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

051199 Other

05119985 Other

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	Type

Part II: Certification

II. Health information		

II. Health information

I, the undersigned official veterinarian, hereby certify that:

II.1. The semen of porcine animals described in Part I:

II.1.1. has been collected, processed and stored, and dispatched from the semen collection centre (1) which is approved and kept in a register by the competent authority;

II.1.2. has been collected, processed and stored, and dispatched from the semen collection centre which 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.

(2) ○ either [II.1.3. is dispatched from the semen collection centre or a zone not subject to movement restrictions affecting porcine 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 [II.1.3. is dispatched from the semen collection centre or a zone subject to movement restrictions affecting porcine animals and established for (3), but derogations from movement restrictions have been granted, and:

(2) ☐ [it complies with the requirements set out in (4);]]

(2) ☐ [and in particular, it is (5).]]

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, prior to the commencement of the quarantine referred to in point II.2.8, 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 at least 30 days and in which foot and mouth disease has not been reported during at least 3 months, and:

(2) ○ either [the donor animals were not vaccinated against foot and mouth disease;]

(2) ○ or [the donor animals were vaccinated against foot and mouth disease during 12 months prior to the date of collection of the semen but not during 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 the donor animals 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 in accordance with the requirements laid down in Part 5, Chapter IV, of Annex II to Delegated Regulation (EU) 2020/686;

II.2.2.3. where no clinical, serological, virological or pathological evidence of infection with Aujeszky's disease virus had been detected during at least 12 months;

II.2.2.4. where, during at least 3 months, no animal was vaccinated against infection with porcine reproductive and respiratory syndrome virus and no infection with porcine reproductive and respiratory syndrome virus was detected;

II.2.3. did not show symptoms or clinical signs of transmissible animal diseases on the date of their admission to a semen collection centre and on the date of collection of the semen;

II.2.4. are identified as provided for in Article 52 or Article 54(2) of Commission Delegated Regulation (EU) 2019/2035;

II.2.5. for 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 in establishments situated in a zone not subject to movement restrictions established due to the occurrence of foot and mouth disease, infection with rinderpest virus, classical swine fever or African swine fever, or of an emerging disease relevant for porcine animals;

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II. Health information		

II.2.5.2. were kept in a single establishment where infection with Brucella abortus, B. melitensis and B. suis, infection with rabies virus, anthrax, infection with Aujeszky's disease virus and infection with porcine reproductive and respiratory syndrome virus have not been reported;

II.2.5.3. were not in contact with animals from establishments situated in a zone subject to movement restrictions 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;

II.2.6. have been subjected to a quarantine for 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 date of their admission to the semen collection centre complied with the following conditions:

II.2.6.1. it was situated in a zone not subject to movement restrictions established due to diseases referred to in point II.2.5.1 or it was under derogation as referred to in point II.1.3., if applicable;

II.2.6.2. none of the diseases referred to in point II.2.5.2 has been reported for 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 at least 30 days;

II.2.6.4. has had no outbreak of foot and mouth disease reported during at least 3 months preceding the date of admission of the animals into the semen collection centre;

II.2.6.5. it was free from infection with Brucella abortus, Brucella melitensis and Brucella suis for at least 3 months;

II.2.7. were kept in the semen collection centre:

II.2.7.1. which was situated in a zone not subject to movement restrictions established due to diseases referred to in point II.2.5.1 or it was under derogation as referred to in point II.1.3, if applicable;

II.2.7.2. where none of the diseases referred to in point II.2.5.2 has been reported for at least 30 days prior to the date of collection of the semen, and:

(2) (6) ○ either [at least 30 days following the date of collection;]

(2) (7) ○ or [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 at least 30 days; and

(2) (6) ○ either [free from foot and mouth disease for at least 3 months prior to the date of collection of the semen and 30 days from the date of collection;]

(2) (7) ○ or [free from foot and mouth disease for 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.2.7.4. where no clinical, serological, virological or pathological evidence of infection with Aujeszky's disease virus had been reported for a period comprising at least 30 days prior to the date of admission and at least 30 days immediately prior to the date of collection of the semen;

II.2.8. have been subjected to the following tests, carried out within 30 days prior to the date of commencement of the quarantine referred to in point II.2.6, with negative results, required in accordance with Part 2, Chapter I, point 1(b), of Annex II to Delegated Regulation (EU) 2020/686:

II.2.8.1. as regards infection with Brucella abortus, B. melitensis and B. suis, a buffered Brucella antigen test (rose Bengal test), a competitive ELISA or an indirect ELISA for the detection of antibodies to smooth Brucella species;

II.2.8.2. as regards infection with Aujeszky's disease virus:

(2) ○ either [in the case of non-vaccinated animals, an ELISA to detect antibodies to the whole Aujeszky's disease virus or to glycoprotein B (ADV-gB) or glycoprotein D (ADV-gD) of the virus or a serum neutralisation test;]

(2) ○ or [in the case of animals vaccinated with a gE deleted vaccine, an ELISA to detect antibodies to glycoprotein E (ADV-gE) of Aujeszky's disease virus;]

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(2) ☐ [II.2.8.3. as regards classical swine fever, an antibody ELISA or serum neutralisation test, in case of animals coming from a Member State or zone thereof where classical swine fever has been reported or vaccination against this disease has been practiced for the preceding 12 months;]

II.2.8.4. as regards infection with porcine reproductive and respiratory syndrome virus, a serological test (the immunoperoxidase monolayer assay (IPMA), immunofluorescence assay (IFA), or ELISA);

II.2.9. have been subjected to the following tests, carried out on samples taken at least 21 days after the date of the commencement of the quarantine referred to in point II.2.6, with negative results, required in accordance with Part 2, Chapter I, point 1(c), of Annex II to Delegated Regulation (EU) 2020/686:

II.2.9.1. as regards infection with Brucella abortus, B. melitensis and B. suis, a buffered Brucella antigen test (rose Bengal test), a competitive ELISA or an indirect ELISA for the detection of antibodies to smooth Brucella species;

II.2.9.2. as regards infection with Aujeszky's disease virus:

(2) ○ either [in the case of non-vaccinated animals, an ELISA to detect antibodies to the whole Aujeszky's disease virus or to glycoprotein B (ADV-gB) or glycoprotein D (ADV-gD) of the virus or a serum neutralisation test;]

(2) ○ or [in the case of animals vaccinated with a gE deleted vaccine, an ELISA to detect antibodies to glycoprotein E (ADV-gE) of Aujeszky's disease virus;]

II.2.9.3. as regards infection with porcine reproductive and respiratory syndrome virus, a serological test (IPMA, IFA, or ELISA) and a test for virus genome (reverse-transcription polymerase chain reaction (RT-PCR), nested set RT-PCR, real-time RT-PCR);

II.2.10. have been subjected, at semen collection centre, to the following compulsory routine tests, required in accordance with Part 2, Chapter I, point 2(a), of Annex II to Delegated Regulation (EU) 2020/686:

II.2.10.1. as regards infection with Brucella abortus, B. melitensis and B. suis, a buffered Brucella antigen test (rose Bengal test), a competitive ELISA or an indirect ELISA for the detection of antibodies to smooth Brucella species;

II.2.10.2. as regards infection with Aujeszky's disease virus:

(2) ○ either [in the case of non-vaccinated animals, an ELISA to detect antibodies to the whole Aujeszky's disease virus or to glycoprotein B (ADV-gB) or glycoprotein D (ADV-gD) of the virus or a serum neutralisation test;]

(2) ○ or [in the case of animals vaccinated with a gE deleted vaccine, an ELISA to detect antibodies to glycoprotein E (ADV-gE) of Aujeszky's disease virus;]

(2) ☐ [II.2.10.3. as regards classical swine fever, an antibody ELISA or serum neutralisation test, in case of animals coming from a Member State or zone thereof where classical swine fever has been reported or vaccination against this disease has been practiced for the preceding 12 months;]

II.2.10.4. as regards infection with porcine reproductive and respiratory syndrome virus, a serological test (IPMA, IFA, or ELISA);

II.2.11. have been subjected to the tests referred to in point II.2.10 carried out, in accordance with Part 2, Chapter I, point 2(b), of Annex II to Delegated Regulation (EU) 2020/686, on samples taken from:

(2) ○ either [all animals immediately prior to the date of departure from the semen collection centre, or upon arrival at the slaughterhouse, and in no case later than 12 months from the date of admission to the semen collection centre.]

(2) ○ or [at least 25 % of the animals in the semen collection centre every 3 months to test for infection with Brucella abortus, Brucella melitensis and Brucella suis, infection with Aujeszky's disease virus and classical swine fever and from at least 10 % of the animals in the semen collection centre every month to test for infection with porcine reproductive and respiratory syndrome virus.]

(2) ○ or [at least 10 % of the animals in the semen collection centre every month to test for infection with Brucella abortus, Brucella melitensis and Brucella suis, infection with Aujeszky's disease virus, classical swine fever and infection with porcine reproductive and respiratory syndrome virus.]

II.3. The semen described in Part I:

II.3.1. has been collected, processed and stored in accordance with animal health requirements set out in Part 1, points 1 and 2, of Annex III to Delegated Regulation (EU) 2020/686;

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II. Health information		

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 date of 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;

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

(2) ☐ [II.4. Where an antibiotic or a mixture of antibiotics was added to the semen described in Part I:

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:

(8),

II.4.2. Immediately after the addition of the antibiotic(s), and before any possible freezing, the diluted semen was kept at a temperature of at least 5°C or 15°C for not less than 45 minutes, or under a time-temperature regime with a documented equivalent bactericidal activity.]

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 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 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”: Semen.

“Identification number”: Indicate identification number of each donor animal.

“Identification mark”: Indicate mark on the straw or other packages where the semen of the consignment is placed.

“Date of collection/production”: Indicate the date on which the semen of the consignment was collected.

“Approval or registration number of plant/establishment/centre”: Indicate the unique approval number of the semen collection centre where the semen of the consignment was collected.

“Quantity”: Indicate number of straws or other packages with the same mark.

Part II:

(1) Only semen collection centres approved by the competent authority and included in the register referred to in Article 101(1), point (b), of Regulation (EU) 2016/429 of the European Parliament and of

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the Council and Article 7 of Delegated Regulation (EU) 2020/686.

(2) Delete if not applicable.

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

(4) 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.

(5) 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.

(6) Applicable for frozen semen.

(7) Applicable for fresh and chilled semen.

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

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