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)(2)

has been 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 and no vaccination against these diseases has taken place during the same period.

II.2. The semen described above was collected before 31 December 2004 at the semen collection centre which:

II.2.1. meets the conditions laid down in Chapter I 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 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 the period commencing 30 days prior to the date of collection of the semen to be exported and the 30 days after collection.

II.4. At the time semen described above was collected, all bovine animals standing at the semen collection centre:

II.4.1. came from herds and/or were born to dams which satisfy the conditions of paragraph 1(b) and (c) of Chapter I of Annex B to Directive 88/407/EEC;

II.4.2. had tested negative, within the 30 days preceding the quarantine isolation period, to:

- 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 an ELISA test for infectious bovine rhinotracheitis/infectious pustular vulvo-vaginitis, and
- a virus isolation test (fluorescent antibody test or immunoperoxidase test) for bovine viral diarrhoea, deferred until the animal reached the age of six months in the case of younger animals;

II.4.3. had undergone the 30-day quarantine isolation period and had tested negative 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 the case of a female animal a vaginal mucus agglutination test;

II.4.4. had tested negative, at least once a year, to the routine tests referred to in points 1(a), (b) and (c) of Chapter II of Annex B to Directive 88/407/EEC.

II.5. At the time the semen described above was collected,

II.5.1. all female bovine animals in the centre had tested negative at least once a year to a vaginal mucus agglutination test for Campylobacter fetus infection, and

II.5.2. all bulls used for semen production had tested negative either to an immunofluorescent antibody test or to a culture test for Campylobacter fetus infection on a sample of preputial material or artificial vagina washings carried out in 12 months prior to collection.

II.6. The semen to be exported was obtained from donor bulls which

II.6.1. satisfy the conditions laid down in Annex C of Directive 88/407/EEC;

(1)either ○ [II.6.2. were resident in the exporting country during the six months immediately prior to collection of the semen for export;]

(1)or ○ [II.6.2. were imported from _____.(2) after spending less than six months in the exporting country and at the time of import satisfied the animal health conditions applying to donors of the semen which is intended for export to the European Union;]

II.6.3. stand in a semen collection centre at which:

Part II: Certification

II. Health information

(1) either ○ [all bovine animals were not vaccinated against infectious bovine rhinotracheitis and tested negative at least once a year to a serum neutralisation test or an ELISA test for infectious bovine rhinotracheitis/ infectious pustular vulvo-vaginitis;]

(1) or ○ [bovine animals not vaccinated against infectious bovine rhinotracheitis tested negative, at least once a year, to a serum neutralisation test or an ELISA test for infectious bovine rhinotracheitis/ infectious pustular vulvo-vaginitis, at which testing for infectious bovine rhinotracheitis was not carried out on bulls which had received their first vaccination against infectious bovine rhinotracheitis at the insemination centre after they had tested negative to a serum neutralisation test or an ELISA test for infectious bovine rhinotracheitis/ infectious pustular vulvo-vaginitis and which had been regularly re-vaccinated at intervals of not more than six months since the first vaccination;]

(1)either ○ [II.6.4. have not been vaccinated against infectious bovine rhinotracheitis,]

(1)or ○ [II.6.4. have been vaccinated against infectious bovine rhinotracheitis in accordance with point II.6.3.,]

II.6.5. fulfil the import conditions for bovine semen laid down in the Bluetongue Chapter of the Terrestrial Animal Health Code of the OIE, depending on the status of the country or zone of residence;****

II.6.6. were resident in the country of export in which the following serotypes of epizootic haemorrhagic disease (EHD) exist: _____ : and tested negative on two occasions not more than 12 months apart to an agar-gel immuno-diffusion test(3) and to a virus neutralization test for all above-listed serotypes of EHD, carried out in approved laboratory on samples of blood taken prior to and not less than 21 days following collection of the semen;***

II.6.7. were resident in the country of export in which the following serotypes of epizootic haemorrhagic disease (EHD) exist: _____ : and tested negative, prior to entry and at six-monthly intervals, to an agar-gel immuno-diffusion test(3) and a virus neutralization test for all above-listed serotypes of EHD, carried out in approved laboratory;**

II.6.8. tested negative on two occasions not more than 12 months apart to a serum neutralization test for Akabane virus carried out in approved laboratory on samples of blood taken prior to and not less than 21 days following collection of the semen.*

II.7. 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.8. The semen to be exported was processed, stored and transported under conditions which satisfy the terms of Directive 88/407/EEC prior to its amendment by Directive 2003/43/EC.

Notes

Part I:

Box I.6.: person responsible for the load in EU: 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 where the semen was collected.

Box I.12.: place of destination: this box is to be filled in only if it is a certificate for transit commodity.

Box I.19.: identification of container and seal number shall be indicated.

Box I.18.: fill in according to whether it is a transit or an import certificate.

Box I.20.: fill in according to whether it is a transit or an import certificate.

Box I.21.: number of packages shall correspond to the number of containers.

Box I.22.: donor identity shall correspond to the official identification of the animal.

date of collection shall be prior to 31 December 2004 and indicated in the following format: dd/mm/yyyy.

approval number of the centre shall correspond to the approval number of the approved semen collection centre where the semen was collected.

Part II:

Part II: Certification

II. Health information

(1) Delete as necessary.

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

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

**** To be used only by Australia, Canada and the USA.

*** To be used only by Australia and the USA.

** To be used only by Canada.

* To be used only by Australia.

. 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)(2)

has been 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 and no vaccination against these diseases has taken place during the same period.

II.2. The semen described above was collected before 31 December 2004 at the semen collection centre which:

II.2.1. meets the conditions laid down in Chapter I 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 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 the period commencing 30 days prior to the date of collection of the semen to be exported and the 30 days after collection.

II.4. At the time semen described above was collected, all bovine animals standing at the semen collection centre:

II.4.1. came from herds and/or were born to dams which satisfy the conditions of paragraph 1(b) and (c) of Chapter I of Annex B to Directive 88/407/EEC;

II.4.2. had tested negative, within the 30 days preceding the quarantine isolation period, to:

- 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 an ELISA test for infectious bovine rhinotracheitis/infectious pustular vulvo-vaginitis, and
- a virus isolation test (fluorescent antibody test or immunoperoxidase test) for bovine viral diarrhoea, deferred until the animal reached the age of six months in the case of younger animals;

II.4.3. had undergone the 30-day quarantine isolation period and had tested negative 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 the case of a female animal a vaginal mucus agglutination test;

II.4.4. had tested negative, at least once a year, to the routine tests referred to in points 1(a), (b) and (c) of Chapter II of Annex B to Directive 88/407/EEC.

II.5. At the time the semen described above was collected,

II.5.1. all female bovine animals in the centre had tested negative at least once a year to a vaginal mucus agglutination test for Campylobacter fetus infection, and

II.5.2. all bulls used for semen production had tested negative either to an immunofluorescent antibody test or to a culture test for Campylobacter fetus infection on a sample of preputial material or artificial vagina washings carried out in 12 months prior to collection.

II.6. The semen to be exported was obtained from donor bulls which

II.6.1. satisfy the conditions laid down in Annex C of Directive 88/407/EEC;

(1)either ○ [II.6.2. were resident in the exporting country during the six months immediately prior to collection of the semen for export;]

(1)or ○ [II.6.2. were imported from _____.(2) after spending less than six months in the exporting country and at the time of import satisfied the animal health conditions applying to donors of the semen which is intended for export to the European Union;]

II.6.3. stand in a semen collection centre at which:

Part II: Certification

II. Informace týkající se zdraví

(1) either ○ [all bovine animals were not vaccinated against infectious bovine rhinotracheitis and tested negative at least once a year to a serum neutralisation test or an ELISA test for infectious bovine rhinotracheitis/ infectious pustular vulvo-vaginitis;]

(1) or ○ [bovine animals not vaccinated against infectious bovine rhinotracheitis tested negative, at least once a year, to a serum neutralisation test or an ELISA test for infectious bovine rhinotracheitis/ infectious pustular vulvo-vaginitis, at which testing for infectious bovine rhinotracheitis was not carried out on bulls which had received their first vaccination against infectious bovine rhinotracheitis at the insemination centre after they had tested negative to a serum neutralisation test or an ELISA test for infectious bovine rhinotracheitis/ infectious pustular vulvo-vaginitis and which had been regularly re-vaccinated at intervals of not more than six months since the first vaccination;]

(1)either ○ [II.6.4. have not been vaccinated against infectious bovine rhinotracheitis,]

(1)or ○ [II.6.4. have been vaccinated against infectious bovine rhinotracheitis in accordance with point II.6.3.,]

II.6.5. fulfil the import conditions for bovine semen laid down in the Bluetongue Chapter of the Terrestrial Animal Health Code of the OIE, depending on the status of the country or zone of residence;****

II.6.6. were resident in the country of export in which the following serotypes of epizootic haemorrhagic disease (EHD) exist: _____ : and tested negative on two occasions not more than 12 months apart to an agar-gel immuno-diffusion test(3) and to a virus neutralization test for all above-listed serotypes of EHD, carried out in approved laboratory on samples of blood taken prior to and not less than 21 days following collection of the semen;***

II.6.7. were resident in the country of export in which the following serotypes of epizootic haemorrhagic disease (EHD) exist: _____ : and tested negative, prior to entry and at six-monthly intervals, to an agar-gel immuno-diffusion test(3) and a virus neutralization test for all above-listed serotypes of EHD, carried out in approved laboratory;**

II.6.8. tested negative on two occasions not more than 12 months apart to a serum neutralization test for Akabane virus carried out in approved laboratory on samples of blood taken prior to and not less than 21 days following collection of the semen.*

II.7. 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.8. The semen to be exported was processed, stored and transported under conditions which satisfy the terms of Directive 88/407/EEC prior to its amendment by Directive 2003/43/EC.

Notes

Part I:

Box I.6.: person responsible for the load in EU: 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 where the semen was collected.

Box I.12.: place of destination: this box is to be filled in only if it is a certificate for transit commodity.

Box I.19.: identification of container and seal number shall be indicated.

Box I.18.: fill in according to whether it is a transit or an import certificate.

Box I.20.: fill in according to whether it is a transit or an import certificate.

Box I.21.: number of packages shall correspond to the number of containers.

Box I.22.: donor identity shall correspond to the official identification of the animal.

date of collection shall be prior to 31 December 2004 and indicated in the following format: dd/mm/yyyy.

approval number of the centre shall correspond to the approval number of the approved semen collection centre where the semen was collected.

Part II:

Part II: Certification

II. Informace týkající se zdraví

(1) Delete as necessary.

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

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

**** To be used only by Australia, Canada and the USA.

*** To be used only by Australia and the USA.

** To be used only by Canada.

* To be used only by Australia.

. 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