OMARs with an uncertain status

These OMARs have not been used for a significant period of time. Therefore the requirements may have changed without the Ministry for Primary Industries knowledge.

If an exporter can provide the current import conditions, and the requirements still match, the certificate and the OMARs will be moved back into the published list of export certificates and OMAR's.

Overseas Market Access Requirements Notification - Animal Products Act 1999 – MAF Biosecurity New Zealand

Ref: AE-EC 24/11L Date: 12 June 2007

OMAR B CAOSEMEC.ECU 12.06.07 – CAPRINE AND OVINE SEMEN TO ECUADOR

1. Statutory authority

Pursuant to section 60 of the Animal Products Act 1999, I notify the following overseas market access requirements, entitled caprine and ovine semen to Ecuador.

This notice takes effect from date of signing.

Dated at Wellington this 19th day of July 2007.

Signed: Karen Sparrow Manager Exports Border Standards MAF Biosecurity New Zealand (pursuant to delegated authority)

2. Requirements of Ecuador

Ovine and caprine semen exported from New Zealand to Ecuador must comply with the import requirements of Ecuador listed in this notice as follows:

- 2.1 An import permit is required for the exportation of caprine and ovine semen to Ecuador.
- 2.2 An official veterinarian of New Zealand Ministry of Agriculture and Forestry must certify, after due enquiry, the following:
- 2.2.1 New Zealand is declared officially free of rinderpest, peste des petits ruminants, sheep & goat pox, Rift Valley fever, foot-and-mouth disease, brucellosis (*Brucella abortus* & *Brucella melitensis*), infectious ovine encephalomyelitis ('louping-ill'), bluetongue, Akabane disease, Aino, maedi-visna, enzootic abortion of ewes (*Chlamydophila abortus*), scrapie, vesicular stomatitis, ovine pulmonary adenomatosis ('jaagsiekte'), contagious agalactia, contagious bovine, ovine & caprine pleuropneumonia, Nairobi sheep disease, and Q fever.

- 2.2.2 Vaccinating against these diseases is not permitted in New Zealand.
- 2.2.3 The semen collection centre, at which the semen was collected, is approved by the New Zealand Ministry of Agriculture and Forestry as having facilities suitable for collecting, processing, and storing semen in accordance with the recommendations of the *OIE Terrestrial Animal Health Code*.
- 2.2.4 The semen collection centre:
- 2.2.4.1 is under the direct supervision and sanitary control of a centre veterinarian who is approved by the New Zealand Ministry of Agriculture and Forestry, and who is responsible for the health of the donor animals and the hygiene of the semen
- 2.2.4.2 is regularly inspected by an Official Veterinarian of the New Zealand Ministry of Agriculture and Forestry.
- 2.2.5 The semen collection centre is isolated from other livestock establishments.
- 2.2.6 In the two years preceding the collection of semen, there has been no clinical evidence of contagious ecthyma ('orf'), campylobacteriosis and listeriosis on either the semen collection centre or the farm(s) of origin of the donor animals.
- 2.2.7 In case of permanent residence on the semen collection centre, all donor animals were tested at least every 12 months, for the following diseases, with negative results:
- 2.2.7.1 ovine epididymitis (*Brucella ovis*), using the complement fixation (CF) test or ELISA Test used.

(To be deleted if not applicable)

2.2.8 In case of permanent residence on the semen collection centre, all donor animals were subjected every 6 months to an intramuscular injection of dihydrostreptomycin (at a dose rate of 25 mg/kg) or long-acting oxytetracycline (at a dose rate of 20 mg/kg). Antibiotic used

(To be deleted if not applicable)

- 2.2.9 The donor animals were held in isolation for 30 days prior to the first collection of semen, during which time they were tested for the following diseases, with negative results:
- 2.2.9.1 caprine arthritis-encephalitis, using either the agar-gel immunodiffusion AGID (test) or ELISA (<u>for goats only</u>). Test used
- 2.2.9.2 border disease (ovine pestivirus), using virus isolation
- 2.2.9.3 ovine epididymitis (*Brucella ovis*), using the complement fixation (CF) test or ELISA. Test used.
- 2.2.10 The donor animals were held in isolation for 30 days prior to the first collection of semen, during which time they were subjected to two intramuscular injections of dihydrostreptomycin (at a dose arte of 25 mg/kg), given at an interval of approximately 14 days, or one injection of long-acting oxytetracycline (at a dose rate of 20 mg/kg). Antibiotic used.

- 2.2.11 According to the centre veterinarian's declaration, the donor animals remained healthy during the period from 60 days prior to collection to 30 days after the final collection of semen.
- 2.2.12 The semen was collected, processed, and stored under the supervision of the centre veterinarian in accordance with the OIE *Terrestrial Animal Health Code* Section 3.2, appendix 3.2.1.
- 2.2.13 Biological products of animal origin used in the processing of semen, including additives or diluents, were obtained from sources which present no recognised animal health risk, or were so treated prior to use that such risk is prevented. Antibiotics have been added in accordance with the recommendations of the OIE *Terrestrial Animal Health Code* Section 3.2, appendix 3.2.1.
- 2.2.14 Following the final collection, the semen was stored in liquid nitrogen and kept isolated for at least 30 days prior to export.
- 2.2.15 The straws or ampoules have been marked, such that the dates of collection and freezing can be identified.
- 2.2.16 The semen for export was placed in fresh liquid nitrogen in a container, which was either new or disinfected with a MAF-approved disinfectant. Disinfectant used and active ingredient. (Options for disinfectants include: 10% formalin, 2% available chlorine, Virkon at the manufacturer's recommended rate, or irradiation at 50kGy.)
- 2.2.17 No animals, semen, fertilised ova, products or equipment other than that listed on the Import Permit are included in the shipment.
- 2.2.18 Immediately prior to export, the transportation container was sealed in the presence of a New Zealand Ministry of Agriculture and Forestry Official Veterinarian, using a seal that bears the mark and serial number of the container which must be recorded on the export certificate.

3. 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.

Explanatory note

These overseas market access requirements are based on the export certificate for caprine and ovine semen to Ecuador dated 12 June 2007.

Additional Information on OMAR Notification: OACEMBEC.UK 18.05.07

- 1. This OMAR is based on the import conditions that were received, via an exporter, from Dr Alex Andrade Orlando, Zoosanitary Certification Supervisor, Servicio Ecuatoriano de Sanidad Agropecuaria (SESA), Ecuador; this country being a member of the Andean Community of Nations (which, at present. includes Bolivia, Columbia, Ecuador, Peru and Venezuela). The export certificate was approved by Dr Andrade in an e-mail, dated 12 July 2007. The export certificate of 12 June 2007 was officially translated.
- 2. To import ovine or caprine semen into Ecuador, the importer must send/fax a copy of the completed and signed export certificate to the Servicio Ecuatoriano de Sanidad Agropecuaria (SESA) in Quito. SESA will then issue the Import Permit within 72 hours. Once the Import Permit has been issued, the consignment can be exported to Ecuador, accompanied by the original export certificate. Fax number for SESA is: + 593 2 2228448.
- 3. Transportation containers with broken or altered seals will be rejected on arrival in Ecuador. Seals may only be removed by Official Veterinarians in Ecuador. In case there is a need to inspect or re-charge containers, Veterinary Officials at any transit port are required to notify in writing on the export certificate the reason and the replacement seal number/mark.

Section 61.A of the Animal Products Amendments Act 2005 states that 'The Crown is not liable, and nor is the Director-General or any employee of the Ministry liable, for any loss arising through the refusal or failure of the relevant authority of an overseas market to admit export animal material or animal product to that market'