INTRA

	I.1. Consignor				I.2. IMSOC reference I.2.a. Local reference			ence	
	Name Address						I.3. Central Comp	petent Authority	
	Country ISO Code					I.4. Local Compe	tent Authority		
ent	I.5. Consignee				I.6. Operator conducting assembly operations independently of an establishment				
m	Name Address				Name				
\mathbf{sig}	Country		ISO Code		Address				
ons					Approval Number				
$^{ m c}$					Country ISO Code				
tion (I.7. Country of ori	gin	ISO Code	9	I.9. Country of destination	on		ISO Code	
Part I: Description of consignment	I.8. Region of origin Code				I.10. Region of destination	on		Code	
)es	I.11. Place of dispatch				I.12. Place of destination	າ			
I: I	Name				Name				
ırt	Address Approval				Address Approval				
Ρĉ	Number				Number				
	Country		ISO Code		Country		ISO Code		
	I.13. Place of loadi	ing			I.14. Date and time of de	eparture			
	Name								
_	Approval								
	Approval Number								
	Country	Country ISO Code							
	I.15. Means of Tra		- 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		I.16. Transporter				
	Mode	International transport	Identification		Name				
		document			Address Approval				
					Number		100 0 1		
					Country		ISO Code		
					I.17. Accompanying doc	uments			
					Document Type Accompanying document				
				Date of Issue Country Place of issue					
	=	.18. Transport conditions							
	Chilled Ambient Ambient			Frozen 🗆					
	I.19. Container No / Seal No								
	I.20. Certified as								
	Germinal product	s L							
	I.21. For transit through a third country								
	Third country Exit point Entry point			ISO Code					
				BCP code BCP code					
	I.22. For transit through Member State(s)				I.23. For export				
	Member State ISO Code I.24. Estimated journey time			Third country	Third country ISO Code				
				Exit point BCP code I.25. Journey Log					
	I.26. Total number		I.27. Total quan	ntity	I.28. Total gross weight				
	I.30. Description o		1 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	,		1	- 0 .		
	_	_	, NOT ELSEWHERE SPECIFIE	ED OR IN	CLUDED				
	0511 Animal products not elsewhere specified or included; dead animals of Chapter 1 or 3, unfit for human consumption 051199 Other								
	05119985 Other								

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_	ION	1	T	T		INIK
	#1. Commodity	Identification Number	Quantity	Nature of commodity	Identification Mark	
	Species	Package count	Date of collection / production	Plant / Establishment / Centre	Туре	
5						
tar transferred to conditions transferred						
b						
1						
I						
I						
I						

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	II. Health information								
	I, the undersigned official veterinarian, hereby certify that:								
	II.1.								
		(2) approved by the competent authority and processed in an appropriate laboratory;							
	II.2.	□ [Ova] (1)	□ [Ova] (1) □ [embryos] (1) were collected from donor mares which:						
Part II: Certification		II.2.1.	on the day of collection have been located in premises situated on the territory or in the case of regionalisation in a part of the territory of a Member State which is not considered to be infected with African horse sickness in accordance with Article 5(2), poins (a) and (b) of Directive 2009/156/EC (3),						
		II.2.2.			er veterinary supervision which on the day of collection of Directive 2009/156/EC,				
		II.2.3.	have been kept price equine metritis for		ion in holdings	free from clin	ical signs of contagious		
		II.2.4.	have not been used for natural breeding during the period of 30 days prior to the collection of \Box [ova] (1) \Box [embryos] (1),						
		II.2.5.	to the best of my knowledge and as far as I could ascertain, have not been in contact with equidae suffering from an infectious or contagious disease during the 15 days immediate preceding the collection of \square [ova] (1) \square [embryos] (1),						
		II.2.6.	have on the day of	collection not s	hown clinical s	igns of an infe	ctious or contagious disease;		
	II.3.	□ [Ova] (1) □ [embryos] (1):							
	II.3.1.	were collected, processed, stored and transported under conditions which comply with the requirements of Annex D of Directive 92/65/EEC;							
	II.3.2.	are dispatched:							
	(1) either	o [by an □ [embryo collection team] (1) □ [embryo production team] (1) or from a zone not subject to movement restrictions affecting equine animals and established for reasons of listed diseases relevant for those species or diseases subject to emergency measures relevant for those species, or those restrictions do not apply to these ova or embryos because they were collected before the restrictions were established, and they have not been in contact with other ova or embryos of a lower health status for an adequate period.]							
	(1) or \circ [by an \square [embryo collection team] (1) \square [embryo production team] (1) or from a zone subject to movement restrictions affecting equine animals and established for (4), but derogations from movement restrictions have been granted, and:								
	(1)	☐ [they co	mply with the requir	rements set out	in	(5);]]			
	(1) 🗆 [and	in particula	r, they are	(6).]]					
	(7) (1)	□ [II.4.	The semen used for requirements of Di			the donor mar	res complies with the		
	(1)	□ [II.5.	The ova used for th Directive 92/65/EEC	_	action of embry	os comply wit	h the requirements of		
	Notes								
	In accordance with the Agreement on the withdrawal of the United Kingdom of Great Britain and Northern Ireland from the European Union and the European Atomic Energy Community, and in particular Article 5(4) of the Protocol on Ireland/Northern Ireland in conjunction with Annex 2 to that Protocol, references to the Union in this animal health certificate include the United Kingdom in respect of Northern Ireland.								
	This animal health certificate shall be completed in accordance with the notes for the completion of certificates provided for in Chapter 2 of Annex I to Commission Implementing Regulation (EU) 2020/2235.								
	Part I:								
	Box Place of dispatch shall correspond to the embryo collection team of ova/embryos collection. reference I.11:								
	Box Place of destination shall correspond to the embryo collection team, embryo production team or to the reference holding of ova/embryos destination								

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		,							
	II. Health info	rmation							
	I.12:								
	Box reference I.26:	Box Total number of packages shall correspond to the number of containers. reference							
	Box reference I.19:	Seal number shall be indicated.							
	Box reference I.30:	"Type": Specify if: in vivo derived embryos, in vivo derived oocytes, in vitro produced embryos or micromanipulated embryos.							
Irt I		Donor identity shall correspond to the official identification of the animal.							
Pa		Date of collection shall be indicated in the following format:dd/mm/yyyy.							
		Approval number of the team shall correspond to the embryo collection team of ova/embryos collection.							
	Part II:	'art II:							
	(1)	Delete if not applicable.							
	(2)	Only embryo collection teams approved by the competent authority and listed in accordance with Article 11(4) of Council Directive 92/65/EEC.							
	(3)	OJ L 192, 23.7.2010, p. 1.							
	(4)	Insert the name of the disease(s).							
	(5)	Insert the specific reference to the article(s), title, and number of the relevant legal act(s) adopted by the Commission providing for those requirements.							
	(6)	Insert the specific attestation(s) provided for in and required by the relevant legal act(s) adopted by the Commission, as referred to in Article 159(2), points (a), (b) and (c), of Regulation (EU) 2016/429 of the European Parliament and of the Council.							
	(7)	Does not apply to ova.							
	Certifying Offi Name (in capi	icer/Official veterinarian tal letters)							
	Date of declar Stamp	· · · · · · · · · · · · · · · · · · ·							
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