2. Embryos (except embryos subject to penetration of the zona pellucida)
Health Certificate

New Zealand — Veterinary certificate to the EU

Part I : Details of dispatched consignment

I.1. Consignor Name Address Country		I.2. Certificate reference No	I.2.a. TRACES reference number:
		I.3. Central competent authority	
		I.4. Local competent authority	
I.5. Consignee Name Address Country		I.6. Person responsible for the consignment in EU	
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 Name / Approval number Address		I.12. Place of destination Name / Approval number Address Postal code	
I.13. Place of loading Name Postal code/ Region		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 BIP in EU Name / BIP unit no. I.17.	
I.18. Temperature of product Frozen ☐		I.19. Total Gross Weight	I.20. Total number of packages
I.21. Seal/Container numbers			
I.22. Commodities certified for: Artificial reproduction ☐			
I.23. For transit through EU to 3rd Country ☐ 3rd country / ISO code		I.24. For import or temporary admission into EU ☐	
I.25. Identification of the commodities Custom code and title: Species (scientific name) / Breed / Breed/category (semen) / Donor identity / Date of collection Approval number of the team / Quantity			

NEW-ZEALAND **2. Embryos (except embryos subject to penetration of the zona pellucida)**

Part II: Certification

II. Health attestation	II.a. Certificate reference No	II.b.

The live animal(s) or animal product(s) herein described, complies/y with the relevant New Zealand standards and requirements which have been recognised as equivalent to the European Union standards and requirements as prescribed in the European Union/New Zealand Agreement on sanitary measures (Council Decision 97/132/EC).

(1)[**III. Additional health attestation**

Either

(1)[III.1. The live animal(s) or animal product(s) herein described, complies/y with the additional conditions laid down in Chapter 29.B. of Section 5 of Annex V to Decision 97/132/EC in the event of the occurrence of a specific disease:

(3)[III.1.1. The in vivo derived embryos herein described were derived from donors that:

III.1.1.1. Were free of clinical signs of FMD, at the time of collection; and from which the embryos were conceived by artificial insemination using semen collected, processed and stored in semen collection centres approved by the competent authority in conformity with OIE standards. In addition the embryos have been collected, processed and stored in accordance with standards laid down by the competent authority; and

III.1.1.2. The donor animals from which the embryos were collected originate from a herd(s) that was/were not located within a protection or surveillance zone. Embryos collected within the protection and surveillance zones have been clearly identified and detained under official supervision.]

(3)[III.1.2. The in vivo derived embryos herein described were derived from donors that:

III.1.2.1. Were free of clinical signs of BT at the time of collection and from which the embryos were conceived by artificial insemination using semen collected, processed and stored in semen collection centres approved by the competent authority in conformity with the OIE standards; and

III.1.2.2. The embryos were collected, processed and stored in accordance with standards laid down by the competent authority.]

(3)[III.1.3. The in vivo derived embryos herein described were derived from donors that:

Were kept for 21 days prior to, and during, collection in an establishment where no case of VS was reported during that period and were subject to a diagnostic test for VS, with negative results, within 21 days prior to embryo collection. In addition the embryos were collected, processed and stored in conformity with OIE notified standards; and the establishment was not located within a protection or surveillance zone. Embryos collected within protection and surveillance zones has been clearly identified and detained under official supervision]

(3)[III.1.4. The in vivo derived embryos herein described were derived from donors that:

either

(2)[1.4.1. have not been vaccinated against CBPP and were subjected to the complement fixation test for CBPP with negative results, on two occasions, with an interval of not less than 21 days and not more than 30 days between each test, the second test being performed within 14 days prior to collection; and were isolated from other domestic bovidae from the day of the first complement fixation test until collection;]

or

(2)[1.4.1. were vaccinated using a vaccine complying with the standards described in the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals not more than 4 months prior to collection;] and

1.4.2. showed no clinical sign of CBPP on the day of collection of the embryos; and were kept since birth, or for the past 6 months, in a herd(s) where no case of CBPP was reported during that period, and that the herd(s) was/were not situated in a CBPP infected zone; and the embryos were collected,

II. Health attestation	II.a. Certificate reference No	II.b.

processed and stored in accordance with standards laid down by the competent authority.]]

Notes

This health certificate is for veterinary purposes only.

Part I

Box I.6.: Complete only in case of transit throught the Union.

Box I.8.: Region of origin: if applicable, otherwise must be crossed out: for animal species or for products affected by the regionalisation measures or by the setting up of approved zones in accordance with Union decisions.

Box I.12.: Complete only in case of storage of products in transit: name and address (street, town and postal code) and the approval or registration number of the warehouse in a free zone, the customs warehouse or the ship supplier.

Box I.14.: For animal products: indicate the date of departure of the means of transport (aeroplane, ship, railway or road vehicle).

Box I.18.: Complete only in case of animal products.

Box I.19.: Enter the 'Total gross weight (kg)' and 'Total net weight (kg)'.

Box I.21.: If applicable, enter the identification number of the container and the seal number.

Box I.22.: Enter the intended use for animal products (the available options will vary in accordance with the specific certificate in the Union import requirements).

Box I.23.: Complete only in case of transit through the Union.

Box I.24.: Complete only in case of importation or temporary admission to the Union.

Box I.25.: Use the appropriate Harmonised System (HS) code under the following headings: 05119985

Part II

(1) Only to be completed if special conditions apply. Otherwise delete.
(2) Delete as appropriate.
(3) Keep if appropriate.

Official veterinarian

Name (in capital letters): Qualification and title:

Date: Signature:

Stamp: