Terrestrial Animal Health Standards Commission Report

October 2008

NEW

CHAPTER 4.5.

COLLECTION AND PROCESSING OF BOVINE, SMALL RUMINANT AND PORCINE SEMEN

Article 4.5.1.

General considerations

The purposes of official sanitary control of semen production are to:

- 1. maintain the health of *animals* on an *artificial insemination centre* at a level which permits the international distribution of semen with a negligible *risk* of infecting other *animals* or humans with pathogens transmissible by semen;
- 2. ensure that semen is hygienically collected, processed and stored.

Artificial insemination centres should comply with recommendations in Chapter 4.6.

Standards for diagnostic tests are described in the Terrestrial Manual.

Article 4.5.2.

Conditions applicable to testing of bulls and teaser animals

Bulls and teaser animals should only enter an artificial insemination centre if they fulfil the following requirements.

1. <u>Pre-quarantine</u>

The *animals* should comply with the following requirements prior to entry into isolation at the *quarantine station*.

a) Bovine brucellosis

The *animals* should comply with point 3 or 4 of Article 11.3.5.

b) Bovine tuberculosis

The animals should comply with point 3 or 4 of Article 11.7.5.

c) Bovine viral diarrhoea-mucosal disease (BVD-MD)

The *animals* should be subjected to the following tests:

- i) a virus isolation test or a test for virus antigen, with negative results;
- ii) a serological test to determine the serological status of every animal.
- d) Infectious bovine rhinotracheitis-infectious pustular vulvovaginitis

If the *artificial insemination centre* is to be considered as infectious bovine rhinotracheitis-infectious pustular vulvovaginitis free (IBR/IPV), the *animals* should either:

- i) come from an IBR/IPV free herd as defined in Article 11.12.3.; or
- ii) be subjected, with negative results, to a serological test for IBR/IPV on a blood sample.

e) Bluetongue

The *animals* should comply with Articles 8.3.5., 8.3.6. or 8.3.7., depending on the bluetongue status of the country of origin of the *animals*.

2. Testing in the quarantine station prior to entering the semen collection facilities

Prior to entering the semen collection facilities of the *artificial insemination centre*, bulls and teaser animals should be kept in a *quarantine station* for at least 28 days. The *animals* should be subjected to diagnostic tests as described below a minimum of 21 days after entering the *quarantine station*, except for *Campylobacter fetus* subsp. *venerealis* and *Trichomonas foetus*, for which testing may commence after 7 days in quarantine. All the results should be negative except in the case of BVD-MD antibody serological testing (see point 2b)i) below).

a) Bovine brucellosis

The animals should be subjected to a serological test with negative results.

b) BVD-MD

i) All *animals* should be tested for viraemia as described in point 1c) above.

Only when all the *animals* in quarantine test negative for viraemia, may the *animals* enter the semen collection facilities upon completion of the 28-day quarantine period.

- ii) After 21 days in quarantine, all *animals* should be subjected to a serological test to determine the presence or absence of BVD-MD antibodies.
- iii) Only if no sero-conversion occurs in the *animals* which tested seronegative before entry into the *quarantine station*, may any *animal* (seronegative or seropositive) be allowed entry into the semen collection facilities.
- iv) If sero-conversion occurs, all the *animals* that remain seronegative should be kept in quarantine over a prolonged time until there is no more seroconversion in the group for

a period of 3 weeks. Serologically positive *animals* may be allowed entry into the semen collection facilities.

c) Campylobacter fetus subsp. venerealis

- i) Animals less than 6 months old or kept since that age only in a single sex group prior to quarantine should be tested once on a preputial specimen, with a negative result.
- ii) Animals aged 6 months or older that could have had contact with females prior to quarantine should be tested three times at weekly intervals on a preputial specimen, with a negative result in each case.

d) Trichomonas foetus

- i) Animals less than 6 months old or kept since that age only in a single sex group prior to quarantine, should be tested once on a preputial specimen, with a negative result.
- ii) Animals aged 6 months or older that could have had contact with females prior to quarantine should be tested three times at weekly intervals on a preputial specimen, with a negative result in each case.

e) IBR-IPV

If the *artificial insemination centre* is to be considered as IBR/IPV free, the *animals* should be subjected, with negative results, to a diagnostic test for IBR/IPV on a blood sample. If any *animal* tests positive, the *animal* should be removed immediately from the *quarantine station* and the other *animals* of the same group should remain in quarantine and be retested, with negative results, not less than 21 days after removal of the positive *animal*.

f) Bluetongue

The *animals* should comply with Articles 8.3.5., 8.3.6. or 8.3.7., depending on the bluetongue status of the country of origin of the *animals*.

3. Testing for BVD-MD prior to the initial dispatch of semen from each serologically positive bull

Prior to the initial dispatch of semen from BVD-MD serologically positive bulls, a semen sample from each *animal* should be subjected to a virus isolation or virus antigen test for BVD-MD. In the event of a positive result, the bull should be removed from the centre and all of its semen destroyed.

4. Testing of frozen semen for IBR/IPV in artificial insemination centres not considered as IBR/IPV free

Each aliquot of frozen semen should be tested as per Article 11.12.7.

5. Testing programme for bulls and teasers resident in the semen collection facilities

All bulls and teasers resident in the semen collection facilities should be tested at least annually for the following *diseases*, with negative results, where the country of origin is not free:

a) Bovine brucellosis

b) Bovine tuberculosis

c) BVD-MD

Animals negative to previous serological tests should be retested to confirm absence of antibodies.

Should an *animal* become serologically positive, every ejaculate of that *animal* collected since the last negative test should be either discarded or tested for virus with negative results.

d) Campylobacter fetus subsp. venerealis

- i) A preputial specimen should be cultured.
- ii) Only bulls on semen production or having contact with bulls on semen production need to be tested. Bulls returning to collection after a lay off of more than 6 months should be tested not more than 30 days prior to resuming production.

e) Bluetongue

The *animals* should comply with the provisions referred to in Articles 8.3.5., 8.3.6. or 8.3.7., depending on the bluetongue status of the country of origin of the *animals*.

f) Trichomonas foetus

- i) A preputial specimen should be cultured.
- ii) Only bulls on semen production or having contact with bulls on semen production need to be tested. Bulls returning to collection after a lay off of more than 6 months should be tested not more than 30 days prior to resuming production.

g) IBR-IPV

If the *artificial insemination centre* is to be considered as IBR/IPV free, the *animals* should comply with the provisions in point 2)c) of Article 11.12.3.

Article 4.5.3.

Conditions applicable to testing of rams/bucks and teaser animals

Rams/bucks and teaser animals should only enter an artificial insemination centre if they fulfil the following requirements.

1. <u>Pre-quarantine</u>

The *animals* should comply with the following requirements prior to entry into isolation at the *quarantine station*.

a) Caprine and ovine brucellosis

The animals should comply with Article 14.1.6.

b) Ovine epididymitis

The animals should comply with Article 14.7.3.

c) Contagious agalactia

The animals should comply with points 1 and 2 of Article 14.3.1.

d) Peste des petits ruminants

The animals should comply with points 1, 2, and 4 or 5 of Article 14.8.7.

e) Contagious caprine pleuropneumonia

The *animals* should comply with Article 14.4.5. or Article 14.4.7., depending on the contagious caprine pleuropneumonia status of the country of origin of the *animals*.

f) Paratuberculosis

The animals should be free from clinical signs for the past 2 years.

g) Scrapie

If the *animals* do not originate from a scrapie free country or zone as defined in Article 14.9.3., the *animals* should comply with Article 14.9.6.

h) Maedi-visna

The *animals* should comply with Article 14.6.2.

i) Caprine arthritis/encephalitis

In the case of goats, the *animals* should comply with Article 14.2.2.

j) Bluetongue

The *animals* should comply with Articles 8.3.5., 8.3.6. or 8.3.7., depending on the bluetongue status of the country of origin of the *animals*.

k) Tuberculosis

In the case of goats, the *animals* should be subject to a single or comparative tuberculin test, with negative results.

Border disease

The animals should be subject to a viral agent isolation test with negative results.

2. Testing in the quarantine station prior to entering the semen collection facilities

Prior to entering the semen collection facilities of the *artificial insemination centre*, rams/bucks and teasers should be kept in a *quarantine station* for at least 28 days. The *animals* should be subjected to diagnostic tests as described below a minimum of 21 days after entering the *quarantine station*, with negative results.

a) Caprine and ovine brucellosis

The animals should be subject to testing as described in point 1c) of Article 14.1.8.

b) Ovine epididymitis

The *animals* and semen should be subject to testing as described in points 1d) and 2 of Article 14.7.4.

c) Maedi-visna and caprine arthritis/encephalitis

The animals and semen should be subjected to a serological test.

d) Bluetongue

The *animals* should comply with the provisions referred to in Articles 8.3.5., 8.3.6. or 8.3.7., depending on the bluetongue status of the country of origin of the *animals*.

3. Testing programme for rams/bucks and teasers resident in the semen collection facilities

All rams/bucks and teasers resident in the semen collection facilities should be tested at least annually for the following *diseases*, with negative results, where the country of origin is not free:

- a) caprine and ovine brucellosis;
- b) ovine epididymitis;
- c) Maedi-visna and caprine arthritis/encephalitis;
- d) tuberculosis (for goats only);
- e) bluetongue.

Article 4.5.4

Conditions applicable to testing of boars

Boars should only enter an artificial insemination centre if they fulfil the following requirements.

1. Pre-quarantine

The *animals* should be clinically healthy, physiologically normal and comply with the following requirements within 30 days prior to entry into isolation at the *quarantine station*.

a) Porcine brucellosis

The *animals* should comply with Article 15.4.3.

b) Foot and mouth disease

The animals should comply with Article 8.5.10., 8.5.11. or 8.5.12.

c) Aujeszky's disease

The *animals* should comply with Article 8.2.8. or 8.2.9.

d) Teschovirus encephalomyelitis

The *animals* should comply with Article 15.6.5. or 15.6.7.

e) Transmissible gastroenteritis

The animals should comply with Article 15.7.2.

f) Swine vesicular disease

The animals should comply with Article 15.5.5. or 15.5.7.

g) African swine fever

The *animals* should comply with Article 15.1.5. or 15.1.6.

h) Classical swine fever

The animals should comply with Article 15.3.7., 15.3.8. or 15.3.9.

i) Porcine reproductive and respiratory syndrome

The *animals* should be subject to the test complying with the standards in the *Terrestrial Manual*.

2. Testing in the quarantine station prior to entering the semen collection facilities

Prior to entering the semen collection facilities of the *artificial insemination centre*, boars should be kept in a *quarantine station* for at least 28 days. The animals should be subjected to diagnostic tests as described below a minimum of 21 days after entering the *quarantine station*, with negative results.

a) Porcine brucellosis

The animals should comply with Article 15.4.5.

b) Foot and mouth disease

The animals should comply with Article 8.5.13., 8.5.14., 8.5.15. or 8.5.16.

c) Aujeszky's disease

The animals should comply with Article 8.2.12., 8.2.13. or 8.2.14.

d) Teschovirus encephalomyelitis

The *animals* should comply with Article 15.6.9. or 15.6.10.

e) Transmissible gastroenteritis

The animals should comply with Article 15.7.4.

f) Swine vesicular disease

The animals should comply with Article 15.5.9. or 15.5.10.

g) African swine fever

The *animals* should comply with Article 15.1.8. or 15.1.9.

h) Classical swine fever

The animals should comply with Article 15.3.11., 15.3.12. or 15.3.13.

i) Porcine reproductive and respiratory syndrome

The *animals* should be subject to the test complying with the standards in the *Terrestrial Manual*.

3. Testing programme for boars resident in the semen collection facilities

All boars resident in the semen collection facilities should be tested at least annually for the following diseases, with negative results, where the *compartment/zone* or country is not free:

a) Porcine brucellosis

The *animals* should comply with Article 15.4.5.

b) Foot and mouth disease

The animals should comply with Article 8.5.13., 8.5.14., 8.5.15. or 8.5.16.

c) Aujeszky's disease

The *animals* should comply with Article 8.2.12., 8.2.13. or 8.2.14. regarding testing every four months.

d) Teschovirus encephalomyelitis

The animals should comply with Article 15.6.9. or 15.6.10.

e) Transmissible gastroenteritis

The animals should comply with Article 15.7.4.

f) Swine vesicular disease

The animals should comply with Article 15.5.9. or 15.5.10.

g) African swine fever

The *animals* should comply with Article 15.1.8. or 15.1.9. Routine test to be applied at least every six months.

h) Classical swine fever

The animals should comply with Article 15.3.11., 15.3.12. or 15.3.13.

i) Porcine reproductive and respiratory syndrome

The *animals* should be subject to the test complying with the standards in the *Terrestrial Manual*.

Article 4.5.5.

General considerations for hygienic collection and handling of semen

Observation of the recommendations described in the Articles below will very significantly reduce the likelihood of the semen being contaminated with common bacteria which are potentially pathogenic.

Article 4.5.6.

Conditions applicable to the collection of semen

- 1. The floor of the mounting area should be easy to clean and to disinfect. A dusty floor should be avoided.
- 2. The hindquarters of the teaser, whether a dummy or a live teaser animal, should be kept clean. A dummy should be cleaned completely after each period of collection. A teaser animal should have its hindquarters cleaned carefully before each collecting session. The dummy or hindquarters of the teaser animal should be sanitized after the collection of each ejaculate. Disposable plastic covers may be used.
- 3. The hand of the person collecting the semen should not come into contact with the *animal*'s penis. Disposable gloves should be worn by the collector and changed for each collection.
- 4. The artificial vagina should be cleaned completely after each collection where relevant. It should be dismantled, its various parts washed, rinsed and dried, and kept protected from dust. The inside of the body of the device and the cone should be disinfected before re-assembly using approved disinfection techniques such as those involving the use of alcohol, ethylene oxide or steam. Once re-assembled, it should be kept in a cupboard which is regularly cleaned and disinfected.
- 5. The lubricant used should be clean. The rod used to spread the lubricant should be clean and should not be exposed to dust between successive collections.
- 6. The artificial vagina should not be shaken after ejaculation, otherwise lubricant and debris may pass down the cone to join the contents of the collecting tube.

- 7. When successive ejaculates are being collected, a new artificial vagina should be used for each mounting. The vagina should also be changed when the *animal* has inserted its penis without ejaculating.
- 8. The collecting tubes should be sterile, and either disposable or sterilised by autoclaving or heating in an oven at 180°C for at least 30 minutes. They should be kept sealed to prevent exposure to the environment while awaiting use.
- 9. After semen collection, the tube should be left attached to the cone and within its sleeve until it has been removed from the collection room for transfer to the laboratory.

Article 4.5.7.

Conditions applicable to the handling of semen and preparation of semen samples in the laboratory

1. Diluents

- a) All receptacles used should have been sterilised.
- b) Buffer solutions employed in diluents prepared on the premises should be sterilized by filtration (0.22 μm) or by autoclaving (121°C for 30 minutes) or be prepared using sterile water before adding egg yolk (if applicable) or equivalent additive and antibiotics.
- c) If the constituents of a diluent are supplied in commercially available powder form, the water used must have been distilled or demineralised, sterilized (121°C for 30 minutes or equivalent), stored correctly and allowed to cool before use.
- d) Whenever milk, egg yolk or any other animal protein is used in preparing the semen diluent, the product must be free of pathogens or sterilised; milk heat-treated at 92°C for 3-5 minutes, eggs from SPF flocks when available. When egg yolk is used, it should be separated from eggs using aseptic techniques. Alternatively, commercial egg yolk prepared for human consumption or egg yolk treated by, for example, pasteurisation or irradiation to reduce bacterial contamination, may be used. Other additives must also be sterilized before use.
- e) Diluent should not be stored for more than 72 hours at +5°C before use. A longer storage period is permissible for storage at -20°C. Storage vessels should be stoppered.
- f) A mixture of antibiotics should be included with a bactericidal activity at least equivalent to that of the following mixtures in each ml of frozen semen: gentamicin (250 μg), tylosin (50 μg), lincomycin-spectinomycin (150/300 μg); penicillin (500 IU), streptomycin (500 μg), lincomycin-spectinomycin (150/300 μg); or amikacin (75μg), divekacin (25μg).

The names of the antibiotics added and their concentration should be stated in the international veterinary certificate.

2. Procedure for dilution and packing

a) The tube containing freshly collected semen should be sealed as soon as possible after collection, and kept sealed until processed.

- b) After dilution and during refrigeration, the semen should also be kept in a stoppered container.
- c) During the course of filling receptacles for dispatch (such as insemination straws), the receptacles and other disposable items should be used immediately after being unpacked. Materials for repeated use should be disinfected with alcohol, ethylene oxide, steam or other approved disinfection techniques.
- d) If sealing powder is used, care should be taken to avoid its being contaminated.

3. Conditions applicable to the storage of semen

Semen for export should be stored separately from other genetic material not meeting these recommendations in fresh liquid nitrogen in sterilised/sanitised flasks before being exported.

Semen straws should be sealed and code marked in line with the international standards of the International Committee for Animal Recording (ICAR)¹.

Prior to export, semen straws or pellets should be identified and placed into new liquid nitrogen in a new or sterilised flask or container under the supervision of an *Official Veterinarian*. The contents of the container or flask should be verified by the *Official Veterinarian* prior to sealing with an official numbered seal before export and accompanied by an *international veterinary certificate* listing the contents and the number of the official seal.