Export Health Certificate

	I.1. Consignor				I.2. IMSOC Re	ference			
	Name				Specimen not	to be used for exports	from El	U	
	Address				I.2.a. Local Reference				
	Country		ISO Code						
ŀ	I.5. Consignee				I.3. Central competent authority				
I	Name				I.4. Local competent authority				
E	Address								
Part I : Details of consignment	Country		ISO Code						
<u></u>									
ns	I.7. Country of origin			ISO Code	I.9. Country of destination ISO C		ISO Code		
잉									
of	I.8. Region of origin			Code	I.10. Region of				Code
i;	I.11. Place of Dispatch				I.12. Place of o	destination			
멅	Name				Name				
Ă۱	Address				Address				
ᆲ	Approval Number				Approval Number				
ar T	Country		ISO Code		Country			ISO Code	
ם	I.13. Place of Loading				I.14. Date and	time of departure			
	Name					•			
	Address								
	Approval Number								
	Country		ISO Code						
- }	145 M				TACE II. D.				
	I.15. Means of Transport		I.16 Entry Poi	nt					
	Mode Internation		Identification						
	documen	t							
					-				
ŀ	I.18. Transport conditions				I.17. Accompanying documents				
	Ambient	1	Controlled _ Fro	ozen 🗆	Commercial				
		•	temperature \square	document Date of issu			of issue		
					reference		Dlaga	of	
					Country		Place of issue	01	
	I.19. Container No / Seal No								
	700 0		7//						
	I.20. Certified as		Artificial reproduction	,					
	Breeding		Artificial reproduction	u Ш					
İ	I.21. For transit through a th	ird cour	ntry		I.22. For trans	sit through Member Sta	te(s)		
	Country		ISO Code						
	EU Exit		DCD l.						
	EU Exit Authority BCP code				Country		ICO C-	odo	
	Authority				Country		ISO Co	ode	
	Authority EU Entry Authority		BCP code		j				
	AuthorityEU Entry	es			Country I.25. Total net	weight		ode otal gross weigh	t
-	Authority EU Entry Authority I.23. Total number of packag		BCP code		j	weight			t
-	Authority EU Entry Authority I.23. Total number of packag I.28. Description of consignm	nent	BCP code I.24. Total quantity	ECIFIED OF TV	I.25. Total net	weight			t
-	Authority EU Entry Authority I.23. Total number of packag I.28. Description of consignm 1.05 PRODUCTS OF ANIMAI	nent L ORIGII	BCP code I.24. Total quantity N, NOT ELSEWHERE SP		I.25. Total net		I.25. T	otal gross weigh	t
-	Authority EU Entry Authority I.23. Total number of packag I.28. Description of consignm 1.05 PRODUCTS OF ANIMAI 0511 Animal products not	nent L ORIGII	BCP code I.24. Total quantity N, NOT ELSEWHERE SP		I.25. Total net		I.25. T	otal gross weigh	t
•	Authority EU Entry Authority I.23. Total number of packag I.28. Description of consignm 1.05 PRODUCTS OF ANIMAI	nent L ORIGII	BCP code I.24. Total quantity N, NOT ELSEWHERE SP		I.25. Total net		I.25. T	otal gross weigh	t
-	Authority EU Entry Authority I.23. Total number of packag I.28. Description of consignm 1. 05 PRODUCTS OF ANIMAI 0511 Animal products not 051199 Other 05119985 Other	nent L ORIGII	BCP code I.24. Total quantity N, NOT ELSEWHERE SP tere specified or include	ed; dead anima	I.25. Total net	or 3, unfit for human	I.25. T	otal gross weigh	
•	Authority EU Entry Authority I.23. Total number of packag I.28. Description of consignm 1. 05 PRODUCTS OF ANIMAI 0511 Animal products not 051199 Other	nent L ORIGII	BCP code I.24. Total quantity N, NOT ELSEWHERE SP tere specified or include		I.25. Total net		I.25. T	otal gross weigh	
•	Authority EU Entry Authority I.23. Total number of packag I.28. Description of consignm 1. 05 PRODUCTS OF ANIMAI 0511 Animal products not 051199 Other 05119985 Other Commodity	ent L ORIGII elsewh	BCP code I.24. Total quantity N, NOT ELSEWHERE SP tere specified or include	ed; dead anima Quantity	I.25. Total net CLUDED ls of Chapter 1	or 3, unfit for human	I.25. T	otal gross weigh nption Manufacturing	plant
•	Authority EU Entry Authority I.23. Total number of packag I.28. Description of consignm 1. 05 PRODUCTS OF ANIMAI 0511 Animal products not 051199 Other 05119985 Other	ent L ORIGII elsewh	BCP code I.24. Total quantity N, NOT ELSEWHERE SP tere specified or include	ed; dead anima	I.25. Total net CLUDED ls of Chapter 1	or 3, unfit for human	I.25. T	otal gross weigh	plant
•	Authority EU Entry Authority I.23. Total number of packag I.28. Description of consignm 1. 05 PRODUCTS OF ANIMAI 0511 Animal products not 051199 Other 05119985 Other Commodity Cold store	ent L ORIGII elsewh	BCP code I.24. Total quantity N, NOT ELSEWHERE SPacere specified or include es	ed; dead anima Quantity	I.25. Total net CLUDED ls of Chapter 1	or 3, unfit for human Batch number Date of production	I.25. T	otal gross weigh nption Manufacturing Date of slaught	plant
•	Authority EU Entry Authority I.23. Total number of packag I.28. Description of consignm 1. 05 PRODUCTS OF ANIMAI 0511 Animal products not 051199 Other 05119985 Other Commodity	ent L ORIGII elsewh	BCP code I.24. Total quantity N, NOT ELSEWHERE SP tere specified or include	ed; dead anima Quantity	I.25. Total net CLUDED ls of Chapter 1	or 3, unfit for human Batch number Date of production	I.25. T	otal gross weigh nption Manufacturing	plant
•	Authority EU Entry Authority I.23. Total number of packag I.28. Description of consignm 1. 05 PRODUCTS OF ANIMAI 0511 Animal products not 051199 Other 05119985 Other Commodity Cold store	ent L ORIGII elsewh	BCP code I.24. Total quantity N, NOT ELSEWHERE SPacere specified or include es	ed; dead anima Quantity	I.25. Total net CLUDED ls of Chapter 1	or 3, unfit for human Batch number Date of production	I.25. T	otal gross weigh nption Manufacturing Date of slaught	plant
•	Authority EU Entry Authority I.23. Total number of packag I.28. Description of consignm 1. 05 PRODUCTS OF ANIMAI 0511 Animal products not 051199 Other 05119985 Other Commodity Cold store	ent L ORIGII elsewh	BCP code I.24. Total quantity N, NOT ELSEWHERE SPacere specified or include es	ed; dead anima Quantity	I.25. Total net CLUDED ls of Chapter 1	or 3, unfit for human Batch number Date of production	I.25. T	otal gross weigh nption Manufacturing Date of slaught	plant
•	Authority EU Entry Authority I.23. Total number of packag I.28. Description of consignm 1. 05 PRODUCTS OF ANIMAI 0511 Animal products not 051199 Other 05119985 Other Commodity Cold store	ent L ORIGII elsewh	BCP code I.24. Total quantity N, NOT ELSEWHERE SPacere specified or include es	ed; dead anima Quantity	I.25. Total net CLUDED ls of Chapter 1	or 3, unfit for human Batch number Date of production	I.25. T	otal gross weigh nption Manufacturing Date of slaught	plant
•	Authority EU Entry Authority I.23. Total number of packag I.28. Description of consignm 1. 05 PRODUCTS OF ANIMAI 0511 Animal products not 051199 Other 05119985 Other Commodity Cold store	ent L ORIGII elsewh	BCP code I.24. Total quantity N, NOT ELSEWHERE SPacere specified or include es	ed; dead anima Quantity	I.25. Total net CLUDED ls of Chapter 1	or 3, unfit for human Batch number Date of production	I.25. T	otal gross weigh nption Manufacturing Date of slaught	plant
	Authority EU Entry Authority I.23. Total number of packag I.28. Description of consignm 1. 05 PRODUCTS OF ANIMAI 0511 Animal products not 051199 Other 05119985 Other Commodity Cold store	ent L ORIGII elsewh	BCP code I.24. Total quantity N, NOT ELSEWHERE SPacere specified or include es	ed; dead anima Quantity	I.25. Total net CLUDED ls of Chapter 1	or 3, unfit for human Batch number Date of production	I.25. T	otal gross weigh nption Manufacturing Date of slaught	plant

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Part II: Certification

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:

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)

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

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

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

III. Additional health attestation

- III.1. The animal product is eligible for intra-Union trade without restriction.
- 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/;
- III.3. For Q-Fever:

Donors have never been confirmed positive for Q-Fever.

and

- (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.]
- (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.]
 - III.4. For bovine viral diarrhea:
- (2) either III.4.1. [The donor animal was subjected to an antigen detection ELISA or virus isolation test for BVDV, with a negative result, within 30 days prior to the entry into the embryo collection centre and has been on the embryo collection centre for more than 6 months prior to embryo collection for this consignment and has remained isolated from other animals that have not been tested negative.]
- (2) or III.4.2. [The donor animal has had a sample of the unfiltered collection fluid or an embryo from the collection for export to New Zealand tested for BVDV2 with virus isolation or real-time RT PCR with negative results.]
 - III.5. To manage Leptospira interrogans, 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/.
 - III.6. For Mycoplasma bovis, either:
- (2) \circ either III.6.1. [The embryos for export to New Zeland were subjected to the following treatment: After being washed 10 times, the embryos were subjected to incubation in tylosin (200 μ g/mL) at 37°C for a minimum of 4 hours.]

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EUROPEAN UNION (NZ) Bovine Embryos

Eſ	JROPEAN	UNION			(NZ) Bovine Embryos
	II. Health inf	ormation			
	(2)	o or	III.6.2.	testing as	oryos for export to New Zeland underwent DNA extraction and PCR described in MPI-STD-TVTL, found here: vw.mpi.govt.nz/dmsdocument/2040/.]
	(1)(3)	□ III.7.	[The comr	-	ein described, complies/y with the additional conditions in the event of isease:
Part II: Certification			III.7.1.	nodity herein described was kept separate from all other commodities not meet the requirements during all stages of production, storage and and all necessary precautions were taken to prevent contamination of nodity with any potential source of the relevant virus.	
ဗီ		(1)	□ III.7.2.		l Mouth Disease:
Ħ				The in viv	vo derived embryos herein described were derived from donors that:
Paı				III.7.2.1.	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;
				III.7.2.2.	and 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
		(1)		[Dluston	official supervision.]
		(1)	□ III.7.3.	[Blueton; III.7.3.1.	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
			C	III.7.3.2.	the embryos were collected, processed and stored in accordance with standards laid down by the competent authority.]
		(1)	□ III.7.4.	[Vesicula	r Disease:
				III.7.4.1.	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.7.5.	[Lumpy S	Skin Disease:
				III.7.5.1.	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.7.6.	[Rift Vall	ey Fever:

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II. He	ealth information			
			III.7.6.1.	The donor must not show any clinical signs of RVF within the perio from 14 days prior to and 14 days following germplasm collection; And
	(2)	o either	III.7.6.2.	[The donor must be vaccinated against RVF in accordance with MP STD-TVTL at least 14 days prior to collection.]
	(2)	o or	III.7.6.3.	[The donor must be demonstrated to be seropositive on the day of collection using a test listed in MPI-STD-TVTL.]
	(2)	o or	III.7.6.4.	[Testing of paired samples using a test listed in MPI-STD-TVTL mus demonstrate that seroconversion did not occur between germplasm collection and 14 days after.]]
	(1)	□ III.7.7.	[Contagio	ous Bovine Pleuropneumonia:
1			The in viv	vo derived embryos herein described were derived from donors that:
	(2)	o either	III.7.7.1.	[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)	o or	III.7.7.2.	[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.7.7.3.	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 store in accordance with standards laid down by the competent authority.]]]
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en / cs **4** / 10

EU	IROPEAN U	JNION		(NZ) Bovine Embryos
	II. Health info	rmation		
	Notes			
		certificate is for veterinary purposes only.		
	Part I	r certificate is for veterifiary purposes offry.		
_	Box I.6.:			
tioī	Box I.8.:	Region of origin: if applicable, otherwise must	he crossed out: for animal en	acies or for products
ertifica	BOX 1.0	affected by the regionalisation measures or by Union decisions.		
Part II: Certification	Box I.11:	Place of Origin shall correspond to the approv in accordance with Article 9(2) of Directive 88/ http://ec.europa.eu/food/animal/semen_ova/bo	407/EEC on the Commission v	•
	Box I.14.:	For animal products: indicate the date of depa or road vehicle).	rture of the means of transpo	rt (aeroplane, ship, railway
	Box I.18.:	Complete only in case of animal products.		
	Box I.19.:	Enter the 'Total gross weight (kg)' and 'Total ne	et weight (kg)'.	
	Box I.21.:	Enter the identification number of the contain	er and the official seal number	er.
	Box I.22.:	Enter the intended use for animal products (the specific certificate in the Union import requires		in accordance with the
	Box I.24.:	Complete only in case of importation or temporation	orary admission to the Union.	
	Box I.25.:	Use the appropriate Harmonised System (HS) 85	code under the following (nor	nexclusive) heading: 0511 99
	Part II			
	(1)	Only to be completed if special conditions app	ly. Otherwise delete.	
	(2)	Delete as appropriate.		
	(3)	In vivo derived embryos only (except embryos pellucida).	s that have been subjected to p	penetration of the zona
	Certifying Offi			
	Name (in cap Date of signa Stamp		Qualification and title Signature	
		5		

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	I.1. Odesílatel					I.2. Referenční číslo IMSOC				
	Název					Specimen not to be used for exports from EU				
	Adresa Země			Kód ISO		I.2.a. Local Re	ference			
	I.5. Příjemce					I 2 Tatžadní nějalužný ongán				
	Název					I.3. Ústřední příslušný orgán I.4. Local competent authority				
	Nazev Adresa					1.4. Local com	peterit authority			
	Země			Kód ISO						
•	I.7. Země původu				Kód ISO	I.9. Country of	destination		Kód ISO)
ĭt I	I.8. Region of origi	n			Kód	I.10. Region ur	rčení		Kód	
	I.11. Place of Dispa					I.12. Místo urč				
	Název					Název				
	Adresa					Adresa				
	Číslo schválení					Číslo schváler	ní			
	Země			Kód ISO		Země			Kód ISO	
	I.13. Místo nakládl	ку				I.14. Date and	time of departure			
	Název									
	Adresa									
	Číslo schválení									
	Země			Kód ISO						
	I.15. Dopravní prostředky					I.16 Entry Poir	nt			
	Тур	Doklad		Identifikace						
	Тур	DORIGU		Identifikace		.^/				
	I.18. Transport cor	nditions				I.17. Průvodní doklady				
		Chlazený [Controlled _ Zm	razené 🗆	Referenční	autuuy			
		Critazon,		temperature \square	-	číslo obchodního do k ladu		Datum vydání		
						Země		Místo vydán	ú	
	I.19. Č. kontejneru	/ č. plomby	7							
	I.20. Certified as Breeding \square			Umělé rozmnožování						
ŀ	I.21. For transit th	rough a thi	rd cour	try		I 22 For trans	it through Member Sta	te(s)		
	Země	i ougii d tilli	i a couft	Kód ISO		1.44. FUI (Falls	n un ough member sta	ic(5)		
	EU Exit									
	Authority			BCP code		Země Kód ISO				
	EU Entry Authority			BCP code						
	I.23. Celkový počet	balení		I.24. Celkové množstv	í	I.25. Celková č	fistá hmotnost	I.25. C	elková hrubá hmotnost	
ŀ	I.28. Description of	f consignm	ent	I		l				
	-	_		DU, JINDE NEUVEDENI	έ ΔΝΙ ΝΕ 7 ΔΗΡΙ	MITÉ				
				•			pitol 1 nebo 3, nezpůs	ohilá k	lidskému nožívání	
	051199 Ostat		paroac	i, jiride rieuvederie urii	riezarii riace, rie	reva zvirata naj	prior 1 nebo o, nezpus	obna k	naskema pozivam	
	05119985 (
			Danish		Mr. a × atra-f		Číala žauža		V/	
	Komodita		Druh		Množství		Číslo šarže		Výrobní zařízení	
			1		<u> </u>		<u> </u>		<u> </u>	
	Chladírenské zaří	zení	Bourá	rna	Datum zmraze	ení	Datum výroby		Datum porážky	
	Čistá hmotnost			Product Description		Počet balení		Identi	fikační značka	
				<u></u> _						

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		7 7	
_	II. Informace týkající se zdraví		

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:

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)

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

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

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

III. Additional health attestation

- III.1. The animal product is eligible for intra-Union trade without restriction.
- 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/;
- III.3. For Q-Fever:

Donors have never been confirmed positive for Q-Fever.

and

Part II: Certification

- (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.]
- (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.]
 - III.4. For bovine viral diarrhea:
- (2) either III.4.1. [The donor animal was subjected to an antigen detection ELISA or virus isolation test for BVDV, with a negative result, within 30 days prior to the entry into the embryo collection centre and has been on the embryo collection centre for more than 6 months prior to embryo collection for this consignment and has remained isolated from other animals that have not been tested negative.]
- (2) or III.4.2. [The donor animal has had a sample of the unfiltered collection fluid or an embryo from the collection for export to New Zealand tested for BVDV2 with virus isolation or real-time RT PCR with negative results.]
 - III.5. To manage Leptospira interrogans, 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/.
 - III.6. For Mycoplasma bovis, either:
- (2) \circ either III.6.1. [The embryos for export to New Zeland were subjected to the following treatment: After being washed 10 times, the embryos were subjected to incubation in tylosin (200 μ g/mL) at 37°C for a minimum of 4 hours.]

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EVROPSKÁ UNIE (NZ) Bovine Embryos

							(,
	II. Informace t	ýkající se zdra	ví				
	(2)	o or	III.6.2.	testing as d	lescribed in	ort to New Zeland underwent MPI-STD-TVTL, found here: nz/dmsdocument/2040/.]	DNA extraction and PCR
	(1)(3)	□ III.7.		nodity herei		complies/y with the addition	al conditions in the event of
Part II: Certification			III.7.1.	that did no transport a	ot meet the reand all neces	described was kept separate equirements during all stages sary precautions were taken by potential source of the rele	of production, storage and to prevent contamination of
ပ္		(1)	□ III.7.2.	[Foot and	Mouth Disea	se:	
ᄪ				The in vivo	derived em	bryos herein described were	derived from donors that:
Paı				III.7.2.1.	which the e semen colle approved b standards.	f clinical signs of FMD, at the embryos were conceived by a ected, processed and stored in the competent authority in addition the embryos have in accordance with standards	rtificial insemination using a semen collection centres conformity with OIE
				III.7.2.2.	from a hero surveillanc	nnimals from which the embral(s) that was/were not located e zone. Embryos collected wire zones have been clearly ide ervision.]	within a protection or thin the protection and
		(1)	□ III.7.3.	[Bluetong	ue:		
			4	III.7.3.1.	which the e	of clinical signs of BT at the tire embryos were conceived by a ected, processed and stored in any the competent authority in	rtificial insemination using a semen collection centres
			C	III.7.3.2.	,	s were collected, processed a ards laid down by the compet	
		(1)	□ III.7.4.	[Vesicular	Disease:		
				III.7.4.1.	an establish period and results, wit embryos w OIE notified a protection protection	os were kept for 21 days prion nament where no case of VS wa were subject to a diagnostic t hin 21 days prior to embryo c ere collected, processed and s d standards and the establish n or surveillance zone. Embry and surveillance zones has be nder official supervision.]	as reported during that test for VS, with negative collection. In addition the stored in conformity with ment was not located within yos collected within
		(1)	□ III.7.5.	[Lumpy Sl	kin Disease:		
				III.7.5.1.	clinical sign the followin country for centre whe period, and or buffer zo	os were derived from donor an of LSD on the day of collecting 28 days; and the animals we the 28 days prior to collection re no case of LSD was official the centre was not situated if one and any semen from buffund controlled.]	on of the embryos and for vere kept in the exporting n, in an embryo collection ly reported during that n either a LSD infected zone
		(1)	□ III.7.6.	[Rift Valle	y Fever:		

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II. Inform	nace týkající se zo	draví		
			III.7.6.1.	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)	o either	III.7.6.2.	[The donor must be vaccinated against RVF in accordance with MP STD-TVTL at least 14 days prior to collection.]
	(2)	\circ or	III.7.6.3.	[The donor must be demonstrated to be seropositive on the day of collection using a test listed in MPI-STD-TVTL.]
	(2)	o or	III.7.6.4.	[Testing of paired samples using a test listed in MPI-STD-TVTL mus demonstrate that seroconversion did not occur between germplasm collection and 14 days after.]]
	(1)	□ III.7.7.	[Contagio	ous Bovine Pleuropneumonia:
			The in viv	o derived embryos herein described were derived from donors that:
	(2)	∘ either	III.7.7.1.	[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)	o or	III.7.7.2.	[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.7.7.3.	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 store in accordance with standards laid down by the competent authority.]]]
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EVROPSKÁ UNIE (NZ) Bovine Embryos

EVROPSKÁ U	UNIE		(NZ) Bovine Embry							
II. Informace	týkající se zdraví									
Notes										
This health certificate is for veterinary purposes only.										
Part I	recording to the control purposes of the									
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.									
Box I.6.: Box I.8.: Box I.11:										
Box I.14.:	I.14.: For animal products: indicate the date of departure of the means of transport (aeroplane, ship, railway or road vehicle).									
Box I.18.:										
Box I.19.:										
Box I.21.:	Enter the identification number of the contain	er and the official seal number	er.							
Box I.22.:	Enter the intended use for animal products (the specific certificate in the Union import require		in accordance with the							
Box I.24.:	Complete only in case of importation or temporation	orary admission to the Union.								
Box I.25.:	Use the appropriate Harmonised System (HS) code under the following (nonexclusive) heading: 0511 99 85									
Part II										
(1)	Only to be completed if special conditions app	ly. Otherwise delete.								
(2)	Delete as appropriate.									
(3)	In vivo derived embryos only (except embryos pellucida).	s that have been subjected to p	penetration of the zona							
Certifying Of										
Name (in ca Datum podr Razítko	pital letters) pisu	Qualification and title Podpis								

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