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.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
Ambient ☐ Chilled ☐ 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.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	Type

Part II: Certification

II. Health information

I, the undersigned official veterinarian, hereby certify that:

(1) ☐ [II.1. The semen of ovine(1)/ caprine(1) animals described in Part I has been collected, processed and stored, and dispatched from the semen collection centre(2) which

II.1.1. is approved and kept in a register by the competent authority;

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.]

(1) ☐ [II.1. The semen of ovine(1)/ caprine(1) animals described in Part I has been collected, processed and stored, and dispatched from the establishment where the donor animals are kept as referred to in Article 13 of Delegated Regulation (EU) 2020/686, and

II.1.1. the operator obtained the prior consent of the competent authority of the Member State of destination to accept the consignment;

II.1.2. the donor animals have been clinically examined by a veterinarian prior to semen collection;

II.1.3. the operator keeps records at the establishment which include at least the information provided for in Article 8(1)(a) of Delegated Regulation (EU) 2020/686.]

(1) ○ either [II.1.4. was collected from animals which have been kept continuously since birth on a holding or holdings recognised as having a negligible or controlled risk of classical scrapie according to point 1 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001, except during the period when they were kept at a semen collection centre that complied during that period with the conditions set out in the four indents of point 1.3.(c)(iv) of that Section;]

(1) ○ or [II.1.4. was collected from animals which have been kept continuously for the last three years before the collection on a holding or holdings which has/have complied for the last three years before the collection with the requirements laid down in points 1.3. (a) to (f) of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001, except during the period when they were kept at a semen collection centre that complied during that period with the conditions set out in the four indents of point 1.3.(c)(iv) of that Section;]

(1) ○ or [II.1.4. was collected from animals which have been kept continuously since birth in a Member State or zone of a Member State listed in point 2.3. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001 as having a negligible risk status for classical scrapie;]

(1) ○ or [II.1.4. was collected from ovine animals of the ARR/ARR prion protein genotype;]

II.2. 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;

II.2.2. come, before the commencement of the quarantine referred to in point II.2.6., from establishments in 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, and

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

(1) ○ 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 Brucella abortus, B. melitensis and B. suis and have never been kept previously in any establishment of a lower health status;

(1)(3) ☐ [II.2.2.3. in which infection with Mycobacterium tuberculosis complex (M. bovis, M. caprae and M. tuberculosis) has not been reported during the last 42 days;]

Part II: Certification

II. Health information

(1)(4) ☐ [II.2.2.3. in which surveillance for infection with Mycobacterium tuberculosis complex (M. bovis, M. caprae and M. tuberculosis) has been carried out on the caprine animals kept on the establishments during at least the 12 month period, as referred to in Article 15(3) of Commission Delegated Regulation (EU) 2020/688, and in case, during this period, infection with Mycobacterium tuberculosis complex (M. bovis, M. caprae and M. tuberculosis) has been reported in caprine animals kept on the establishment, measures were taken in accordance with Part 1(3) of Annex II to that Delegated Regulation;]

II.2.2.4. in which surra (Trypanosoma evansi) has not been reported during the 30 day period, and

(1) ○ either [surra has not been reported in the establishments during the last 2 years;]

(1) ○ 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 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;]

(1)(3) ☐ [II.2.2.5. in which ovine epididymitis (Brucella ovis) has not been reported during the 12 month period;]

(1)(8) ☐ [II.2.2.6. where, during the period of 60 days prior to their stay in the quarantine accommodation referred to in point II.2.6., they have been subjected, with negative results, to a serological test for ovine epididymitis (Brucella ovis), or any other test for ovine epididymitis (Brucella ovis) of an equivalent documented sensitivity and specificity, as required in accordance with point 1(b) of Chapter I of Part 3 of Annex II to Delegated Regulation (EU) 2020/686;]

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;

II.2.4. are individually identified as provided for in Article 45(2) or (4), or Article 46(1) of Commission Delegated Regulation (EU) 2019/2035;

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

II.2.5.1. were kept on establishments not situated in a restricted zone established due to the occurrence of foot-and-mouth disease, infection with rinderpest virus, infection with Rift Valley fever virus, infection with peste des petits ruminants virus, sheep pox and goat pox or contagious caprine pleuropneumonia, or of an emerging disease relevant for ovine and caprine animals;

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), infection with epizootic haemorrhagic disease virus, infection with bluetongue virus (serotypes 1-24) and, in case of ovine animals and those caprine animals which are kept together with ovine animals, ovine epididymitis (Brucella ovis) have not been reported;

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.;

II.2.5.4. were not used for natural breeding;

Part II: Certification

II. Health information

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, which on the day of their admission to the semen collection centre 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.;

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;

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;

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;

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.;

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, and

(1)(3) ☐ [at least 30 days following the date of the collection;]

(1)(4) ☐ [until the date of dispatch of the consignment of semen to another Member State;]

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; and

(1)(3) ☐ [free from foot-and-mouth disease for a period of at least 3 months prior to the date of collection of the semen and 30 days from the date of collection;]

(1)(4) ☐ [free from foot-and-mouth disease for a period of at least 3 months prior to the date of collection of the semen and until the date of dispatch of the consignment of semen to another Member State and the donor animals have been kept at that semen collection centre for a continuous period of at least 30 days immediately prior to the date of collection of the semen;]

II.2.8. comply with at least one of the following conditions as regards infection with bluetongue virus (serotypes 1-24):

(1) ☐ 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 or zone thereof 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;]

(1) ☐ 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);]

(1) ☐ 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;]

(1) ☐ 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;]

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II. Health information

(1) ☐ 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;]

(1) ☐ 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;]

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):

(1) ☐ 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;]

(1) ☐ 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;]

(1) ☐ 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:

(1) ☐ 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;]]

(1) ☐ 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.]]

(1)(5) ☐ [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, required in accordance with point 1(c) of Chapter I of Part 3 of Annex II to Delegated Regulation (EU) 2020/686:

II.2.10.1. 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;

(1)(8) ☐ [II.2.10.2. for ovine epididymitis (Brucella ovis), a serological test or any other test with an equivalent documented sensitivity and specificity;]

II.2.11. have been subjected to the following tests, carried out on blood samples taken within a period of at least 21 days after the commencement of the quarantine referred to in point II.2.6., with negative results, required in accordance with point 1(d) of Chapter I of Part 3 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;

(1)(8) ☐ [II.2.11.2. for ovine epididymitis (Brucella ovis), a serological test or any other test with an equivalent documented sensitivity and specificity;]

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 3 of Annex II to Delegated Regulation (EU) 2020/686:

II.2.12.1. 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;

(1)(8) ☐ [II.2.12.2. for ovine epididymitis (Brucella ovis), a serological test or any other test with an equivalent documented sensitivity and specificity.]]

(1)(9) ☐ [II.2.13. have been subjected to the following tests, carried out on blood samples taken within the period of 30 days prior to collection of the semen, with negative results:

Part II: Certification

II. Health information

II.2.13.1. 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;

(1)(8) ☐ [II.2.13.2. for ovine epididymitis (Brucella ovis), a serological test or any other test with an equivalent documented sensitivity and specificity;]]

II.3. The semen described in Part I

(1)(5) ☐ [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;

(1)(6) ☐ [II.3.3.3. has been filled in with the cryogenic agent which not have been previously used for other products.]

(1)(10) ☐ [II.4. The semen is preserved by the addition of antibiotics as follows:

II.4.1. The following antibiotic or mixture of antibiotics 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:

(1) ○ either [gentamicin (250 µg);]

(1) ○ or [a mixture of penicillin (500 IU) and streptomycin (500 µg);]

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

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

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

(1) ○ or [an antibiotic or a mixture of antibiotics(11) , with a bactericidal activity at least equivalent to one of the following mixtures:

- gentamicin (250 µg);
- penicillin (500 IU) and streptomycin (500 µg);
- 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.]

Part II: Certification

II. Health information		

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 European Union in this certificate include the United Kingdom in respect of Northern Ireland.

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 reference I.11:	"Place of dispatch": Indicate the unique approval number and the name and address of the semen collection centre or, in case of an establishment as referred in Article 13 of Delegated Regulation (EU) 2020/686, the unique registration number and address of the establishment of dispatch of the consignment of semen.
Box reference I.12:	"Place of destination": Indicate the address and unique registration or approval number of the establishment of destination of the consignment of semen.
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:	"Type": Indicate semen.
	"Species": Select amongst "Ovis aries" or "Capra hircus" as appropriate.
	"Identification number": Indicate the identification number of each donor animal.
	Identification mark: Indicate the mark on the straw or other packages where semen of the consignment is placed.
	"Date of collection/production": Indicate the date on which semen of the consignment was collected.
	"Approval or registration number of plant/establishment/centre": Indicate the unique approval number of the semen collection centre or, in the case of an establishment as referred in Article 13 of Delegated Regulation (EU) 2020/686, the unique registration number of the establishment where the semen was collected.
	"Quantity": Indicate the number of straws or other packages with the same mark.
	"Type": Indicate for BTV-test: II.2.8.5. and/or II.2.8.6., and/or for EHD-test: II.2.9.3.1. and/or II.2.9.3.2., if relevant.

Part II:

(1)	Delete if not applicable.
(2)	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.
(3)	Applicable for ovine animals.
(4)	Applicable for caprine animals.
(5)	Applicable for semen collected at a semen collection centre.
(6)	Applicable for frozen semen.
(7)	Applicable for fresh and chilled semen.
(8)	Applicable for ovine animals and for those caprine animals which are kept together with ovine animals.
(9)	Applicable for semen collected at an establishment where the donor animals are kept as referred to in Article 13 of Delegated Regulation (EU) 2020/686.
(10)	Mandatory attestation in case antibiotics were added.

Part II: Certification

II. Health information		

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

Certifying Officer/Official veterinarian

Name (in capital letters)	Authority name
Date of signature	Signature
Stamp	