Part I: Description of consignment

I.1. Consignor
Name
Address
Country ISO Code

I.2. IMSOC reference

I.2.a. Local reference

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Consignee
Name
Address
Country ISO Code

I.6. Operator conducting assembly operations independently of an establishment
Name
Address
Approval Number
Country ISO Code

I.7. Country of origin ISO Code

I.9. Country of destination ISO Code

I.8. Region of origin Code

I.10. Region of destination Code

I.11. Place of dispatch
Name
Address
Approval Number
Country ISO Code

I.12. Place of destination
Name
Address
Approval Number
Country ISO Code

I.13. Place of loading
Name
Address
Approval Number
Country ISO Code

I.14. Date and time of departure

I.15. Means of Transport

Mode	International transport document	Identification

I.16. Transporter
Name
Address
Activity ID
Country ISO Code

I.17. Accompanying documents
Commercial document reference
Date of issue
Country
Place of issue

I.18. Transport conditions
Frozen ☐ Chilled ☐ Ambient ☐

I.19. Container No / Seal No

I.20. Certified as
Germinal products ☐

I.21. For transit through a third country ☐
Third country ISO Code
Exit point BCP code
Entry point BCP code

I.22. For transit through Member State(s) ☐
Member State ISO Code

I.23. For export ☐
Third country ISO Code
Exit point BCP code

I.25. Journey Log

I.26. Total number of packages

I.27. Total quantity

I.28. Total gross weight

I.30. Description of consignment

Commodity	Species	Identification Number	Quantity	Nature of commodity

Identification Mark	Package count	Date of collection / production	Plant / Establishment / Centre

Part II: Certification

II. Health information

II.1. Animal health attestation

I, the undersigned official veterinarian, hereby certify that:

II.1.1. The semen described in Part I was collected before the date of 31 December 2004 on a semen collection centre(1) which:

(a) was approved under conditions laid down in Chapter I of Annex A to Council Directive 88/407/EEC;

(b) was operated and supervised under conditions laid down in Chapter II of Annex A to Directive 88/407/EEC.

II.1.2. At the time the semen described in Part I was collected, all bovine animals at the semen collection centre:

(a) came from herds and/or were born to dams which satisfy the conditions of points 1(b) and (c) in Chapter I of Annex B to Directive 88/407/EEC;

(b) have, within the 30 days preceding the quarantine isolation period, undergone, with negative results:

- the tests referred to in points 1(d)(i), (ii) and (iii) of Chapter I of Annex B to Directive 88/407/EEC, and
- a serum neutralization test or ELISA test for infectious bovine rhinotracheitis/infectious pustular vulvovaginitis, and
- a virus isolation test (fluorescent antibody test or immunoperoxidase test) for bovine viral diarrhoea, which in the case of an animal less than six months of age has been deferred until that age was reached;

(c) have satisfied the quarantine isolation period of 30 days and have been subjected with the required negative results to the following health tests:

- a serological test for brucellosis carried out in accordance with the procedure described in Annex C to Directive 64/432/EEC;
- either an immunofluorescent antibody test or a culture test for Campylobacter fetus infection on a sample of preputial material or artificial vagina washings, or, in the case of a female animal, a vaginal mucus agglutination test;
- a microscopic examination and culture test for Trichomonas foetus on a sample of preputial material or artificial vagina washings, or in case of a female animal a vaginal mucus agglutination test;

(d) have undergone, at least once a year, with negative results, the routine tests referred to in points 1(a), (b) and (c) in Chapter II of Annex B to Directive 88/407/EEC.

II.1.3. At the time the semen described in Part I was collected,

(a) all female bovine animals in the centre have undergone, at least once a year, a vaginal mucus agglutination test for Campylobacter fetus infection with negative results, and

(b) all bulls used for semen production have undergone with negative result either an immunofluorescent antibody test or a culture test for Campylobacter fetus infection on a sample of preputial material or artificial vagina washings carried out within 12 months prior to collection.

II.1.4. The semen described in Part I was collected from bulls standing in a semen collection centre in which:

(2) ○ either [all bovine animals have not been vaccinated against infectious bovine rhinotracheitis and have undergone at least once a year with negative result a serum neutralisation test or an ELISA test for infectious bovine rhinotracheitis/infectious pustular vulvovaginitis;]

Part II: Certification

II. Health information

(2) ○ or [bovine animals not vaccinated against infectious bovine rhinotracheitis have undergone, at least once a year, with negative result a serum neutralisation test or ELISA test for infectious bovine rhinotracheitis/infectious pustular vulvovaginitis, and testing for infectious bovine rhinotracheitis is not carried out on bulls which have received a first vaccination against infectious bovine rhinotracheitis at the insemination centre after they have been tested with negative result in a serum neutralisation test or ELISA test for infectious bovine rhinotracheitis/infectious pustular vulvovaginitis and which since the first vaccination have been regularly re-vaccinated with an interval of not more than six months;].

II.1.5. The semen described in Part I was collected from bulls which:

II.1.5.1.

(2) ○ either [have not been vaccinated against foot-and-mouth disease within 12 months prior to collection;]

(2) ○ or [have been vaccinated against foot-and-mouth disease less than 12 months and more than 30 days prior to collection, and 5% of doses of the semen from each collection, with a minimum of five straws, have been submitted to a virus isolation test for foot-and-mouth disease, carried out with negative results in the laboratory ((3)), situated in or designated by the Member State of destination;]

II.1.5.2.

(2) ○ either [have not been vaccinated against infectious bovine rhinotracheitis,]

(2) ○ or [have been vaccinated against infectious bovine rhinotracheitis in accordance with point II.1.4,].

II.1.6. The semen described in Part I was stored in approved conditions for a minimum period of 30 days immediately following collection(4).

II.1.7. The semen described in Part I was sent to the place of loading in a sealed container and bearing the number detailed in Box I.19.

Notes

This animal health certificate shall be completed according to the notes for the completion of certificates provided for in Chapter 2 of Annex I to Commission Implementing Regulation (EU) 2020/2235.

Part I:

Box I.11: Place of dispatch shall correspond to the semen collection centre (as defined in Article 2(b) first indent of Directive 88/407/EEC) where the semen was collected.

Box I.12: Place of destination shall correspond to the semen collection or storage centre (as defined in Article 2(b) of Directive 88/407/EEC), or to the holding of semen destination.

Box I.19: Identification of container and Seal number shall be indicated.

Box I.30: Donor identity shall correspond to the official identification of the animal.

Date of collection shall be indicated in the following format: dd/mm/yyyy and shall be earlier than 31 December 2004.

Approval number of the centre shall correspond to the approval number of the semen centre indicated in Box I.11. where the semen was collected.

Part II:

(1) Only semen collection centres approved by the competent authority and listed in accordance with Article 5(2) of Council Directive 88/407/EEC.

(2) Delete as appropriate.

(3) Name of the laboratory.

(4) May be deleted for fresh semen.

Certifying Officer/Official veterinarian

Name (in capital letters) Authority name

Date of signature Signature

Stamp