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 for/products certified for:

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

I.27. Transit through Member states ☐

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

Part II: Certification

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

I, the undersigned official veterinarian, hereby certify that:

(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 in vitro 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 in vivo derived embryos described above meet the requirements of Chapter III(II)(1) of Annex D to Directive 92/65/EEC;]

(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;]

(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 meet the requirements of Chapter III(II)(4) of Annex D to Directive 92/65/EEC;]

II.3. the ova or embryos described above come from donor 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 equidae satisfying the conditions laid down in Articles 4 and 5 or Articles 12 to 16 of Directive 2009/156/EC have been admitted;

II.3.2. meet the additional requirements of Chapter IV(4) of Annex D to Directive 92/65/EEC;

II.3.3. have not been used for natural breeding during at least 30 days prior to the date of collection of ova or embryos and between the date of the first sample referred to in points II.3.4. and II.3.5. and the date of the collection of ova and embryos;

II.3.4. have been subjected with negative result to an agar-gel immuno-diffusion test (Coggins test) or an ELISA for equine infectious anaemia carried out on a blood samples taken on (3), being during the past 30 days prior to the date of the first collection of ova or embryos and the last test was carried out on a sample of blood taken on (3); being not more than 90 days before the ova and embryos were collected;

II.3.5. have been subjected to an agent identification test for contagious equine metritis by isolation of Taylorella equigenitalis after a cultivation of 7 to 14 days carried out with negative results in each case on samples taken during the past 30 days prior to the date of the first collection of ova or embryos from mucosal surfaces of the clitoral fossa and clitoral sinuses on two consecutives oestrus periods on (3) and on (3), and on an additional culture specimen taken during one of the oestrus periods from the endometrial cervix on (3);

(1)either [II.4. the embryos described above were conceived as a result of artificial insemination of the donor mares with semen which was collected, processed, stored and transported under conditions which comply with the requirements of Chapters I(I), II(I) and III(I) of Annex D to Directive 92/65/EEC;]

(1) or [II.4. the embryos described above were conceived as a result of in vitro fertilisation of ova complying with the conditions in point 2 of Chapter III(II) of Annex D to Directive 92/65/EEC with semen which was collected, processed, stored and transported under conditions which comply with the requirements of Chapters I(I), II(I) and III(I) of Annex D to Directive 92/65/EEC;]

(1) or [II.4. the ova have not been in contact with semen of the equine species;]

II.5. the ova or embryos described above were sent to the place of loading in a sealed container in accordance with point 6 of Chapter III(II) of Annex D to Directive 92/65/EEC and bearing the number detailed in Box I.23.

Notes

Part I:

Box I.12.: place of origin shall correspond to the embryo collection team or embryo production team of ova/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.

Part II:

(1) Delete as appropriate.

(2) Only approved embryo collection or production teams listed in accordance with Article 11(4) of Council Directive 92/65/EEC on the Commission website:

http://ec.europa.eu/food/animal/approved_establishments/establishments_vet_field_en.htm.

(3) Insert date.

(4) OJ L 192, 23.7.2010, p. 1.

· 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