DECISIONS

COMMISSION DECISION

of 26 August 2010

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

(notified under document C(2010) 5779)

(Text with EEA relevance)

(2010/470/EU)

THE EUROPEAN COMMISSION,

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

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

Whereas:

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

concerning semen storage centres and detailed conditions for their approval and supervision. It also sets out detailed conditions for the approval and supervision of embryo collection and production teams, for the collection and processing of *in vivo* derived embryos and the production and processing of *in vitro* fertilised embryos and micromanipulated embryos. Annex D, as thus amended, also amended the conditions to be applied to the donor animals of semen, ova and embryos of animals of the equine, ovine and caprine species and of ova and embryos of porcine species.

- (4) It is necessary to establish new model health certificates for trade within the Union of the commodities taking into account the animal health requirements set out in Annex D to Directive 92/65/EEC, as amended by Regulation (EU) No 176/2010.
- (5) In addition, provision should be made for existing stocks of commodities in the Union that comply with the provisions of Directive 92/65/EEC established prior to the entry into force of the amendments introduced by Regulation (EU) No 176/2010. Accordingly, it is necessary to set out separate model health certificates for trade within the Union in semen, ova and embryos of animals of the equine, ovine and caprine species and trade in ova and embryos of animals of the porcine species collected or produced, processed and stored in accordance with Annex D to Directive 92/65/EEC prior to 1 September 2010.
- (6) The long lasting stocking capabilities for such commodities make it impossible at present to fix a date for the exhaustion of the existing stocks. Therefore, it is not possible to fix a date for the termination of the use of those model health certificates for the existing stocks.

⁽¹⁾ OJ L 268, 14.9.1992, p. 54.

⁽²⁾ OJ L 52, 3.3.2010, p. 14.

- (7) In the interests of consistency and simplification of Union legislation, the model health certificates should be set out in a single decision and take account of Commission Regulation (EC) No 599/2004 of 30 March 2004 concerning the adoption of a harmonised model certificate and inspection report linked to intra-Community trade in animals and products of animal origin (1).
- (8) In order to ensure full traceability of the commodities, model health certificates should be set out in this Decision for trade within the Union in semen of animals of the equine, ovine and caprine species collected in approved semen collection centres and dispatched from an approved semen storage centre, whether or not the latter constitutes part of a semen collection centre approved under a different approval number.
- In the interests of clarity of Union legislation, the Union acts setting out model health certificates for trade within the Union in the commodities concerned should be expressly repealed. Accordingly, Commission Decision 95/294/EC of 24 July 1995 determining the specimen animal health certificate for trade in ova and embryos of the equine species (2), Commission Decision 95/307/EC of 24 July 1995 determining the specimen animal health certificate for trade in semen of the equine species (3), Commission Decision 95/388/EC of 19 September 1995 determining the specimen certificate for intra-Community trade in semen, ova and embryos of the ovine and caprine species (4) and Commission Decision 95/483/EC of 9 November 1995 determining the specimen certificate for intra-Community trade in ova and embryos of swine (5) should be repealed.
- (10) The measures provided for in this Decision are in accordance with the opinion of the Standing Committee on the Food Chain and Animal Health,

HAS ADOPTED THIS DECISION:

Article 1

Subject matter

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

- (a) semen of animals of the equine species;
- (b) ova and embryos of animals of the equine species;
- (c) semen of animals of the ovine and caprine species;
- (d) ova and embryos of animals of the ovine and caprine species;
- (1) OJ L 94, 31.3.2004, p. 44.
- (2) OJ L 182, 2.8.1995, p. 27.
- (3) OJ L 185, 4.8.1995, p. 58.
- (4) OJ L 234, 3.10.1995, p. 30.
- (5) OJ L 275, 18.11.1995, p. 30.

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

Article 2

Trade in semen of animals of the equine species

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

- (a) model health certificate IA as set out in Part A, for consignments of semen collected after 31 August 2010 and dispatched from an approved semen collection centre of origin of the semen;
- (b) model health certificate IB as set out in Part B, for consignments of stocks of semen collected, processed and stored before 1 September 2010 and dispatched after 31 August 2010 from an approved semen collection centre of origin of the semen;
- (c) model health certificate IC as set out in Part C, for consignments of semen and stocks of semen referred to in (a) and (b) dispatched from an approved semen storage centre.

Article 3

Trade in ova and embryos of animals of the equine species

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

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

Article 4

Trade in semen of animals of the ovine and caprine species

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

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

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

Article 5

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

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

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

Article 6

Trade in ova and embryos of the porcine species

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

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

Article 7

Repeals

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

Article 8

Applicability

This Decision shall apply from 1 September 2010.

Article 9

Addressees

This Decision is addressed to the Member States.

Done at Brussels, 26 August 2010.

For the Commission

John DALLI

Member of the Commission

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

EUR	OPEA	N UNION							Intra t	rade certificate	
	1.1.	•			I.2. C	I.2. Certificate reference No I.2.a. Local reference No					
		Name			13 0	I.3. Central competent authority					
		Address Postal code					entrar competent a	authonty			
nted							ocal competent au	ithority			
esei	1.5.	Consignee			1.6.						
t p		Name									
nen		Address									
gi		Postal code									
of consignment presented	1.8.	Country of origin ISO code	I.9. Region	of origin	Code		Country of destination	ISO code	I.11. Region of destination	Code	
Part I: Details of	1.12.	Place of origin				113	Place of destinatio	l n			
<u> </u>		Semen ce	entre 🗆			1.10.		_	Holding		
a z			_				Semen cent	ie 🖂	Holding \square		
۳		Name Address	Approval n	umber			Name		Approval num	nber	
							Address				
		Postal code				_	Postal code				
	1.14.					l.15.					
	1.16.	Means of transport				1.17.	-				
		Aeroplane Ship		Railway wag	on \square						
		. – .	er 🗆	nanway wag	011						
			21 LJ								
		Identification									
	I.18. 	Description of commodity					I.19. Co	•	ode (HS code)		
									5 11 99 85		
								1.20.	Quantity		
	1.21.	Temperature of product									
		Ambient	Chille	ed 🔲			Frozen 🔲	1.22.	Number of package	S	
	1.23.	Seal/container No						1.24.	Type of packaging		
		0 100 100 10									
	1.25.	Commodities certified for: Artificial reproduction									
	1.26.	Transit through a third count	ry []		1.27.	Fransit through Me	mber State	s		
		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 commod	ities								
		Species (scientific name)	Breed	Donor ider	ntity [Date of o	collection	Approval nu of the te		Quantity	

Part II: Certification

	FUDODE	AN LINION				Familia comos Dort A				
		AN UNION Health infor	mation		II.a. Certificate reference No	Equine semen — Part A				
		Tioditi' iiiioii	madon		n.a. Ooranoate reference 140	11.0.				
	I, the un	dersigned o	fficial veter	inarian, hereby certify that:						
	II.1.	The semen collection centre (2), in which the semen described above was collected, processed and stored for trade is approved and supervised by the competent authority in accordance with Chapter I(I)(1) and Chapter I(II)(1) of Annex D to Directive 92/65/EEC;								
	II.1.1.	during the period commencing 30 days prior to the date of first collection of the semen described above until the date the fresh or chilled semen was dispatched or until the 30 days storage period for frozen semen elapsed, the semen collection centre;								
	II.1.1.1.			erritory or in the case of regionalisation in a norse sickness in accordance with Article						
	II.1.1.2.	fulfilled the	conditions	for a holding laid down in Article 4(5) of	Directive 2009/156/EC;					
•	II.1.1.3.	contained of	only equida	ae which were free of clinical signs of equ	uine viral arteritis and contagious equir	ne metritis.				
	II.2.	Only equidathe centre.	ae satisfyin	g the conditions laid down in Articles 4 an	d 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	e centre and on the day the semen				
	II.3.2.			days prior to the date of semen collection equine metritis during that period;	n in holdings where no equine has sho	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		tion and from the dates of the first				
	II.3.4.	Vaccines for	or Terrestri	ollowing tests, which meet at least the red al Animals of the OIE, carried out on san recognised by the competent authority:						
		(1) either	[II.3.4.1.	an agar-gel immuno-diffusion test (Cogg	ins test) for equine infectious anaemia	(EIA) with negative result;]				
		(¹) or	[II.3.4.1.	an ELISA for equine infectious anaemia	(EIA) with negative result;]					
	and	(1) either	[II.3.4.2.	a serum neutralisation test for equine vi	ral arteritis (EVA) with negative result	at a serum dilution of one in four;]				
		(¹) or	[II.3.4.2.	a virus isolation test for equine viral arter of the donor stallion;]	itis (EVA) carried out with negative resu	ılt on an aliquot of the entire semen				
	and		[II.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 each	on of <i>Taylorella equigenitalis</i> after a cl d from genital swabs taken at least fr	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 s follows:	case to at least one of the test progra	mmes (4) detailed in points II.3.5.1,				
	II.3.5.1.	during the	period of co	continuously resident on the semen colle ollection of the semen described above an th status than the donor stallion;						

the tests described in point II.3.4 have been carried out on samples taken (5) prior to the first semen collection and at least 14 days following the date of the commencement of the residence period of at least 30 days;

EUROPEAN UNION	Equine semen — Parl	. A

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

II.3.5.2. the donor stallion was resident on the semen collection centre for at least 30 days prior to the date of the first collection and during the period of collection of the semen described above, but has left the centre under the responsibility of the centre veterinarian for a continuous period of less than 14 days, and/or other equidae on the collection centre came into direct contact with equidae of lower health status:

the tests described in point II.3.4 have been carried out on samples taken (5) prior to the first semen collection of the breeding season or collection period in the year the semen described above was collected and at least 14 days following the date of the commencement of the residence period of at least 30 days;

and the test described in point II.3.4.1 for equine infectious anaemia was last carried out on a sample of blood taken (5) not more than 90 days before the semen described above was collected;

and (1) either [one of the tests described in point II.3.4.2 for equine viral arteritis was last carried out on a sample taken (5) not more than 30 days before the semen described above was collected;]

(1) or [a virus isolation test for equine viral arteritis was carried out with negative result on an aliquot of the entire semen of the donor stallion taken (5) not more than 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. The tests described in point II.3.4 have been carried out on samples taken (5) prior to the first semen collection of the breeding season or collection period in the year the semen described above was collected,

and the tests described in point II.3.4 were last carried out on samples taken (5) not less than 14 days and not more than 90 days after the collection of the semen described above.

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

_		Start date	e (⁵)		D	ate of sampling for h	ealth tests (5)	
Identification of semen	Test programme	Donor residence	Semen collection	EIA II.3.4.1	E II.3	VA 3.4.2	CE II.3	EM .4.3
Identi of s	T				Blood Semen sample sample		1. sample	2. sample

(')	either	[11.4	No	antibiotics	were	added	to	the	semen;	
-----	--------	-------	----	-------------	------	-------	----	-----	--------	--

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

II.5. The semen described above was:

- II.5.1. collected, processed, stored and transported under conditions which comply with the requirements of Chapters II(I)(1) and III(I) of Annex D to Directive 92/65/EEC;
- II.5.2. sent to the place of loading in a sealed container in accordance with point 1.4 of Chapter III(I) of Annex D to Directive 92/65/EEC and bearing the number indicated in Box I.23.

EUROPEAN UNION	Equine semen — Part A
----------------	-----------------------

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

Notes

Part I:

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

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

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

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

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

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

Part II:

Guidance for the completion of Table in II.3.6:

Abbreviations:

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

EIA-2 EIA testing second occasion

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

EVA-B2 EVA testing on blood sample second occasion

EVA-S1 EVA testing on semen sample first occasion

EVA-S2 EVA testing on semen sample second occasion

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

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

CEM-21 CEM testing second occasion first sample

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

Instructions:

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

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

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

EUROP	EAN UN	ION				Equine semen — Part A
II.	Health	information		II.a. Certifi	cate reference No	II.b.
ے	Start date (5)			Dat	e of sampling for health	tests (5)

							The same of the sa	
_	0	Start date	; (⁵)		D	ate of sampling for he	ealth tests (5)	
Identification of semen	Test gramme	Donor residence			_	EVA II.3.4.2		EM .4.3
Ident of s	T				Blood sample	Semen sample	1. sample	2. sample
_	В	6	_	EIA-1	EVA-B1	EVA-S1	CEM-11	CEM-12
A	В	С	D	EIA-2	EVA-B2	EVA-S2	CEM-21	CEM-22

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

Official veterinarian (*)								
	Name (in capital letters):	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 on the certificate.

PART B

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

URO	PEA	N UNION	Intra trade certificate
	l.1.	Consignor	I.2. Certificate reference No I.2.a. Local reference No
		Name	
		Address	I.3. Central competent authority
ted		Postal code	I.4. Local competent authority
esen	l.5.	Consignee	1.6.
ğ		Name	
ient		Address	1.7.
gnr		Postal code	
of consignment presented	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
Part I: Details	140	Discourse and addition	
) C	1.12.	Place of origin	I.13. Place of destination
-		Semen centre	Semen centre Holding
2		Name Approval number	Name Approval number
		Address	Address
		Postal code	Postal code
	1.14.		1.15.
	I.16.	Means of transport	I.17.
		Road vehicle Other	
		Identification	
	I.18.	Description of commodity	I.19. Commodity code (HS code) 05 11 99 85
			I.20. Quantity
	I.21.	Temperature of product	
		Ambient Chilled Chilled	Frozen I.22. Number of packages
	1.23.	Seal/container No	I.24. Type of packaging
	1.25.	Commodities certified for:	
		Artificial reproduction	
	1.26.	Transit through 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 Date (scientific name)	te of collection Approval number Quantity of the team

	EUROPE	AN UNION		Equine semen — Part E					
	II.	Health information		II.a. Certificate reference No	II.b.				
	I, the ur	dersigned official veterir	narian, hereby certify that:						
Part II: Certification	II.1.	The semen collection centre (2), in which the semen described above was collected, processed and stored for trade:							
	II.1.1.	is approved and super	vised by the competent authority accor	ding to the conditions of Chapter I o	f Annex D to Directive 92/65/EEC;				
	II.1.2.	is situated on the territory or in the case of regionalisation in a part of the territory (1) of a Member State which was on the day semen was collected until the date the semen was dispatched as fresh/chilled (1) semen or until the 30 days mandatory storage period for frozen semen elapsed (1) not considered to be infected with African horse sickness in accordance with Article 5(2)(a) and (b) of Directive 2009/156/EC (3);							
Part II: C	II.1.3.	fulfilled during the period commencing 30 days prior to the date of semen collection until the date the semen was dispatched as fresh/chilled (¹) semen or until the 30 days mandatory storage period for frozen semen elapsed (¹), the conditions of Article 4 of Directive 2009/156/EC;							
	II.1.4.	contained during the period commencing 30 days prior to the date of semen collection until the date the semen was dispatched as fresh/chilled (¹) semen or until the 30 days mandatory storage period for frozen semen elapsed (¹) only equidae which were free of clinical signs of equine viral arteritis and contagious equine metritis;							
	II.2.	All equidae have been	admitted into the centre under the prov	isions of Article 4 and 5 of Directive 2	2009/156/EC.				
	II.3.	The semen described above was collected from donor stallions, which:							
	II.3.1.	on the day the semen was collected have not shown clinical signs of an infectious or contagious disease;							
	II.3.2.	during at least 30 days	s prior to collection of the semen have n	ot been used for natural service;					
	II.3.3.	during the last 30 days arteritis;	prior to collection of the semen have bee	n kept on holdings where no equidae	showed clinical signs of equine viral				
	II.3.4.	during the last 60 days prior to collection of the semen have been kept on holdings where no equidae showed clinical signs of contagious equine metritis;							
	II.3.5.	to the best of my knowledge and as far as I could ascertain, have not been in contact with equidae suffering from an infectious or contagious disease during 15 days immediately preceding collection of the semen;							
	II.3.6.	have undergone the following animal health tests, carried out in a laboratory recognised by the competent authority, in accordance with a test programme as specified in point II.3.7;							
		[II.3.6.1.	an agar gel immunodiffusion test (Cog	gins test) for equine infectious anaem	ia with negative result;]				
	and	(¹) either [II.3.6.2.	a serum neutralisation test for equine	viral arteritis with negative result at a	serum dilution of one in four; and]				
		(¹) or [II.3.6.2.	a virus isolation test for equine viral art the donor stallion;]	eritis carried out with negative result of	on 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 <i>Taylorella equi</i>	genitalis from pre-ejaculatory fluid or				
	II.3.7.	have been subject to t	he one of the following test programmes	s (⁴):					
	II.3.7.1.		continuously resident in the collection of o equidae in the collection centre came of						
		contagious equine met	point II.3.6 have been carried out on sar ritis on a second sample taken on above residence period and at least at	(⁵), being	at least 14 days after the				

EUROPE	AN UNION			Equine semen — Part B		
II.	Health inforn	ation	II.a. Certificate reference No	II.b.		
II.3.7.2.		stallion was not continuously resident in the collection lower health status than the donor stallion;	on centre or other equidae in the colle	ection centre came into contact with		
	contagious	escribed in point II.3.6 have been carried out on sa equine metritis on a second sample taken on men collection and at least at the beginning of the	(⁵), being			
and		the test described in point II.3.6.1 for equine infectious anaemia was last carried out on a sample of blood taken on				
and (1) either [one of the tests described in point II.3.6.2 for equine viral arteritis was last carried out on a sample c						
	(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 above	the donor stallion collected on	d by a virus isolation test which was		
II.3.7.3.	14 days af	lescribed in point II.3.6 have been carried out during ter the collection of the semen on samples taken of ad sample taken on	on (5) and in the			
II.4.		described above was collected, processed, stored and III of Annex D to Directive 92/65/EEC.	I and transported under conditions wh	ich comply with the requirements of		
Notes						
Part I:						
Box I.12	2: place of or	igin shall correspond to the semen collection centre	e of origin of the semen.			
Box I.13	3: place of de	estination shall correspond to the semen collection	or storage centre or to the holding of	semen destination.		
Box I.23	3: identificatio	n of container and seal number shall be indicated.				
Box I.31	: donor iden	tity shall correspond to the official identification of t	the animal.			
	date of col	lection shall be indicated in the following format: do	d/mm/yyyy.			
	approval no collected.	umber of the centre shall correspond to the approva	al number of the semen centre indicate	d in Box I.12 where the semen was		
Part II:						
` '	te as appropi					
		men collection centres listed in accordance with A u/food/animal/approved_establishments/establishmer		5/EEC on the Commission website:		
	. 192, 23.7.20					
' '		gramme(s) that do(es) not apply to the consignmen	nt.			
(⁵) Inser						
		r official inspector (*)				
	me (in capital	,	Qualification and title:			
Loc	al veterinary	unit:	LVU No:			
Dai	te:		Signature:			
Sta	mp:					
(*) The c	olour 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

=UKC	ROPEAN UNION Intra trade certificate									
	l.1.	Consignor				I.2. Certificate	reference N	0	I.2.a. Local ref	ference No
		Name Address Postal code Consignee Name			I.3. Central co	I.3. Central competent authority				
ъ					I.4. Local com	npetent autho	rity			
ente	1.5.				I.6. No(s) of re	elated origina	l	No(s) of accompa	nvina	
pres					certificates			documents		
ent		Address				1.7.				
gum		Postal code								
of consignment presented	1.8.	Country of origin	ISO code	I.9. Region of origin	Code	I.10. Country destination		ISO code	I.11. Region of destination	Code
siis	110	Place of origin				I do Divisió	-1			
Part I: Details	1.12.	Place of origin				I.13. Place of				
-		Semen centre				emen centre	Ш	Holding \square		
Pal		Name Address		Approval number		Name Address			Approval nu	mber
		Postal code					مام			
	1.14.					Postal co	oue			
						1.10.				
	I.16.	Means of transport				l.17.			_	
		Aeroplane \square	Ship [Railway wa	igon 🔲					
		Road vehicle	Other							
		Identification								
	I.18.	8. Description of commodity		I.19. Commodity code (HS code) 05 11 99 85						
									Quantity	
	1.01	Temperature of prod	duat							
	1.21.	Ambient	Juct	Chilled		Frozen		es		
	1.23.	Seal/container No				I.24. Type of packaging		<u> </u>		
	1.25.	Commodities certific								
	1.26.	Transit through a th				I.27. Transit th	rough Memb	er State	 S	
		Third country	ISO co			Member	State		ISO code	
		Exit point	Code			Member			ISO code	
		Entry point	BIP N	0		Member	State		ISO code	
	1.28.	Export	100	_ 🗆		1.29.				
		Third country Exit point	ISO co Code	ode						
	1.00									
	1.30.									
	I.31.	Identification of the	commoditie	es						
		Species (scientific name)		Breed Donor ide	entity	Date of collec	tion		oval number the team	Quantity

Part II: Certification

EUROPEAN UNION	Equine semen — F	Part C

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

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

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

- (1) either [Part A of Annex I to Decision 2010/470/EU;]
- (1) or [Part B of Annex I to Decision 2010/470/EU;]
- (1) or [Decision 95/307/EC;]
- (1) or [II.1. was collected, processed and stored for a minimum period of 30 days immediately following collection in an approved semen collection centre (2) situated in the European Union and operated and supervised in accordance with Chapter I(I)(1) and Chapter I(II)(1) of Annex D to Directive 92/65/EEC, and was moved to the semen storage centre detailed in Box I.12 in accordance with:
 - (1) either [Part A of Annex I to Decision 2010/470/EU;]
 - (1) or [Part B of Annex I to Decision 2010/470/EU;]
 - (1) or [Part C of Annex I to Decision 2010/470/EU;]
 - (1) or [Decision 95/307/EC;]
- (1) or [II.1. was collected, processed and stored in an approved semen collection centre (2) situated in a third country or part(s) thereof listed in columns 2 and 4 of Annex I to Commission Decision 2004/211/EC which is operated and supervised in accordance with Chapter I(I)(1) and Chapter I(II)(1) of Annex D to Directive 92/65/EEC, and was imported into the European Union under the conditions of Article 4 of Decision 2004/211/EC in accordance with:
 - (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) indicated in Box I.12, which is operated and supervised in accordance with Chapter I(I)(2) and Chapter I(II)(2) of Annex D to Directive 92/65/EEC;
 - II.3. was sent to the place of loading in a sealed container in accordance with point 1.4 of Chapter III(I) of Annex D to Directive 92/65/EEC and bearing the number indicated in Box I.23.

Notes

Part I:

Box I.6: shall correspond to the serial number of the individual official document(s) or health certificate(s) [either INTRA or CVED] that accompanied the semen described above from the approved semen collection centre of its origin to the described above semen storage centre.

The original(s) of this/these document(s) or certificate(s) or the officially endorsed copy/copies thereof must be attached to this certificate.

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

EUROPE	EUROPEAN UNION Equine semen — Part						
II.	Health information	II.a. Certificate reference No	II.b.				
Box I.13	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 1.23	Box I.23: identification of container and seal number shall be indicated.						
Box I.31	Box I.31: donor identity shall correspond to the official identification of the animal.						
	date of collection shall be indicated in the following format: do	il/mm/yyyy.					
	approval number of the centre shall correspond to the approv	al number of the semen collection cer	ntre of origin of the semen.				
Part II:							
(1) Delet	e as appropriate.						
	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				
1 '	/ec.europa.eu/food/animal/approved_establishments/establishmer /ec.europa.eu/food/animal/semen_ova/equine/index_en.htm	nts_vet_field_en.htm					
Official v	veterinarian or official inspector (*)						
	Name (in capital letters):	Qualification and title:					
	Local veterinary unit:	LVU No:					
	Date: Signature:						
	Stamp:						
(*) The co	(*) The colour of the stamp and signature must be different from that of the other particulars in the certificate.						
() The colour of the stamp and signature files be different from that of the other particulars in the certificate.							

ANNEX II

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

PART A

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

		N UNION	Intra trade certificat			
	l.1.	· ·	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			
esent	l.5.	Consignee	1.6.			
t pr		Name	17			
men		Address	1.7.			
ign		Postal code				
s of consignment presented	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			
taile	1.12.	Place of origin	I.13. Place of destination			
٠٠		Embryo team □	Holding ☐ Embryo team ☐			
Part I: Details		Annual multi-				
4		Name Approval number Address	Name Approval number			
		Postal code	Address			
			Postal code			
	1.14.		I.15.			
	l.16.	Means of transport	1.17.			
		Aeroplane Ship Railway wagon				
		Road vehicle Other				
		Identification				
	I.18.	Description of commodity	I.19. Commodity code (HS code)			
			05 11 99 85			
			I.20. Quantity			
	1.21.	Temperature of products	I.22. Number of packages			
		Ambient Chilled 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				
	1.00	<u>'</u>				
	1.30.					
	I.31.	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 — Part A					
II. He	alth info	rmation	II.a. Certificate reference No	II.b.				
I, the under	rsigned o	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 embryo collection team (2) approved and supervised in accordance with Chapter I(III)(1) of Annex D to Directive 92/65/EEC;]							
(¹) or	[II.1. the <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;]							
(¹) either	[II.2.	the <i>in vivo</i> derived embryos described above meet the requirements of Chapter III(II)(1) of Annex D to Directive 92/65/EEC;]						
(¹) either	[II.2.	the <i>in vivo</i> derived ova described above me 92/65/EEC;]	et the requirements of Chapter III	(II)(2) of Annex D to Directive				
(¹) or	[II.2.	the <i>in vitro</i> produced embryos described above 92/65/EEC;]	meet the requirements of Chapter	III(II)(3) of Annex D to Directive				
(¹) or	[II.2.	the micromanipulated embryos described above 92/65/EEC;]	meet the requirements of Chapter	III(II)(4) of Annex D to Directive				
	II.3.	the ova or embryos described above come from de	onor mares which:					
II.3.1. coming from holdings fulfilling the conditions laid down in Article 4(5) of Directive 2009/156/EC (4) onto which only satisfying the conditions laid down in Articles 4 and 5 or Articles 12 to 16 of Directive 2009/156/EC have been a								
	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 let 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	(3), being during the page last test was carried out of	ast 30 days prior to the date of the n a sample of blood taken				
	II.3.5.	have been subjected to an agent identification test froultivation of 7 to 14 days carried out with negative redate of the first collection of ova or embryos frounds consecutives oestrus periods on	results in each case on samples taken om mucosal surfaces of the clitoral to and on	during the past 30 days prior to the fossa and clitoral sinuses on two				
(¹) either	[II.4.	the embryos described above were conceived as a collected, processed, stored and transported under III(I) of Annex D to Directive 92/65/EEC;]						
(¹) or	[II.4.	the embryos described above were conceived as a of Chapter III(II) of Annex D to Directive 92/65/EEC conditions which comply with the requirements of C	with semen which was collected, proce	ssed, stored and transported under				
(¹) or	[11.4.	the ova have not been in contact with semen of th	e equine species;]					
II.5. the ova or embryos described above were sent to the place of loading in a sealed container in accordance with point 6 or Chapter III(II) of Annex D to Directive 92/65/EEC and bearing the number detailed in Box I.23.								
Notes								
Part I:								
Box I.12: F	Place of o	origin shall correspond to the embryo collection team	n or embryo production team of ova/er	nbryos collection/production.				
	Place of Jestinatio	destination shall correspond to the embryo collect	tion team, embryo production team	or to the holding of ova/embryos				
		ion of container and seal number shall be indicated.						

EUROP	EAN UNION	Equine ova and embryos — Part A					
II.	Health information	II.a. Certificate reference No	II.b.				
Box I.3	31: Category: specify if: in vivo derived embryos, in vivo derived	ova, <i>in vitro</i> produced embryos or mic	romanipulated embryos.				
	Donor identity shall correspond to the official identification of the animal.						
	Date of collection shall be indicated in the following format: d	d/mm/yyyy.					
	Approval number of the team shall correspond to the collection/production.	embryo collection team or embryo	production team of ova/embryos				
Part II	:						
(1) De	lete as appropriate.						
	ly approved embryo collection or production teams listed in accord bsite:	ance with Article 11(4) of Council Direct	ctive 92/65/EEC on the Commission				
http	o://ec.europa.eu/food/animal/approved_establishments/establishme	nts_vet_field_en.htm					
(3) Ins	ert date.						
(4) OJ	L 192, 23.7.2010, p. 1.						
Officia	veterinarian or official inspector (*)						
Na	me (in capital letters):	Qi	ualification and title:				
Loc	cal veterinary unit:	L\	/U No:				
Da	te:	Si	gnature:				
Sta	Stamp:						
(*) The							
(c) the	(*) The colour of the stamp and signature must be different from that of the other particulars in the certificate.						

PART B

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

EUR	UROPEAN UNION Intra trade certificate							
	l.1.	Consignor		I.2. Certificate	e reference	No	I.2.a. Local refere	ence No
		Name	I.3. Central competent authority					
		Address	1.3. Central c	ompetent a	uthority			
		D		I.4. Local cor	mpetent auti	nority		
pa		Postal code			·			
sent	1.5.	Consignee		I.6.				
pres		Name						
Ţ		Address	1.7.					
E		Postal code		***************************************				
consignment presented	1.8.	Country of origin ISO code I.9. Region of origin Code		I.10. Country	of	ISO	I.11. Region of	Code
co	1.0.			destinat	ion .	code	destination	
of of								
Part I: Details	1.12.	Place of origin		I.13. Place of	destination			
De		Embryo team [П		Holdi	ng 🔲	Embryo team [7
벌					rioidi	ng 🗀	Embryo team [_
Pa			Approval number	Name			Approval numb	er
		Address		Address				
		Postal code		Postal c	ode			
	1.14.			I.15.				
	I.16. Means of transport		_	l.17.				
		Aeroplane Ship	Railway wagon 🗌					
		Road vehicle Other		_				
		Identification						
	I.18.	Description of commodity			I.19. Com		de (HS code)	
						1.20. Qu	5 11 99 85	
					1.20. Qu	arrity		
	1.21.	Temperature of products				1.22. Nur	mber of packages	
		Ambient	Chilled	Frozen				
	1.23.	Seal/Container No				1.24. Tvr	pe of packaging	
	1.25.	Commodities certified for:						
		Artificial reproduction		I				
	1.26.	Transit through third country		I.27. Transit t	•	nber State		
		Third country	SO code	Member	r State		ISO code	
		Exit point C	Code	Member	State		ISO code	
		Entry point B	BIP No	Member	r State		ISO code	
	1.28.	Export		1.29.				
		Third country IS	SO code					
		Exit point C	Code					
	1.30.							
	1,31.	Identification of the commodities	-					
		Species Breed (Scientific name)	Category Donor identity	y Date o	of collection	A	approval number of the team	Quantity

II: Certification

Part

EUROPEAN UNION Equine ova and embryos — Part B

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

- I, the undersigned official veterinarian, hereby certify that:
- II.1. Ova/embryos (1) described above were collected by a collection team (2) approved by the competent authority and processed in an appropriate laboratory;
- II.2. Ova/embryos (1) were collected from donor mares which:
- II.2.1. on the day of collection have been located in premises situated on the territory or in the case of regionalisation in a part of the territory of a Member State which is not considered to be infected with African horse sickness in accordance with Article 5(2)(a) and (b) of Directive 2009/156/EC (3);
- II.2.2. have been located in holdings under veterinary supervision which on the day of collection fulfilled the conditions of Article 4 of Directive 2009/156/EC;
- II.2.3. have been kept prior to the collection in holdings free from clinical signs of contagious equine metritis for 60 days;
- II.2.4. have not been used for natural breeding during the period of 30 days prior to the collection of ova/embryos (1);
- II.2.5. to the best of my knowledge and as fare as I could ascertain, have not been in contact with equidae suffering from an infectious or contagious disease during the 15 days immediately preceding the collection of ova/embryos (1);
- II.2.6. have on the day of collection not shown clinical signs of an infectious or contagious disease;
- II.3. Ova/embryos (1) were collected, processed, stored and transported under conditions which comply with the requirements of Annex D of Directive 92/65/EEC:
- II.4. The semen used for the artificial insemination of the donor mares complies with the requirements of Directive 92/65/EEC (4) (1);
- II.5. The ova used for the in vivo production of embryos comply with the requirements of Directive 92/65/EEC (1).

Notes

Part I:

- Box I.12: Place of origin shall correspond to the embryo collection team of ova/embryos collection.
- Box I.13: Place of destination shall correspond to the embryo collection team, embryo production team or to the holding of ova/embryos destination.
- Box I.23: Identification of container and seal number shall be indicated.
- Box I.31: Category: specify if: in vivo derived embryos, in vivo derived ova, in vitro produced embryos or micromanipulated embryos.

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

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

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

Part II:

- (1) Delete as appropriate.
- (2) Only approved embryo collection teams listed in accordance with Article 11(4) of 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) Does not apply to ova.

EUROPEAN UNION	Equi	ne ova and embryos — Part B				
II. Health information	II.a. Certificate reference No	II.b.				
Official veterinarian or official inspector (*)						
Name (in capital letters):	Qual	fication and title:				
Local veterinary unit:	LVU	No:				
Date:	Signa	ature:				
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

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	EUROPEAN UNION Intra trade certifica							
	l.1.	3	I.2. Certificate reference No I.2.a. Local reference No					
		Name Address	I.3. Central competent authority					
٦		Postal code	I.4. Local competent authority					
ente	1.5.	Consignee	1.6.					
Les		Name						
벌		Address	1.7.	1.7.				
June		Postal code						
of consignment presented	1.8.	Country ISO code I.9. Region of origin Code of origin	I.10. Country of destination	I.11. Region of Code destination				
ails	1.12.	Place of origin	I.13. Place of destination	·				
Part I: Details		Semen centre	Semen centre	Holding				
±		Name Approval number	Name	Approval number				
Pa		Address	Address	Approvar number				
		Postal code	Postal code					
	1.14.		1.15.					
	1.16.	Means of transport	1.17.					
		Aeroplane						
	1.18.	Description of commodity	I.19. Commodity	code (HS code)				
				05 11 99 85				
			I.	20. Quantity				
	1.21.	Temperature of products		22. Number of packages				
		Ambient Chilled Chilled	Frozen	T.ZZ. Nambor 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 St	ates				
		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)		roval number Quantity the centre				

EUROPEAN UNION

Ovine and caprine semen — Part A

	II. He	alth infor	mation	II.a. Certificate reference No	II.b.				
	I, the under	signed o	official veterinarian, hereby certify that:						
		II.1.	. the semen described above:						
		II.1.1.	 was collected, processed and stored in a semen collection centre (2) approved and supervised by the competent authority in accordance with Chapter I(I)(1) and Chapter I(II)(1) of Annex D to Directive 92/65/EEC; 						
tion		II.1.2.	comes from the donor animals which meet the requ	uirements of Chapter II(II) of Annex D	to Directive 92/65/EEC;				
Part II: Certification		II.1.3.	was collected, processed, stored and transported un III(I) of Annex D to Directive 92/65/EEC;	nder conditions which comply with the	requirements of Chapters II(II) and				
‡ ∺	(¹) either	[II.1.4.	meets the requirements of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001;]						
Par	(¹) or	[II.1.4.	meets the requirements of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and is destined for a Member State which benefits, for all or part of its territory, from the provisions laid down in point (b) or (c) of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and the donor animals comply regarding scrapie with the guarantees provided for by the programmes referred to in that point and with the guarantees (3) requested by the Member State of destination;]						
		II.1.5.	was sent to the place of loading in a sealed container in accordance with point 1.4 of Chapter III(I) of Annex D to Directive 92/65/EEC and bearing the number detailed in Box I.23;						
	(¹) either	[II.2.	no antibiotics or no mixture of antibiotics were adde	ed to the semen;]					
	(¹) or	[II.2.	the following antibiotic or combination of antibiotics less than (4):	was added to produce a concentrati	_				
	Mataa								
	Notes Part I:								
		lace of a	origin shall correspond to the semen collection centre	e of origin of the semen					
			destination shall correspond to the semen collection	•	semen destination				
			ion of container and seal number shall be indicated.	or delayer control or to the ficially of	comen assumation.				
			entity shall correspond to the official identification of t	he animal.					
			ollection shall be indicated in the following format: do						
		pproval ollected.	number of the centre shall correspond to the approva	I number of the semen centre indicate	d in Box I.12 where the semen was				
	Part II:								
	(1) Delete a	s appro	priate.						
	()		semen collection centres listed in accordance with A eu/food/animal/approved_establishments/establishmer	,	5/EEC on the Commission website:				
	(3) Addition	al guara	ntees as laid down in Article 2 of Regulation (EC) No	o 546/2006 (OJ L 94, 1.4.2006, p. 28	3).				
	(4) Insert na	ames an	d concentrations.						
	Official veterinarian or official inspector (*)								
	Name (in capital letters): Qualification and title:								
	Local ve	eterinary	unit:	L	/U No:				
	Date:			S	gnature:				
	Stamp:								
	(*) The colour of the stamp and signature must be different from that of the other particulars in the certificate.								

PART B

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

EUR	OPEA	N UNION					Intra trade	certificate		
	l.1.	Consignor Name		I.2. Certificate re	eference No	1.2	.a. Local reference	e No		
		Address Postal code		I.3. Central com	I.3. Central competent authority					
ted		rostal code	I.4. Local compe	etent authorit	у					
presen	1.5.	Consignee Name		1.6.						
ent		Address		1.7.						
guu		Postal code								
Part I: Details of consignment presented	1.8.	Country ISO code I.9. Region of or of origin	igin Code	I.10. Country of destination	ISO co	ode I.11.	Region of destination	Code		
etail	140	Place Collete								
ă	1.12.	Place of origin		I.13. Place of de			_			
ar		Semen centre		Seme	n centre 🗌		Holding 🔲			
۵		Name Approval n Address	umber	Name Address			Approval numbe	r		
		Postal code		Postal code	€					
	1.14.			I.15.						
	116	Means of transport		1.17.						
		·								
	Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐									
	140	Identification								
	1.18.	Description of commodity		I.19. Commodity code (HS code) 05 11 99 85						
						I.20. Quanti	ty			
	1.21.	Temperature of products Ambient ☐ Chilled ☐		Frozen		I.22. Numbe	r of packages			
	1.23.	Seal/Container No				I.24. Type o	f packaging			
	1.25.	Commodities certified for:								
		Artificial reproduction								
	1.26.	Transit through third country		I.27. Transit thro	ugh Member	States				
		Third country ISO code		Member St	ate	IS	O code			
		Exit point Code		Member St			O code			
		Entry point BIP No		Member St	ate	IS	O code			
	1.28.	Export		1.29.						
		Third country ISO code Exit point Code								
	1.30.									
	1.31.	Identification of the commodities								
			r identity	Date of collection	Ap _l	proval numbe f the centre	r Quanti	ity		

	EUROPEAN UNION Ovine and caprine semen — Par								
	II. H	lealth info	rmation	II.a. Certificate reference No	II.b.				
	I, the undersigned official veterinarian, hereby certify that the semen described above:								
	II.1. was collected, processed and stored in a semen collection centre (2) approved and supervised by the competent authority in accordance with Chapter I(I) and Chapter I(II) of Annex D to Directive 92/65/EEC;								
		II.2.	comes from the donor animals which meet the req	uirements of Chapter II(II) of Annex D	to Directive 92/65/EEC;				
el ullicado		II.3.	was collected, processed, stored and transported u III of Annex D to Directive 92/65/EEC;	nder conditions which comply with the	requirements of Chapters II(II) and				
:	(1) either	[11.4.	meets the requirements of Chapter A(I) of Annex V	'III to Regulation (EC) No 999/2001;]					
ב	(1) or [II.4. meets the requirements of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and is destined for a Member State which benefits, for all or part of its territory, from the provisions laid down in point (b) or (c) of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and the donor animals comply regarding scrapie with the guarantees provided for by the programmes referred to in that point and with the guarantees (3) requested by the Member State of destination.]								
_	Notes								
	Part I:								
	Box I.12:	Place of	origin shall correspond to the semen collection centre	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:	Identificat	tion of container and seal number shall be indicated.						
	Box I.31:	Donor ide	entity shall correspond to the official identification of t	he animal.					
		Date of c	collection shall be indicated in the following format: do	d/mm/yyyy.					
		Approval collected	number of the centre shall correspond to the approva	I number of the semen centre indicate	d in Box I.12 where the semen was				
	Part II:								
	(1) Delete	as appro	priate.						
			semen collection centres listed in accordance with A eu/food/animal/approved_establishments/establishmer		E/EEC on the Commission website:				
	(³) Additio	onal guara	antees as laid down in Article 2 of Regulation (EC) N	o 546/2006 (OJ L 94, 1.4.2006, p. 28).				
	Official ve	terinarian	or official inspector (*)						
	Name	(in capita	ıl letters):	Qı	ualification and title:				
	Local	veterinary	unit:	LV	/U No:				
	Date:			Si	gnature:				
	Stamp	:							
	(*) The colour of the stamp and signature must be different from that of the other particulars in the certificate.								

PART C

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

EUR	OPEA	N UNION	Intra trade certificate			
	l.1.	3	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			
consignment presented	1.5.	Consignee Name	I.6. No(s) of related original certificates	No(s) of accompanying documents		
nment		Address Postal code	1.7.			
₽	1.8.	Country ISO code I.9. Region of origin Code of origin	I.10. Country of destination	I.11. Region of Code destination		
Part I: Details	l.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.			
	I.16.	Means of transport	1.17.			
		Aeroplane Ship Railway wagon				
		Road vehicle Other				
		Identification				
	I.18.	Description of commodity	I.19. Commodity coc 05	de (HS code) 11 99 85		
			1.20.	I.20. Quantity		
	I.21.	Temperature of products Ambient ☐ Chilled ☐	Frozen 🗆	Number of packages		
	1.23.	Seal/Container No	1.24.	Type of packaging		
	1.25.	Commodities certified for:	1			
		Artificial reproduction				
	1.26.	Transit through third country	I.27. Transit through Member States			
		Third country ISO code	Member State Member State	ISO code ISO code		
		Exit point 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 r of the ce			

ı	EUROPEA	N UNION		Ovi	ine and caprine semen — Part C
	II. I	Health informa	ation	II.a. Certificate reference No	II.b.
	I, the und	dersigned offic	cial veterinarian, hereby certify that the semen des	scribed above:	
	(¹) either	[II.1.	was collected, processed and stored for a minir semen collection centre (2) situated in the Me accordance with Chapter I(I)(1) and Chapter I(II) moved to the semen storage centre detailed in P animal health and veterinary certification at least	ember State of origin of the semen (1) of Annex D to Directive 92/65/EEC Part I.12 situated in the same Member	and operated and supervised in C, and from where the semen was
ation		(1) either	[Part A of Annex III to Decision 2010/470/EU;]		
Part II: Certification		(1) or	[Part B of Annex III to Decision 2010/470/EU;]		
art II: ((¹) or	[Decision 95/388/EC;]]		
۵	(¹) or	[II.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/Eaccordance with:	ean Union and operated and supervis	ed 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	[II.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	untry or part(s) thereof listed in Annex oter I(I)(1) and Chapter I(II)(1) of Annex	I to Decision 2010/472/EU which is D to Directive 92/65/EEC, and was
		(1) either	[Section A of Part 2 of Annex II to Decision 2010	0/472/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		ated 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		hapter III(I) of Annex D to Directive
	Notes				
	Part I:				
	Box I.6:	panied the s	pond to the serial number of the individual official emen described above from the approved semen of of this/these document(s) or certificate(s), or the o	collection centre of its origin to the desc	ribed above semen storage centre.

- The original of this/these document(s) or certificate(s), or the officially endorsed copy/copies thereof must be attached to this certificate.
- Box I.12: Place of origin shall correspond to the semen storage centre of dispatch of the semen.
- Box I.13: Place of destination shall correspond to the semen collection or storage centre or to the holding of destination of the semen.
- Box I.23: Identification of container and seal number shall be indicated.
- Box I.31: Donor identity shall correspond to the official identification of the animal.

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

Approval number of the centre shall correspond to the approval number of the semen collection centre of the semen origin.

EUROPEAN UNION	Equi	ne ova and embryos — Part C						
II. Health information	II.a. Certificate reference No	II.b.						
Part II:								
(¹) Delete as appropriate.								
(2) Only approved semen collection or storage centres listed in 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/establishmehttp://ec.europa.eu/food/animal/semen_ova/ovine/index_en.htm	http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm http://ec.europa.eu/food/animal/semen_ova/ovine/index_en.htm							
Official veterinarian or official inspector (*)								
Name (in capital letters):	Qual	ification and title:						
Local veterinary unit:	LVU	No:						
Date:	Date: Signature:							
Stamp:								
(*) The colour of the stamp and signature must be different from that of the other pa	articulars in the certificate.							

ANNEX IV

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

PART A

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

EUR	OPE/	AN UNION							In	tra trade ce	ertificate
	l.1.	Consignor				I.2. Certificat	e reference No		I.2.a. Loca	l reference l	No
		Name				I.3. Central competent authority					
		Address				1.3. Central c	competent author	ority			
ted		Postal code				I.4. Local co	mpetent authori	ty			
seu	1.5.	Consignee				I.6.					
pre		Name									
Jen		Address				1.7.					
guu		Postal code									
of consignment presented		Country of origin	ISO code	I.9. Region of origin	Code	I.10. Country destinat	of tion	ISO code	I.11. Re des	gion of stination	Code
sis.											
Part I: Details	1.12.	Place of origin		_		I.13. Place o		_			_
# #		N	Embryo tea				Hold	ing 🔲		nbryo team [
Par		Name Address		Approval number		Name			Ар	proval numb	oer
						Address					
		Postal code				Postal o	code				
	1.14					I.15.					_
	I.16.	Means of transport				1.17.					
		Aeroplane	Ship 🔲	Railway wagon				_			
		Road vehicle	Other \square								
		Identification									
	I.18.	Description of commo	dity				I.19. Commod				
									05 11 99 8). Quantity		
								1.20	7. Quantity		
	1.21.	Temperature of produc	cts					1.22	2. Number	of packages	;
		Ambient		Chilled		Frozen					
	1.23.	Seal/Container No						1.24	I. Type of	packaging	
	1.25.	Commodities certified	for:								
		Artificial reproduction [
	1.26.	Transit through third co	ountry			I.27. Transit t	hrough Member	r States			
		Third country	·	ISO code		Member	-			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 cor	mmodities								
		Species (Scientific name)	Breed	Category	Donor io		Date of collection		al number e team	Qu	antity

EUROPEAN UNION

Ovine and caprine ova/embryos — Part A

				•		na capinio crajonisi yoo ir are A
		II.	Health in	formation	II.a. Certificate reference No	II.b.
Γ		I, the und	dersigned	official veterinarian, hereby certify that:		
		(¹) either	[II.1.	the <i>in vivo</i> derived embryos/ <i>in vivo</i> derived ova (1) collection team (2) approved and supervised in acco		
 -	ation	(¹) or	[II.1.	the <i>in vitro</i> produced embryos/micromanipulated emembryo production team (²) approved and supervise 92/65/EEC;]		
	92/65/EEC;] (1) either [II.2. the in vivo derived embryos described above meet the requirements of Chapter III(II)(1) of Annex D to Directive 92					
;	Part II: ((¹) or	[II.2.	the in vivo derived ova described above meet the	requirements of Chapter III(II)(2) of	Annex D to Directive 92/65/EEC;]
	۵	(¹) or	[II.2.	the in vitro produced embryos described above meet	the requirements of Chapter III(II)(3)	of Annex D to Directive 92/65/EEC;]
		(¹) or	[II.2.	the micromanipulated embryos described above 92/65/EEC;]	meet the requirements of Chapter	III(II)(4) of Annex D to Directive
			II.3.	the ova or embryos described above:		
		(¹) either	[II.3.1.	meet the requirements of Chapter A(I) of Annex VIII	to Regulation (EC) No 999/2001;]	
		(¹) or	[II.3.1.	meet the requirements of Chapter A(I) of Annex VIII which benefits, for all or part of its territory, from the Regulation (EC) No 999/2001 and the donor anim programmes referred to in that point and with the g	e provisions laid down in point (b) or als comply regarding scrapie with t	(c) of Chapter A(I) of Annex VIII to he guarantees provided for by the
			II.3.2.	come from female donors of the ovine/caprine spe Directive 92/65/EEC;	ecies (1) which meet the requiremen	ts of Chapter IV(3) of Annex D to
		(¹) either	[II.4.	the embryos described above were conceived as a r collected, produced, stored and transported under co of Annex D to Directive 92/65/EEC;]		
		(¹) or	[II.4.	the embryos described above were conceived as a re III(II)(2) of Annex D to Directive 92/65/EEC with a conditions which comply with the requirements of C	semen which was collected, proces	sed, stored and transported under
		(¹) or	[II.4.	the ova have not been in contact with semen of the	ovine and caprine species;]	
			II.5.	the ova or embryos described above were sent to chapter III(II) of Annex D to Directive 92/65/EEC and		

Notes

Part I:

- Box I.12: Place of origin shall correspond to the embryo collection team or embryo production team of embryos collection/production.
- Box I.13: Place of destination shall correspond to the embryo collection team, embryo production team or to the holding of ova/embryos destination.
- Box I.23: Identification of container and seal number shall be indicated.
- Box I.31: Category: specify if: in vivo derived embryos, in vivo derived ova, in vitro produced embryos or micromanipulated embryos.

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

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

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

					M

Ovine and caprine ova/embryos — Part A

II. Health information	II.a. Certificate reference No	II.b.					
Part II:							
 Delete as appropriate. Only approved embryo collection or production teams listed in accordance with Article 11(4) of Council Directive 92/65/EEC on the Commission website: http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm Additional guarantees as laid down in Article 2 of Regulation (EC) No 546/2006 (OJ L 94, 1.4.2006, p. 28). 							
Official veterinarian or official inspector (*)							
Name (in capital letters):		Qualification and title:					
Local veterinary unit:		LVU No:					
Date:		Signature:					
Stamp:							
(*) The colour of the stamp and signature must be different from that of the other particulars in the certificate.							

PART 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

OPE/	AN UNION			Intra trade certificate
	Consignor		I.2. Certificate reference No	I.2.a. Local reference No
	Name Address		I.3. Central competent authority	
<i>'</i>	Address		. ,	
	Postal code		I.4. Local competent authority	
l.5. (Consignee		1.6.	
	Name			
	Address		1.7.	
	Postal code			
	Country of ISO code origin	I.9. Region of origin Code	I.10. Country of ISO destination code	I.11. Region of Code destination
1.12	Place of origin		I.13. Place of destination	
	Embryo tear	n 🗆	Holding	Embryo team
	Name	Approval number	Name	Approval number
	Address		Address	
	Postal code		Postal code	
1.14			I.15.	
l.16.	. Means of transport		l.17.	
	Aeroplane Ship	Railway wagon 🗌		
	Road vehicle Other			
	Identification			
l.18.	. Description of commodity		I.19. Commodity code	
				05 11 99 85 20. Quantity
			1.	20. Quantity
l.21.	. Temperature of products			22. Number of packages
	Ambient	Chilled	Frozen	
1.23.	. Seal/Container No		1.	24. Type of packaging
1.25.	. Commodities certified for:			
	Artificial reproduction			
1.26.	. Transit through third country		I.27. Transit through Member States	
	Third country	ISO code	Member State	ISO code
ĺ	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 id		oval number Quantity
	(Scientific name)		collection of	tne team
		Successive Bonories		the team

Part II: Certification

EN

EUROPEAN UNION Ovine and caprine ova/embryos - Part B Health information II.a. Certificate reference No I, the undersigned official veterinarian, hereby certify that the ova/embryos (1) described above: II.1. were collected, processed and stored under conditions which meet the requirements of Directive 92/65/EEC; 11.2. come from female donors of the ovine/caprine species (1) which meet the requirements of Chapter IV of Annex D to Directive (1) either [II.3. meet the requirements of Chapter III of Annex D to Directive 92/65/EEC and of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001.] (1) or [II.3. meet the requirements of Chapter III of Annex D to Directive 92/65/EEC and of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and are destined for a Member State which benefits, for all or part of its territory, from the provisions laid down in point (b) or (c) of Chapter A(l) of Annex VIII to Regulation (EC) No 999/2001 and the donor animals comply regarding scrapie with the guarantees provided for by the programmes referred to in that point and with the guarantees (2) requested by the Member State of destination.] (1) either [II.4. in the case of embryos, the semen used for fertilisation meets the requirements of Directive 92/65/EEC and the requirements of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001.] (1) or [11.4. in the case of embryos, the semen used for fertilisation meets the requirements of Directive 92/65/EEC and the requirements of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and is destined for a Member State which benefits, for all or part of its territory, from the provisions laid down in point (b) or (c) of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001 and the donor animals comply regarding scrapie with the guarantees provided for by the programmes referred to in that point and with the guarantees (2) requested by the Member State of destination.] Notes Part I: Box I.12: Place of origin shall correspond to the embryo collection team of ova/embryos collection. Place of destination shall correspond to the embryo collection team, embryo production team or to the holding of ova/embryos Box I.13: destination. Box I.23: Identification of container and seal number shall be indicated. Box I.31: Category: specify if: in vivo derived embryos, in vivo derived ova, in vitro produced embryos or micromanipulated embryos. Donor identity shall correspond to the official identification of the animal. Date of collection shall be indicated in the following format: dd/mm/yyyy. Approval number of the team shall correspond to the embryo collection team of ova/embryos collection indicated in Box I.12. Part II: Delete as appropriate. (2) Additional guarantees as laid down in Article 2 of Regulation (EC) No 546/2006 (OJ L 94, 1.4.2006, p. 28).

Official veterinarian or official inspector (*)

Name (in capital letters):

Qualification and title:

Local veterinary unit:

LVU No:

Date:

Signature:

Stamp:

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

ANNEX V

Model health certificates for trade within the Union in 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

EUF	ROPEAN UNION Intra trade certific						
	1.1.	Consignor	I.2. Certificate reference No I.2.a. Local reference No				
		Name					
		Address	I.3. Central competent authority				
consignment presented		Postal code	I.4. Local competent authority				
rese	1.5.	Consignee	1.6.				
t p		Name					
mer		Address	1.7.				
ign		Postal code					
Suo	1.8.	Country of ISO code I.9. Region of origin Code	I.10. Country of ISO I.11. Region of Code				
₽		origin	destination code destination				
Si S	L						
Part I: Details	1.12	. Place of origin Embryo team □	I.13. Place of destination Holding ☐ Embryo team ☐				
=		Name Approval number	Name Approval number				
Par		Address	Address				
		Postal code	Postal code				
	1.1	4.	1.15.				
	1.16	. Means of transport	1.17.				
		Aeroplane ☐ Ship ☐ Railway wagon ☐					
		Road vehicle Other O					
		Identification					
	1.18	. Description of commodity	I.19. Commodity code (HS code)				
			05 11 99 85				
			I.20. Quantity				
	1.21	. Temperature of products	I.22. Number of packages				
		Ambient Chilled From From From From From From From From	ozen 🗆				
	1.23	. Seal/Container No	I.24. Type of packaging				
	1.25	. Commodities certified for:	I				
		Artificial reproduction					
	100	<u> </u>	LOZ. Transit through Marchae Chates				
	1.26	. Transit through third country SO code	I.27. Transit through Member States Member State ISO code				
		Exit point Code	Member State ISO code Member State ISO code				
		Entry point BIP No					
	1.28	. Export	1.29.				
		Third country ISO code					
		Exit point Code					
	1.30						
	1.31	. Identification of the commodities Species Breed Category Donor ider (Scientific name)	ntity Date of Approval number Quantity collection of the team				

Part II: Certification

EUROPE/	AN UNION			Porcine ova/embryos — Part A
II.	Health in	formation	II.a. Certificate reference No	II.b.
I, the und	dersigned (official veterinarian, hereby certify that the ova/embry	os (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 in
	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 Ch	apter IV(2) of Annex D to Directive
(¹) either	[II.4.	are in vivo derived embryos which:		
	II.4.1.	were conceived as a result of artificial insemination	with semen meeting the requiremen	nts 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 c 2008/185/EC;]	lestined for a Member State or regior	n thereof listed in Annex I to Decision
	(¹) or	[listed in Annex I to Decision 2008/185/EC and are of Decision 2008/185/EC;]	destined for a Member State or regio	n thereof not listed in Annex I or II to
	(¹) or	[listed in Annex II to Decision 2008/185/EC and are a 2008/185/EC and have been washed with trypsin;]	destined for a Member State or region	n thereof listed in Annex I to Decision
	(¹) or	[listed in Annex II to Decision 2008/185/EC and are c 2008/185/EC;]	lestined for a Member State or regior	n thereof listed in Annex II to Decision
	(¹) or	[not listed in Annex I or II to Decision 2008/185/EC a to Decision 2008/185/EC and have been washed w		or region thereof listed in Annex I or II
	(¹) or	[not listed in Annex I or II to Decision 2008/185/EC a or II to Decision 2008/185/EC;]]	and are destined for a Member State	or region thereof not listed in Annex I
(¹) or	[II.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 c 2008/185/EC;]	lestined for a Member State or regior	n thereof listed in Annex I to Decision
	(¹) or	[listed in Annex I to Decision 2008/185/EC and are of Decision 2008/185/EC;]	destined for a Member State or regio	n thereof not listed in Annex I or II to
	(¹) or	[listed in Annex II to Decision 2008/185/EC and are of 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;]	lestined for a Member State or regior	n thereof listed in Annex II to Decision
	(¹) or	[not listed in Annex I or II to Decision 2008/185/EC a to Decision 2008/185/EC and 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 Annex I

EUROPEAN UNION Porcine ova/embryos — Part A

II.	Health info	rmation	II.a. Certificate reference No	II.b.							
(¹) or	[11.4.	[II.4. are in vivo derived ova which originate from a Member State or region thereof:									
	(¹) either	(i) 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	[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	(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 to Decision 2008/185/EC and which come from donor females complying with the conditions of Article 1 of Decision 2008/185/EC;]										
	(1) or [not listed in Annex I or II to Decision 2008/185/EC and are destined for a Member State or region thereof not listed in Annex or II to Decision 2008/185/EC;]]										
	II.5.	II.5. were sent to the place of loading in a sealed container under conditions complying with point 6 of Chapter III(II) of Annex D to Directive 92/65/EEC and bearing the number detailed in Box I.23.									
Notes											
Part I:											
Box I.12	: place of	place of origin shall correspond to the embryo collection team or embryo production team of ova/embryos collection/production.									
Box I.13	: place of	place of destination shall correspond to the embryo collection team, embryo production team or to the holding of ova/embryos destination.									
Box I.23	: identification of container and seal number shall be indicated.										
Box I.31	: category: specify if: <i>in vivo</i> derived embryos, <i>in vivo</i> derived ova, <i>in vitro</i> produced embryos or micromanipulated embryos. donor identity shall correspond to the official identification of the animal. date of collection shall be indicated in the following format: dd/mm/yyyy.										
	approval	approval number of the team shall correspond to the embryo collection team or embryo production team of ova/embryos collection/production indicated in Box I.12.									
Part II:											
(1) Delete as appropriate. (2) Only approved embryo collection or production teams listed in accordance with Article 11(4) of Council Directive 92/65/EEC on the Commission website: http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm											
Official veterinarian or official inspector (*)											
Na	me (in cap	ital letters):	Qualification and title:								
Lo	cal veterina	ry unit:	LVU No:								
Da Sta	te: amp:		Signature:								
(*) The c	olour of the s	stamp and signature must be different from that of the other p	articulars 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

UROPEAN UNION Intra trade certificate														
		Consignor				1.2.	Certificat	e reference No		I.2.a. Loca	al reference l	No		
		Name					12 Control competent outbouits							
		Address				I.3. Central competent authority								
ıtea		Postal code					I.4. Local Competent Authority							
eser	l.5.	I.5. Consignee				1.6.								
i bi	Name													
ails of co		Address				1.7.								
		Postal code												
		Country of origin	ISO code	I.9. Region of origin	Code	I.10.	Country destinat		ISO code	I.11. Re de	egion of estination	Code		
	1.40	DI () ;				1.40	D.							
3	1.12.	Place of origin	Embryo tear	_m □		1.13.	Place of	f destination Hold	ing 🔲	Fr	mbryo team	П		
		Name	Approval number				riola			proval numb				
-		Address	• •			Name Approval numb								
		Postal code				Postal code								
	1.14													
			1.15.											
	I.16.	Means of transport	Ship 🔲	Railway wagon	_	1.17.								
		Aeroplane 🗌												
		Road vehicle Other Identification												
								1						
	I.18.	I.18. Description of commodity					I.19. Commodity code (HS code) 05 11 99 85							
									I.20. Qu		00			
							100 N 1 1							
	1.21.	Temperature of produc	1.22. Nu 			I.22. Nui	umber of packages							
		Ambient	- Cr	nilled	Fro	ozen L								
	1.23.	Seal/Container No	I.24. Type of packaging					ging						
	1.25.	Commodities certified for:												
		Artificial reproduction												
	1.26.	Transit through third c	ountry			1.27.	Transit t	hrough 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 co	mmodities											
		Species	Breed	Category D	Donor ide	ntity		ate of		val number	C	uantity		
		(Scientific name)					col	llection	of t	he team				

Part II: Certification

EUROPEAN UNION Porcine ova/embryos — Part B 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: II.1.1. were collected, processed and stored under conditions which meet the requirements of Directive 92/65/EEC; II.1.2. come from donor female swine which meet the requirements of Chapter IV of Annex D to Directive 92/65/EEC; meet the requirements of Chapter III of Annex D to Directive 92/65/EEC. II.1.3. (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; 11.2.2. the embryos have been washed with trypsin (2).] (1) or [II.2. In the case of ova, the ova comes from a donor female swine which meets the conditions of Article 1 of Decision 2008/185/EC (2).] Notes Part I: Box I.12: place of origin shall correspond to the embryo collection team of ova/embryos collection. Box I.13: place of destination shall correspond to the embryo collection team, embryo production team or to the holding of ova/embryos destination. Box I.23: identification of container and seal number shall be indicated. Box I.31: category: specify if: in vivo derived embryos, in vivo derived ova, in vitro produced embryos or micromanipulated embryos. donor identity shall correspond to the official identification of the animal. date of collection shall be indicated in the following format: dd/mm/yyyy. approval number of the team shall correspond to the embryo collection team of ova/embryos collection indicated in Box I.12. Part II: (1) Delete as appropriate. (2) This condition applies only to ova and embryos which originate in the Member States or regions thereof not listed in Annexes I and II to Decision 2008/185/EC (OJ L 59, 4.3.2008, p. 19) and destined to the Member States or regions thereof so listed. It shall also apply to movements from Member States or regions thereof listed in Annex 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 (*) 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.