Guidance for BOV-SEM-A-INTRA until 02/11/2026

MODEL ANIMAL HEALTH CERTIFICATE FOR THE MOVEMENT BETWEEN MEMBER STATES OF CONSIGNMENTS OF SEMEN OF BOVINE ANIMALS COLLECTED, PROCESSED AND STORED IN ACCORDANCE WITH REGULATION (EU) 2016/429 AND DELEGATED REGULATION (EU) 2020/686 AFTER 20 APRIL 2021, DISPATCHED FROM THE SEMEN COLLECTION CENTRE WHERE THE SEMEN WAS COLLECTED

EUROPEAN UNION **Certificate**
model BOV-SEM-A-INTRA

EUROPEAN UNION

Part I: Description of consignmen

I.1 Consignor enter accurate details
Name
Address including accurate post code
Country
ISO country code XI

I.2 IMSOC reference insert TRACES certificate number
QR CODE N/A

I.2a Local reference enter aPVP unique stamp code

I.3 Central Competent Authority **DEFRA**

I.4 Local Competent Authority **DAERA**

I.5 Consignee enter accurate details
Name
Address
https://www.nationsonline.org/oneworld/country_code_list.htm
Country
.
ISO country code code of receiving country e.g. IE (Ireland)

I.6 Operator conducting assembly operations independently establishment N/A
Name
Registration No
Address
Country
ISO country code

I.7 Country of origin **N. Ireland**
ISO country code XI

I.9 Country of destination As in 1.5
ISO country code as in

I.8 Region of origin N/A
Code N/A

I.10 Region of destination N/A
Code N/A

I.11 Place of dispatch I.1
Name
Address
Country
Registration/Approval No Centre approval number https://www.daera-ni.gov.uk/publications/approved-germinal-product-establishments-northern-ireland
ISO country code XI

I.12 Place of destination
Name
Registration/Approva
Address
Country
ISO country code as in

I.13 Place of loading I.1

I.14 Date and time of departure

I.15 Means of transport
□ Vessel
□ Aircraft
□ Railway
□ Road vehicle X
Identification Reg no
□ Other
Document

I.16 Transporter N/A
Name
Registration/Authorisation
Address
Country
ISO country code

I.17 N/A
Type
Code
Country
ISO country code
Commercial document reference

I.18	**Transport conditions**	□ Ambient	□ Chilled □ Frozen select condition which applies

I.19 Container number/Seal number N/A if no seal

Container No

Seal No

I.20 Certified as or for

□ Further keeping	□ Slaughter	□ Confined establishment	X□ Germinal products
□ Registered equine animal	□ Travelling circus/animal act	□ Exhibition	□ Event or activity near borders
□ Release into the wild	□ Dispatch centre	□ Relaying area/purification centre	□ Ornamental aquaculture establishment
□ Further processing	□ Organic fertilizers and soil improvers	□ Technical use	□ Quarantine or similar establishment
□ Products for human consumption	□ Pollination	□ Live aquatic animals for human consumption	□ Other

I.21 □ For transit through a third country if via GB

Third country

ISO country code GB

Exit point

BCP code

Entry point

BCP code

I.22	**□ For transit through Member State(s) e.g. IRL**		**I.23**	**□ For export N/A**	
	Member State e.g. Ireland	ISO country code e.g. IE		Third country	ISO country code
	Member State	ISO country code		Exit point	BCP code
	Member State	ISO country code			

I.24	**Estimated journey time N/A**		**I.25**	**Journey log**	□ yes □ no X
I.26	**Total number of packages number of containers**		**I.27**	**Total quantity number of straws**	
I.28	**Total net weight/gross weight (kg)**	Estimated weight	**I.29**	**Total space foreseen for the consignment** N/A	

I.30 Description of consignment

CN code	Species	Subspecies/Category	~~Sex~~	~~Identification system~~	Identification number	~~Age~~	Quantity Type
~~Region of origin~~		~~Cold store~~		Identification mark	~~Type of packaging~~		~~Net weight~~
~~Slaughterhouse~~		~~Treatment type~~		~~Nature of commodity~~	~~Number of packages~~		~~Batch N~~
		Date of collection/production		~~Manufacturing plant~~	Approval or registration number of plant/establishment/centre	~~Test~~	

NOTE October 2026: Purple text below indicates requirements that are current but no longer apply from 02/11/2026.

The aPVP may need to privately request support certification:

1. If the authorising PVP completing this certificate does not have knowledge of donor movement or disease or test histories they must personally obtain veterinary certification as needed from the relevant veterinary surgeon to allow them to certify the statements below.
2. For Notifiable disease clearances: to apply for a Veterinary Service Support Certificate (VSSPT), see application here Exports of consignments of semen of bovine animals to EU Member States collected, processed and stored after 20 April 2021 | Department of Agriculture, Environment and Rural Affairs (daera-ni.gov.uk)

A DAERA VSSPT, (veterinary service support certificate), is valid for 10 days after the date of its certification unless the disease situation changes after its issue.

Reg (EU) 2020/686: https://eur-lex.europa.eu/eli/reg_del/2020/686/oj/eng
Select *current consolidated version.*

DAERA GUIDANCE ON INTRA TEMPLATE FOR HEALTH ATTESTATIONS:

I, the undersigned official veterinarian, hereby certify that:

II.1. The semen of bovine animals described in Part I has been collected, processed and stored, and dispatched from the semen collection centre (1) which

II.1.1. is approved and kept in a register by the competent authority; aPVP to confirm. Check DAERA website https://www.daera-ni.gov.uk/publications/approved-germinal-product-establishments-northern-ireland

II.1.2. complies with requirements as regards responsibilities, operational procedures, facilities and equipment set out in Part 1 of Annex I to Commission Delegated Regulation (EU) 2020/686 aPVP to confirm.

The semen described in Part I is intended for artificial reproduction and was obtained from **donor animals which**

II.2.1. have been born and remained since birth in the Union, or have entered the Union in accordance with the requirements for entry into the Union; aPVP to confirm through movement history.

II.2.2**. come**, before the commencement of the quarantine referred to in point II.2.6., **from establishments i**n a Member State or zone thereof, or from establishments under official

control by the competent authority in a third country or territory, or a zone thereof

II.2.2.1. situated **in an area where foot-and-mouth disease has not been reported within a 10-km radius centred on the establishment for a period of at least 30 days** and in which foot-and-mouth disease has not been reported during a period of at least 3 months, Requires VSSPT

and aPVP to select the relevant option below depending on movement/import history and disease status.

(2) ○ either [they were not vaccinated against foot-and-mouth disease;]

(2) ○ or [they were vaccinated against foot-and-mouth disease during the period of 12 months prior to the date of collection of the semen but not during the period of the last 30 days immediately prior to the date of collection of the semen, and 5 % (with a minimum of five straws) of each quantity of semen taken from a donor animal at any time is submitted to a virus isolation test for foot-and-mouth disease with negative results;]

II.2.2.2. **free from infection with Mycobacterium tuberculosis complex** (M. bovis, M. caprae and M. tuberculosis), current status provided by VSSPT and they have never been kept previously in any establishment of a lower health status; aPVP to establish previous history.

II.2.2.3. **free from infection with Brucella abortus, B. melitensis and B. suis**, and they have never been kept previously in any establishment of a lower health status; VSSPT to provide for current establishment; aPVP to establish previous history.

(2) ○ either [II.2.2.4. **free from enzootic bovine leukosis**, and they have never been kept previously in any establishment of a lower health status;] VSSPT to provide for current establishment; aPVP to establish previous history and select accordingly.

(2) ○ or [II.2.2.4. not free from enzootic bovine leukosis and the donor animals are younger than 2 years of age and have been produced by dams which have been subjected, with negative results, to a serological test for enzootic bovine leukosis after removal of the animal from the dam;]

(2) ○ or [II.2.2.4. not free from enzootic bovine leukosis and the donor animals have reached the age of 2 years and have been subjected, with a negative result, to a serological test for enzootic bovine leukosis;]

(2) ○ either [II.2.2.5. **free from infectious bovine rhinotracheitis/infectious pustular vulvovaginitis,** and they have never been kept previously in any establishment of a lower health status;] aPVP to establish previous IBR/IPV disease history and select accordingly.

(2) ○ or [II.2.2.5. not free from infectious bovine rhinotracheitis/infectious pustular

vulvovaginitis and the donor animals have been subjected, with a negative

result, to a serological test (whole virus) on a blood sample;]

II.2.2.6. in which **surra (Trypanosoma evansi) has not been reported during the 30 day**

period, and

(2) ○ either [**surra has not been reported in the establishments during the last 2 years**.] VSSPT to provide. Provided free, select this and delete next option.

(2) ○ or [surra has been reported in the establishments during the last 2 years and following the last outbreak the establishments have remained under movement restrictions until

- the infected animals have been removed from the establishment, and

- the remaining animals on the establishment have been

subjected to a test for surra (Trypanosoma evansi) with

one of the diagnostic methods provided for in Part 3 of

Annex I to Commission Delegated Regulation (EU)

2020/688, carried out, with negative results, on samples taken at least 6 months after the infected animals have been removed from the establishment.]

Donor animals which

II.2.3. did not show symptoms or clinical signs of transmissible animal diseases on the day of their

admission to a semen collection centre and on the day of collection of the semen; aPVP to establish

II.2.4. are individually identified as provided for in Article 38 of Commission Delegated Regulation

(EU) 2019/2035; aPVP to establish

II.2.5. **for a period of at least 30 days prior to the date of first collection of the semen and during the collection period** **For frozen semen, aPVP must provide these dates to DAERA so DAERA can find disease status for the VSSPT for 30 days before collection of frozen semen; for fresh semen, VSSPT to be provided based on calculating 30 days from the date of issue of VSSPT.**

II.2.5.1. were **kept** on **establishments not situated in a restricted zone** established due to aPVP to establish movement history; VSSPT required to provide disease freedoms below.

the occurrence of **foot-and-mouth** disease, infection with **rinderpest** virus,

infection with **Rift Valley fever** virus, **contagious bovine pleuropneumonia** or

lumpy skin disease, or of an **emerging disease relevant for bovine animals**;

for a period of at least 30 days prior to the date of first collection of the semen and during the collection period:

II.2.5.2. were kept on a **single establishment** where infection with **Brucella abortus, B. melitensis and B. suis, infection with Mycobacterium tuberculosis complex (M. bovis, M. caprae and M. tuberculosis), rabies, anthrax, surra (Trypanosoma evansi), enzootic bovine leukosis, infectious bovine rhinotracheitis/infectious pustular vulvovaginitis, bovine viral diarrhoea, [infection with epizootic haemorrhagic disease virus, infection with bluetongue virus (serotypes 1-24),]**

OCTOBER 2026: The conditions in purple above apply until 02/11/2026; after that establishment freedom from EHD and BTV is no longer required. Donor requirements remain and are laid out at II.2.8 and II.2.9.

bovine genital campylobacteriosis and trichomonosis have not been reported;

VSSPT for some diseases; aPVP to provide for movement history and diseases not covered in VSSPT.

For a period of at least 30 days prior to the date of first collection of the semen and during the collection period:

II.2.5.3. were not in contact with animals from establishments situated in a restricted zone due to the occurrence of diseases referred to in point II.2.5.1. or from establishments which do not meet the conditions referred to in point II.2.5.2.; aPVP to provide

II.2.5.4. were not used for natural breeding; aPVP to provide based on Operator Declaration/centre vet.

II.2.6. have been subjected to a quarantine for a period **of at least 28 days** in **quarantine accommodation,** where only other cloven-hoofed animals with at least the same health status were present, aPVP to provide

which on the day of their admission to the semen collection centre **aPVP must provide to DAERA for VSSPT: details of quarantine accommodation and date of entry to the SCC**

complied with the following conditions:

II.2.6.1. it **was not situated in a restricted zone established due to diseases referred to in point II.2.5.1.;** aPVP and VSSPT

II.2.6.2. **none of the diseases referred to in point II.2.5.2. has been reported for a period of at least 30 days;** aPVP and VSSPT

II.2.6.3. it was situated in an **area where foot-and-mouth disease has not been reported within a 10-km radius centred on the quarantine accommodation for a period of at least 30 days; VSSPT**

II.2.6.4. has had no outbreak of foot-and-mouth disease reported during a period of at **least 3 months preceding the date of admission of the animals into the semen collection centre; VSSPT; aPVP** must provide date of entry to SCC to DAERA to check 3 months preceding date of admission for VSSPT;

II.2.7. were kept in the **semen collection centre**

II.2.7.1. **which was not situated in a restricted zone established due to diseases referred to in point II.2.5.1.;** aPVP and VSSPT

II.2.7.2. where **none of the diseases referred to in point II.2.5.2. has been reported for a period of at least 30 days prior to the date of collection of the semen**, aPVP and VSSPT

and

(2)(3) ☐ [at least 30 days following the date of the collection;] delete for fresh; keep for frozen

(2)(4) ☐ [until the date of dispatch of the consignment of semen to another Member State;] Select for fresh/chilled

II.2.7.3. situated in an **area where foot-and-mouth disease has not been reported** within **a 10-km radius centred on the semen collection centre for a period of at least 30 days;** Request VSSPT

and

(The semen described in Part I is intended for artificial reproduction and was obtained from **donor animals which)**

II.2.8. comply with at least one of the following conditions as regards infection with **bluetongue** virus (serotypes 1-24): aPVP to select according to animal movement history

(2) ☐ either [II.2.8.1. they have been kept for a period of at least 60 days prior to and during collection

of the semen in a **Member State <u>or zone thereof</u> free from infection with**

bluetongue virus (serotypes 1-24) where no case of infection with bluetongue

virus (serotypes 1-24) has been confirmed during the last 24 months in the

targeted animal population.

This option not applicable as BTV was confirmed in NI 29/11/25.
Choose relevant option below; in effect, testing options.

(2) ☐ and/or [II.2.8.2. they have been kept in a seasonally disease-free zone, during the seasonally disease-free period, for a period of at least 60 days prior to and during collection of the semen, in a Member State or zone thereof with an approved eradication programme against infection with bluetongue virus (serotypes 1-24);]

(2) ☐ and/or [II.2.8.3. they have been kept in a seasonally disease-free zone, during the seasonally disease-free period, for a period of at least 60 days prior to and during collection of the semen, in a Member State or zone thereof where the competent authority of the place of origin of the consignment of semen has obtained the prior written consent of the competent authority of the Member State of destination to the conditions for establishment of that seasonally disease-free zone and to accept the consignment of semen;]

(2) ☐ and/or [II.2.8.4. they have been kept in a vector-protected establishment for a period of at least

60 days prior to and during collection of the semen;]

(2) ☐ and/or [II.2.8.5. they have been subjected to a serological test to detect antibodies to the bluetongue virus serogroup 1-24, with negative results, between 28 and 60 days from the date of each collection of the semen;]

(2) ☐ and/or [II.2.8.6. they have been subjected to an agent identification test for bluetongue virus (serotypes 1-24), with negative results, on blood samples taken at commencement and final collection of the semen and during collection of the semen at intervals of at least every 7 days, in the case of the virus isolation test, or of at least every 28 days, in the case of PCR;]

(The semen described in Part I is intended for artificial reproduction and was obtained from **donor animals which:)**

II.2.9. comply with at least one of the following conditions as regards infection with epizootic haemorrhagic disease virus (serotypes 1-7) (EHDV 1-7):

(2) ☐ either [II.2.9.1. they have been kept for a period of at least 60 days prior to and during collection of the semen in a **Member State or zone thereof where EHDV 1-7 has not been reported for a period of at least the preceding 2 years within a radius of 150 km of the establishment;]** **This option relevant if VSSPT assures EHD freedom; aPVP to confirm movement history.**

If NI not free of EHD, choose relevant option below:

(2) ☐ and/or [II.2.9.2. they have been kept in a vector-protected establishment for a period of at least 60 days prior to and during collection of the semen;]

(2) ☐ and/or [II.2.9.3. were resident in the Member State in which according to official findings the following serotypes of EHDV exist: and have been subjected with negative results in each case to the following tests carried out in an official laboratory:

(2) ☐ either [II.2.9.3.1. a serological test to detect antibodies to EHDV 1-7, with negative results, at least every 60 days throughout the collection period and between 28 and 60 days from the date of the final collection of the semen;]]

(2) ☐ and/or [II.2.9.3.2. an agent identification test for EHDV 1-7, with negative results, on blood samples taken at the commencement and final collection of the semen and during the collection of the semen at intervals of at least every 7 days, in the case of virus isolation test, or of at least every 28 days, in the case of PCR.

(The semen described in Part I is intended for artificial reproduction and was obtained from donor animals which:)

The aPVP certifies the statements below provided they have own knowledge or supporting veterinary certificate/s.

II.2.10 have been subjected to the following tests, carried out on blood samples taken within the period of 30 days prior to the commencement of the quarantine referred to in point II.2.6., with negative results, except for the bovine viral diarrhoea antibody test referred to in point II.2.10. required in accordance with point 1(b) of Chapter I of Part 1 of Annex II to Delegated Regulation (EU) 2020/686

II.2.10.5.2., required in accordance with point 1(b) of Chapter I of Part 1 of Annex II to Delegated Regulation (EU) 2020/686:

II.2.10.1. for infection with Mycobacterium tuberculosis complex (M. bovis, M. caprae and M. tuberculosis), an intradermal tuberculin test referred to in point 1 of Part 2 of Annex I to Delegated Regulation (EU) 2020/688;

II.2.10.2. for infection with Brucella abortus, B. melitensis and B. suis, a serological test referred to in point 1 of Part 1 of Annex I to Delegated Regulation (EU) 2020/688;

(2)(5) ☐ for enzootic bovine leukosis, a serological test referred to in point (a) of Part 4 of Annex I to Delegated Regulation (EU) 2020/688;]

II.2.10.4. for infectious bovine rhinotracheitis/infectious pustular vulvovaginitis, a serological test (whole virus) on a blood sample if the animals do not come from an establishment free from infectious bovine rhinotracheitis/infectious pustular vulvovaginitis;

II.2.10.5. for bovine viral diarrhoea:

II.2.10.5.1. a virus isolation test, a test for virus genome or a test for virus antigen, and

II.2.10.5.2. a serological test to determine the presence or absence of antibodies;

II.2.11. have been subjected to the following tests, carried out on blood samples taken within a period of at least 21 days, or 7 days in the case of the tests referred to in points II.2.11.4. and II.2.11.5., after the commencement of the quarantine referred to in point II.2.6., with negative results, except for the bovine viral diarrhoea antibody test referred to in point II.2.11.3.2., required in accordance with point 1(c) of Chapter I of Part 1 of Annex II to Delegated Regulation (EU) 2020/686:

II.2.11.1. for infection with Brucella abortus, B. melitensis and B. suis, a serological test referred to in point 1 of Part 1 of Annex I to Delegated Regulation (EU) 2020/688;

II.2.11.2. for infectious bovine rhinotracheitis/infectious pustular vulvovaginitis, a serological test (whole virus) on a blood sample;

II.2.11.3. for bovine viral diarrhoea:

II.2.11.3.1. a virus isolation test, a test for virus genome or a test for virus antigen, and

II.2.11.3.2. a serological test to determine the presence or absence of antibodies;

II.2.11.4. for bovine genital campylobacteriosis (Campylobacter fetus ssp. venerealis):

(2) ○ either [II.2.11.4.1

.

a single test carried out on a sample of artificial vagina washings or preputial specimen, in the case of animals less than 6 months old or kept since that age in a single sex group without contact with females prior to the quarantine referred to in point II.2.6.;]

(2) ○ or [II.2.11.4.2.

tests carried out on samples of artificial vagina washings or preputial specimens taken on three occasions at intervals of at least 7 days;]

II.2.11.5. for trichomonosis (Trichomonas foetus):

(2) ○ either [II.2.11.5.1. a single test carried out on a sample of preputial specimen, in the case of animals less than 6 months old or kept since that age in a single sex group without contact with females prior to the quarantine referred to in point II.2.6.;]

(2) ○ or [II.2.11.5.2. tests carried out on preputial specimens taken on three occasions at intervals of at least 7 days;]

II.2.12. have been subjected at semen collection centre, at least once a year, to the following compulsory routine tests, required in accordance with point 2 of Chapter I of Part 1 of Annex II to Delegated Regulation (EU) 2020/686:

II.2.12.1. for infection with Mycobacterium tuberculosis complex (M. bovis, M. caprae and M. tuberculosis), an intradermal tuberculin test referred to in point 1 of Part 2 of Annex I to Delegated Regulation (EU) 2020/688;

II.2.12.2. for infection with Brucella abortus, B. melitensis and B. suis, a serological test referred to in point 1 of Part 1 of Annex I to Delegated Regulation (EU) 2020/688;

II.2.12.3. for enzootic bovine leukosis, a serological test referred to in point (a) of Part 4 of Annex I to Delegated Regulation (EU) 2020/688;

II.2.12.4. for infectious bovine rhinotracheitis/infectious pustular vulvovaginitis, a serological test (whole virus) on a blood sample;

(2)(6) ☐

[II.2.12.5.

for bovine viral diarrhoea, a serological test for detection of an antibody;]

(2)(7) [II.2.12.6. ☐ for bovine genital campylobacteriosis (Campylobacter fetus ssp. venerealis), a test on a sample of preputial specimen;]

(2)(7) ☐

[II.2.12.7.

for trichomonosis (Trichomonas foetus), a test on a sample of preputial specimen;]

II.3. The semen described in Part I

II.3.1. has been collected, processed and stored in accordance with animal health requirements set out in points 1 and 2 of Part 1 of Annex III to Delegated Regulation (EU) 2020/686;

II.3.2. is placed in straws or other packages on which the mark is applied in accordance with requirements provided for in Article 10 of Delegated Regulation (EU) 2020/686 and that mark is indicated in Box I.30;

II.3.3. is transported in a container which:

II.3.3.1. was sealed and numbered prior to the dispatch from the semen collection centre under responsibility of the centre veterinarian, or by an official veterinarian, and the seal bears the number as indicated in Box I.19;

II.3.3.2. has been cleaned and either disinfected or sterilised before use, or is single-use container;

(2)(3) ☐

[II.3.3.3.

has been filled in with the cryogenic agent which not have been previously used for other products.]

II.4. The semen is preserved by the addition of antibiotics as follows:

II.4.1. The following antibiotic or mixture of antibiotics, effective in particular against campylobacters, leptospires and mycoplasmas, has been added to the semen after final dilution, or is contained in the used semen diluents, to reach the indicated concentration per ml of semen: Select appropriate option and if appropriate, describe the mixture used:

(2) ○ either [a mixture of gentamicin (250 µg), tylosin (50 µg) and lincomycin-spectinomycin (150/300 µg);]

(2) ○ or [a mixture of lincomycin-spectinomycin (150/300 µg), penicillin (500 IU) and streptomycin (500 µg);]

(2) ○ or [a mixture of amikacin (75 µg) and divekacin (25 µg);]

(2) ○ or [an antibiotic or a mixture of antibiotics (8) **Describe** , with a bactericidal activity at least equivalent to one of the following mixtures:

- gentamicin (250 µg), tylosin (50 µg) and lincomycin-spectinomycin (150/300 µg);
- lincomycin-spectinomycin (150/300 µg), penicillin (500 IU) and streptomycin (500 µg);
- amikacin (75 µg) and divekacin (25 µg).]

II.4.2. Immediately after the addition of the antibiotics, and before any possible freezing, the diluted semen was kept at a temperature of at least 5°C for a period of not less than 45 minutes, or under a time-temperature regime with a documented equivalent bactericidal activity.

Provided NI is given freedom from Lumpy Skin Disease on VSSPT, for donors from NI, delete both options below. For imported animals, check the incoming health certificate and select accordingly.

(2) ○ either [II.5

Germinal products (semen, ova and/or embryos, indicate as appropriate) obtained from bovine animals kept in vaccination zone I in relation to emergency protective vaccination against lumpy skin disease, in compliance with Article 13(3) of, and Annex IX, Part 3, point (3.4.1), to Commission Delegated

(2) ○ or [II.5

Germinal products (semen, ova and/or embryos, indicate as appropriate) obtained from bovine animals kept in vaccination zone II in relation to emergency protective vaccination against lumpy skin disease, in compliance with Article 13(3) of, and Annex IX, Part 3, point (3.4.2), to Commission Delegated Regulation (EU) 2023/361.

FOOTNOTES:

Part II:

(1) Only semen collection centres approved by the competent authority and included in the register

referred to in Article 101(1)(b) of Regulation (EU) 2016/429 and Article 7 of Delegated Regulation (EU)

2020/686.

(2) Delete if not applicable.

(3) Applicable for frozen semen.

(4) Applicable for fresh and chilled semen.

(5) Not applicable to animals which come from an establishment not free from enzootic bovine leukosis and which are less than 2 years of age as referred to in Article 20(2)(a) of Delegated Regulation (EU)

2020/686.

(6) Applicable only to seronegative animals.

Applicable only to bulls in semen production or having contact with bulls in semen production. Bulls

returning to collection after a lay-off period of more than 6 months shall be tested during a period of 30 days prior to resuming production.

(8) Insert the name(s) of the antibiotic(s) added and its (their) concentration or the commercial name of the semen diluent containing antibiotics.

Name (in capital letters)	Qualification and title
Local Control Unit name Local DVO name	Local Control Unit code Obtain from local DVO
Date	
Stamp	Signature