

Part I : Details of consignment	I.1. Consignor Name Address Country		ISO Code	I.2. IMSOC Reference <b>Specimen not to be used for exports from EU</b>		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		I.16 Entry Point					
	Mode	International transport document	Identification					
	I.18. Transport conditions Ambient <input type="checkbox"/> Controlled temperature <input type="checkbox"/> Frozen <input type="checkbox"/> Chilled <input type="checkbox"/>		I.17. Accompanying documents Commercial document reference Country		Date of issue Place of issue			
	I.19. Container No / Seal No							
I.20. Certified as Transhumance <input type="checkbox"/> Pollination <input type="checkbox"/> Animal Feedingstuff <input type="checkbox"/> Pharmaceutical use <input type="checkbox"/> Game Restocking <input type="checkbox"/> Fattening <input type="checkbox"/> Registered equidae <input type="checkbox"/> Quarantine <input type="checkbox"/> Artificial reproduction <input type="checkbox"/> Training <input type="checkbox"/> Further process <input type="checkbox"/> Racing <input type="checkbox"/> Production of petfood <input type="checkbox"/> Breeding and production <input type="checkbox"/> Consignments according to Regulation No 999/2001 <input type="checkbox"/> Competition <input type="checkbox"/> Storage <input type="checkbox"/> Breeding <input type="checkbox"/> Ornamental use/research <input type="checkbox"/> Laboratory <input type="checkbox"/> Rodent food <input type="checkbox"/> Pets <input type="checkbox"/> Other <input type="checkbox"/> Production <input type="checkbox"/> Technical use <input type="checkbox"/> Ornamental bird food <input type="checkbox"/> Relaying <input type="checkbox"/> Slaughter <input type="checkbox"/> Approved Bodies <input type="checkbox"/> Human consumption <input type="checkbox"/> Pet food <input type="checkbox"/> Sales <input type="checkbox"/> Organic fertilizers <input type="checkbox"/> Unregistered equidae <input type="checkbox"/> Circus exhibition <input type="checkbox"/> Category 3 fish oil/fish meal for detoxification according to Regulation 2015/786 <input type="checkbox"/>								
I.21. For transit through a third country <input type="checkbox"/> Country _____ ISO Code _____ EU Exit Authority _____ BCP code _____ EU Entry Authority _____ BCP code _____				I.22. For transit through Member State(s) <input type="checkbox"/> 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				
	<b>1. 05 PRODUCTS OF ANIMAL ORIGIN, NOT ELSEWHERE SPECIFIED OR INCLUDED</b>				
	<b>0511</b> Animal products not elsewhere specified or included; dead animals of Chapter   1   or 3, unfit for human consumption				
	<b>051199</b> Other				
	<b>05119985</b> 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	
SPECIMEN					

## 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 collection 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 collection;

(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 collection and did not carry out vaccination against foot-and-mouth disease or lumpy skin disease during that period.]

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

- the embryos were not subjected to penetration of the zona pellucida,

- the embryos were stored under approved conditions for at least 30 days immediately after their collection,

- 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 embryos were collected.]

II.1.2. were collected 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 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.1.3. were collected and processed 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 dispatch to Great Britain, in the case of fresh embryos, or during the 30 days after collection, in the case of embryos subject to a mandatory storage for at least 30 days in accordance with point II.1.1.2.

II.1.4. from the time of collection until 30 days thereafter or, in the case of fresh embryos until the day of their dispatch to Great Britain, they 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.1.5. were collected from the donor females, which:

II.1.5.1. were located, during the 30 days immediately prior to collection, on premises situated in an area of at least 10-km radius on 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.1.5.2. showed no clinical signs of disease on the day of collection;

II.1.5.3. 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,

<b>Part II: Certification</b>	II. Health information			
	<p style="margin-left: 150px;">- which were free from enzootic bovine leukosis or in which no bovine animal showed clinical signs of enzootic bovine leukosis during the previous three years,</p> <p style="margin-left: 150px;">- in which no bovine animal showed clinical signs of infectious bovine rhinotracheitis/infectious pustular vulvo-vaginitis during the previous 12 months;</p> <p>II.1.6. The embryos to be exported were conceived by artificial insemination 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 part thereof listed in Annex 1 to implementing Decision 2011/630/EU (4) or by the competent authority of Great Britain.</p> <p>Notes</p> <p>(*) Those countries subject to the transitional import arrangements include: an EU member State; Liechtenstein; Norway and Switzerland.</p> <p>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).</p> <p>References to Great Britain in this certificate include Channel Islands and Isle of Man.</p> <p>Part I:</p> <p>Box I.6.: Person responsible for the load in Great Britain: this box is to be filled in only if it is a certificate for transit commodity.</p> <p>Box I.11.: Place of origin shall correspond to the embryo production team from which the embryos are dispatch to Great Britain and listed in accordance with Article 8(2) of Directive 89/556/EEC.</p> <p>Box I.20.: Number of packages shall correspond to the number of containers.</p> <p>Box I.21.: identification of container and seal number shall be indicated.</p> <p>Box I.23.: fill in according to whether it is a transit or an import certificate.</p> <p>Box I.24.: fill in according to whether it is a transit or an import certificate.</p> <p>Box I.25.: Species: select amongst "Bos taurus", "Bison bison" or "Bubalus bubalis" as appropriate.</p> <p style="margin-left: 20px;">Category: select "in vitro derived embryos".</p> <p style="margin-left: 20px;">Dam identity shall correspond to the official identification of the animal.</p> <p style="margin-left: 20px;">Sire identity shall correspond to the official identification of the animal.</p> <p style="margin-left: 20px;">Date of freezing shall be indicated in the following format: dd.mm.yyyy</p> <p style="margin-left: 20px;">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 .</p> <p>Part II:</p> <p>(1) Delete as appropriate.</p> <p>(2) Only third countries listed in Annex I to Decision 2006/168/EC.</p> <p>(3) Only embryo production teams listed in accordance with Article 8(2) of Directive 89/556/.</p> <p style="margin-left: 20px;">The signature and the stamp must be in a different colour to that of the printing.</p>			
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 <b>Specimen not to be used for exports from EU</b> 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		I.16. Entry Point																
	<table border="1"> <thead> <tr> <th>Typ</th> <th>Doklad</th> <th>Identifikace</th> </tr> </thead> <tbody> <tr><td> </td><td> </td><td> </td></tr> <tr><td> </td><td> </td><td> </td></tr> <tr><td> </td><td> </td><td> </td></tr> <tr><td> </td><td> </td><td> </td></tr> </tbody> </table>		Typ	Doklad	Identifikace														
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I.18. Transport conditions Okolní <input type="checkbox"/> Controlled temperature <input type="checkbox"/> Zmrazené <input type="checkbox"/> Chlazený <input type="checkbox"/>		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 <input type="checkbox"/> Pollination <input type="checkbox"/> Krmivo pro zvířata <input type="checkbox"/> Použití pro farmaceutické účely <input type="checkbox"/> Game Restocking <input type="checkbox"/> Výkrmová <input type="checkbox"/> Evidování koňovití <input type="checkbox"/> Karanténa <input type="checkbox"/> Umělé rozmnožování <input type="checkbox"/> Training <input type="checkbox"/> Další postup <input type="checkbox"/> Racing <input type="checkbox"/> Production of petfood <input type="checkbox"/> Breeding and production <input type="checkbox"/> Consignments according to Regulation No 999/2001 <input type="checkbox"/> Competition <input type="checkbox"/> Storage <input type="checkbox"/> Breeding <input type="checkbox"/> Ornamental use/research <input type="checkbox"/> Laboratory <input type="checkbox"/> Rodent food <input type="checkbox"/> Zvířata v zájmovém chovu <input type="checkbox"/> Jiné <input type="checkbox"/> Production <input type="checkbox"/> Technické použití <input type="checkbox"/> Ornamental bird food <input type="checkbox"/> Sádkování <input type="checkbox"/> Porážka <input type="checkbox"/> Schválené orgány <input type="checkbox"/> Lidská spotřeba <input type="checkbox"/> Krmivo pro domácí zvířata <input type="checkbox"/> Sales <input type="checkbox"/> Organic fertilizers <input type="checkbox"/> Unregistered equidae <input type="checkbox"/> Cirkus/výstava <input type="checkbox"/> Category 3 fish oil/fish meal for detoxification according to Regulation 2015/786 <input type="checkbox"/>																			
I.21. For transit through a third country <input type="checkbox"/> Země _____ Kód ISO _____ EU Exit Authority _____ BCP code _____ EU Entry Authority _____ BCP code _____		I.22. For transit through Member State(s) <input type="checkbox"/> Země _____ Kód ISO _____																	
I.23. Celkový počet balení	I.24. Celkové množství	I.25. Celková čistá hmotnost	I.25. Celková hrubá hmotnost																

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	

Část I

SPECIMEN

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 collection 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 collection;

(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 collection and did not carry out vaccination against foot-and-mouth disease or lumpy skin disease during that period.]

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

- the embryos were not subjected to penetration of the zona pellucida,

- the embryos were stored under approved conditions for at least 30 days immediately after their collection,

- 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 embryos were collected.]

II.1.2. were collected 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 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.1.3. were collected and processed 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 dispatch to Great Britain, in the case of fresh embryos, or during the 30 days after collection, in the case of embryos subject to a mandatory storage for at least 30 days in accordance with point II.1.1.2.

II.1.4. from the time of collection until 30 days thereafter or, in the case of fresh embryos until the day of their dispatch to Great Britain, they 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.1.5. were collected from the donor females, which:

II.1.5.1. were located, during the 30 days immediately prior to collection, on premises situated in an area of at least 10-km radius on 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.1.5.2. showed no clinical signs of disease on the day of collection;

II.1.5.3. 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,

<b>Part II: Certification</b>	II. Informace týkající se zdraví		
	<p>- which were free from enzootic bovine leukosis or in which no bovine animal showed clinical signs of enzootic bovine leukosis during the previous three years,</p> <p>- in which no bovine animal showed clinical signs of infectious bovine rhinotracheitis/infectious pustular vulvo-vaginitis during the previous 12 months;</p> <p>II.1.6. The embryos to be exported were conceived by artificial insemination 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 part thereof listed in Annex 1 to implementing Decision 2011/630/EU (4) or by the competent authority of Great Britain.</p> <p>Notes</p> <p>(*) Those countries subject to the transitional import arrangements include: an EU member State; Liechtenstein; Norway and Switzerland.</p> <p>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).</p> <p>References to Great Britain in this certificate include Channel Islands and Isle of Man.</p> <p>Part I:</p> <p>Box I.6.: Person responsible for the load in Great Britain: this box is to be filled in only if it is a certificate for transit commodity.</p> <p>Box I.11.: Place of origin shall correspond to the embryo production team from which the embryos are dispatch to Great Britain and listed in accordance with Article 8(2) of Directive 89/556/EEC.</p> <p>Box I.20.: Number of packages shall correspond to the number of containers.</p> <p>Box I.21.: identification of container and seal number shall be indicated.</p> <p>Box I.23.: fill in according to whether it is a transit or an import certificate.</p> <p>Box I.24.: fill in according to whether it is a transit or an import certificate.</p> <p>Box I.25.: Species: select amongst "Bos taurus", "Bison bison" or "Bubalus bubalis" as appropriate.</p> <p>Category: select "in vitro derived embryos".</p> <p>Dam identity shall correspond to the official identification of the animal.</p> <p>Sire identity shall correspond to the official identification of the animal.</p> <p>Date of freezing shall be indicated in the following format: dd.mm.yyyy</p> <p>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.</p> <p>Part II:</p> <p>(1) Delete as appropriate.</p> <p>(2) Only third countries listed in Annex I to Decision 2006/168/EC.</p> <p>(3) Only embryo production teams listed in accordance with Article 8(2) of Directive 89/556/.</p> <p>The signature and the stamp must be in a different colour to that of the printing.</p>		
Certifying Officer			
Name (in capital letters)		Qualification and title	
Datum podpisu		Podpis	
Razítko			