Part I : Details of consignment

I.1. Consignor
Name
Address
Country ISO Code

I.2. IMSOC Reference
I.2.a. Local Reference

I.5. Consignee
Name
Address
Country ISO Code

I.3. Central competent authority
I.4. Local competent authority

I.7. Country of origin ISO Code

I.9. Country of destination ISO Code

I.8. Region of origin Code

~~I.10. Region of destination~~

I.11. Place of Dispatch
Name
Address
Approval Number
Country ISO Code

I.12. Place of destination
Name
Address
Approval Number
Country ISO Code

I.13. Place of Loading
Name
Address
Approval Number
Country ISO Code

I.14. Date and time of departure

I.15. Means of Transport

| Mode | International transport document | Identification |
|---|---|---|
| | | |
| | | |
| | | |
| | | |

I.16 Entry Point

I.18. Transport conditions
Controlled temperature ☐ Chilled ☐ Ambient ☐ Frozen ☐

I.17. Accompanying documents
Commercial document reference
Date of issue
Country
Place of issue

I.19. Container No / Seal No

I.20. Certified as
Artificial reproduction ☐

I.21. For transit through a third country ☐
Country ISO Code
EU Exit Authority BCP code
EU Entry Authority BCP code

I.22. For transit through Member State(s) ☐
Country ISO Code

I.24. Total quantity

I.25. Total gross weight

I.28. Description of consignment

**1. 05** PRODUCTS OF ANIMAL ORIGIN, NOT ELSEWHERE SPECIFIED OR INCLUDED
**0511** Animal products not elsewhere specified or included; dead animals of Chapter|1|or 3, unfit for human consumption
**051110** Bovine semen
**05111000** Bovine semen

| Commodity | Species | Breed/Category | Identification number | Date of collection/production |
|---|---|---|---|---|
| | | | | |

| Manufacturing plant | Quantity |
|---|---|
| | |

Part II: Certification

II. Health information

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

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

Regulation (EU) 2016/429 of the European Parliament and of the Council of 9 March 2016 on transmissible animal diseases and amending and repealing certain acts in the area of animal health (Animal Health Law)

Commission Delegated Regulation (EU) 2020/686 of 17 December 2019 supplementing Regulation (EU) 2016/429 of the European Parliament and of the Council as regards the approval of germinal product establishments and the traceability and animal health requirements for movements within the Union of germinal products of certain kept terrestrial animals

Commission Delegated Regulation (EU) 2020/687 of 17 December 2019 supplementing Regulation (EU) 2016/429 of the European Parliament and the Council, as regards rules for the prevention and control of certain listed diseases

Commission Delegated Regulation (EU) 2020/688 of 17 December 2019 supplementing Regulation (EU) 2016/429 of the European Parliament and of the Council, as regards animal health requirements for movements within the Union of terrestrial animals and hatching eggs

III. Additional health attestation

III.1. The animal product is eligible for intra-Union trade without restriction.

III.2. For diseases not regulated by the EU: All laboratory samples required by this veterinary certificate have been collected, processed, and stored in accordance with the OIE's recommendations or as described in Approved Diagnostic Tests, Vaccines, Treatments and Post-Arrival Testing Laboratories for Animal Import Health Standards, MPI-STD-TVTL, found here: https://www.mpi.govt.nz/dmsdocument/2040/.

III.3. The donor's centres of residence have had no cases of BHV 5 (suspected or diagnosed) in the year prior to semen collection for export to New Zealand.

III.4. The tests for brucellosis must be proven capable of detecting infection with Brucella suis and must be one of the following: CFT, BBAT, I-ELISA, or FPA in accordance with Commission Delegated Regulation (EU) 2020/688.

III.5. To manage Leptospira interrogans, antibiotics have been added in accordance with the OIE Code, or with an approved combination listed in MPI-STD-TVTL.

III.6. Q-Fever:

To the best of my knowledge and as far as I can ascertain, the donors have never been confirmed positive for Q-Fever:

(2) ○ either [The donors were subjected to an ELISA test for Q-fever, on a sample collected between 21 to 120 days after each semen collection (a period of 60 days or less) for export to New Zealand, with negative results.]

(2) ○ or [An aliquot of semen from each collection for export to New Zealand was tested using a laboratory validated Q-fever PCR test which is in accordance with the methods described in the Q-fever Chapter of the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals.]

III.7. Mycoplasma bovis:

(2) ○ either [The semen for export to New Zealand underwent DNA extraction and PCR testing as described in MPI-STD-TVTL, found here: https://www.mpi.govt.nz/dmsdocument/2040/.]

(2) ○ or [The semen for export to New Zealand underwent antibiotic treatment as described in MPI-STD-TVTL, found here: https://www.mpi.govt.nz/dmsdocument/2040/.]

(1) ☐ III.8. [The commodity herein described, complies/y with the additional conditions in the event of the occurrence of a disease:

III.8.1. The commodity herein described was kept separate from all other commodities that did not meet the requirements during all stages of production, storage and transport and all necessary precautions were taken to prevent contamination of the commodity with any potential source of the relevant virus.

Part II: Certification

II. Health information

(1) ☐ III.8.2. [Foot and Mouth Disease:

(2) ○ either [The semen herein described was derived from donor animals which were kept in a semen collection centre where no animals have been added in last the 30 days before collection and FMD has not occurred within 10 kilometres for 30 days before and after collection, and which showed no clinical sign of FMD on the day of collection, have not been vaccinated against FMD and were subjected, not less than 21 days after collection of the semen, to a tests for antibodies against FMD virus, with negative results and no other animals present in the semen collection centre has been vaccinated against FMD. Additionally, the semen was collected from a semen collection centre not located within a protection or surveillance zone and any semen collected within a protection and surveillance zone has been clearly identified and detained under official supervision; and the semen collected, was further processed and stored in conformity with the provisions of Chapter 4.5 or Chapter 4.6 of the OIE Terrestrial Animal Health Code as relevant and was further stored in the country of origin for a period of at least one month following collection, and during this period no animal on the establishment where the donor animals were kept showed any sign of FMD.]

(2) ○ or [The semen herein described has been collected and stored frozen at least 21 days before the estimated date of earliest infection with the foot-and-mouth disease virus on a holding in the protection and surveillance zone; and any semen collected after the date of earliest infection has been stored separately and was only released after all the measures relating to the outbreak of FMD have been removed; and all animals accommodated in the semen collection centre have undergone a clinical examination and samples taken have been subjected to a serological test to substantiate the absence of infection in the centre concerned; and the donor animals have been subjected with negative result to a serological test for the detection of antibodies against the FMD virus on a sample taken not earlier than 28 days after the collection of the semen]]

(1) ☐ III.8.3. [Lumpy Skin Disease:

The semen herein described was derived from donor animals that showed no clinical sign of LSD on the day of collection of the semen and for the following 28 days; and the animals were kept in the exporting country for the 28 days prior to collection, in a semen collection centre where no case of LSD was officially reported during that period, and the centre was not situated in either a LSD infected zone or buffer zone and any semen from buffer zone has been clearly identified and controlled.]

(1) ☐ III.8.4. [Rift Valley Fever:

The donor must not show any clinical signs of RVF within the period from 14 days prior to and 14 days following germplasm collection;

And

(2) ○ either [The donor must be vaccinated against RVF in accordance with MPI-STD-TVTL at least 14 days prior to collection.]

(2) ○ or [The donor must be demonstrated to be seropositive on the day of collection using a test listed in MPI-STD-TVTL.]

(2) ○ or [Testing of paired samples using a test listed in MPI-STD-TVTL must demonstrate that seroconversion did not occur between germplasm collection and 14 days after.]

(1) ☐ III.8.5. [Contagious Bovine Pleuropneumonia (CBPP):

The donors:

Showed no clinical sign of CBPP on the day of collection of the semen; 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 semen was collected, processed and stored in accordance with standards laid down by the competent authority.

Part II: Certification

II. Health information

And

(2) ○ either [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.]

(2) ○ or [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.]]]

Notes

This health certificate is for veterinary purposes only.

Part I

Box I.6.: Complete only in case of transit through 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.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: https://food.ec.europa.eu/animals/semen-oocytes-embryos/bovine_en

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.: Enter the identification number of the container and the official 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.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 (nonexclusive) heading: 0511 99 85.

Part II

(1) Only to be completed if special conditions apply. Otherwise delete.

(2) Delete as appropriate.

Certifying Officer

| | |
|---|---|
| Name (in capital letters) | Qualification and title |
| Date of signature | Signature |
| Stamp | |