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 centre	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 88/407/EEC;

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

II.3. The bovine semen complies with provisions of the Bluetongue Chapter of the OIE Code;

II.4. 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 semen collection period (a period of 60 days or less) for export to New Zealand, with negative results.

Notes

Part I:

Box I.11.: Place of origin shall correspond to the approved semen collection centre or semen storage centre listed in accordance with Article 9(2) of Directive 88/407/EEC on the Commission website:

http://ec.europa.eu/food/animal/semen_ova/bovine/index_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 centre shall correspond to the approval number of the semen collection centre in which the semen was 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