Part I : Details of consignment

I.1. Consignor
Name
Address
Country ISO Code

I.2. IMSOC Reference
Specimen not to be used for exports from EU

I.2.a. Local Reference

I.5. Consignee
Name
Address
Country ISO Code

I.3. Central competent authority

I.4. Local competent authority

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 Entry Point

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

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

I.19. Container No / Seal No

I.20. Certified as

Transhumance ☐	Pollination ☐	Animal Feedingstuff ☐	Pharmaceutical use ☐
Game Restocking ☐	Fattening ☐	Registered equidae ☐	Quarantine ☐
Artificial reproduction ☐	Training ☐	Further process ☐	Racing ☐
Production of petfood ☐	Breeding and production ☐	Consignments according to Regulation No 999/2001 ☐	Competition ☐
Storage ☐	Breeding ☐	Ornamental use/research ☐	Laboratory ☐
Rodent food ☐	Pets ☐	Other ☐	Production ☐
Technical use ☐	Ornamental bird food ☐	Relaying ☐	Slaughter ☐
Approved Bodies ☐	Human consumption ☐	Pet food ☐	Sales ☐
Organic fertilizers ☐	Unregistered equidae ☐	Circus exhibition ☐	Category 3 fish oil/fish meal for detoxification according to Regulation 2015/786 ☐

I.21. For transit through a third country ☐
Country ISO Code
EU Exit Authority BCP code
EU Entry Authority BCP code

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

I.24. Total quantity

I.25. Total gross weight

Part I : Details of consignment

I.28. 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

Commodity	Species	Identification number	Identification mark	Nature of commodity

Quantity	Date of collection/production	Manufacturing plant

Part II: Certification

II. Health information

I, the undersigned official veterinarian, hereby certify that :

II.1. _____ (name of exporting country or part thereof)(2)

was free from rinderpest and foot-and-mouth disease during the 12 months immediately prior to collection of the semen for export and until its date of dispatch to Great Britain and no vaccination against these diseases has taken place during the same period.

II.2. The centre(3) described in Box. I.11. at which the semen to be exported was collected:

II.2.1. meets the conditions laid down in Chapter I(1) of Annex A to Directive 88/407/EEC;

II.2.2. is operated and supervised in accordance with the conditions laid down in Chapter II(1) of Annex A to Directive 88/407/EEC.

II.3. The centre at which the semen to be exported was collected was free from rabies, tuberculosis, brucellosis, anthrax and contagious bovine pleuropneumonia during 30 days prior to the date of collection of the semen to be exported and the 30 days after collection (in the case of fresh semen until the day of dispatch to Great Britain).

II.4. The bovine animals standing at the semen collection centre:

(8)II.4.1. come from herds which satisfy the conditions of paragraph 1(b) of Chapter I of Annex B to Directive 88/407/EEC;

II.4.2. come from herds or were born to dams which comply with the conditions of paragraph 1(c) of Chapter I of Annex B to Directive 88/407/EEC, or were tested at the age of at least 24 months in accordance with paragraph 1(c) of Chapter II of Annex B to that Directive;

II.4.3. underwent the tests required in accordance with paragraph 1(d) of Chapter I of Annex B to Directive 88/407/EEC in the 28 days preceding the quarantine isolation period;

II.4.4. have satisfied the quarantine isolation period and testing requirements laid down in paragraph 1(e) of Chapter I of Annex B to Directive 88/407/EEC;

II.4.5. have undergone, at least once a year, the routine tests referred to in Chapter II of Annex B to Directive 88/407/EEC.

II.5. The semen to be exported was obtained from donor bulls which:

II.5.1. satisfy the conditions laid down in Annex C of Directive 88/407/EEC;

(1)either ○ [II.5.2. have remained in the exporting country for at least the last six months prior to collection of the semen to be exported;]

(1)or ○ [II.5.2. have remained in the exporting country for at least 30 days prior to the collection of the semen since entry and they were imported from _____(2) during the period of less than six months prior to the collection of the semen and satisfied the animal health conditions applying to donors of the semen which is intended for export to Great Britain;]

II.5.3. comply with at least one of the following conditions as regards bluetongue, as detailed in the table in point I.28.:

(1)either ☐ [II.5.3.1. were kept in a bluetongue virus-free country or zone for at least 60 days prior to, and during, collection of the semen;]

(1)and/or ☐ [II.5.3.2. were kept during a bluetongue virus seasonally free period in a seasonally free zone for at least 60 days prior to, and during, collection of the semen;]

(1)and/or ☐ [II.5.3.3. were kept in a vector-protected establishment for at least 60 days prior to, and during, collection of the semen;]

(1)and/or ☐ [II.5.3.4. were subjected to a serological test for the detection of antibody to the bluetongue virus serogroup, carried out in accordance with the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals, with negative results, at least every 60 days throughout the collection period and between 21 and 60 days after the final collection for this consignment of semen;]

(1)and/or ☐ [II.5.3.5. were subjected to an agent identification test for bluetongue virus, carried out in accordance with the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals, with negative results, on blood samples taken at commencement and final collection for this consignment of semen and at least every 7 days (virus isolation test) or at least every 28 days, if carried out as polymerase chain reaction (PCR), during collection for this consignment of semen;]

Part II: Certification

II. Health information

II.5.4. comply with at least one of the following conditions as regards epizootic haemorrhagic disease (EHD), as detailed in the table in point I.28.:

(1)either ☐ [II.5.4.1. were resident in the exporting country which according to official findings is free from epizootic haemorrhagic disease (EHD);]

(1)(5)and/or ☐ [II.5.4.2. were resident in the exporting country in which according to official findings the following serotypes of epizootic haemorrhagic disease (EHD) exist: _____ and were subjected with negative results in each case to the following tests carried out in an approved laboratory:

(1)either ☐ [II.5.4.2.1. a serological test(4) for the detection of antibody to the EHD virus serogroup, carried out on samples of blood taken on two occasions not more than 12 months apart prior to and not less than 21 days following collection for this consignment of semen;]

(1)and/or ☐ [II.5.4.2.2. a serological test(4) for the detection of antibody to the EHD virus serogroup, carried out on samples taken at intervals of not more than 60 days throughout the collection period and between 21 and 60 days after the final collection for this consignment of semen.]

(1)and/or ☐ [II.5.4.2.3. an agent identification test(4) carried out on blood samples collected at commencement and conclusion of, and at least every 7 days (virus isolation test) or at least every 28 days, if carried out as PCR, during collection for this consignment of semen.]]

II.6. The semen to be exported was collected after the date on which the centre was approved by the competent national authorities of the exporting country.

II.7. The semen to be exported was processed, stored and transported under conditions which satisfy the terms of Directive 88/407/EEC.

Notes

(*) Those countries subject to the transitional import arrangements include: an EU member State; Liechtenstein; Norway and Switzerland.

References to European Union legislation within this certificate are references to direct EU legislation which has been retained in Great Britain (retained EU law as defined in the European Union (Withdrawal) Act 2018).

References to Great Britain in this certificate include Channel Islands and Isle of Man.

Part II: Certification

II. Health information

Part I:

Box I.6.: Person responsible for the load in Great Britain: this box is to be filled in only if it is a certificate for transit commodity.

Box I.11.: Place of origin shall correspond to the semen collection centre listed in accordance with Article 9(2) of Directive 88/407/EEC and where the semen was collected.

Box I.16: Do not use this box until the end of the transitional staging period.

Box I.20.: Number of packages shall correspond to the number of containers.

Box I.21.: Identification of container and seal number shall be indicated.

Box I.23.: Fill in according to whether it is a transit or an import certificate.

Box I.24.: Fill in according to whether it is a transit or an import certificate.

Box I.25.: Species: select amongst “Bos taurus”, “Bison bison" or "Bubalus bubalis” as appropriate.

Donor identity shall correspond to the official identification of the animal.

Date of collection shall be indicated in the following format: dd/mm/yyyy.

Quantity shall correspond to the number of straws of semen collected on a particular date from an identified donor bull complying with particular conditions for bluetongue and EHD.

Part II:

(1) Delete as necessary.

(2) Only third countries or parts thereof listed in Annex I to Implementing Decision 2011/630/EU.

(3) Only semen collection centres listed in accordance with Article 9(2) of Directive 88/407/EEC.

(4) Standards for EHD virus diagnostic tests are described in the Bluetongue Chapter (2.1.3) of the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals.

(5) Compulsory for Australia, Canada and the United States.

(6) Referring to each straw or batch of straws indicate applicable condition (for example II.5.3.1.).

(7) Referring to each straw or batch of straws indicate applicable condition (for example II.5.4.1. or II.5.4.2.1.).

(8) For New Zealand, appearing with the entry “XII” in column 6 of Part 1 of Annex I to Commission Regulation (EU) No 206/2010 (OJ L 73, 20.3.2010, p.1), officially tuberculosis-free bovine herds shall be considered equivalent to officially tuberculosis-free bovine herds in the Member States recognised based on the conditions laid down in paragraphs 1 and 2 of Annex A.I to Council Directive 64/432/EEC.

The signature and the stamp must be in a different colour to that of the printing.

Certifying Officer

Name (in capital letters) Qualification and title

Date of signature Signature

Stamp

Část I

I.1. Odesílatel
Název
Adresa
Země Kód ISO

I.2. Referenční číslo IMSOC
Specimen not to be used for exports from EU

I.2.a. Local Reference

I.5. Příjemce
Název
Adresa
Země Kód ISO

I.3. Ústřední příslušný orgán

I.4. Local competent authority

I.7. Země původu Kód ISO

I.9. Country of destination Kód ISO

I.8. Region of origin Kód

I.10. Region určení Kód

I.11. Place of Dispatch
Název
Adresa
Číslo schválení
Země Kód ISO

I.12. Místo určení
Název
Adresa
Číslo schválení
Země Kód ISO

I.13. Místo nakládky
Název
Adresa
Číslo schválení
Země Kód ISO

I.14. Date and time of departure

I.15. Dopravní prostředky

Typ	Doklad	Identifikace

I.16 Entry Point

I.18. Transport conditions
Okolní ☐ Controlled temperature ☐ Zmrazené ☐ Chlazený ☐

I.17. Průvodní doklady
Referenční číslo obchodního do k ladu
Datum vydání
Země
Místo vydání

I.19. Č. kontejneru / č. plomby

I.20. Certified as

Transhumance ☐	Pollination ☐	Krmivo pro zvířata ☐	Použití pro farmaceutické účely ☐
Game Restocking ☐	Výkrmová ☐	Evidovaní koňovití ☐	Karanténa ☐
Umělé rozmnožování ☐	Training ☐	Další postup ☐	Racing ☐
Production of petfood ☐	Breeding and production ☐	Consignments according to Regulation No 999/2001 ☐	Competition ☐
Storage ☐	Breeding ☐	Ornamental use/research ☐	Laboratory ☐
Rodent food ☐	Zvířata v zájmovém chovu ☐	Jiné ☐	Production ☐
Technické použití ☐	Ornamental bird food ☐	Sádkování ☐	Porážka ☐
Schválené orgány ☐	Lidská spotřeba ☐	Krmivo pro domácí zvířata ☐	Sales ☐
Organic fertilizers ☐	Unregistered equidae ☐	Cirkus/výstava ☐	Category 3 fish oil/fish meal for detoxification according to Regulation 2015/786 ☐

I.21. For transit through a third country ☐
Země Kód ISO
EU Exit Authority BCP code
EU Entry Authority BCP code

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

I.24. Celkové množství

I.25. Celková hrubá hmotnost

Část I

I.28. Description of consignment

1. 05 VÝROBKY ŽIVOČIŠNÉHO PŮVODU, JINDE NEUVEDENÉ ANI NEZAHRNUTÉ

0511 Výrobky živočišného původu, jinde neuvedené ani nezahrnuté; mrtvá zvířata kapitol|1|nebo 3, nezpůsobilá k lidskému požívání

051110 Býčí sperma

05111000 Býčí sperma

Komodita	Druh	Identifikační číslo	Identifikační značka	Nature of commodity

Množství	Datum odběru/produkce	Výrobní zařízení

Part II: Certification

II. Informace týkající se zdraví

I, the undersigned official veterinarian, hereby certify that :

II.1. _____ (name of exporting country or part thereof)(2)

was free from rinderpest and foot-and-mouth disease during the 12 months immediately prior to collection of the semen for export and until its date of dispatch to Great Britain and no vaccination against these diseases has taken place during the same period.

II.2. The centre(3) described in Box. I.11. at which the semen to be exported was collected:

II.2.1. meets the conditions laid down in Chapter I(1) of Annex A to Directive 88/407/EEC;

II.2.2. is operated and supervised in accordance with the conditions laid down in Chapter II(1) of Annex A to Directive 88/407/EEC.

II.3. The centre at which the semen to be exported was collected was free from rabies, tuberculosis, brucellosis, anthrax and contagious bovine pleuropneumonia during 30 days prior to the date of collection of the semen to be exported and the 30 days after collection (in the case of fresh semen until the day of dispatch to Great Britain).

II.4. The bovine animals standing at the semen collection centre:

(8)II.4.1. come from herds which satisfy the conditions of paragraph 1(b) of Chapter I of Annex B to Directive 88/407/EEC;

II.4.2. come from herds or were born to dams which comply with the conditions of paragraph 1(c) of Chapter I of Annex B to Directive 88/407/EEC, or were tested at the age of at least 24 months in accordance with paragraph 1(c) of Chapter II of Annex B to that Directive;

II.4.3. underwent the tests required in accordance with paragraph 1(d) of Chapter I of Annex B to Directive 88/407/EEC in the 28 days preceding the quarantine isolation period;

II.4.4. have satisfied the quarantine isolation period and testing requirements laid down in paragraph 1(e) of Chapter I of Annex B to Directive 88/407/EEC;

II.4.5. have undergone, at least once a year, the routine tests referred to in Chapter II of Annex B to Directive 88/407/EEC.

II.5. The semen to be exported was obtained from donor bulls which:

II.5.1. satisfy the conditions laid down in Annex C of Directive 88/407/EEC;

(1)either ○ [II.5.2. have remained in the exporting country for at least the last six months prior to collection of the semen to be exported;]

(1)or ○ [II.5.2. have remained in the exporting country for at least 30 days prior to the collection of the semen since entry and they were imported from _____(2) during the period of less than six months prior to the collection of the semen and satisfied the animal health conditions applying to donors of the semen which is intended for export to Great Britain;]

II.5.3. comply with at least one of the following conditions as regards bluetongue, as detailed in the table in point I.28.:

(1)either ☐ [II.5.3.1. were kept in a bluetongue virus-free country or zone for at least 60 days prior to, and during, collection of the semen;]

(1)and/or ☐ [II.5.3.2. were kept during a bluetongue virus seasonally free period in a seasonally free zone for at least 60 days prior to, and during, collection of the semen;]

(1)and/or ☐ [II.5.3.3. were kept in a vector-protected establishment for at least 60 days prior to, and during, collection of the semen;]

(1)and/or ☐ [II.5.3.4. were subjected to a serological test for the detection of antibody to the bluetongue virus serogroup, carried out in accordance with the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals, with negative results, at least every 60 days throughout the collection period and between 21 and 60 days after the final collection for this consignment of semen;]

(1)and/or ☐ [II.5.3.5. were subjected to an agent identification test for bluetongue virus, carried out in accordance with the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals, with negative results, on blood samples taken at commencement and final collection for this consignment of semen and at least every 7 days (virus isolation test) or at least every 28 days, if carried out as polymerase chain reaction (PCR), during collection for this consignment of semen;]

Part II: Certification

II. Informace týkající se zdraví

II.5.4. comply with at least one of the following conditions as regards epizootic haemorrhagic disease (EHD), as detailed in the table in point I.28.:

(1)either ☐ [II.5.4.1. were resident in the exporting country which according to official findings is free from epizootic haemorrhagic disease (EHD);]

(1)(5)and/or ☐ [II.5.4.2. were resident in the exporting country in which according to official findings the following serotypes of epizootic haemorrhagic disease (EHD) exist: _____ and were subjected with negative results in each case to the following tests carried out in an approved laboratory:

(1)either ☐ [II.5.4.2.1. a serological test(4) for the detection of antibody to the EHD virus serogroup, carried out on samples of blood taken on two occasions not more than 12 months apart prior to and not less than 21 days following collection for this consignment of semen;]

(1)and/or ☐ [II.5.4.2.2. a serological test(4) for the detection of antibody to the EHD virus serogroup, carried out on samples taken at intervals of not more than 60 days throughout the collection period and between 21 and 60 days after the final collection for this consignment of semen.]

(1)and/or ☐ [II.5.4.2.3. an agent identification test(4) carried out on blood samples collected at commencement and conclusion of, and at least every 7 days (virus isolation test) or at least every 28 days, if carried out as PCR, during collection for this consignment of semen.]]

II.6. The semen to be exported was collected after the date on which the centre was approved by the competent national authorities of the exporting country.

II.7. The semen to be exported was processed, stored and transported under conditions which satisfy the terms of Directive 88/407/EEC.

Notes

(*) Those countries subject to the transitional import arrangements include: an EU member State; Liechtenstein; Norway and Switzerland.

References to European Union legislation within this certificate are references to direct EU legislation which has been retained in Great Britain (retained EU law as defined in the European Union (Withdrawal) Act 2018).

References to Great Britain in this certificate include Channel Islands and Isle of Man.

Part II: Certification

II. Informace týkající se zdraví

Part I:

Box I.6.: Person responsible for the load in Great Britain: this box is to be filled in only if it is a certificate for transit commodity.

Box I.11.: Place of origin shall correspond to the semen collection centre listed in accordance with Article 9(2) of Directive 88/407/EEC and where the semen was collected.

Box I.16: Do not use this box until the end of the transitional staging period.

Box I.20.: Number of packages shall correspond to the number of containers.

Box I.21.: Identification of container and seal number shall be indicated.

Box I.23.: Fill in according to whether it is a transit or an import certificate.

Box I.24.: Fill in according to whether it is a transit or an import certificate.

Box I.25.: Species: select amongst “Bos taurus”, “Bison bison" or "Bubalus bubalis” as appropriate.

Donor identity shall correspond to the official identification of the animal.

Date of collection shall be indicated in the following format: dd/mm/yyyy.

Quantity shall correspond to the number of straws of semen collected on a particular date from an identified donor bull complying with particular conditions for bluetongue and EHD.

Part II:

(1) Delete as necessary.

(2) Only third countries or parts thereof listed in Annex I to Implementing Decision 2011/630/EU.

(3) Only semen collection centres listed in accordance with Article 9(2) of Directive 88/407/EEC.

(4) Standards for EHD virus diagnostic tests are described in the Bluetongue Chapter (2.1.3) of the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals.

(5) Compulsory for Australia, Canada and the United States.

(6) Referring to each straw or batch of straws indicate applicable condition (for example II.5.3.1.).

(7) Referring to each straw or batch of straws indicate applicable condition (for example II.5.4.1. or II.5.4.2.1.).

(8) For New Zealand, appearing with the entry “XII” in column 6 of Part 1 of Annex I to Commission Regulation (EU) No 206/2010 (OJ L 73, 20.3.2010, p.1), officially tuberculosis-free bovine herds shall be considered equivalent to officially tuberculosis-free bovine herds in the Member States recognised based on the conditions laid down in paragraphs 1 and 2 of Annex A.I to Council Directive 64/432/EEC.

The signature and the stamp must be in a different colour to that of the printing.

Certifying Officer

Name (in capital letters) Qualification and title

Datum podpisu Podpis

Razítko