EUROPEAN UNION **Intra trade certificate**

Part I : Details of dispatched consignment

I.1. Consignor
Name
Address
Country

I.2. Certificate reference number

I.2.a.Local reference number::

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Consignee
Name
Address
Country

I.6. No.(s) of related original certificates

No.(s) of accompanying documents

I.7. Dealer
Name
Approval number

I.8.Country of origin ISO code

I.9. Region of origin Code

I.10. Country of destination ISO code

I.11. Region of destination Code

I.12. Place of origin/Place of harvest
Holding ☐ Assembly centre ☐ Dealer's premise ☐
Approved body ☐ Semen centre ☐ Approved aquaculture holding ☐
Embryo team ☐ Establishment ☐ Other ☐
Name
Approval number
Address
Postal code / Region

I.13. Place of destination
Holding ☐ Assembly centre ☐ Dealer's premise ☐
Approved body ☐ Semen centre ☐ Approved aquaculture holding ☐
Embryo team ☐ Establishment ☐ Other ☐
Name
Approval number
Address
Postal code / Region

I.14. Place of loading
Postal code / Region

I.15. Date and time of departure

I.16. Means of transport
Aeroplane ☐ Ship ☐ Railway wagon ☐
Road vehicle ☐ Other ☐
Identification::
Number(s):

I.17. Transporter
Name
Approval number
Address
Postal code / Region Member state

I.21 Temperature of products
Ambient ☐ Chilled ☐ Frozen ☐

I.20. Number/Quantity

I.22. Number of packages

I.23. Identification of container/Seal number

I.25. Animals certified as/products certified for:
Artificial reproduction ☐

I.26. Transit through 3rd country ☐
3rd country ISO code
3rd country ISO code
3rd country ISO code
Exit point Code
Entry point BIP unit no.:

I.27. Transit through Member states ☐
Member state ISO code
Member state ISO code
Member state ISO code

I.28. Export ☐
3rd country ISO code
Exit point Code

I.29. Estimated journey time

I.30. Route plan
Yes ☐ No ☐

I.31. Identification of the animals

Species	Donor identity	Date(s) of collection	Quantity	Type of packaging

Part II: Certification

II. Health information	II.a. Certificate reference number	II.b.Local reference number:

I, the undersigned official veterinarian, hereby certify that:

II.1. The semen collection centre(2), in which the semen described above was collected, processed and stored for trade is approved and supervised by the competent authority in accordance with Chapter I(I)(1) and Chapter I(II)(1) of Annex D to Directive 92/65/EEC;

II.1.1. during the period commencing 30 days prior to the date of first collection of the semen described above until the date the fresh or chilled semen was dispatched or until the 30 days storage period for frozen semen elapsed, the semen collection centre:

II.1.1.1. was situated on the territory or in the case of regionalisation in a part of the territory(1) of a Member State which was not considered to be infected with African horse sickness in accordance with Article 5(2)(a) and(b) of Directive 2009/156/EC(3);

II.1.1.2. fulfilled the conditions for a holding laid down in Article 4(5) of Directive 2009/156/EC(3);

II.1.1.3. contained only equidae which were free of clinical signs of equine viral arteritis and contagious equine metritis;

II.2. Only equidae satisfying the conditions laid down in Articles 4 and 5 or Articles 12 to 16 of Directive 2009/156/EC(3) have been admitted onto the centre.

II.3. The semen described above was collected from donor stallions, which:

II.3.1. have not shown any clinical sign of an infectious or contagious disease at the time of admission onto the centre and on the day the semen was collected;

II.3.2. have been kept for 30 days prior to the date of semen collection in holdings where no equine has shown any clinical sign of equine viral arteritis or contagious equine metritis during that period;

II.3.3. have not been used for natural mating during at least 30 days prior to the date of first semen collection and from the dates of the first sample referred to in points II.3.5.1., II.3.5.2. or II.3.5.3. until the end of the collection period;

II.3.4. have undergone the following tests , which meet at least the requirements of the relevant Chapter of the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals of the OIE, carried out on samples taken in accordance with one of the programmes specified in point II.3.5 in a laboratory recognised by the competent authority:

(1)either [II.3.4.1. an agar-gel immuno-diffusion test (Coggins test) for equine infectious anaemia (EIA) with negative result;]

(1)or [II.3.4.1. an ELISA for equine infectious anaemia (EIA) with negative result;]

and (1)either [II.3.4.2. a serum neutralisation test for equine viral arteritis (EVA) with negative result at a serum dilution of one in four;]

(1)or [II.3.4.2. a virus isolation test for equine viral arteritis (EVA) carried out with negative result on an aliquot of the entire semen of the donor stallion;]

and II.3.4.3. an agent identification test for contagious equine metritis (CEM) carried out on two occasions on samples taken with an interval of seven days by isolation of Taylorella equigenitalis after a cultivation of 7 to 14 days from pre-ejaculatory fluid or a semen sample and from genital swabs taken at least from the penile sheath, urethra and urethral fossa with negative result in each case;

II.3.5. have been subjected with the results specified in II.3.4. in each case to at least one of the test programmes(4) detailed in points II.3.5.1., II.3.5.2. and II.3.5.3. as follows:

II.3.5.1. The donor stallion was continuously resident on the semen collection centre for at least 30 days prior to the date of the first collection and during the period of collection of the semen described above and no equidae on the semen collection centre came into direct contact with equidae of lower health status than the donor stallion.

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

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:

Identification of semen	Test programme	Start date(5)		Date of sampling for health tests(5)				
		Donor residence	Semen collection	EIA II.3.4.1.	EVA II.3.4.2.	CEM II.3.4.3.		
					Blood sample	Semen sample	1. Sample	2. Sample

Part II: Certification

II. Health information	II.a. Certificate reference number	II.b.Local reference number:

(1)either [II.4. No antibiotics were added to the semen;]

r

(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.

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

Contagious equine metritis (CEM) testing first occasion first sample

Part II: Certification

II. Health information	II.a. Certificate reference number	II.b.Local reference number:

CEM-11

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.

Identification of semen	Test programme	Start date(5)		Date of sampling for health tests(5)				
		Donor residence	Semen collection	EIA II.3.4.1	EVA II.3.4.2		CEM II.3.4.3	
					Blood sample	Semen sample	1. Sample	2. Sample
A	B	C	D	EIA-1	EVA-B1	EVA-S1	CEM-11	CEM-12
				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.

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

Official veterinarian or official inspector

Name (in Capital): Qualification and title:

Local Veterinary Unit: LVU N°:

Date: Signature:

Stamp

Part III: Control

III.1. Date of the inspection

III.2. Certificate Reference Number::

III.3. Documentary Check:: No Yes

EU Standard Satisfactory Not satisfactory

Additional guarantees Satisfactory Not satisfactory

National requirements Satisfactory Not satisfactory

III.4. Identity Check:: No Yes

Satisfactory Not satisfactory

III.5. Physical Check:: No Total animals checked

Satisfactory Not satisfactory

III.6. Laboratory Tests:: No Yes

Date:

Tested for::

Random Suspicion

Results:: Pending Satisfactory Not satisfactory

III.7. Welfare check No Yes

Satisfactory Not satisfactory

III.8.Infringement of welfare regulation::

III.8.1.Transporter authorisation invalid

III.8.2.Non-compliance of the means of transport

III.8.3.Stocking density exceeded Average space

III.8.4.Travel times exceeded

III.8.5.Watering and feeding not fulfilled

III.8.6.Mishandling or negligence to the animals

III.8.7.Supplementary measures for the journeys of long duration

III.8.8.Certificate of proficiency of the driver

III.8.9.Data registered in the log book

III.8.10.Other

III.9. Infringement of health legislation

III.9.1. Absence/Invalid certificate

III.9.2. Mis-match with documents

III.9.3. Non authorised country

III.9.4. Non approved region/ zone

III.9.5. Prohibited species

III.9.6. Absence of additional guarantee

III.9.7. Non approved holding

III.9.8. Diseased or suspect animals

III.9.9. Unsatisfactory tests

III.9.10. Absence or non legal identification

III.9.11.National requirements not fulfilled

III.9.12. Address of destination invalid

III.9.13. Other

III.10. Impact of the transport on animals

Number of dead animals:: Estimation:

Number of unfit animals :: Estimation:

Number of birth or abortion:

III.11. Corrective action

III.11.1. Delayed departure

III.11.2. Transfer procedure

III.11.3. Quarantine

III.11.4. Humane killing/Euthanasia

III.11.5. Destruction of carcasses/products

III.II.6. Return of consignment

III.11.7. Treatment of products

III.11.8.7. Use of products for other purpose

Identification:

III.12. Follow-up of quarantine

III.12.1.Humanely killing/Euthanasia

III.12.2.Release

III.13. Place of inspection

Establishment Holding Assembly centre

Dealer's premise Approved body Semen centre

Port Airport Exit point

Enroute Other

III.14. Official veterinarian or official inspector

Local Veterinary Unit LVU N°

Name (in Capital):

Qualification and title

Date: Signature:

PLANNING

1.1. ORGANISER name and address (a) (b)	1.2. Name of the person in charge of the journey
	1.3. Telephone / Fax

2. TOTAL EXPECTED DURATION (hours / days)	
3.1. Place and country of DEPARTURE	4.1. Place and country of DESTINATION

3.2. Date	3.3. Time	4.2. Date	4.3. Time

5.1. Species	5.2. Number of animals	5.3. Veterinary certificate(s) number(s)

5.4. Estimated total weight of the consignment (in kg)	5.5. Total space foreseen for the consignment (in m²)

6. LIST OF FORESEEN RESTING, TRANSFER OR EXIT POINTS

6.1. Name of the places where animals are to be rested, or transferred (including exit points)	6.2. Arrival		6.3. Length (in hours)	6.4. Transporter name and authorisation N° (if different from the organiser)	6.5 identification
	Date	Time			

7. I, the organiser, hereby declare that I am responsible for the organisation of the above-mentioned journey and I have made suitable arrangements to safeguard the welfare of the animals throughout the journey in accordance with the provisions of Council Regulation 1/2005
8. Signature of the organiser

(a) Organiser: see definition laid down in Article 2(q) of Council Regulation 1/2005

(b) If the organiser is a transporter the authorisation number shall be specified