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 country)(2)

II.1.1. has been free from rinderpest, peste des petits ruminants, sheep and goat pox, contagious caprine pleuropneumonia and Rift Valley fever during the 12 months immediately prior to collection of the semen to be exported and until its date of dispatch to Great Britain and no vaccination against these diseases took place during that period;

II.1.2. has been free from foot-and-mouth disease during the 12 months immediately prior to collection of the semen to be exported and until its date of dispatch to Great Britain and no vaccination against this disease took place during that period.

II.2. The semen collection centre described in Box I.11. and at which the semen to be exported was collected and stored:

II.2.1. meets the conditions for the approval of semen collection centres laid down in Chapter I(I)(1) of Annex D to Directive 92/65/EEC;

II.2.2. is operated and supervised in accordance with the conditions applicable to semen collection centres and storage centres laid down in Chapter I(II)(1) of Annex D to Directive 92/65/EEC.

II.3. The ovine/caprine(1) animals standing at the semen collection centre:

II.3.1. prior to their stay in the quarantine accommodation described in point II.3.3.,

(1)(4)either ○ [II.3.1.1. originate from the territory described in Box I.8, which has been recognised as officially brucellosis (B. melitensis)-free,]

(1)or ○ [II.3.1.1. have belonged to a holding which has obtained and maintained its officially brucellosis (B. melitensis)-free status in accordance with Directive 91/68/EEC,]

(1)or ○ [II.3.1.1. originate from a holding, where in respect of brucellosis (B. melitensis) all susceptible animals have been free from clinical or any signs of this disease for the last 12 months, none of the ovine and caprine animals have been vaccinated against this disease, save those vaccinated with Rev. 1 vaccine more than two years ago, and all ovine and caprine animals over six months of age have been subjected to at least two tests(3), carried out with negative results on samples taken on _____ (date) and on _____ (date) at least six months apart, the latter being within 30 days of entry into the quarantine accommodation,]

and have not been kept previously in a holding of a lower status;

II.3.1.2. have been kept continuously for at least 60 days on a holding where no case of contagious epididymitis (Brucella ovis) has been diagnosed in the last 12 months,

(1)and ☐ [they are animals of the ovine species and have undergone during the 60 days prior to their stay in the quarantine accommodation described in point II.3.3. a complement fixation test, or any other test with an equivalent documented sensitivity and specificity, to detect contagious epididymitis with result of less than 50 ICFTU/ml;]

II.3.1.3. to the best of my knowledge do not come from holdings and have not been in contact with animals of a holding, in which based on the official notification system and according to the written declaration made by the owner any of the following diseases has been clinically detected within the periods referred to in (a) to (d) prior to their stay in the quarantine accommodation described in point II.3.3.:

(a) contagious agalactia of sheep or goats (Mycoplasma agalactiae, Mycoplasma capricolum, Mycoplasma mycoides var. mycoides "large colony"), within the last six months,

(b) paratuberculosis and caseous lymphadenitis, within the last 12 months,

(c) pulmonary adenomatosis, within the last three years;

(1)either ○ [(d) Maedi/Visna for sheep or caprine viral arthritis/encephalitis for goats, within the last three years;]

Part II: Certification

II. Health information

(1)or ○ [(d) Maedi/Visna for sheep or caprine viral arthritis/encephalitis for goats, within the last 12 months, and all the infected animals were slaughtered and remaining animals subsequently reacted negatively to two tests carried out at least six months apart;]

II.3.2. have undergone the following tests carried out on a blood sample collected within the 28 days preceding the commencement of the period of quarantine specified in point II.3.3.for:

– brucellosis (B. melitensis), with negative results in each case in accordance with Annex C to Directive 91/68/EEC;

- contagious epididymitis (B. ovis), in the case of sheep only, with negative results in each case in accordance with Annex D to Directive 91/68/EEC, or any other test with an equivalent documented sensitivity and specificity;

– border disease in accordance with point 1.4 (c) of Chapter II(II) of Annex D to Directive 92/65/EEC;

II.3.3. have satisfied the quarantine isolation period of at least 28 days in a quarantine accommodation specifically approved for the purpose by the competent authority and during that period

II.3.3.1. only animals of at least the same health status were present in the quarantine accommodation;

II.3.3.2. the animals have undergone the following tests, carried out by the laboratory approved by the competent authority of the exporting country on samples taken not earlier than 21 days after the animals were admitted to the quarantine accommodation, for:

– brucellosis (B. melitensis) with negative results in accordance with Annex C to Directive 91/68/EEC;

– contagious epididymitis (Brucella ovis), in the case of sheep only, with negative results in accordance with Annex D to Directive 91/68/EEC, or any other test with an equivalent documented sensitivity and specificity;

– border disease in accordance with point 1.6 of Chapter II(II) of Annex D to Directive 92/65/EEC;

II.3.4. have undergone at least once a year the routine tests for:

– brucellosis (B. melitensis) with negative results in each case in accordance with Annex C to Directive 91/68/EEC;

– contagious epididymitis (Brucella ovis), in the case of sheep only, with negative results in each case in accordance with Annex D to Directive 91/68/EEC, or any other test with an equivalent documented sensitivity and specificity;

– border disease in accordance with point 5(c) of Chapter II(II) of Annex D to Directive 92/65/EEC.

II.4. The semen to be exported was obtained from donor rams/bucks (1) which:

II.4.1. were admitted to the approved semen collection centre with the express permission of the centre veterinarian.

II.4.2. show no clinical signs of disease on the day of admission to the approved semen collection centre and on the day the semen was collected;

(1)either ○ [II.4.3. have not been vaccinated against foot-and-mouth disease during the 12 months prior to collection of the semen;]

(1)or ○ [II.4.3. have been vaccinated against foot-and-mouth disease at least 30 days prior to the collection, and 5 % (with a minimum of five straws) of each collection have been submitted to a virus isolation test for foot-and-mouth disease with negative results;]

II.4.4. have been kept at an approved semen collection centre for a continuous period of at least 30 days immediately prior to collection of the semen, in the case of collections of fresh semen;

Part II: Certification

II. Health information

II.4.5. have not served naturally after their entry to the quarantine accommodation described in point II.3.3. and up to and including the day of semen collection;

II.4.6. have been kept at the approved semen collection centres:

II.4.6.1. which have been free from foot-and-mouth disease for at least three months prior to collection of the semen and 30 days after collection or, in the case of fresh semen, until the date of dispatch, and which are situated in the centre of an area of 10 kilometres radius in which there has been no case of foot-and-mouth disease for at least 30 days prior to collection of the semen;

II.4.6.2. which have been free, during the period commencing 30 days prior to collection and ending 30 days after collection of the semen or, in the case of fresh semen, until the date of dispatch, from brucellosis (B. melitensis), contagious epididymitis (Brucella ovis), anthrax and rabies;

(1)either ○ [II.4.7. have remained in the exporting country for at least the past six months prior to collection of the semen to be exported;]

(1)or ○ [II.4.7. during the last six months prior to collection of the semen they complied with the animal health conditions applying to donors of the semen which is intended for export to Great Britain and they have been imported into the exporting country at least 30 days prior to collection of the semen from_____(2);]

(1)either ○ [II.4.8. were kept in a bluetongue virus-free country or zone for at least 60 days prior to, and during, collection of the semen;]

(1)or ○ [II.4.8. 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)or ○ [II.4.8. were kept in a vector-protected establishment for at least 60 days prior to, and during collection of the semen;]

(1)or ○ [II.4.8. were subjected to a serological test for the detection of antibody to the bluetongue virus group, carried out in accordance with the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals, with negative results, on blood samples taken 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)or ○ [II.4.8. were subjected to an agent identification test for bluetongue virus, carried out in accordance with the 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 (PCR test) during collection for this consignment of semen;]

(1)(5)either ○ [II.4.9. were resident in the exporting country which according to official findings is free from epizootic haemorrhagic disease (EHD);]

(1)or ○ [II.4.9. 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:]

(1)either ○ [a serological test(6) for the detection of antibody to the EHDV group carried out in an approved laboratory on samples of blood taken on two occasions not more than 12 months apart prior to and not less than 21 days after the final collection for this consignment of semen.]]

(1)or ○ [a serological test(6) for the detection of antibody to the EHDV group, carried out in an approved laboratory on samples of blood 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)or ○ [an agent identification test(6) carried out in an approved laboratory on samples of blood taken at commencement and conclusion of, and at least every 7 days (virus isolation test) or at least every 28 days (PCR test) during collection for this consignment of semen.]]

II.4.10. have been kept continuously since birth in a country where the following conditions are fulfilled:

II.4.10.1. classical scrapie is compulsorily notifiable;

Part II: Certification

II. Health information

	II.4.10.2.	an awareness, surveillance and monitoring system is in place;
	II.4.10.3.	ovine and caprine animals affected with classical scrapie are killed and completely destroyed;
	II.4.10.4.	the feeding to ovine and caprine animals of meat-and-bone meal, or greaves of ruminant origin, as defined in the OIE Terrestrial Animal Health Code, has been banned and effectively enforced in the whole country for a period of at least the last seven years;
(1)either	○ [II.4.11.	have been kept continuously for a period of the last three years preceding the date of before the collection of the semen to be exported in a holding or holdings which has/have fulfilled during that period all the requirements set out in points 1.3. (a) to (f) of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001, except during the period when they were kept at a semen collection centre that complied during that period with the conditions set out in the four indents of point 1.3.(c)(iv) of that Section;]
(1)or	○ [II.4.11.	are ovine animals of ARR/ARR prion protein genotype.]
II.5.	The semen to be exported:	
	II.5.1.	was collected after the date on which the semen collection centre was approved by the competent authority of the exporting country;
	II.5.2.	was collected, processed, preserved, stored and transported in accordance with the requirements applicable to semen laid down in Chapter III(I) of Annex D to Directive 92/65/EEC;
	II.5.3.	was sent to the place of loading in a sealed container in accordance with the requirements for semen to be subject to trade laid down in point 1.4 of Chapter III(I) of Annex D to Directive 92/65/EEC and bearing the number indicated in Box I.23.
(1)either	○ [II.6.	No antibiotics were added to the semen.]
(1)or	○ [II.6.	The following antibiotic or combination of antibiotics was added to produce a concentration in the final diluted semen of not less than(7):_____]

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 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 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 approved semen collection centre in which the semen was collected and listed in accordance with Article 17(3)(b) of Directive 92/65/EEC.

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 “Ovis aries” or “Capra hircus” 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.

Approval number of the centre shall correspond to the approval number of the semen collection centre indicated in Box I.11.

Part II:

(1) Delete as necessary.

(2) Only third countries listed in Annex I to Decision 2010/472/EU.

(3) Tests shall be carried out in accordance with Annex C to Directive 91/68/EEC.

(4) Only for the territory appearing with the entry "V" in column 6 of Part 1 of Annex I to Commission Regulation (EU) No 206/2010.

(5) See remarks for exporting country concerned in Annex I to Decision 2010/472/EU.

(6) Standards for EHD virus diagnostic tests are described in Chapter 2.1.7. of the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals.

(7) Insert names and concentrations.

- 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 country)(2)

II.1.1. has been free from rinderpest, peste des petits ruminants, sheep and goat pox, contagious caprine pleuropneumonia and Rift Valley fever during the 12 months immediately prior to collection of the semen to be exported and until its date of dispatch to Great Britain and no vaccination against these diseases took place during that period;

II.1.2. has been free from foot-and-mouth disease during the 12 months immediately prior to collection of the semen to be exported and until its date of dispatch to Great Britain and no vaccination against this disease took place during that period.

II.2. The semen collection centre described in Box I.11. and at which the semen to be exported was collected and stored:

II.2.1. meets the conditions for the approval of semen collection centres laid down in Chapter I(I)(1) of Annex D to Directive 92/65/EEC;

II.2.2. is operated and supervised in accordance with the conditions applicable to semen collection centres and storage centres laid down in Chapter I(II)(1) of Annex D to Directive 92/65/EEC.

II.3. The ovine/caprine(1) animals standing at the semen collection centre:

II.3.1. prior to their stay in the quarantine accommodation described in point II.3.3.,

(1)(4)either ○ [II.3.1.1. originate from the territory described in Box I.8, which has been recognised as officially brucellosis (B. melitensis)-free,]

(1)or ○ [II.3.1.1. have belonged to a holding which has obtained and maintained its officially brucellosis (B. melitensis)-free status in accordance with Directive 91/68/EEC,]

(1)or ○ [II.3.1.1. originate from a holding, where in respect of brucellosis (B. melitensis) all susceptible animals have been free from clinical or any signs of this disease for the last 12 months, none of the ovine and caprine animals have been vaccinated against this disease, save those vaccinated with Rev. 1 vaccine more than two years ago, and all ovine and caprine animals over six months of age have been subjected to at least two tests(3), carried out with negative results on samples taken on _____ (date) and on _____ (date) at least six months apart, the latter being within 30 days of entry into the quarantine accommodation,]

and have not been kept previously in a holding of a lower status;

II.3.1.2. have been kept continuously for at least 60 days on a holding where no case of contagious epididymitis (Brucella ovis) has been diagnosed in the last 12 months,

(1)and ☐ [they are animals of the ovine species and have undergone during the 60 days prior to their stay in the quarantine accommodation described in point II.3.3. a complement fixation test, or any other test with an equivalent documented sensitivity and specificity, to detect contagious epididymitis with result of less than 50 ICFTU/ml;]

II.3.1.3. to the best of my knowledge do not come from holdings and have not been in contact with animals of a holding, in which based on the official notification system and according to the written declaration made by the owner any of the following diseases has been clinically detected within the periods referred to in (a) to (d) prior to their stay in the quarantine accommodation described in point II.3.3.:

(a) contagious agalactia of sheep or goats (Mycoplasma agalactiae, Mycoplasma capricolum, Mycoplasma mycoides var. mycoides "large colony"), within the last six months,

(b) paratuberculosis and caseous lymphadenitis, within the last 12 months,

(c) pulmonary adenomatosis, within the last three years;

(1)either ○ [(d) Maedi/Visna for sheep or caprine viral arthritis/encephalitis for goats, within the last three years;]

Part II: Certification

II. Informace týkající se zdraví

(1)or ○ [(d) Maedi/Visna for sheep or caprine viral arthritis/encephalitis for goats, within the last 12 months, and all the infected animals were slaughtered and remaining animals subsequently reacted negatively to two tests carried out at least six months apart;]

II.3.2. have undergone the following tests carried out on a blood sample collected within the 28 days preceding the commencement of the period of quarantine specified in point II.3.3.for:

– brucellosis (B. melitensis), with negative results in each case in accordance with Annex C to Directive 91/68/EEC;

- contagious epididymitis (B. ovis), in the case of sheep only, with negative results in each case in accordance with Annex D to Directive 91/68/EEC, or any other test with an equivalent documented sensitivity and specificity;

– border disease in accordance with point 1.4 (c) of Chapter II(II) of Annex D to Directive 92/65/EEC;

II.3.3. have satisfied the quarantine isolation period of at least 28 days in a quarantine accommodation specifically approved for the purpose by the competent authority and during that period

II.3.3.1. only animals of at least the same health status were present in the quarantine accommodation;

II.3.3.2. the animals have undergone the following tests, carried out by the laboratory approved by the competent authority of the exporting country on samples taken not earlier than 21 days after the animals were admitted to the quarantine accommodation, for:

– brucellosis (B. melitensis) with negative results in accordance with Annex C to Directive 91/68/EEC;

– contagious epididymitis (Brucella ovis), in the case of sheep only, with negative results in accordance with Annex D to Directive 91/68/EEC, or any other test with an equivalent documented sensitivity and specificity;

– border disease in accordance with point 1.6 of Chapter II(II) of Annex D to Directive 92/65/EEC;

II.3.4. have undergone at least once a year the routine tests for:

– brucellosis (B. melitensis) with negative results in each case in accordance with Annex C to Directive 91/68/EEC;

– contagious epididymitis (Brucella ovis), in the case of sheep only, with negative results in each case in accordance with Annex D to Directive 91/68/EEC, or any other test with an equivalent documented sensitivity and specificity;

– border disease in accordance with point 5(c) of Chapter II(II) of Annex D to Directive 92/65/EEC.

II.4. The semen to be exported was obtained from donor rams/bucks (1) which:

II.4.1. were admitted to the approved semen collection centre with the express permission of the centre veterinarian.

II.4.2. show no clinical signs of disease on the day of admission to the approved semen collection centre and on the day the semen was collected;

(1)either ○ [II.4.3. have not been vaccinated against foot-and-mouth disease during the 12 months prior to collection of the semen;]

(1)or ○ [II.4.3. have been vaccinated against foot-and-mouth disease at least 30 days prior to the collection, and 5 % (with a minimum of five straws) of each collection have been submitted to a virus isolation test for foot-and-mouth disease with negative results;]

II.4.4. have been kept at an approved semen collection centre for a continuous period of at least 30 days immediately prior to collection of the semen, in the case of collections of fresh semen;

Part II: Certification

II. Informace týkající se zdraví

II.4.5. have not served naturally after their entry to the quarantine accommodation described in point II.3.3. and up to and including the day of semen collection;

II.4.6. have been kept at the approved semen collection centres:

II.4.6.1. which have been free from foot-and-mouth disease for at least three months prior to collection of the semen and 30 days after collection or, in the case of fresh semen, until the date of dispatch, and which are situated in the centre of an area of 10 kilometres radius in which there has been no case of foot-and-mouth disease for at least 30 days prior to collection of the semen;

II.4.6.2. which have been free, during the period commencing 30 days prior to collection and ending 30 days after collection of the semen or, in the case of fresh semen, until the date of dispatch, from brucellosis (B. melitensis), contagious epididymitis (Brucella ovis), anthrax and rabies;

(1)either ○ [II.4.7. have remained in the exporting country for at least the past six months prior to collection of the semen to be exported;]

(1)or ○ [II.4.7. during the last six months prior to collection of the semen they complied with the animal health conditions applying to donors of the semen which is intended for export to Great Britain and they have been imported into the exporting country at least 30 days prior to collection of the semen from_____(2);]

(1)either ○ [II.4.8. were kept in a bluetongue virus-free country or zone for at least 60 days prior to, and during, collection of the semen;]

(1)or ○ [II.4.8. 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)or ○ [II.4.8. were kept in a vector-protected establishment for at least 60 days prior to, and during collection of the semen;]

(1)or ○ [II.4.8. were subjected to a serological test for the detection of antibody to the bluetongue virus group, carried out in accordance with the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals, with negative results, on blood samples taken 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)or ○ [II.4.8. were subjected to an agent identification test for bluetongue virus, carried out in accordance with the 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 (PCR test) during collection for this consignment of semen;]

(1)(5)either ○ [II.4.9. were resident in the exporting country which according to official findings is free from epizootic haemorrhagic disease (EHD);]

(1)or ○ [II.4.9. 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:]

(1)either ○ [a serological test(6) for the detection of antibody to the EHDV group carried out in an approved laboratory on samples of blood taken on two occasions not more than 12 months apart prior to and not less than 21 days after the final collection for this consignment of semen.]]

(1)or ○ [a serological test(6) for the detection of antibody to the EHDV group, carried out in an approved laboratory on samples of blood 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)or ○ [an agent identification test(6) carried out in an approved laboratory on samples of blood taken at commencement and conclusion of, and at least every 7 days (virus isolation test) or at least every 28 days (PCR test) during collection for this consignment of semen.]]

II.4.10. have been kept continuously since birth in a country where the following conditions are fulfilled:

II.4.10.1. classical scrapie is compulsorily notifiable;

Part II: Certification

II. Informace týkající se zdraví

II.4.10.2. an awareness, surveillance and monitoring system is in place;

II.4.10.3. ovine and caprine animals affected with classical scrapie are killed and completely destroyed;

II.4.10.4. the feeding to ovine and caprine animals of meat-and-bone meal, or greaves of ruminant origin, as defined in the OIE Terrestrial Animal Health Code, has been banned and effectively enforced in the whole country for a period of at least the last seven years;

(1)either ○ [II.4.11. have been kept continuously for a period of the last three years preceding the date of before the collection of the semen to be exported in a holding or holdings which has/have fulfilled during that period all the requirements set out in points 1.3. (a) to (f) of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001, except during the period when they were kept at a semen collection centre that complied during that period with the conditions set out in the four indents of point 1.3.(c)(iv) of that Section;]

(1)or ○ [II.4.11. are ovine animals of ARR/ARR prion protein genotype.]

II.5. The semen to be exported:

II.5.1. was collected after the date on which the semen collection centre was approved by the competent authority of the exporting country;

II.5.2. was collected, processed, preserved, stored and transported in accordance with the requirements applicable to semen laid down in Chapter III(I) of Annex D to Directive 92/65/EEC;

II.5.3. was sent to the place of loading in a sealed container in accordance with the requirements for semen to be subject to trade laid down in point 1.4 of Chapter III(I) of Annex D to Directive 92/65/EEC and bearing the number indicated in Box I.23.

(1)either ○ [II.6. No antibiotics were added to the semen.]

(1)or ○ [II.6. The following antibiotic or combination of antibiotics was added to produce a concentration in the final diluted semen of not less than(7):_____]

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 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 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 approved semen collection centre in which the semen was collected and listed in accordance with Article 17(3)(b) of Directive 92/65/EEC.

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 “Ovis aries” or “Capra hircus” 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.

Approval number of the centre shall correspond to the approval number of the semen collection centre indicated in Box I.11.

Part II:

(1) Delete as necessary.

(2) Only third countries listed in Annex I to Decision 2010/472/EU.

(3) Tests shall be carried out in accordance with Annex C to Directive 91/68/EEC.

(4) Only for the territory appearing with the entry "V" in column 6 of Part 1 of Annex I to Commission Regulation (EU) No 206/2010.

(5) See remarks for exporting country concerned in Annex I to Decision 2010/472/EU.

(6) Standards for EHD virus diagnostic tests are described in Chapter 2.1.7. of the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals.

(7) Insert names and concentrations.

- 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