

I.	IDENTIFICATION OF SEMEN			
1)	Identification of donors			
	onor identity (species, Breed e, date of birth, number)	Number (in words) and code-mark of semen straws	Date of collection	Notes
2)	Identification of semen consignment			
	(Fresh, chilled, frozen semen)			
3)	Seal number of semen container			
II.	Origin and destination of semen			
1)	Name, address and approval number of Semen Collection Centre			
		•••••••		
2)	The semen consignment is to be sent from			
		of loading)		
	Directly to:			
	(place o	of destination)		
3)	Means of transportation			

(Indicate means of transport and registration marks)

#### 4) Name and address of consignor:

5) Name and address of consignee:

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# **III.** Health information

It the undersigned, certify that the semen described in Paragraph I, meets the following requirements:

- 1) the United Kingdom is free from foot and mouth disease and rinderpest in accordance with the Office International des Epizooties (OIE) Terrestrial Animal Health Code;
- 2) in the 30 day period prior to the bovine animals entering the isolation facility, they have been subjected to the following tests, with negative results:
  - a) an intradermal comparative tuberculin test;

  - c) an agar gel immunodiffusion test (AGIDT) or an enzyme linked immunosorbent assay (ELISA) for enzotic bovine leucosis;
  - d) a serum-neutralization test (SNT) or an enzyme linked immunosorbent assay (ELISA) for infectious bovine rhinotracheitis/infectious pustular vulvovaginitis (IBR/IPV);
  - e) a virus isolation test (fluorescent antibody test or immunoperoxidase test) for bovine viral diarrhoea. In the case of an animal less than six months old the test has been deferred until that age has been reached;
- 3) prior to entering the approved Semen Collection Centre, the animals have been subjected to a period of 30 days isolation in special accommodation approved by the competent veterinary authority and have been subjected to the following tests, with negative results:

  - b) an immunofluorescent antibody test or a culture test for Campylobacter foetus infection on a sample of preputial material;
  - c) microscopic examination and culture test for Trichomonas foetus on sample of preputial washings;
  - a serum neutalization test or an enzyme linked immunosorbent assay (ELISA) for infectious bovine rhinotracheitis/infectious pustular vulvo vaginitis (IBR/IPV);
- all tests have been carried out at a laboratory approved by the competent veterinary authority;
- 5) the semen described above is derived from donors which:
  - a) have not shown any disease symptoms on the day of collection;
  - b) have not been used for natural service while at the Centre;
  - c) have not been vaccinated against foot and mouth disease, infectious bovine rhinotracheitis/infectious pustular vulvo-vaginitis (IBR/IPV) and epizootic haemorrhagic disease (EHD);

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- d) originate from the centre of an area of 10 kilometres radius in which there has been no case of foot and mouth disease for at least 30 days prior to collection;
- e) originate from a centre under permanent veterinary supervision where bovine animals are tested periodically for tuberculosis, brucellosis, campylobacteriosis, infectious bovine rhinotracheitis/infectious pustular vulvo-vaginitis (IBR/IPV) and bovine leucosis by the competent veterinary authority;

in respect of bluetongue virus (BTV), the semen was obtained from donor bull(s) which comply with at least one of the following conditions:

- they were kept in a BTV free country or zone for a period of at least 60 days before commencement of, and during, collection of the semen; or
- they have been protected from attack from *Culicoides* likely to be competent BTV vectors for a period of at least 60 days before commencement of, and during, collection of the semen; or
- they have been subjected to a serological test according to the OIE Terrestrial Manual to detect antibodies to the BTV group, with negative results, at least every 60 days during the collection period and between 21 and 60 days following the final collection for this consignment; or
- \*iv. they have been subjected, with negative results, to an agent identification test for BTV according to the OIE Terrestrial Manual carried out on blood samples collected:

  (a) at commencement and final collection; and
  (b) during the period of semen collection for this consignment:

  i. at least every seven days, in the case of a virus isolation test,
  ii. at least every 28 days, in the case of a polymerase chain reaction (PCR) test;

#### 7) the semen described above:

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\*ii.

iii

- a) has been stored in an approved Semen Collection Centre for at least 30 days before dispatch;
- b) has been treated during processing by the addition of antibiotics to produce these concentrations in the final diluted semen:

### EITHER\*

not less than: 500 IU per ml streptomycin, 500 IU per ml penicillin, 150 µg per ml lincomycin, 300 µg per ml spectinomycin;

OR\*

an alternative combination of antibiotics with an equivalent effect against Campylobacters, Leptospires and Mycoplasmas, namely.....

c) has been transported in previously cleaned and disinfected container which before dispatch was officially sealed and placed in approved storage.

# \* Delete as appropriate

Official Stamp

Signed ..... RCVS Official Veterinarian

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Name in block letters

Date	Address
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