

**Ref:** AE-FK05L

**Date:** 12.12.01

## **OVERSEAS MARKET ACCESS REQUIREMENTS NOTIFICATION**

### **ANIMAL PRODUCTS ACT 1999**

### **BIOSECURITY AUTHORITY**

**OMAR B BOVSEMEC.FAL -BOVINE SEMEN to the FALKLAND ISLANDS**

#### **Statutory authority**

Pursuant to section 60 of the Animal Products Act 1999, I notify the following overseas market access requirements and specifications, entitled bovine semen to Falkland Islands.

This notice takes effect from date of signing.

Dated at Wellington this 16<sup>th</sup> day of July 2002.

Signed by Carolyn Hini  
National Manager International Animal Trade  
MAF Biosecurity Authority  
(pursuant to delegated authority)

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#### **IMPORT PERMIT**

An import permit is required for the exportation of bovine semen to the Falkland Islands.

#### **EXPORT REQUIREMENTS.**

An Official Veterinarian of the New Zealand Ministry of Agriculture and Forestry must certify, after due enquiry, in regards to the donor animal and semen identified, the following requirements.

#### **Country Freedom**

New Zealand must be free from African swine fever, Akabane, bluetongue, brucellosis (*Brucella abortus*), bovine spongiform encephalopathy, contagious bovine pleuropneumonia, ephemeral fever, epizootic haemorrhagic disease of deer, foot and mouth disease, lumpy skin disease, rabies, rinderpest and vesicular stomatitis.

Vaccination against these diseases must be prohibited in New Zealand.

## **Semen Collection Centre**

The semen collection centre at which the semen is to be collected must be:

- approved by the New Zealand Ministry of Agriculture and Forestry as having facilities suitable for isolating animals and collecting, processing and storing semen in accordance with *Bovine Semen* Appendix 3.2.1 of the OIE *International Animal Health Code, mammals, birds and bees*
- under the direct supervision and sanitary control of a veterinarian who is responsible for the hygiene of the centre and the health of the animals
- regularly inspected by a veterinary officer accredited by the New Zealand Ministry of Agriculture and Forestry

## **Semen Collection Centre Status**

In the 12 months preceding the date of export of the semen, there must be no confirmed diagnoses of the following diseases on the semen collection centre: bovine genital campylobacteriosis, bovine tuberculosis, bovine viral diarrhoea/mucosal disease (BVD/MD), enzootic bovine leucosis, infectious bovine rhinotracheitis/infectious pustular vulvovaginitis (IBR/IPV), Johne's disease (*Mycobacterium paratuberculosis*), leptospirosis and trichomoniasis.

## **Pre-Collection Period**

The donor bull must be resident in a semen collection centre, under veterinary supervision, for at least 30 days prior to first collection of semen.

During the pre-collection period the donor bull must not perform natural service.

During the pre-collection and collection period the donor bull and all in-contact animals must remain healthy.

## **Testing on the Centre**

The donor bulls must be resident in a semen collection centre and be tested, with negative results, for the following diseases within the 12-month period immediately prior to semen collection:

- Leptospirosis either using the microagglutination test for serotypes *L hardjo*, and *L pomona* (negative is less than 50% agglutination at 1:200 dilution), or each donor bull must be subjected to an intramuscular injection of dihydrostreptomycin at a dose rate of 25 mg/kg bodyweight, on two occasions, at an interval of 14 days. The dates of injection must be recorded
- *Campylobacter fetus* subsp. *venerealis* using culture examination of preputial washings

- *Trichomonas fetus* using direct microscopic examination and culture examination of preputial washings
- Bovine tuberculosis using an intradermal test applied to either the caudal fold or cervical area using bovine tuberculin
- Enzootic bovine leucosis using either the AGID test or ELISA.

The donor bull, or the semen for export, must be tested for infectious bovine rhinotracheitis as follows:

- either negative semen culture carried out within the 12 months prior to collection of the semen, where the donor bull having given a negative result to either an IBR ELISA or SNT is routinely vaccinated with an inactivated IBR vaccine
- or negative to either an ELISA or a SNT carried out within the 12 months prior to collection of the semen for export
- or negative semen culture on each batch of semen for export when the donor bull is IBR seropositive.

The donor bull, or its semen, must be tested with negative results for bovine virus diarrhoea virus as follows:

- either an antigen ELISA
- or culture of a serum sample within the 12 months prior to collection of the semen for export
- or culture of the semen within the 12 months prior to export of the semen.

### **Semen Collection and Storage**

The semen must be collected, processed, packaged and stored in accordance with the recommendations of Appendix 3.2.1 of the OIE *International Animal Health Code, mammals, birds and bees*. The ingredients of the diluent must be recorded along with the name and concentration of antibiotics added to the diluent.

Prior to export, the transportation flasks must be sealed under veterinary supervision.

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### **Certification**

Certification is to be provided on the appropriate export certificate form determined to be an official assurance under section 62(1) of the Animal Products Act 1999.

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## **Definitions**

For the purposes of this document:

Any term or expression that is defined in the Animal Products Act 1999 and used, but not defined in this document, has the same meaning as in this Act.

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## ***Explanatory note***

*These overseas market access requirements are drafted on the current export certificate for the exportation of bovine semen to the Falkland Islands dated 12 December 2001.*

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## **Disclaimer**

This information is intended for use as guidance only and should not be taken as definitive or exhaustive. MAF endeavours to keep this information current and accurate. However, it may be subject to change without notice. Exporters should make their own enquiries in relation to import requirements. MAF will not accept liability for any loss resulting from reliance on this information.