EUROPEAN UNION

INTRA

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.8. Region of origin Code

I.9. Country of destination ISO 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
Approval Number
Country ISO Code

I.17. Accompanying documents
Document Type
Accompanying document reference
Date of Issue
Country
Place of issue

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

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.24. Estimated journey time

I.25. Journey Log

I.26. Total number of packages

I.27. Total quantity

I.28. Total gross weight

I.30. Description of consignment

1. 05 PRODUCTS OF ANIMAL ORIGIN, NOT ELSEWHERE SPECIFIED OR INCLUDED

0511 Animal products not elsewhere specified or included; dead animals of Chapter|1|or 3, unfit for human consumption

051110 Bovine semen

05111000 Bovine semen

Part I: Description of consignment

#1.	Commodity	Identification Number	Quantity	Nature of commodity	Identification Mark
Species		Package count	Date of collection / production	Plant / Establishment / Centre	

Part II: Certification

II. Health information

I, the undersigned official veterinarian, hereby certify that:

II.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.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 6 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.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.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.]

(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 6 months.]

II.5. The semen described in Part I was collected from bulls which:

II.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

Part II: Certification

II. Health information

the laboratory ( ) (3), situated in or designated by the Member State of destination;]

II.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.6. The semen described in Part I was stored in approved conditions for a minimum 30 days immediately following collection (4).

II.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.

II.8. The semen described in Part I is dispatched from:

(2) ○ either [a semen collection center or a zone not subject to movement restrictions affecting bovine animals and established for reasons of listed diseases relevant for those species or diseases subject to emergency measures relevant for those species or those restrictions do not apply to this semen because it was collected before the restrictions were established, and the semen has not been in contact with other semen of a lower health status for an adequate period.]

(2) ○ or [a semen collection center or a zone subject to movement restrictions affecting bovine animals and established for (5), but derogations from movement restrictions have been granted, and:

(2) ☐ [it complies with the requirements set out in (6);]]

(2) ☐ [and in particular, it is (7).]]

Notes

In accordance with the Agreement on the withdrawal of the United Kingdom of Great Britain and Northern Ireland from the European Union and the European Atomic Energy Community, and in particular Article 5(4) of the Protocol on Ireland/Northern Ireland in conjunction with Annex 2 to that Protocol, references to the Union in this animal health certificate include the United Kingdom in respect of Northern Ireland.

This animal health certificate shall be completed in accordance with 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 reference I.11: Place of dispatch shall correspond to the semen collection centre (as defined in Article 2, second paragraph, point (b), first indent, of Directive 88/407/EEC) where the semen was collected.

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

Box reference I.19: Seal number shall be indicated.

Box reference I.26: Total number of packages shall correspond to the number of containers.

Box reference 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 Directive 88/407/EEC.

(2) Delete if not applicable.

II. Health information		

(3) Name of the laboratory.

(4) May be deleted for fresh semen.

(5) Insert the name of the disease(s).

(6) Insert the specific reference to the article(s), title, and number of the relevant legal act(s) adopted by the Commission providing for those requirements.

(7) Insert the specific attestation(s) provided for in and required by the relevant legal act(s) adopted by the Commission, as referred to in Article 159(2), points (a), (b) and (c), of Regulation (EU) 2016/429 of the European Parliament and of the Council.

Part II: Certification

Certifying Officer/Official veterinarian	
Name (in capital letters)	Qualification and title
Date of declaration	Signature
Stamp	