Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I ision 2008/185/EC;]]								

EUROPEAN UNION

Porcine ova/embryos - Part A

ONOFL	AN UNION		Porcine ova/em	Porcine ova/embryos — Part						
II.	Health info	rmation	II.a. Certificate reference No II.b.							
1) or	[II.4. are in vivo derived ova which originate from a Member State or region thereof:									
	(¹) either	[listed in Annex I to Decision 2008/185/EC a 2008/185/EC;]	nd are destined for a Member State or region thereof listed in Anr	nex I to Decisio						
	(¹) or	[listed in Annex I to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex I or II to Decision 2008/185/EC;]								
	(1) or	[listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I to Decision 2008/185/EC and which come from donor females complying with the conditions of Article 1 of Decision 2008/185/EC.]								
	(¹) or	[listed in Annex II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex II to Decisio 2008/185/EC;]								
	(1) or	[not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof listed in Annex I or to Decision 2008/185/EC and which come from donor females complying with the conditions of Article 1 of Decisio 2008/185/EC;]								
	(1) or	 [not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annor II to Decision 2008/185/EC;]] 								
	II.5.	were sent to the place of loading in a sealed container under conditions complying with point 6 of Chapter III(II) of Annex D to Directive 92/65/EEC and bearing the number detailed in Box I.23.								
Votes										
Part I:										
3ox 1.12	: place of	origin shall correspond to the embryo collecti	on team or embryo production team of ova/embryos collection/pr	oduction.						
3ox 1.13	: place of	destination shall correspond to the embryo col	ection team, embryo production team or to the holding of ova/emb	ryos destinatio						
3ox 1.23	: identifica	tion of container and seal number shall be in	dicated.							
Box I.31	I: category: specify if: in vivo derived embryos, in vivo derived ova, in vitro produced embryos or micromanipulated embryos. donor identity shall correspond to the official identification of the animal. date of collection shall be indicated in the following format: dd/mm/yyyy.									
	approval number of the team shall correspond to the embryo collection team or embryo production team of ova/embryos collecti production indicated in Box I.12.									
Part II:										
(²) Only			accordance with Article 11(4) of Council Directive 92/65/EEC on	the Commissio						
webs http:/		eu/food/animal/approved_establishments/estat	vlishments_vet_field_en.htm							
Official v	/eterinarian	or official inspector (*)								
Na	me (in capi	tal letters):	Qualification and title:							
Lo	cal veterina	ry unit:	LVU No:							
Da	te:		Signature:							
Sta	amp:									
(*) The o	olour of the s	tamp and signature must be different from that of th	other particulars in the certificate.							

PART B

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

UR	OPE/	AN UNION									Intra trade	e certificate	
	1.1.	I.1. Consignor						I.2. Certificate reference No I.2.a. Local reference No					
		Name											
	Address					1.3.	I.3. Central competent authority						
nted		Postal code					I.4. Local Competent Authority						
Part I: Details of consignment presented	1.5.	Consignee Name				1.6.							
		Address				1.7.							
		Postal code											
		Country of ISO origin		ode I.9. Region of origin C	Code	l.10.	I.10. Country of destination		ISO code	I.11. Re de	egion of estination	Code	
	112	Place of origin				I.13. Place of destination							
		· · · · · · · · · · · · · · · · · · ·	Embryo tea	m 🔲 Approval number					lding 🔲	ding 🗌 Embryo team			
art		Name							Approva		proval num	l number	
ä		Address				Address							
	Postal code				Postal code								
	1.14	1.14.				l.15.							
	1.16.	.16. Means of transport				1.17.							
		Aeroplane 🔲	Ship 🗖	Railway wagor									
		Road vehicle Other											
		Identification											
	I.18. Description of commodity					I.19. Commodity code (HS code)							
							05 11 99 85						
									1.20. Qu	antity			
	I.21. Temperature of products						I.22. Number of packages						
	Ambient Chilled Fro]						
	I.23. Seal/Container No					I.24. Type of packaging							
	I.25. Commodities certified for:												
	Artificial reproduction												
	1.26.	Transit through third c	country			1.27.	Transit ti	nrough Memb	er 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 co	mmodities										
		Species (Scientific name)	Breed	Category I	Donor i d e	entity		ate of lection		val number he team		Quantity	

EUROPEAN UNION Porcine ova/embryos - Part B Ш Health information II.a. Certificate reference No II.b. I, the undersigned official veterinarian, hereby certify that: II.1. The ova/embryos (1) described above: 11.1.1. were collected, processed and stored under conditions which meet the requirements of Directive 92/65/EEC; come from donor female swine which meet the requirements of Chapter IV of Annex D to Directive 92/65/EEC; II.1.2. II.1.3. meet the requirements of Chapter III of Annex D to Directive 92/65/EEC. Certification (¹) either [II.2. In the case of embryos. II.2.1. the semen used for fertilisation meets the requirements of Directive 90/429/EEC: Part II: (11.2.2. the embryos have been washed with trypsin (2).] In the case of ova, the ova comes from a donor female swine which meets the conditions of Article 1 of Decision 2008/185/EC (²).] (1) or [11.2. Notes Part I: Box I.12: place of origin shall correspond to the embryo collection team of ova/embryos collection. Box I.13: place of destination shall correspond to the embryo collection team, embryo production team or to the holding of ova/embryos destination. Box I.23: identification of container and seal number shall be indicated. Box 1.31: category: specify if: in vivo derived embryos, in vivo derived ova, in vitro produced embryos or micromanipulated embryos. donor identity shall correspond to the official identification of the animal. date of collection shall be indicated in the following format: dd/mm/yyyy. approval number of the team shall correspond to the embryo collection team of ova/embryos collection indicated in Box I.12. Part II: (1) Delete as appropriate. (2) This condition applies only to ova and embryos which originate in the Member States or regions thereof not listed in Annexes I and II to Decision 2008/185/EC (OJ L 59, 4.3.2008, p. 19) and destined to the Member States or regions thereof so listed. It shall also apply to movements from Member States or regions thereof listed in Annex II of Decision 2008/185/EC to Member States or regions thereof listed in Annex I of Decision 2008/185/EC to 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 to 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 to 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 to 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 to 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 to 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 to 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 to 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 to 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 to 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 to Member States or regions thereof li 2008/185/EC. Official veterinarian or official inspector (*) Name (in capital letters): Qualification and title: LVU No: Local veterinary unit: Date: Signature: Stamp:

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