

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				I.16 Entry Point			
Mode	International transport document	Identification					
I.18. Transport conditions Chilled <input type="checkbox"/> Controlled temperature <input type="checkbox"/> Frozen <input type="checkbox"/> Ambient <input type="checkbox"/>				I.17. Accompanying documents Commercial document reference Date of issue Country Place of issue			
I.19. Container No / Seal No							
I.20. Certified as Artificial reproduction <input type="checkbox"/> Breeding <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	
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		
	<p>The live animal(s) or animal product(s) herein described, complies/y with the relevant European Union standards and requirements which have been recognised as equivalent to the New Zealand standards and requirements as prescribed in the European Union/New Zealand Agreement on sanitary measures (Council Decision 97/132/EC), as last amended, specifically, in accordance with:</p> <p>Regulation (EU) 2016/429 of the European Parliament and of the Council of 9 March 2016 on transmissible animal diseases and amending and repealing certain acts in the area of animal health (Animal Health Law)</p> <p>Commission Delegated Regulation (EU) 2020/686 of 17 December 2019 supplementing Regulation (EU) 2016/429 of the European Parliament and of the Council as regards the approval of germinal product establishments and the traceability and animal health requirements for movements within the Union of germinal products of certain kept terrestrial animals</p> <p>Commission Delegated Regulation (EU) 2020/687 of 17 December 2019 supplementing Regulation (EU) 2016/429 of the European Parliament and the Council, as regards rules for the prevention and control of certain listed diseases</p> <p>Commission Delegated Regulation (EU) 2020/688 of 17 December 2019 supplementing Regulation (EU) 2016/429 of the European Parliament and of the Council, as regards animal health requirements for movements within the Union of terrestrial animals and hatching eggs</p>		
	<p>III. Additional health attestation</p> <p>III.1. The animal product is eligible for intra-Union trade without restriction.</p> <p>III.2. For diseases not regulated by the EU: All laboratory samples required by this veterinary certificate have been collected, processed, and stored in accordance with the OIE's recommendations or as described in Approved Diagnostic Tests, Vaccines, Treatments and Post-Arrival Testing Laboratories for Animal Import Health Standards, MPI-STD-TVTL, found here: https://www.mpi.govt.nz/dmsdocument/2040/;</p> <p>III.3. Q-Fever: Donors have never been confirmed positive for Q-Fever. and</p> <p>(2) ○ either III.3.1. [The donors were subjected to an ELISA test for Q fever, on a sample collected between 21 to 120 days after each embryo collection for export to New Zealand, with negative results.]</p> <p>(2) ○ or III.3.2. [A sample of embryos/oocytes, collection fluids and/or washing fluids from each germplasm collection for export to New Zealand was tested using a laboratory validated Q-Fever PCR test which is in accordance with the methods described in the Q-Fever Chapter of the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals.]</p> <p>III.4. To manage <i>Leptospira interrogans</i>, antibiotics have been added in accordance with the OIE Code, or with an approved combination listed in MPI-STD-TVTL, found here: https://www.mpi.govt.nz/dmsdocument/2040/.</p> <p>III.5. For <i>Mycoplasma bovis</i>, either:</p> <p>(2) ○ either III.5.1. [The embryos for export to New Zealand were subjected to the following treatment: After being washed 10 times, the embryos were subjected to incubation in tylosin (200 µg/mL) at 37°C for a minimum of 4 hours.]</p> <p>(2) ○ or III.5.2. [The embryos for export to New Zealand underwent DNA extraction and PCR testing as described in MPI-STD-TVTL, found here: https://www.mpi.govt.nz/dmsdocument/2040/.]</p> <p>(1)(3) <input type="checkbox"/> III.6. [The commodity herein described, complies/y with the additional conditions in the event of the occurrence of a disease:</p> <p> III.6.1. The commodity herein described was kept separate from all other commodities that did not meet the requirements during all stages of production, storage and transport and all necessary precautions were taken to prevent contamination of the commodity with any potential source of the relevant virus.</p> <p>(1) <input type="checkbox"/> III.6.2. [Foot and Mouth Disease: The in vivo derived embryos herein described were derived from donors that:</p>		

II. Health information

- III.6.2.1. were free of clinical signs of FMD, at the time of collection and from which the embryos were conceived by artificial insemination using semen collected, processed and stored in semen collection centres approved by the competent authority in conformity with OIE standards. In addition the embryos have been collected, processed and stored in accordance with standards laid down by the competent authority;
- and
- III.6.2.2. the donor animals from which the embryos were collected originate from a herd(s) that was/were not located within a protection or surveillance zone. Embryos collected within the protection and surveillance zones have been clearly identified and detained under official supervision.]
- (1) ☐ III.6.3. [Bluetongue:
- III.6.3.1. were free of clinical signs of BT at the time of collection and from which the embryos were conceived by artificial insemination using semen collected, processed and stored in semen collection centres approved by the competent authority in conformity with the OIE standards.
- and
- III.6.3.2. the embryos were collected, processed and stored in accordance with standards laid down by the competent authority.]
- (1) ☐ III.6.4. [Vesicular Disease:
- III.6.4.1. The embryos were kept for 21 days prior to, and during, collection in an establishment where no case of VS was reported during that period and were subject to a diagnostic test for VS, with negative results, within 21 days prior to embryo collection. In addition the embryos were collected, processed and stored in conformity with OIE notified standards and the establishment was not located within a protection or surveillance zone. Embryos collected within protection and surveillance zones has been clearly identified and detained under official supervision.]
- (1) ☐ III.6.5. [Lumpy Skin Disease:
- III.6.5.1. The embryos were derived from donor animals that showed no clinical sign of LSD on the day of collection of the embryos and for the following 28 days; and the animals were kept in the exporting country for the 28 days prior to collection, in an embryo collection centre where no case of LSD was officially reported during that period, and the centre was not situated in either a LSD infected zone or buffer zone and any semen from buffer zone has been clearly identified and controlled.]
- (1) ☐ III.6.6. [Rift Valley Fever:
- III.6.6.1. The donor must not show any clinical signs of RVF within the period from 14 days prior to and 14 days following germplasm collection;
- And
- (2) ☐ either III.6.6.2. [The donor must be vaccinated against RVF in accordance with MPI-STD-TVTL at least 14 days prior to collection.]
- (2) ☐ or III.6.6.3. [The donor must be demonstrated to be seropositive on the day of collection using a test listed in MPI-STD-TVTL.]
- (2) ☐ or III.6.6.4. [Testing of paired samples using a test listed in MPI-STD-TVTL must demonstrate that seroconversion did not occur between germplasm collection and 14 days after.]]
- (1) ☐ III.6.7. [Contagious Bovine Pleuropneumonia:

Part II: Certification	II. Health information									
	<p>The in vivo derived embryos herein described were derived from donors that:</p>									
	(2)	○ either	III.6.7.1.	[have not been vaccinated against CBPP and were subjected to the complement fixation test for CBPP with negative results, on two occasions, with an interval of not less than 21 days and not more than 30 days between each test, the second test being performed within 14 days prior to collection; and were isolated from other domestic bovidae from the day of the first complement fixation test until collection;]						
	(2)	○ or	III.6.7.2.	[were vaccinated using a vaccine complying with the standards described in the OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals not more than 4 months prior to collection;						
	and		III.6.7.3.	showed no clinical sign of CBPP on the day of collection of the embryos; and were kept (2) [since birth], or (2) [for the past 6 months], in a herd(s) where no case of CBPP was reported during that period, and that the herd(s) was/were not situated in a CBPP infected zone; and the embryos were collected, processed and stored in accordance with standards laid down by the competent authority.]]						
<p>Notes</p> <p>This health certificate is for veterinary purposes only.</p> <p>Part I</p> <p>Box I.6.: </p> <p>Box I.8.: Region of origin: if applicable, otherwise must be crossed out: for animal species or for products affected by the regionalisation measures or by the setting up of approved zones in accordance with Union decisions.</p> <p>Box I.11: Place of Origin shall correspond to the approved semen collection centre or semen storage centre listed in accordance with Article 9(2) of Directive 88/407/EEC on the Commission website: http://ec.europa.eu/food/animal/semen_ova/bovine/index_en.htm.</p> <p>Box I.14.: For animal products: indicate the date of departure of the means of transport (aeroplane, ship, railway or road vehicle).</p> <p>Box I.18.: Complete only in case of animal products.</p> <p>Box I.19.: Enter the 'Total gross weight (kg)' and 'Total net weight (kg)'.</p> <p>Box I.21.: Enter the identification number of the container and the official seal number.</p> <p>Box I.22.: Enter the intended use for animal products (the available options will vary in accordance with the specific certificate in the Union import requirements).</p> <p>Box I.24.: Complete only in case of importation or temporary admission to the Union.</p> <p>Box I.25.: Use the appropriate Harmonised System (HS) code under the following (nonexclusive) heading: 0511 99 85</p> <p>Part II</p> <p>(1) Only to be completed if special conditions apply. Otherwise delete.</p> <p>(2) Delete as appropriate.</p> <p>(3) In vivo derived embryos only (except embryos that have been subjected to penetration of the zona pellucida).</p>										
<p>Certifying Officer</p> <table border="0"> <tr> <td>Name (in capital letters)</td> <td>Qualification and title</td> </tr> <tr> <td>Date of signature</td> <td>Signature</td> </tr> <tr> <td>Stamp</td> <td></td> </tr> </table>					Name (in capital letters)	Qualification and title	Date of signature	Signature	Stamp	
Name (in capital letters)	Qualification and title									
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Stamp										

I. rész	I.1. Feladó Név Cím Ország ISO-kód			I.2. IMSOC hivatkozási szám Specimen not to be used for exports from EU	
				I.2.a. Local Reference	
	I.5. Címzett Név Cím Ország ISO-kód			I.3. Központi illetékes hatóság	
				I.4. Local competent authority	
	I.7. Származási ország ISO-kód		I.9. Country of destination ISO-kód		
	I.8. Region of origin Kód		I.10. Rendeltetési régió Kód		
	I.11. Place of Dispatch Név Cím Jóváhagyási szám Ország ISO-kód			I.12. Rendeltetési hely Név Cím Jóváhagyási szám Ország ISO-kód	
	I.13. Berakodás helye Név Cím Jóváhagyási szám Ország ISO-kód			I.14. Date and time of departure	
	I.15. Szállítóeszköz			I.16 Entry Point	
	Típus	Okmány	Azonosítás		
	I.18. Transport conditions Hűtött <input type="checkbox"/> Controlled temperature <input type="checkbox"/> Fagyasztva <input type="checkbox"/> Környezet <input type="checkbox"/>			I.17. Kísérőokmányok A kereskedelmi okmány hivatkozási száma Ország Kiállítás dátuma Kiállítás helye	
	I.19. Konténerszám/plombaszám				
I.20. Certified as Mesterséges szaporítás <input type="checkbox"/> Breeding <input type="checkbox"/>					
I.21. For transit through a third country <input type="checkbox"/> Country EU Exit Authority EU Entry Authority ISO-kód BCP code BCP code			I.22. For transit through Member State(s) <input type="checkbox"/> Country ISO-kód		
I.23. Csomagok teljes száma	I.24. Teljes mennyiség	I.25. Nettó össztömeg	I.25. Bruttó össztömeg		
I.28. Description of consignment 1. 05 MÁSHOVA NEM SOROLT, VAGY MÁSHOL FEL NEM TŰNTETETT ÉLELMEZÉSI CÉLRA ALKALMAS ÁLLATI EREDETŰ TERMÉK 0511 Másutt nem említett állati termék; az 1. vagy a 3. árucsoportba tartozó, emberi fogyasztásra nem alkalmas élettelen állat 051199 Más 05119985 Más					
Áru	Faj	Mennyiség	Tételszám	Gyártóüzem	
Hűtőház	Darabolóüzem	Fagyasztás dátuma	Az előállítás dátuma	Vágás dátuma	
Nettó tömeg	Product Description	Csomagok darabszáma	Azonosító jelölés		

Part II: Certification	II. Egészségügyi információk		
	<p>The live animal(s) or animal product(s) herein described, complies/y with the relevant European Union standards and requirements which have been recognised as equivalent to the New Zealand standards and requirements as prescribed in the European Union/New Zealand Agreement on sanitary measures (Council Decision 97/132/EC), as last amended, specifically, in accordance with:</p> <p>Regulation (EU) 2016/429 of the European Parliament and of the Council of 9 March 2016 on transmissible animal diseases and amending and repealing certain acts in the area of animal health (Animal Health Law)</p> <p>Commission Delegated Regulation (EU) 2020/686 of 17 December 2019 supplementing Regulation (EU) 2016/429 of the European Parliament and of the Council as regards the approval of germinal product establishments and the traceability and animal health requirements for movements within the Union of germinal products of certain kept terrestrial animals</p> <p>Commission Delegated Regulation (EU) 2020/687 of 17 December 2019 supplementing Regulation (EU) 2016/429 of the European Parliament and the Council, as regards rules for the prevention and control of certain listed diseases</p> <p>Commission Delegated Regulation (EU) 2020/688 of 17 December 2019 supplementing Regulation (EU) 2016/429 of the European Parliament and of the Council, as regards animal health requirements for movements within the Union of terrestrial animals and hatching eggs</p>		
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II. Egészségügyi információk

- III.6.2.1. were free of clinical signs of FMD, at the time of collection and from which the embryos were conceived by artificial insemination using semen collected, processed and stored in semen collection centres approved by the competent authority in conformity with OIE standards. In addition the embryos have been collected, processed and stored in accordance with standards laid down by the competent authority;
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