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 of _____(exporting country)(2) certify that:

II.1. The embryos to be exported

II.1.1. were produced in the exporting country, which according to official findings:

II.1.1.1 was free from rinderpest during the 12 months immediately prior to their production;

(1)either ○ [II.1.1.2. was free from foot-and-mouth disease and lumpy skin disease during the 12 months immediately prior to their production and did not carry out vaccination against foot-and-mouth disease or lumpy skin disease during that period.]

(1)or ○ [II.1.1.2. was not free from foot-and-mouth disease or lumpy skin disease during the 12 months immediately prior to their production or carried out vaccination against foot-and-mouth disease or lumpy skin disease during that period, and

- the embryos were produced without penetration of the zona pellucida,
- the embryos were stored under approved conditions for at least 30 days immediately after their production,
- the donor females come from holdings on which no animal was vaccinated against foot-and-mouth disease or lumpy skin disease during the 30 days prior to collection and no animal of a susceptible species showed clinical signs of foot-and-mouth disease or lumpy skin disease during the 30 days prior to, and at least the 30 days after, the oocytes were collected.]

II.1.2. were produced by the embryo production team(3) which:

- has been approved in accordance with Chapter I of Annex A to Directive 89/556/EEC;
- carried out the production, processing, storing and transport of the embryos in accordance with Chapter II of Annex A to Directive 89/556/EEC;
- is subject to inspection by an official veterinarian at least twice a year.

II.2. The oocytes used in the production of the embryos to be exported were collected on premises situated in an area of at least 10 km radius centred on them, on which according to official findings there was no occurrence of foot-and-mouth disease, epizootic haemorrhagic disease, vesicular stomatitis, Rift Valley fever, contagious bovine pleuropneumonia or lumpy skin disease in the 30 days immediately prior to their collection and until their dispatch to Great Britain, in case of fresh embryos, or during the 30 days after collection, in case of embryos subject to a mandatory storage for at least 30 days in accordance with point II.2.2.

II.3. From the time of collection of the oocytes until 30 days thereafter or, in the case of fresh embryos, until the day of dispatch, the embryos to be exported were stored on premises situated in an area of at least 10 km radius centred on them, on which according to official findings there was no occurrence of foot-and-mouth disease, vesicular stomatitis, Rift Valley fever, contagious bovine pleuropneumonia or lumpy skin disease.

II.4. The donors of oocytes used in the production of the embryos to be exported:

II.4.1. were located, during the 30 days immediately prior to collection of the oocytes, on premises within a 10-km radius of which, according to official findings, there was no occurrence of foot-and-mouth disease, bluetongue, epizootic haemorrhagic disease, vesicular stomatitis, Rift Valley fever, contagious bovine pleuropneumonia or lumpy skin disease;

II.4.2. showed no clinical signs of disease on the day of collection;

II.4.3. spent the six months immediately prior to collection within the territory of the exporting country in no more than two herds:

- which, according to official findings, were free from tuberculosis during that time,
- which, according to official findings, were free from brucellosis during that time,
- which were free from enzootic bovine leukosis or in which no animal showed clinical signs of enzootic bovine leukosis during the previous three years,
- in which no bovine animal showed clinical signs of infectious bovine rhinotracheitis/infectious pustular vulvo-vaginitis during the previous 12 months.

(1)either ○ [II.4.4. were kept in a bluetongue virus-free country or zone for at least 60 days prior to, and during, collection of the oocytes.]

Part II: Certification

II. Health information

(1)or ○ [II.4.4. were kept during a seasonally free period or protected from the vector for at least 60 days prior to, and during, the collection of the oocytes, and the embryos were produced without penetration of the zona pellucida, except if the donors underwent a serological test to detect antibodies to the bluetongue virus group, carried out in accordance with the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals between 21 and 60 days after collection and giving negative results and the embryos were stored for at least 30 days.]

(1)or ○ [II.4.4. underwent a serological test to detect antibodies to the bluetongue virus group, carried out in accordance with the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals between 21 and 60 days after collection and giving negative results, and the embryos were stored for at least 30 days.]

(1)or ○ [II.4.4. underwent an agent identification test, carried out in accordance with the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals on a blood sample taken on the day of collection or the day of slaughtering and giving negative results – the embryos having been produced, in the latter case, without penetration of the zona pellucida.]

II.5. The embryos to be exported were conceived by in vitro fertilisation using semen coming from semen collection or storage centres approved for the collection, processing and/or storage of semen by the competent authority of a third country or a part thereof listed in Annex I to Implementing Decision 2011/630/EU(4) or by the competent authority of a Member State.

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 embryo production team from which the embryos are dispatch to the Union and listed in accordance with Article 8(2) of Directive 89/556/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 “Bos taurus”, “Bison bison" or "Bubalus bubalis” as appropriate.

Category: select "in vitro produced embryos".

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

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

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

Approval number of the team: shall correspond to the embryo production team by which the embryos were produced, processed and stored; and listed in accordance with Article 8(2) of Directive 89/556/EEC

Part II:

(1) Delete as appropriate.

(2) Only third countries listed in Annex I to Decision 2006/168/EC.

(3) Only embryo production teams listed in accordance with Article 8(2) of Directive 89/556/EEC

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

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 of _____(exporting country)(2) certify that:

II.1. The embryos to be exported

II.1.1. were produced in the exporting country, which according to official findings:

II.1.1.1 was free from rinderpest during the 12 months immediately prior to their production;

(1)either ○ [II.1.1.2. was free from foot-and-mouth disease and lumpy skin disease during the 12 months immediately prior to their production and did not carry out vaccination against foot-and-mouth disease or lumpy skin disease during that period.]

(1)or ○ [II.1.1.2. was not free from foot-and-mouth disease or lumpy skin disease during the 12 months immediately prior to their production or carried out vaccination against foot-and-mouth disease or lumpy skin disease during that period, and

- the embryos were produced without penetration of the zona pellucida,
- the embryos were stored under approved conditions for at least 30 days immediately after their production,
- the donor females come from holdings on which no animal was vaccinated against foot-and-mouth disease or lumpy skin disease during the 30 days prior to collection and no animal of a susceptible species showed clinical signs of foot-and-mouth disease or lumpy skin disease during the 30 days prior to, and at least the 30 days after, the oocytes were collected.]

II.1.2. were produced by the embryo production team(3) which:

- has been approved in accordance with Chapter I of Annex A to Directive 89/556/EEC;
- carried out the production, processing, storing and transport of the embryos in accordance with Chapter II of Annex A to Directive 89/556/EEC;
- is subject to inspection by an official veterinarian at least twice a year.

II.2. The oocytes used in the production of the embryos to be exported were collected on premises situated in an area of at least 10 km radius centred on them, on which according to official findings there was no occurrence of foot-and-mouth disease, epizootic haemorrhagic disease, vesicular stomatitis, Rift Valley fever, contagious bovine pleuropneumonia or lumpy skin disease in the 30 days immediately prior to their collection and until their dispatch to Great Britain, in case of fresh embryos, or during the 30 days after collection, in case of embryos subject to a mandatory storage for at least 30 days in accordance with point II.2.2.

II.3. From the time of collection of the oocytes until 30 days thereafter or, in the case of fresh embryos, until the day of dispatch, the embryos to be exported were stored on premises situated in an area of at least 10 km radius centred on them, on which according to official findings there was no occurrence of foot-and-mouth disease, vesicular stomatitis, Rift Valley fever, contagious bovine pleuropneumonia or lumpy skin disease.

II.4. The donors of oocytes used in the production of the embryos to be exported:

II.4.1. were located, during the 30 days immediately prior to collection of the oocytes, on premises within a 10-km radius of which, according to official findings, there was no occurrence of foot-and-mouth disease, bluetongue, epizootic haemorrhagic disease, vesicular stomatitis, Rift Valley fever, contagious bovine pleuropneumonia or lumpy skin disease;

II.4.2. showed no clinical signs of disease on the day of collection;

II.4.3. spent the six months immediately prior to collection within the territory of the exporting country in no more than two herds:

- which, according to official findings, were free from tuberculosis during that time,
- which, according to official findings, were free from brucellosis during that time,
- which were free from enzootic bovine leukosis or in which no animal showed clinical signs of enzootic bovine leukosis during the previous three years,
- in which no bovine animal showed clinical signs of infectious bovine rhinotracheitis/infectious pustular vulvo-vaginitis during the previous 12 months.

(1)either ○ [II.4.4. were kept in a bluetongue virus-free country or zone for at least 60 days prior to, and during, collection of the oocytes.]

Part II: Certification

II. Informace týkající se zdraví

(1)or ○ [II.4.4. were kept during a seasonally free period or protected from the vector for at least 60 days prior to, and during, the collection of the oocytes, and the embryos were produced without penetration of the zona pellucida, except if the donors underwent a serological test to detect antibodies to the bluetongue virus group, carried out in accordance with the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals between 21 and 60 days after collection and giving negative results and the embryos were stored for at least 30 days.]

(1)or ○ [II.4.4. underwent a serological test to detect antibodies to the bluetongue virus group, carried out in accordance with the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals between 21 and 60 days after collection and giving negative results, and the embryos were stored for at least 30 days.]

(1)or ○ [II.4.4. underwent an agent identification test, carried out in accordance with the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals on a blood sample taken on the day of collection or the day of slaughtering and giving negative results – the embryos having been produced, in the latter case, without penetration of the zona pellucida.]

II.5. The embryos to be exported were conceived by in vitro fertilisation using semen coming from semen collection or storage centres approved for the collection, processing and/or storage of semen by the competent authority of a third country or a part thereof listed in Annex I to Implementing Decision 2011/630/EU(4) or by the competent authority of a Member State.

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 embryo production team from which the embryos are dispatch to the Union and listed in accordance with Article 8(2) of Directive 89/556/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 “Bos taurus”, “Bison bison" or "Bubalus bubalis” as appropriate.

Category: select "in vitro produced embryos".

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

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

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

Approval number of the team: shall correspond to the embryo production team by which the embryos were produced, processed and stored; and listed in accordance with Article 8(2) of Directive 89/556/EEC

Part II:

(1) Delete as appropriate.

(2) Only third countries listed in Annex I to Decision 2006/168/EC.

(3) Only embryo production teams listed in accordance with Article 8(2) of Directive 89/556/EEC

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

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