EUROPEAN UNION INTRA

	I.1. Consignor					I.2. IMSOC ref	erence	I.2.a. Local re	ference	
	_							I 3 Central Co	ompetent Authority	
	Name									
	Address							1.4. Local Con	npetent Authority	
	Country ISO Code									
_	I.5. Consignee					I.6. Operator conducting assembly operations independently of an				
Part I: Description of consignment	Name					establishment				
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en 1/3

EUROPEAN UNION

	II. Health info	rmation					
		I, the unde	rsigned offi	cial veterinarian, hereby	v certify that:		
	(1)	o either	[II.1.	the in vivo derived emb	oryos(1)/in vivo derived ova(1 d stored by an embryo collec ace with Chapter I(III)(1) of An	tion team(2) approved and	
art in cormication	(1)	o or	[II.1.	Part I were produced, p	mbryos(1)/micromanipulated processed and stored by an er ed in accordance with Chapto 	nbryo production team(2)	
3	(1)	o either	[II.2.	the in vivo derived embryos described in Part I meet the requirements of Chapter III(II)(1) of Annex D to Directive 92/65/EEC;]			
10.1	(1)	o or	[II.2.	the in vivo derived ova III(II)(2) of Annex D to 2	described in Part I meet the Directive 92/65/EEC;]	requirements of Chapter	
	(1)	o or	[II.2.	_	mbryos described in Part I m nex D to Directive 92/65/EEC;]	-	
(1) or [II.2. the micromanipulated embryos described in Chapter III(II)(4) of Annex D to Directive 92/6							
			□ [II.3.		ets of embryos of the ovine or ing conditions as regards clas		
		(1)	∘ either	birth on a holding or h	om animals which have been oldings recognised as having in accordance with point 1 o on (EC) No 999/2001;]	a negligible or a controlled	
		(1)	o or	last three years before complied for the last th	om animals which have been the collection on a holding or ree years before collection w of point 1.3. of Section A of 0 /2001;]	holdings which have ith the requirements laid	
		(1)	o or	birth in a Member State for classical scrapie ap	om animals which have been e or zone of a Member State v proved in accordance with th pter A of Annex VIII to Regula	vith a negligible risk status e first subparagraph of poin	
		(1)	o or	[they were collected fro	om ovine animals and		
			(1)	o either [are of the A	ARR/ARR prion protein genot	ype;]	
			(1)	o or [carry at lease January 20]	ast one ARR allele and were c 15;]]	ollected after the date of 1	
			II.4.	_	scribed in Part I come from fe s(1) which meet the requiren 2/65/EEC;		
	(1)	o either	[II.5.	insemination of the dos stored and transported	in Part I were conceived as a nor females with semen whic under conditions which com II(I) of Annex D to Directive 9	h was collected, processed, ply with the requirements o	
	(1)	o or	[II.5.	fertilisation of ova com to Directive 92/65/EEC transported under cond	in Part I were conceived as a plying with the conditions in with semen which was collect ditions which comply with the nex D to Directive 92/65/EEC;	Chapter III(II)(2) of Annex I ted, processed, stored and e requirements of Chapters	
	(1)	\circ or	[II.5.	the ova have not been i	n contact with semen of the	ovine and caprine species;]	
			II.6.	sealed container in acc	scribed in Part I were sent to ordance with point 6 of Chap d bearing the number detaile	ter III(II) of Annex D to	

en 2/3

EUROPEAN UNION

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:

Box I.11:

Place of dispatch shall correspond to the embryo collection team or embryo production team of embryos collection/production.

Part II: Certification Box I.12:

Place of destination shall correspond to the embryo collection team, embryo production team, germinal product processing establishment, germinal product storage centre or to the establishment of ova/embryos destination.

Box I.19:

Identification of container and Seal number shall be indicated.

Box I.30:

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

Identification number shall correspond to the official identification of the animal.

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 or embryo production team of ova/embryos collection/production.

Part II:

(1) Delete as appropriate.

Only embryo collection or production teams approved by the competent authority and listed in (2) accordance with Article 11(4) of Directive 92/65/EEC

Certifying Officer/Official veterinarian

Name (in capital letters)

Date of signature

Stamp

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Qualification and title

Signature

3 / 3