EUROPEAN UNION INTRA

	I.1. Consignor				I.2. IMSOC ref	erence	I.2.a. Local reference	
	_						I.3. Central Competent Authority	
	Name							
	Address						I.4. Local Competent Authority	
	Country		ISO Code					
	100 code							
	I.5. Consignee					I.6. Operator conducting assembly operations independently of an		
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1 9	Name				Nome			
ਖ਼	Address				Name			
g	Country		ISO Code		Address			
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—	I.8. Region of origin			Code	I.10. Region of		Code	
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	I.13. Place of loadi	ng			I.14. Date and	time of departure		
		0				or acparture		
	Name							
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	I.15. Means of Tran	nsport			I.16. Transpor	ter		
	Mode	International	Identification		Name			
		transport			Address			
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	II. Health info	rmation							
	I, the undersigned official veterinarian, hereby certify that:								
	II.1.		erminal product storage centre(1) described in Box I.11. at which the semen(2)/ oocytes(2)/ in vivo ed embryos(2)/ in vitro produced embryos(2)/ micromanipulated embryos(2) was/were stored:						
		II.1.1.	I.1.1. is approved and kept in a register by the competent authority;						
ication		II.1.2.			ards responsibilities, operation Annex I to Commission Deleg				
Certif	II.2.				ryos(2)/ in vitro produced em for artificial reproduction an				
	(2) □ either	[II.2.1.	has/have been collected or produced, processed and stored in a semen collection centre(2)(3)/ by an embryo collection team(2)(3)/ by an embryo production team(2)(3), and/or processed and stored in a germinal product processing establishment(2)(3), and/or stored in a germinal product storage centre(2)(3) situated in the Member State of its/their collection or production and complying with requirements as regards responsibilities, operational procedures, facilities and equipment set out in Part 1(2)/Part 2(2)/Part 3(2)/Part 4(2)/Part 5(2) of Annex I to Delegated Regulation (EU) 2020/686, and was/were moved to the germinal product storage centre indicated in Box I.11. situated in the Member State of its/their collection or production under animal health certification requirements at least as strict as those provided for in:						
		(2)	\square either	[Model EQUI-SEM-A-IN	TRA(4);]				
		(2)	\square and/or	[Model EQUI-SEM-B-IN	TRA(4);]				
		(2)	\square and/or	[Model EQUI-SEM-C-IN	TRA(4);]				
		(2)	□ and/or [Model EQUI-SEM-D-INTRA(4);]						
		(2)	\square and/or	[Model EQUI-OOCYTES	-EMB-A-INTRA(4);]				
		(2)	□ and/or [Model EQUI-OOCYTES-EMB-B-INTRA(4);]						
		(2)	\square and/or	[Model EQUI-OOCYTES	-EMB-C-INTRA(4);]				
		(2)	\square and/or	[Model EQUI-OOCYTES	-EMB-D-INTRA(4);]				
		(2)	\square and/or	[Model EQUI-GP-PROCE	ESSING-INTRA(4);]				
		(2)	\square and/or	[Model EQUI-GP-STORA	AGE-INTRA(4);]]				
		(2)	\square and/or	[Model IA in Part A of A	Annex I to Decision 2010/470/	EU(4);]			
		(2)	\square and/or	[Model IB in Part B of A	Annex I to Decision 2010/470/I	EU(4);]			
		(2)	\square and/or	[Model IC in Part C of A	annex I to Decision 2010/470/F	EU(4);]			
		(2)	\square and/or	[Model ID in Part D of A	Annex I to Decision 2010/470/	EU(4);]			
		(2)	\square and/or	[Model in Commission	Decision 95/307/EC(4);]				
	(2) □ and/or	[II.2.1.	centre(2)(3 and/or pro stored in a collection of operational 4(2)/Part 50 germinal p)/ by an embryo collection cessed and stored in a go germinal product storage or production and comp Il procedures, facilities a (2) of Annex I to Delegato	ed, processed and stored in a on team(2)(3)/ by an embryo perminal product processing ege centre(2)(3) situated in the lying with requirements as rend equipment set out in Part ed Regulation (EU) 2020/686, adicated in Box I.11. situated it cordance with:	production team(2)(3), stablishment(2)(3), and/or Member State of its/their egards responsibilities, 1(2)/Part 2(2)/Part 3(2)/Part and was/were moved to the			
		(2)	□ either	[Model EQUI-SEM-A-IN					
		(2)	□ and/or	[Model EQUI-SEM-B-IN					
		(2)		[Model EQUI-SEM-C-IN]					
		(2)		[Model EQUI-SEM-D-IN					
		(2)	□ and/or	[Model EQUI-OOCYTES					
		(2)		[Model EQUI-OOCYTES					
		(2)	□ and/or	[Model EQUI-OOCYTES	-EMB-C-INTRA(4);]				

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The shift information		ROTEEN CIVICIA			510MdL-MilM)		
Company Comp		II. Health info	rmation				
Bandor Model EQUI-GP-STORAGE-INTRA(d):]			(2)	□ and/or	[Model EQUI-OOCYTES	-EMB-D-INTRA(4);]	
The part of the product of the pro				□ and/or	•		
Care			(2)	□ and/or	[Model EQUI-GP-STOR	AGE-INTRA(4);]]	
Care			(2)	□ and/or	[Model IA in Part A of A	Annex I to Decision 2010/470/	EU(4);]
(2)	u			□ and/or			
and/or and/or centre(2)(3)/ by an embryo collection team(2)(3)/ by an embryo production team(2)(3)/ by an embryo collection team(2)(3)/ by an embryo production team(2)(3), and/or processed and stored in a germinal product storage centre(2)(3) situated in a third country, territory or zone thereof listed in Annex XII to Commission Implementing Regulation (EU) [C(2021)1800) and complying with requirements as regards responsibilities, operational procedures, facilities and equipment set out in Part 12()Part 3(2)Part 4(2)Part 4(2)Pa	atio			•			
and/or and/or centre(2)(3)/ by an embryo collection team(2)(3)/ by an embryo production team(2)(3)/ by an embryo collection team(2)(3)/ by an embryo production team(2)(3), and/or processed and stored in a germinal product storage centre(2)(3) situated in a third country, territory or zone thereof listed in Annex XII to Commission Implementing Regulation (EU) [C(2021)1800) and complying with requirements as regards responsibilities, operational procedures, facilities and equipment set out in Part 12()Part 3(2)Part 4(2)Part 4(2)Pa	ific			□ and/or			
and/or and/or centre(2)(3)/ by an embryo collection team(2)(3)/ by an embryo production team(2)(3)/ by an embryo collection team(2)(3)/ by an embryo production team(2)(3), and/or processed and stored in a germinal product storage centre(2)(3) situated in a third country, territory or zone thereof listed in Annex XII to Commission Implementing Regulation (EU) [C(2021)1800) and complying with requirements as regards responsibilities, operational procedures, facilities and equipment set out in Part 12()Part 3(2)Part 4(2)Part 4(2)Pa	ert						
(2)	Part II: ((2)	[II.2.1.	centre(2)(3 and/or pro stored in a zone there and compl facilities at to Delegate	s)/ by an embryo collecticessed and stored in a germinal product storator of listed in Annex XII to ying with requirements and equipment set out in the degulation (EU) 2020/	on team(2)(3)/ by an embryo perminal product processing ege centre(2)(3) situated in a the Commission Implementing Ras regards responsibilities, of Part 1(2)/Part 2(2)/Part 3(2)/Part 3(2)/Part 3(2)/Part 2(2)/Part 3(2)/Part 2(2)/Part 3(2)/Part 2(2)/Part 3(2)/Part 2(2)/Part 3(2)/Part 3(2	production team(2)(3), stablishment(2)(3), and/or nird country, territory or egulation (EU) [C(2021)1800] perational procedures, art 4(2)/Part 5(2) of Annex I
C2			(2)	\square either	[Model EQUI-SEM-A-EN	NTRY(4);]	
C2			(2)	□ and/or	[Model EQUI-SEM-B-EN	VTRY(4);]	
C2			(2)	□ and/or	[Model EQUI-SEM-C-EN	JTRY(4);]	
(2) □ and/or [Model EQUI-OOCYTES-EMB-B-ENTRY(4);] (2) □ and/or [Model EQUI-OOCYTES-EMB-C-ENTRY(4);] (2) □ and/or [Model EQUI-GP-PROCESSING-ENTRY(4);] (2) □ and/or [Model EQUI-GP-STORAGE-ENTRY(4);] (2) □ and/or [Model 1 in Section A of Part 1 of Annex III to Regulation (EU) 2018/659(4);] (2) □ and/or [Model 2 in Section B of Part 1 of Annex III to Regulation (EU) 2018/659(4);] (2) □ and/or [Model 3 in Section C of Part 1 of Annex III to Regulation (EU) 2018/659(4);] (2) □ and/or [Model 4 in Section D of Part 1 of Annex III to Regulation (EU) 2018/659(4);] (2) □ and/or [Model 1 in Section A of Part 2 of Annex II to Decision 2010/471/EU(4);] (2) □ and/or [Model 2 in Section B of Part 2 of Annex II to Decision 2010/471/EU(4);] (2) □ and/or [Model 3 in Section C of Part 2 of Annex II to Decision 2010/471/EU(4);] (2) □ and/or [Model in Annex to Commission Decision 96/539/EC(4);] II.2.2. has/have been collected, processed and stored in accordance with animal health requirements set out in Annex III to Delegated Regulation (EU) 2020/686; II.2.3. is/are placed in straws or other packages on which the mark is applied in accordance with requirements provided for in Article 10 of Delegated Regulation (EU) 2020/686 and/or Article 83(a) of Commission Delegated Regulation (EU) 2020/692 and that mark is indicated in Box 1.30; II.2.4. is/are transported in a container which: II.2.4.1. was sealed and numbered prior to the dispatch from the germinal product storage centre under responsibility of the centre veterinarian, or by an official veterinarian, and the seal bears the number as indicated in Box 1.19; II.2.4.2. has been cleaned and either disinfected or sterilised before use, or is single-use container; (2)(5) □			(2)	□ and/or	[Model EQUI-SEM-D-EN	NTRY(4);]	
C2			(2)	\square and/or	[Model EQUI-OOCYTES	-EMB-A-ENTRY(4);]	
C2			(2)	\square and/or	[Model EQUI-OOCYTES	-EMB-B-ENTRY(4);]	
(2) □ and/or [Model EQUI-GP-STORAGE-ENTRY(4);] (2) □ and/or [Model 1 in Section A of Part 1 of Annex III to Regulation (EU) 2018/659(4);] (2) □ and/or [Model 2 in Section B of Part 1 of Annex III to Regulation (EU) 2018/659(4);] (2) □ and/or [Model 3 in Section C of Part 1 of Annex III to Regulation (EU) 2018/659(4);] (2) □ and/or [Model 4 in Section D of Part 1 of Annex III to Regulation (EU) 2018/659(4);] (2) □ and/or [Model 1 in Section A of Part 2 of Annex II to Decision 2010/471/EU(4);] (2) □ and/or [Model 2 in Section B of Part 2 of Annex II to Decision 2010/471/EU(4);] (2) □ and/or [Model 3 in Section C of Part 2 of Annex II to Decision 2010/471/EU(4);] (2) □ and/or [Model in Annex to Commission Decision 96/539/EC(4);] II.2.2. has/have been collected, processed and stored in accordance with animal health requirements set out in Annex III to Delegated Regulation (EU) 2020/686; II.2.3. is/are placed in straws or other packages on which the mark is applied in accordance with requirements provided for in Article 10 of Delegated Regulation (EU) 2020/686 and/or Article 83(a) of Commission Delegated Regulation (EU) 2020/692 and that mark is indicated in Box I.30; II.2.4. is/are transported in a container which: II.2.4.1. was sealed and numbered prior to the dispatch from the germinal product storage centre under responsibility of the centre veterinarian, or by an official veterinarian, and the seal bears the number as indicated in Box I.19; II.2.4.2. has been cleaned and either disinfected or sterilised before use, or is single-use container; has been filled in with the cryogenic agent which not have been previously used for other products;]			(2)	\square and/or	[Model EQUI-OOCYTES	-EMB-C-ENTRY(4);]	
(2)			(2)	\square and/or	[Model EQUI-GP-PROC	ESSING-ENTRY(4);]	
(2) □ and/or [Model 2 in Section B of Part 1 of Annex III to Regulation (EU) 2018/659(4);] (2) □ and/or [Model 3 in Section C of Part 1 of Annex III to Regulation (EU) 2018/659(4);] (2) □ and/or [Model 1 in Section D of Part 1 of Annex III to Regulation (EU) 2018/659(4);] (2) □ and/or [Model 1 in Section A of Part 2 of Annex II to Decision 2010/471/EU(4);] (2) □ and/or [Model 2 in Section B of Part 2 of Annex II to Decision 2010/471/EU(4);] (2) □ and/or [Model 3 in Section C of Part 2 of Annex II to Decision 2010/471/EU(4);] (2) □ and/or [Model in Annex to Commission Decision 96/539/EC(4);] II.2.2. has/have been collected, processed and stored in accordance with animal health requirements set out in Annex III to Delegated Regulation (EU) 2020/686; II.2.3. is/are placed in straws or other packages on which the mark is applied in accordance with requirements provided for in Article 10 of Delegated Regulation (EU) 2020/686 and/or Article 83(a) of Commission Delegated Regulation (EU) 2020/692 and that mark is indicated in Box I.30; II.2.4. is/are transported in a container which: II.2.4.1. was sealed and numbered prior to the dispatch from the germinal product storage centre under responsibility of the centre veterinarian, or by an official veterinarian, and the seal bears the number as indicated in Box I.19; II.2.4.2. has been cleaned and either disinfected or sterilised before use, or is single-use container; (2)(5) □ has been filled in with the cryogenic agent which not have been previously used for other products;]			(2)	\square and/or	[Model EQUI-GP-STOR	AGE-ENTRY(4);]	
(2) □ and/or [Model 3 in Section C of Part 1 of Annex III to Regulation (EU) 2018/659(4);] (2) □ and/or [Model 4 in Section D of Part 1 of Annex III to Regulation (EU) 2018/659(4);] (2) □ and/or [Model 1 in Section A of Part 2 of Annex II to Decision 2010/471/EU(4);] (2) □ and/or [Model 2 in Section B of Part 2 of Annex II to Decision 2010/471/EU(4);] (2) □ and/or [Model 3 in Section C of Part 2 of Annex II to Decision 2010/471/EU(4);] (2) □ and/or [Model in Annex to Commission Decision 96/539/EC(4);] II.2.2. has/have been collected, processed and stored in accordance with animal health requirements set out in Annex III to Delegated Regulation (EU) 2020/686; II.2.3. is/are placed in straws or other packages on which the mark is applied in accordance with requirements provided for in Article 10 of Delegated Regulation (EU) 2020/686 and/or Article 83(a) of Commission Delegated Regulation (EU) 2020/692 and that mark is indicated in Box I.30; II.2.4. is/are transported in a container which: II.2.4.1. was sealed and numbered prior to the dispatch from the germinal product storage centre under responsibility of the centre veterinarian, or by an official veterinarian, and the seal bears the number as indicated in Box I.19; II.2.4.2. has been cleaned and either disinfected or sterilised before use, or is single-use container; (2)(5) □ [II.2.4.3. is been filled in with the cryogenic agent which not have been previously used for other products;]			(2)	□ and/or	[Model 1 in Section A o	of Part 1 of Annex III to Regula	ntion (EU) 2018/659(4);]
(2) □ and/or [Model 4 in Section D of Part 1 of Annex III to Regulation (EU) 2018/659(4);] (2) □ and/or [Model 2 in Section B of Part 2 of Annex II to Decision 2010/471/EU(4);] (2) □ and/or [Model 3 in Section C of Part 2 of Annex II to Decision 2010/471/EU(4);] (2) □ and/or [Model 3 in Section C of Part 2 of Annex II to Decision 2010/471/EU(4);] (2) □ and/or [Model in Annex to Commission Decision 96/539/EC(4);] II.2.2. has/have been collected, processed and stored in accordance with animal health requirements set out in Annex III to Delegated Regulation (EU) 2020/686; II.2.3. is/are placed in straws or other packages on which the mark is applied in accordance with requirements provided for in Article 10 of Delegated Regulation (EU) 2020/686 and/or Article 83(a) of Commission Delegated Regulation (EU) 2020/692 and that mark is indicated in Box I.30; II.2.4. is/are transported in a container which: II.2.4.1. was sealed and numbered prior to the dispatch from the germinal product storage centre under responsibility of the centre veterinarian, or by an official veterinarian, and the seal bears the number as indicated in Box I.19; II.2.4.2. has been cleaned and either disinfected or sterilised before use, or is single-use container; (2)(5) □ II.2.4.3. has been filled in with the cryogenic agent which not have been previously used for other products;]			(2)	\square and/or	[Model 2 in Section B o	f Part 1 of Annex III to Regula	ation (EU) 2018/659(4);]
(2) □ and/or [Model 1 in Section A of Part 2 of Annex II to Decision 2010/471/EU(4);] (2) □ and/or [Model 2 in Section B of Part 2 of Annex II to Decision 2010/471/EU(4);] (2) □ and/or [Model 3 in Section C of Part 2 of Annex II to Decision 2010/471/EU(4);] (2) □ and/or [Model in Annex to Commission Decision 96/539/EC(4);] II.2.2. has/have been collected, processed and stored in accordance with animal health requirements set out in Annex III to Delegated Regulation (EU) 2020/686; II.2.3. is/are placed in straws or other packages on which the mark is applied in accordance with requirements provided for in Article 10 of Delegated Regulation (EU) 2020/686 and/or Article 83(a) of Commission Delegated Regulation (EU) 2020/692 and that mark is indicated in Box I.30; II.2.4. is/are transported in a container which: II.2.4.1. was sealed and numbered prior to the dispatch from the germinal product storage centre under responsibility of the centre veterinarian, or by an official veterinarian, and the seal bears the number as indicated in Box I.19; II.2.4.2. has been cleaned and either disinfected or sterilised before use, or is single-use container; (2)(5) □ has been filled in with the cryogenic agent which not have been previously used for other products;]			(2)	\square and/or	[Model 3 in Section C o	f Part 1 of Annex III to Regula	tion (EU) 2018/659(4);]
(2) □ and/or [Model 2 in Section B of Part 2 of Annex II to Decision 2010/471/EU(4);] (2) □ and/or [Model in Annex to Commission Decision 96/539/EC(4);] II.2.2. has/have been collected, processed and stored in accordance with animal health requirements set out in Annex III to Delegated Regulation (EU) 2020/686; II.2.3. is/are placed in straws or other packages on which the mark is applied in accordance with requirements provided for in Article 10 of Delegated Regulation (EU) 2020/686 and/or Article 83(a) of Commission Delegated Regulation (EU) 2020/692 and that mark is indicated in Box I.30; II.2.4. is/are transported in a container which: II.2.4.1. was sealed and numbered prior to the dispatch from the germinal product storage centre under responsibility of the centre veterinarian, or by an official veterinarian, and the seal bears the number as indicated in Box I.19; II.2.4.2. has been cleaned and either disinfected or sterilised before use, or is single-use container; (2)(5) □ II.2.4.3. has been filled in with the cryogenic agent which not have been previously used for other products;]			(2)	\square and/or	[Model 4 in Section D o	of Part 1 of Annex III to Regula	ation (EU) 2018/659(4);]
(2) □ and/or [Model 3 in Section C of Part 2 of Annex II to Decision 2010/471/EU(4);] (2) □ and/or [Model in Annex to Commission Decision 96/539/EC(4);] II.2.2. has/have been collected, processed and stored in accordance with animal health requirements set out in Annex III to Delegated Regulation (EU) 2020/686; II.2.3. is/are placed in straws or other packages on which the mark is applied in accordance with requirements provided for in Article 10 of Delegated Regulation (EU) 2020/686 and/or Article 83(a) of Commission Delegated Regulation (EU) 2020/692 and that mark is indicated in Box I.30; II.2.4. is/are transported in a container which: II.2.4.1. was sealed and numbered prior to the dispatch from the germinal product storage centre under responsibility of the centre veterinarian, or by an official veterinarian, and the seal bears the number as indicated in Box I.19; II.2.4.2. has been cleaned and either disinfected or sterilised before use, or is single-use container; (2)(5) □ has been filled in with the cryogenic agent which not have been previously used for other products;]			(2)	\square and/or	[Model 1 in Section A o	of Part 2 of Annex II to Decisio	n 2010/471/EU(4);]
(2)			(2)	\square and/or	[Model 2 in Section B o	f Part 2 of Annex II to Decisio	n 2010/471/EU(4);]
II.2.2. has/have been collected, processed and stored in accordance with animal health requirements set out in Annex III to Delegated Regulation (EU) 2020/686; II.2.3. is/are placed in straws or other packages on which the mark is applied in accordance with requirements provided for in Article 10 of Delegated Regulation (EU) 2020/686 and/or Article 83(a) of Commission Delegated Regulation (EU) 2020/692 and that mark is indicated in Box I.30; II.2.4. is/are transported in a container which: II.2.4.1. was sealed and numbered prior to the dispatch from the germinal product storage centre under responsibility of the centre veterinarian, or by an official veterinarian, and the seal bears the number as indicated in Box I.19; II.2.4.2. has been cleaned and either disinfected or sterilised before use, or is single-use container; (2)(5) [II.2.4.3. has been filled in with the cryogenic agent which not have been previously used for other products;]			(2)	\square and/or	[Model 3 in Section C o	f Part 2 of Annex II to Decisio	n 2010/471/EU(4);]
requirements set out in Annex III to Delegated Regulation (EU) 2020/686; II.2.3. is/are placed in straws or other packages on which the mark is applied in accordance with requirements provided for in Article 10 of Delegated Regulation (EU) 2020/686 and/or Article 83(a) of Commission Delegated Regulation (EU) 2020/692 and that mark is indicated in Box I.30; II.2.4. is/are transported in a container which: II.2.4.1. was sealed and numbered prior to the dispatch from the germinal product storage centre under responsibility of the centre veterinarian, or by an official veterinarian, and the seal bears the number as indicated in Box I.19; II.2.4.2. has been cleaned and either disinfected or sterilised before use, or is single-use container; (2)(5) Implicate the product of the dispatch from the germinal product storage centre under responsibility of the centre veterinarian, or by an official veterinarian, and the seal bears the number as indicated in Box I.19; II.2.4.2. has been cleaned and either disinfected or sterilised before use, or is single-use container; (2)(5) III.2.4.3. has been filled in with the cryogenic agent which not have been previously used for other products;]			(2)	\square and/or	[Model in Annex to Co	mmission Decision 96/539/EC(4);]
requirements provided for in Article 10 of Delegated Regulation (EU) 2020/686 and/or Article 83(a) of Commission Delegated Regulation (EU) 2020/692 and that mark is indicated in Box I.30; II.2.4. is/are transported in a container which: II.2.4.1. was sealed and numbered prior to the dispatch from the germinal product storage centre under responsibility of the centre veterinarian, or by an official veterinarian, and the seal bears the number as indicated in Box I.19; II.2.4.2. has been cleaned and either disinfected or sterilised before use, or is single-use container; (2)(5)			II.2.2.				
II.2.4.1. was sealed and numbered prior to the dispatch from the germinal product storage centre under responsibility of the centre veterinarian, or by an official veterinarian, and the seal bears the number as indicated in Box I.19; II.2.4.2. has been cleaned and either disinfected or sterilised before use, or is single-use container; (2)(5)			II.2.3.	requireme 83(a) of Co	nts provided for in Artic	cle 10 of Delegated Regulation	(EU) 2020/686 and/or Article
storage centre under responsibility of the centre veterinarian, or by an official veterinarian, and the seal bears the number as indicated in Box I.19; II.2.4.2. has been cleaned and either disinfected or sterilised before use, or is single-use container; (2)(5) [II.2.4.3. has been filled in with the cryogenic agent which not have been previously used for other products;]			II.2.4.	is/are trans	sported in a container w	hich:	
(2)(5) Container; has been filled in with the cryogenic agent which not have been previously used for other products;]				II.2.4.1.	storage centre under r	esponsibility of the centre vet	erinarian, or by an official
(2)(5) — has been filled in with the cryogenic agent which not have been previously used for other products;]				_		either disinfected or sterilised	before use, or is single-use
(2)(6) [II.2.5. is/are placed in straws or other packages which are securely and hermetically sealed;			(2)(5)	_		the cryogenic agent which no	t have been previously used
		(2)(6) \square [II.2.5. is/are placed in straws or other		ed in straws or other pa	ckages which are securely an	d hermetically sealed;	

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	II. Health information			
	II.2.6.	is/are transported in a container w	there they are congreted from	anch other by physical
	11.2.0.	is/are transported in a container w compartments or by being placed	in secondary protective hags	each other by physical
		comparaments of by being placed.	in occorianty protective bags.	
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Part II: Certification				
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II. Health information

Notes

This animal health certificate shall be completed according to the notes for the completion of certificates provided for in Chapter 2 of Annex I to Commission Implementing Regulation (EU) 2020/2235.

Part I:

Certification Box reference I.11:

"Place of dispatch": Indicate the unique approval number and the name and address of the germinal product storage centre of dispatch of the consignment of semen, oocytes, and/or embryos. Only germinal product storage centres approved by the competent authority and included in the register referred to in Article 101(1)(b) of Regulation (EU) 2016/429 and Article 7 of Delegated Regulation (EU) 2020/686.

Box reference I.12:

Part II:

"Place of destination": Indicate the address and unique registration or approval number of the establishment of destination of the consignment of semen, oocytes, and/or embryos.

Box reference

"Accompanying documents": Number(s) of related original certificate(s) shall correspond to the serial number of the individual official document(s) or health certificate(s) that accompanied the semen, oocytes and/or embryos described in Part I from the semen collection centre where the semen was collected, and/or the embryo collection and/or production team by which the oocytes and/or embryos were collected or produced, and/or the germinal product processing establishment where the semen, oocytes or embryos were processed and stored, and/or the germinal product storage centre where the semen, oocytes or embryos were stored to the germinal product storage centre described in Box I.11. The original(s) of those document(s) or those certificate(s) or the officially endorsed copies thereof must be attached to this certificate.

Seal number shall be indicated. Box

reference I.19:

Box reference I.26:

Total number of packages shall correspond to the number of containers.

reference I.30:

"Type": Specify if semen, in vivo derived embryos, in vivo derived oocytes, in vitro produced embryos or micromanipulated embryos.

"Identification number": Indicate identification number of each donor animal.

"Identification mark": indicate mark on the straw or other packages where semen, oocytes and/or embryos of the consignment are placed.

"Date of collection/production": indicate the date on which semen, oocytes and/or embryos of the consignment was/were collected or produced.

"Approval or registration number of plant/establishment/centre": Indicate the unique approval number of the semen collection centre where the semen was collected, and/or of the embryo collection and/or production team by which the oocytes or embryos were collected or produced.

"Quantity": Indicate number of straws or other packages with the same mark.

Part II:

- (1) Only germinal product storage centres approved by the competent authority and included in the register referred to in Article 101(1)(b) of Regulation (EU) 2016/429 and Article 7 of Delegated Regulation (EU) 2020/686.
- Delete if not applicable. (2)
- (3) Only germinal product establishments approved by the competent authority and included in the register referred to in Article 101(1)(b) of Regulation (EU) 2016/429 and Article 7 of Delegated Regulation (EU) 2020/686.

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	II. Health information	
l: Certification	that accompanied the semen, oocytes or embry where the semen was collected, and/or the eml and/or embryos were collected or produced, an where the semen, oocytes or embryos were pro centre where the semen, oocytes or embryos w semen, oocytes and/or embryos dispatch descr. (5) Applicable for frozen semen, oocytes or embry (6) Applicable for the consignment where in one of	n certificate(s) or the officially endorsed copies of thereof yos described in Part I from the semen collection centre bryo collection or production team by which the oocytes ind/or the germinal product processing establishment ocessed and stored, and/or the germinal product storage were stored to the germinal product storage centre of the libed in Box I.11 must be attached to this certificate.
Part II:		O all'Eurita della
Pal	Name (in capital letters)	Qualification and title
_	Date of signature Stamp	Signature

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