EUROPEAN UNION **Export Certificate**

Part I : Details of dispatched consignment

I.1. Consignor
Name
Address
Country

I.2. Certificate reference number

I.2.a.TRACES 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 Country of origin ISO code

I.8. Region of origin

I.9. Country of destination ISO code

I.10. Region of destination

I.11 Place of origin

I.12. Place of destination

I.13. Place of loading

I.14. Date and time of departure

I.15. Means of transport
Aeroplane ☐ Ship ☐ Railway wagon ☐
Road vehicle ☐ Other ☐
Identification::
Number(s):

I.16. Entry Point

I.17. CITES

I.18 Temperature of products

I.19. Number/Quantity

I.20. Total number of packages

I.21. Seal/Container number

I.22. Commodities certified for :
Artificial reproduction ☐

I.23. Transit through 3rd country

I.24. For Export ☐

I.25. Identification of the commodities

Species	Breed	Donor identity	Date(s) of collection	Approval number of the team	Quantity

Part II: Certification

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

I, the undersigned, official veterinarian of the (Member State of the EU) certify that:

II.1. The animal products herein described, comply with the relevant European Union animal health standards and requirements which have been recognised as equivalent to the New Zealand standards and requirements as prescribed in Council Decision 97/132/EC, as last amended, specifically, in accordance with Council Directive 89/556/EEC;

II.2. The animal products are eligible for intra-Union trade without restriction;

II.3. To the best of my knowledge and as far as I can ascertain, the donors have never recorded a positive test for Q fever.

The donors were subjected to a complement fixation test (CFT) (negative being no fixation of complement at a dilution of 1:10 or higher) or ELISA test for Q fever, on a sample collected between 21 and 120 days after each embryo collection period (a period of 60 days or less) for export to New Zealand, with negative results.

(1)either [II.4. The donor animal was subjected to an antigen detection ELISA or virus isolation test for BVDV, with a negative result, within 30 days prior to entry into the embryo collection centre and has been on the embryo collection centre for more than 6 months prior to embryo collection for this consignment and has remained isolated from other animals that have not been tested negative.]

(1)or [II.4. The donor animal has had either a pooled sample of non-viable oocytes/embryos and washing fluid (as per the OIE Code appendix for in vivo derived embryos) or an embryo, from the first embryo collection for this consignment subjected to either virus isolation or PCR for BVDV with negative results.]

Notes

Part I:

Box I.11.: Place of origin shall correspond to the embryo collection team listed in accordance with Article 8(2) of Directive 89/556/EEC on the Commission website: http://ec.europa.eu/food/animal/semen_ova/bovine/ova_embryos_en.htm.

Box I.20.: Number of packages shall correspond to the number of containers.

Box I.21.: Identification of container and seal number shall be indicated.

Box I.25.: Species: indicate “Bos taurus”, “Bison bison" or "Bubalus bubalis” as appropriate.

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 approval number of the embryo collection team by which the embryos were collected,.

Part II:

· The signature and the stamp must be in a different colour to that of the printing.

Official veterinarian or official inspector

Name (in Capital):

Qualification and title:

Local Veterinary Unit:

LVU N°:

Date:

Signature:

Stamp