EUROPEAN UNION **Export Health Certificate**

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.23. Total number of packages	I.24. Total quantity	I.25. Total net weight	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

051199 Other

05119985 Other

Commodity	Species	Quantity	Batch number	Manufacturing plant

Cold store	Cutting plant	Date of freezing	Date of production	Date of slaughter

Net weight	Product Description	Package count	Identification mark

Part II: Certification

II. Health information

I, the undersigned, official veterinarian, hereby certify that:

II.1. the exporting country _____ (name of exporting county) (2)

(1) either ○ [II.1.1. has during the past 12 months been free of foot-and-mouth disease, classical swine fever and African swine fever,

and that no vaccinations have been carried out against any of these diseases during the past 12 months;]

(1) or ○ is recognised as free of foot-and-mouth disease without vaccination by the World Organisation for Animal Health (OIE) and free of classical swine fever and African swine fever, in accordance with the recommendations laid down in the OIE Terrestrial Animal Health Code;]

II.2. the semen collection centre in which the semen in this consignment was collected:

II.2.1. is approved for export to Great Britain by the veterinary services of _____ (name of third country (2)) and complies with the condition for approval and supervision set out in Chapter I and Chapter II of Annex A to Directive 90/429/EEC;

II.2.2. was, during the period commencing three months prior to the date of collection of the semen in this consignment until the date of its dispatch, situated in an area not restricted due to an outbreak of foot-and-mouth disease, classical swine fever, African swine fever, swine vesicular disease, and vesicular stomatitis;

II.2.3. was, during the period commencing 30 days prior to the date of collection of the semen in this consignment until the date of its dispatch, free from brucellosis and Aujeszky's disease;

(1) either ☐ [II.2.4. contains only animals that have not been vaccinated against Aujeszky's disease and meet the requirements of Annex B to Directive 90/429/EEC.]

(1)(3) and/or ☐ [II.2.4. is a centre in which some or all of the animals have been vaccinated against Aujeszky's disease using a gE deleted vaccine and meet the requirements of Annex B to Directive 90/429/EEC.]

Conditions for the admission of animals to the semen collection centre

II.3. Prior to be admitted to the semen collection centre, all animals:

II.3.1. were subjected to a period of quarantine of at least 30 days in accommodation specifically approved for the purpose by the competent authority, and where only animals having at least the same health status were present (quarantine accommodation);

II.3.2. prior to entering the quarantine accommodation, were chosen from herds or holdings:

II.3.2.1. which were free of brucellosis in accordance with the Chapter on porcine brucellosis of the Terrestrial Animal Health Code of the World Organisation or Animal Health (OIE);

II.3.2.2. in which no animal vaccinated against foot-and-mouth disease was present in the preceding 12 months;

II.3.2.3. which were not situated in a restricted area defined under the provisions of the national legislation due to an outbreak of foot-and-mouth disease, classical swine fever, African swine fever, swine vesicular disease, vesicular stomatitis and Aujeszky's disease;

II.3.2.4. in which no clinical, serological, virological or pathological evidence of Aujeszky's disease was deleted in the preceding 12 months;

II.3.3. prior to entering the quarantine accommodation, were not previously kept in any herd of a lower health status than described in II.3.2;

II.3.4. within 30 days prior to entering the quarantine accommodation referred to in point II.3.1, were subjected to the following tests, performed in accordance with international standards, with negative results;

II.3.4.1. as regards brucellosis, a buffered Brucella antigen test(rose Bengal test), or a cELISA or an iELISA;

II.3.4.2. as regards Aujeszky's disease,

(1)either ○ [II.3.4.2.1. in the case of non-vaccinated animals, a serum neutralisation test or an ELISA for detecting antibodies to the whole Aujeszky's disease virus or to its glycoprotein B (ADV-gB) or glycoprotein D (ADV-gD);]

(2) or ○ [II.3.4.2.1. in the case of animals vaccinated with a gE deleted vaccine, an ELISA for detecting antibodies to glycoprotein E (ADV-gE);]

Part II: Certification

II. Health information

(1)either ○ [II.3.5. were admitted to the centre after all of the animals had reacted with negatives result to a buffered brucella antigen test (rose Bengal test), or a cELISA or an iELISA carried out on samples collected during the last 15 days of the period of quarantine specified in point II.3.1;]

(1) or ○ [II.3.5. were admitted to the centre after not all of the animals had reacted with negative result to a buffered Brucella antigen test (rose Bengal test), or a cELISA or an iELISA carried out on samples collected during the last 15 days of the period of quarantine specified in point II.3.1 and the suspicion of brucellosis was ruled out in accordance with point I.5 of Chapter I of Annex B to Directive 90/429/EEC;]

II.3.6. were subjected to the following tests for Aujeszky's disease carried out on samples collected during the last 15 days of the period of quarantine specified in point II.3.1:

(1) either ○ [II.3.6.1. in the case of non-vaccinated animals, a serum neutralisation test or an ELISA for detecting antibodies to the whole Aujeszky's disease virus or to its glycoprotein B (ADV-gB) or glycoprotein D (ADV-gD);]

(1) or ○ [II.3.6.1. in the case of animals vaccinated with a gE deleted vaccine, an ELISA for detecting antibodies to glycoprotein E (ADV-gE);]

(1) either ○ [II.3.6.2. the tests referred to in point II.3.6.1 were carried out with negative result in each case;]

(1) or ○ [II.3.6.2. the animals that proved positive in a test referred to in point II.3.6.1 were removed immediately from the quarantine accommodation and the competent authority took all necessary measures to ensure that the remaining animals had a satisfactory health status before being admitted to the collection centre in accordance with point II.3;]

II.3.7. All tests were carried out in a laboratory approved by the competent authority;

II.3.8. Animals were only admitted to the semen collection centre with the express permission of the centre veterinarian and all animal movements, entering and exiting the semen collection centre, are recorded;

II.3.9. No animal admitted to the semen collection centre showed any clinical sign of disease on the day of admission; all animals came directly from the quarantine accommodation which, on the day of consignment and during the period of residency of the animals, officially fulfilled the following conditions:

II.3.9.1. it was not situated in a restricted area defined under the provisions of national legislation due to an outbreak of foot-and-mouth disease, classical swine fever, African swine fever, swine vesicular disease, vesicular stomatitis and Aujeszky's disease;

II.3.9.2. no clinical, serological, virological or pathological evidence of foot- and-mouth disease, classical swine fever, African swine fever, swine vesicular disease, vesicular stomatitis and Aujeszky's disease had been recorded for the past 30 days.

Compulsory routine tests for animals kept at the semen collection centre

II.4. All animals kept at the semen collection centre are subjected to the following routine tests carried out in a laboratory approved by the competent authority:

II.4.1. as regards brucellosis, a buffered Brucella antigen test (rose Bengal test), or a cELlSA or an iELlSA;

II.4.2. as regards Aujeszky's disease,

(1) either ○ [II.4.2.1. in the case of non-vaccinated animals, a serum neutralisation test or an ELISA for detecting antibodies to the whole Aujeszky's disease virus or to its glycoprotein B (ADV-gB) or glycoprotein D (ADV-gD);]

(1) or ○ [II.4.2.1. in the case of animals vaccinated with a gE deleted vaccine, an ELISA for detecting antibodies to glycoprotein E (ADV-gE);]

II.4.3. The routine tests referred to in points II.4.1 and II.4.2, are carried out on samples taken in accordance with point 1.2 of Chapter II of Annex B to Directive 90/429/EEC in order to ensure that all animals in the centre have been tested at least once during their stay at that centre and at least every 12 months from the date of admission, if their stay exceeds 12 months;

(1) either ○ [II.4.4. All of the animals have reacted with negative results in the routine tests referred to in points II.4.1 and II.4.2 carried out on samples referred to in point II.4.3.]

(1) or ○ [II.4.4. Not all of the animals have reacted with negative results in the tests referred to in points II.4.1 and II.4.2 carried out on samples referred to in point II.4.3:

(a) the animals which proved positive were isolated,

Part II: Certification

II. Health information

(b) the semen collected from each animal at the centre since the date of that animal's last negative test was held in separate storage from semen eligible for export to Great Britain which was collected before the animal's last negative test or after the health status of the centre had been re-established under responsibility of the competent authority of the exporting country.]

Conditions for semen collected at a semen collection centre and intended for export to Great Britain

II.5. The semen in this consignment was obtained from animals which:

II.5.1. have been resident in _____ (name of third country (2)) for a minimum period of three months immediately prior to collection;

II.5.2. showed no clinical signs of disease on the day the semen was collected;

II.5.3. had not been vaccinated against foot-and-mouth disease;

II.5.4. satisfy the requirements referred to in point II.3;

II.5.5. have not been allowed to serve naturally;

II.5.6. were kept in semen collection centres which were not situated in a restricted area designated under the provisions of the national legislation relating to foot-and-mouth disease, classical swine fever, African swine fever, swine vesicular disease vesicular stomatitis and AujeszkVs disease;

II.5.7. were kept in semen collection centres in which no clinical, serological, virological or pathological evidence of foot-and-mouth disease, classical swine fever, African swine fever, swine vesicular disease, vesicular stomatitis and Aujeszky's disease has been detected in the 30-day period immediately prior to collection.

II.6. An effective combination of antibiotics, in particular against leptospires, was added to the semen in this consignment after final dilution or to the diluent. In the case of frozen semen, antibiotics were added before the semen was frozen.

II.6.1. The combination of antibiotics referred to in point II.6. produced an effect at least equivalent to the following concentration in the final diluted semen:

(a) not less than 500 µ streptomycin per ml final dilution,

(b) not less than 500 IU penicillin per ml final dilution,

(c) not less than 150 µ lincomycin per ml final dilution,

(d) not less than 300 µ spectinomycin per ml final dilution;

II.6.2. Immediately after the addition of the antibiotics the diluted semen was kept at a temperature of at least 15°C for a period of not less than 45 minutes.

II.7. The semen in this consignment:

II.7.1. has been stored as laid down in point 2(d) of Chapter I and point 6(a), (b), (e) and (f) of Chapter II of Annex A to Directive 90/429/EEC prior to dispatch;

II.7.2 is being transported to the country of destination in flasks which were cleaned and disinfected or sterilised before use and which have been sealed prior to dispatch from the approved storage facilities.

Part II: Certification

II. Health information

Notes

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

References made to European Union Legislation forms part of retained EU law in Great Britain.

Part I:

Box 1.6: Person responsible for the load in GB', this box is to be filled in only if it is a certificate for transit commodity.

Box 1.8: Provide the code of the third country as appearing in Annex 1 to Commission Implementing Decision 2012/137/EU.

Box 1.11: Place of origin shall correspond to the semen collection centre of the semen dispatch listed in accordance with Article 8(2) of Directive 90/429/EEC.

Box 1.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity.

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

Box 1.20: Number of packages shall correspond to the number of containers.

Box 1.21: Identification of container and seal number shall be indicated.

Box 1.23: fill in according to whether it is a transit or an import certificate.

Box 1.24: fill in according to whether it is a transit or an import certificate.

Box 1.25: Donor identity shall correspond to the official identification of the animal.

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

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

Part II:

(1) Delete as necessary.1 I to Commission Implementing Decision 2012/137/EU.

(3) This option shall be deleted where the country of destination, or a region thereof, is free of Aujeszky's disease in accordance with Article 10 of 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.23. Celkový počet balení	I.24. Celkové množství	I.25. Celková čistá hmotnost	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

051199 Ostatní

05119985 Ostatní

Komodita	Druh	Množství	Číslo šarže	Výrobní zařízení

Chladírenské zařízení	Bourárna	Datum zmrazení	Datum výroby	Datum porážky

Čistá hmotnost	Product Description	Počet balení	Identifikační značka

Part II: Certification

II. Informace týkající se zdraví

I, the undersigned, official veterinarian, hereby certify that:

II.1. the exporting country _____ (name of exporting county) (2)

(1) either ○ [II.1.1. has during the past 12 months been free of foot-and-mouth disease, classical swine fever and African swine fever,

and that no vaccinations have been carried out against any of these diseases during the past 12 months;]

(1) or ○ is recognised as free of foot-and-mouth disease without vaccination by the World Organisation for Animal Health (OIE) and free of classical swine fever and African swine fever, in accordance with the recommendations laid down in the OIE Terrestrial Animal Health Code;]

II.2. the semen collection centre in which the semen in this consignment was collected:

II.2.1. is approved for export to Great Britain by the veterinary services of _____ (name of third country (2)) and complies with the condition for approval and supervision set out in Chapter I and Chapter II of Annex A to Directive 90/429/EEC;

II.2.2. was, during the period commencing three months prior to the date of collection of the semen in this consignment until the date of its dispatch, situated in an area not restricted due to an outbreak of foot-and-mouth disease, classical swine fever, African swine fever, swine vesicular disease, and vesicular stomatitis;

II.2.3. was, during the period commencing 30 days prior to the date of collection of the semen in this consignment until the date of its dispatch, free from brucellosis and Aujeszky's disease;

(1) either ☐ [II.2.4. contains only animals that have not been vaccinated against Aujeszky's disease and meet the requirements of Annex B to Directive 90/429/EEC.]

(1)(3) and/or ☐ [II.2.4. is a centre in which some or all of the animals have been vaccinated against Aujeszky's disease using a gE deleted vaccine and meet the requirements of Annex B to Directive 90/429/EEC.]

Conditions for the admission of animals to the semen collection centre

II.3. Prior to be admitted to the semen collection centre, all animals:

II.3.1. were subjected to a period of quarantine of at least 30 days in accommodation specifically approved for the purpose by the competent authority, and where only animals having at least the same health status were present (quarantine accommodation);

II.3.2. prior to entering the quarantine accommodation, were chosen from herds or holdings:

II.3.2.1. which were free of brucellosis in accordance with the Chapter on porcine brucellosis of the Terrestrial Animal Health Code of the World Organisation or Animal Health (OIE);

II.3.2.2. in which no animal vaccinated against foot-and-mouth disease was present in the preceding 12 months;

II.3.2.3. which were not situated in a restricted area defined under the provisions of the national legislation due to an outbreak of foot-and-mouth disease, classical swine fever, African swine fever, swine vesicular disease, vesicular stomatitis and Aujeszky's disease;

II.3.2.4. in which no clinical, serological, virological or pathological evidence of Aujeszky's disease was deleted in the preceding 12 months;

II.3.3. prior to entering the quarantine accommodation, were not previously kept in any herd of a lower health status than described in II.3.2;

II.3.4. within 30 days prior to entering the quarantine accommodation referred to in point II.3.1, were subjected to the following tests, performed in accordance with international standards, with negative results;

II.3.4.1. as regards brucellosis, a buffered Brucella antigen test(rose Bengal test), or a cELISA or an iELISA;

II.3.4.2. as regards Aujeszky's disease,

(1)either ○ [II.3.4.2.1. in the case of non-vaccinated animals, a serum neutralisation test or an ELISA for detecting antibodies to the whole Aujeszky's disease virus or to its glycoprotein B (ADV-gB) or glycoprotein D (ADV-gD);]

(2) or ○ [II.3.4.2.1. in the case of animals vaccinated with a gE deleted vaccine, an ELISA for detecting antibodies to glycoprotein E (ADV-gE);]

Part II: Certification

II. Informace týkající se zdraví

(1)either ○ [II.3.5. were admitted to the centre after all of the animals had reacted with negatives result to a buffered brucella antigen test (rose Bengal test), or a cELISA or an iELISA carried out on samples collected during the last 15 days of the period of quarantine specified in point II.3.1;]

(1) or ○ [II.3.5. were admitted to the centre after not all of the animals had reacted with negative result to a buffered Brucella antigen test (rose Bengal test), or a cELISA or an iELISA carried out on samples collected during the last 15 days of the period of quarantine specified in point II.3.1 and the suspicion of brucellosis was ruled out in accordance with point I.5 of Chapter I of Annex B to Directive 90/429/EEC;]

II.3.6. were subjected to the following tests for Aujeszky's disease carried out on samples collected during the last 15 days of the period of quarantine specified in point II.3.1:

(1) either ○ [II.3.6.1. in the case of non-vaccinated animals, a serum neutralisation test or an ELISA for detecting antibodies to the whole Aujeszky's disease virus or to its glycoprotein B (ADV-gB) or glycoprotein D (ADV-gD);]

(1) or ○ [II.3.6.1. in the case of animals vaccinated with a gE deleted vaccine, an ELISA for detecting antibodies to glycoprotein E (ADV-gE);]

(1) either ○ [II.3.6.2. the tests referred to in point II.3.6.1 were carried out with negative result in each case;]

(1) or ○ [II.3.6.2. the animals that proved positive in a test referred to in point II.3.6.1 were removed immediately from the quarantine accommodation and the competent authority took all necessary measures to ensure that the remaining animals had a satisfactory health status before being admitted to the collection centre in accordance with point II.3;]

II.3.7. All tests were carried out in a laboratory approved by the competent authority;

II.3.8. Animals were only admitted to the semen collection centre with the express permission of the centre veterinarian and all animal movements, entering and exiting the semen collection centre, are recorded;

II.3.9. No animal admitted to the semen collection centre showed any clinical sign of disease on the day of admission; all animals came directly from the quarantine accommodation which, on the day of consignment and during the period of residency of the animals, officially fulfilled the following conditions:

II.3.9.1. it was not situated in a restricted area defined under the provisions of national legislation due to an outbreak of foot-and-mouth disease, classical swine fever, African swine fever, swine vesicular disease, vesicular stomatitis and Aujeszky's disease;

II.3.9.2. no clinical, serological, virological or pathological evidence of foot- and-mouth disease, classical swine fever, African swine fever, swine vesicular disease, vesicular stomatitis and Aujeszky's disease had been recorded for the past 30 days.

Compulsory routine tests for animals kept at the semen collection centre

II.4. All animals kept at the semen collection centre are subjected to the following routine tests carried out in a laboratory approved by the competent authority:

II.4.1. as regards brucellosis, a buffered Brucella antigen test (rose Bengal test), or a cELlSA or an iELlSA;

II.4.2. as regards Aujeszky's disease,

(1) either ○ [II.4.2.1. in the case of non-vaccinated animals, a serum neutralisation test or an ELISA for detecting antibodies to the whole Aujeszky's disease virus or to its glycoprotein B (ADV-gB) or glycoprotein D (ADV-gD);]

(1) or ○ [II.4.2.1. in the case of animals vaccinated with a gE deleted vaccine, an ELISA for detecting antibodies to glycoprotein E (ADV-gE);]

II.4.3. The routine tests referred to in points II.4.1 and II.4.2, are carried out on samples taken in accordance with point 1.2 of Chapter II of Annex B to Directive 90/429/EEC in order to ensure that all animals in the centre have been tested at least once during their stay at that centre and at least every 12 months from the date of admission, if their stay exceeds 12 months;

(1) either ○ [II.4.4. All of the animals have reacted with negative results in the routine tests referred to in points II.4.1 and II.4.2 carried out on samples referred to in point II.4.3.]

(1) or ○ [II.4.4. Not all of the animals have reacted with negative results in the tests referred to in points II.4.1 and II.4.2 carried out on samples referred to in point II.4.3:

(a) the animals which proved positive were isolated,

Part II: Certification

II. Informace týkající se zdraví

(b) the semen collected from each animal at the centre since the date of that animal's last negative test was held in separate storage from semen eligible for export to Great Britain which was collected before the animal's last negative test or after the health status of the centre had been re-established under responsibility of the competent authority of the exporting country.]

Conditions for semen collected at a semen collection centre and intended for export to Great Britain

II.5. The semen in this consignment was obtained from animals which:

II.5.1. have been resident in _____ (name of third country (2)) for a minimum period of three months immediately prior to collection;

II.5.2. showed no clinical signs of disease on the day the semen was collected;

II.5.3. had not been vaccinated against foot-and-mouth disease;

II.5.4. satisfy the requirements referred to in point II.3;

II.5.5. have not been allowed to serve naturally;

II.5.6. were kept in semen collection centres which were not situated in a restricted area designated under the provisions of the national legislation relating to foot-and-mouth disease, classical swine fever, African swine fever, swine vesicular disease vesicular stomatitis and AujeszkVs disease;

II.5.7. were kept in semen collection centres in which no clinical, serological, virological or pathological evidence of foot-and-mouth disease, classical swine fever, African swine fever, swine vesicular disease, vesicular stomatitis and Aujeszky's disease has been detected in the 30-day period immediately prior to collection.

II.6. An effective combination of antibiotics, in particular against leptospires, was added to the semen in this consignment after final dilution or to the diluent. In the case of frozen semen, antibiotics were added before the semen was frozen.

II.6.1. The combination of antibiotics referred to in point II.6. produced an effect at least equivalent to the following concentration in the final diluted semen:

(a) not less than 500 µ streptomycin per ml final dilution,

(b) not less than 500 IU penicillin per ml final dilution,

(c) not less than 150 µ lincomycin per ml final dilution,

(d) not less than 300 µ spectinomycin per ml final dilution;

II.6.2. Immediately after the addition of the antibiotics the diluted semen was kept at a temperature of at least 15°C for a period of not less than 45 minutes.

II.7. The semen in this consignment:

II.7.1. has been stored as laid down in point 2(d) of Chapter I and point 6(a), (b), (e) and (f) of Chapter II of Annex A to Directive 90/429/EEC prior to dispatch;

II.7.2 is being transported to the country of destination in flasks which were cleaned and disinfected or sterilised before use and which have been sealed prior to dispatch from the approved storage facilities.

Part II: Certification

II. Informace týkající se zdraví

Notes

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

References made to European Union Legislation forms part of retained EU law in Great Britain.

Part I:

Box 1.6: Person responsible for the load in GB', this box is to be filled in only if it is a certificate for transit commodity.

Box 1.8: Provide the code of the third country as appearing in Annex 1 to Commission Implementing Decision 2012/137/EU.

Box 1.11: Place of origin shall correspond to the semen collection centre of the semen dispatch listed in accordance with Article 8(2) of Directive 90/429/EEC.

Box 1.12: Place of destination: this box is to be filled in only if it is a certificate for transit commodity.

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

Box 1.20: Number of packages shall correspond to the number of containers.

Box 1.21: Identification of container and seal number shall be indicated.

Box 1.23: fill in according to whether it is a transit or an import certificate.

Box 1.24: fill in according to whether it is a transit or an import certificate.

Box 1.25: Donor identity shall correspond to the official identification of the animal.

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

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

Part II:

(1) Delete as necessary.1 I to Commission Implementing Decision 2012/137/EU.

(3) This option shall be deleted where the country of destination, or a region thereof, is free of Aujeszky's disease in accordance with Article 10 of 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