**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 the semen described in Part I:

II.1. was collected, processed and stored in a semen collection centre (1) approved and supervised by the competent authority in accordance with Chapter I(1) and Chapter II(1) of Annex A to Directive 88/407/EEC;

II.2. was collected from bulls, which:

II.2.1. meet the requirements of Chapters I and II of Annex B to Directive 88/407/EEC,

(2) ○ either II.2.2. [have not been vaccinated against foot and mouth disease within 12 months prior to collection;]

(2) ○ or II.2.2. [have been vaccinated against foot and mouth disease less than 12 months and more than 30 days prior to the collection, and 5 % of doses of semen of each collection, with a minimum of five straws, have been submitted to a virus isolation test for foot-an-mouth disease, carried out with negative results in the laboratory ( ) (3) situated in or designated by the Member State of destination;]

II.3. was collected, processed, stored and transported under conditions which comply with the standards laid down in Annex C to Directive 88/407/EEC;

II.4. was stored in approved conditions for a minimum period of 30 days immediately following collection (4).

II.5. 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.

**Part II: Certification**

| II. Health information | | |
|---|---|---|

Approval number of the centre shall correspond to the approval number of the semen centre indicated in box I.12 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.

(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.

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