

Veterinary certificate to EU

Part I : Details of consignment presented

I.1. Consignor Name Address Postal code / Region Country	I.2. Certificate reference number	I.2.a. TRACES reference number :
	I.3. Central Competent Authority	
	I.4. Local Competent Authority	
	I.6 Person responsible for load in EU	
I.5. Consignee Name Address Postal code / Region Country		
I.7. Country of origin, ISO code	I.8. Region of origin, Code	I.9. Country of destination ISO code
I.10. Region of destination Code		I.12. Place of destination
I.11. Place of origin Holding <input type="checkbox"/> Semen centre <input type="checkbox"/> Establishment <input type="checkbox"/> Name Approval number Address Name Approval number Address Name Approval number Address		
I.13 Place of loading Address Approval number		I.14 Date of departure
I.15. Means of transport Aeroplane <input type="checkbox"/> Ship <input type="checkbox"/> Railway wagon <input type="checkbox"/> Road vehicle <input type="checkbox"/> Other <input type="checkbox"/> Identification::: Document:		
I.16. Entry BIP in EU Name BIP unit no.:		I.17. No.(s) of CITES
I.18. Description of commodity		I.19. Commodity code (HS code)
I.21 Temperature of products		I.20. Quantity
I.23. Identification of container/Seal number		I.22. Number of packages
I.25. Commodity certified as: Breeding <input type="checkbox"/> Fattening <input type="checkbox"/> Slaughter <input type="checkbox"/> Approved bodies <input type="checkbox"/> Artificial reproduction <input type="checkbox"/> Quarantine <input type="checkbox"/> Game restocking <input type="checkbox"/> Registered horses <input type="checkbox"/> Pets <input type="checkbox"/> Circus <input type="checkbox"/> Relaying <input type="checkbox"/> Other <input type="checkbox"/> Humane consumption <input type="checkbox"/> Animal feedingstuff <input type="checkbox"/> Further process <input type="checkbox"/> Pharmaceutical use <input type="checkbox"/> Technical use <input type="checkbox"/>		I.24. Type of packaging
I.26. For transit to 3rd Country by EU		I.27. For import or admission into EU Definitive import <input type="checkbox"/> Horses Re-entry <input type="checkbox"/> Temporary admission horses <input type="checkbox"/>
I.28. Identification of the commodity Species (scientific name) Breed/category Identification marks Date of collection Quantity Collection centre approval number Donor identity		

2006/168 Embryos of domestic animals of the bovine species for imports collected or produced before 1st January 2006

Part II: Certification

II. Health information	II.a. Certificat reference number	II.b. TRACES reference number
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11.1, the undersigned official veterinarian of the Government of (insert name of exporting country) certify that:

11.1. the embryo collection (1) / production (1) team identified above:

- is approved in accordance with Chapter I of Annex A to Directive 89/556/EEC,
- carried out the collection, processing, production(1) and storing and transport of the embryos described above in accordance with Chapter II of Annex A to Directive 89/556/EEC,
- is subjected at least twice per year to inspection by an official veterinarian.

11.2. The embryos to be exported were collected(1) or produced(1) in the exporting country, which according to official findings:

11.2.1. has been free from rinderpest during 12 months immediately prior to the collection(1) or production (1) of the embryos;

11.2.2. either has been free from foot-and-mouth disease during the 12 months immediately prior to collection(1) or production (1) of the embryos and has not practiced vaccination against foot-and-mouth disease during this period (1) , or

11.2.2.2 has not been free from foot-and-mouth disease for the 12 months immediately prior to the collection(1) or production (1) of the embryos and/or has practised vaccination against foot-and-mouth disease during this period, and

- the embryos have been stored in approved conditions for a minimum period of 30 days immediately after collection, and
- the donor females and the donors of ovaries, oocytes and other tissues used in the production of embryos come from a holding in which no animal has shown clinical signs of foot-and-mouth disease nor was vaccinated against foot-and-mouth disease during the 30 days prior to collection (1);

11.2.3. either has been free from bluetongue and epizootic haemorrhagic disease (EHD) for the 12 months immediately prior to collection(1) or production (1) of the embryos to be exported and has not practiced vaccination against these diseases during this period (1) , or

11.2.3.2 has not been free from bluetongue and epizootic haemorrhagic disease (EHD) for the 12 months immediately prior to collection(1) or production (1) of the embryos to be exported and/or has practised vaccination against these diseases during this period, and

- the embryos have been stored in approved conditions for a minimum period of 30 days immediately after collection, and
- the donor females and the donors of ovaries, oocytes and other tissues used in the production of embryos were subjected to an agar gel immuno diffusion test and a serum neutralisation test for the detection of antibodies against the epizootic haemorrhagic disease virus carried out with negative results on a blood sample taken not less than 21 days following collection (1);

11.3.1. the premises on which the embryos to be exported or the ovaries, oocytes and other tissues used in the production of embryos to be exported were collected and processed were at the time of collection situated in the centre of an area of 20 km diameter in which according to official findings there had been no incidence of foot-and-mouth disease, bluetongue, epizootic haemorrhagic disease, vesicular stomatitis , Rift Valley fever or contagious bovine pleuropneumonia for 30 days immediately prior to collection and in the case of embryos certified under 11.2.2.2 and 11.2.3.2 for 30 days after collection;

11.3.2. between the time of collection or production of the embryos to be exported and their dispatch, they were stored continuously in approved premises which were situated in the centre of an area of 20 km in diameter in which according to official findings there was no incidence of foot-and-mouth disease, vesicular stomatitis or Rift Valley fever;

11.4. the donor females and the donors of ovaries, oocytes and other tissues used in the production of embryos:

11.4.1. were located during the 30 days immediately prior to collection of the embryos to be exported in premises situated in the centre of an area of 20 km in diameter in which according to official findings there was during this period no incidence of foot-and-mouth disease, bluetongue, epizootic haemorrhagic disease, vesicular stomatitis, Rift Valley fever or contagious bovine pleuropneumonia;

11.4.2. showed no clinical sign of disease on the day of collection;

11.4.3. have spent the six months immediately prior to collection in the territory of the exporting country in a maximum of two herds:

- which, according to official findings, have been free from tuberculosis,
- which, according to official findings, have been free from brucellosis,
- which have been free from enzootic bovine leukosis or in which no bovine animal has shown clinical signs of enzootic bovine leukosis during the previous three years,
- in which no bovine animal has shown clinical signs of infectious bovine rhinotracheitis/infectious pustular vulvo-vaginitis during the previous 12 months.

11.5. The embryos comply with the following additional guarantees (3):

11.5.1. either the embryos to be exported were collected(1) or produced(1) in the exporting country, which according to official findings is free of Akabane disease (1) , or

11.5.2. the embryos to be exported were collected(1) or produced(1) in the exporting country, which according to official findings is not free of Akabane disease (1), and

- the embryos have been stored in approved conditions for a minimum period of 30 days immediately after collection, and
- the donor females and the donors of ovaries, oocytes and other tissues used in the production of embryos were subjected to a serum neutralisation test for Akabane disease carried out with negative results on a blood sample taken not less than 21 days following collection (1).

11.6. The embryos to be exported were conceived as a result of artificial insemination or in vitro fertilisation with semen from a donor sire standing at a semen collection centre approved by the competent authority for the collection, processing and storage of semen or with semen imported from the European Community.

Note for guidance:

- (1) Delete as appropriate.
- (2) Corresponding to the identification of the donor cows and date of collection.
- (3) See the remarks for the exporting country concerned in Annex I to Decision [insert reference to present Decision].
- (4) The signature and the stamp must be in a colour different to that of printing.

Note: This certificate must:

- (a) be drawn up in at least one of the official languages of the Member State of destination and the Member State where the embryos will enter Community territory;
- (b) be made out to a single consignee;
- (c) accompany the embryos in the original;
- (d) not to be used after the date indicated in Article 4 of Decision [insert reference to present Decision].

Information:

In accordance with Article 3 (a) of Council Directive 89/556/EEC, embryos imported under the conditions laid down in this certificate are not eligible for intra-Community trade.

2006/168 Embryos of domestic animals of the bovine species for imports collected or produced before 1st January 2006

Part II: Certification	II. Health information	II.a. Certificat reference number	II.b. TRACES reference number
Official veterinarian or official inspector			
Name (in Capital):		Qualification and title:	
Local Veterinary Unit:		LVU N°:	
Date:		Signature:	
Stamp			