

Part I: Description of consignment	I.1. Consignor		I.2. IMSOC reference		I.2.a. Local reference	
	Name				I.3. Central Competent Authority	
	Address				I.4. Local Competent Authority	
	Country		ISO Code			
	I.5. Consignee			I.6. Operator conducting assembly operations independently of an establishment		
	Name			Name		
	Address			Address		
	Country			Country		
	ISO Code			Approval Number		
				ISO Code		
I.7. Country of origin			I.9. Country of destination		I.8. Region of origin	
ISO Code					Code	
					I.10. Region of destination	
					Code	
I.11. Place of dispatch			I.12. Place of destination			
Name			Name			
Address			Address			
Approval Number			Approval Number			
Country			Country			
ISO Code			ISO Code			
I.13. Place of loading			I.14. Date and time of departure			
Name						
Address						
Approval Number						
Country						
ISO Code						
I.15. Means of Transport			I.16. Transporter			
Mode	International transport document	Identification	Name			
			Address			
			Activity ID			
			Country			
			ISO Code			
			I.17. Accompanying documents			
			Commercial document reference			
			Date of issue			
			Country			
			Place of issue			
I.18. Transport conditions						
Frozen <input type="checkbox"/>		Chilled <input type="checkbox"/>		Ambient <input type="checkbox"/>		
I.19. Container No / Seal No						
I.20. Certified as Germinal products <input type="checkbox"/>						
I.21. For transit through a third country <input type="checkbox"/>						
Third country		ISO Code				
Exit point		BCP code				
Entry point		BCP code				
I.22. For transit through Member State(s) <input type="checkbox"/>			I.23. For export <input type="checkbox"/>			
Member State			Third country			
ISO Code			ISO Code			
			Exit point			
			BCP code			
			I.25. Journey Log			
I.26. Total number of packages		I.27. Total quantity		I.28. Total gross weight		
I.30. Description of consignment						
Commodity	Species	Identification Number	Quantity	Nature of commodity		
Identification Mark	Package count	Date of collection / production	Plant / Establishment / Centre	Type		

Part II: Certification	II. Health information			
	I, the undersigned official veterinarian, hereby certify that:			
	(1)	○ either [II.1.	the in vivo derived embryos(1)/in vivo derived ova(1) described in Part I were collected, processed and stored by an embryo collection team(2) approved and supervised in accordance with Chapter I(III)(1) of Annex D to Directive 92/65/EEC;]	
	(1)	○ or [II.1.	the in vitro produced embryos(1)/micromanipulated embryos(1) described in Part I were produced, processed and stored by an embryo production team(2) approved and supervised in accordance with Chapter I(III)(1) and (2) of Annex D to Directive 92/65/EEC;]	
	(1)	○ 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;]	
	(1)	○ or [II.2.	the in vivo derived ova described in Part I meet the requirements of Chapter III(II)(2) of Annex D to Directive 92/65/EEC;]	
	(1)	○ or [II.2.	the in vitro produced embryos described in Part I meet the requirements of Chapter III(II)(3) of Annex D to Directive 92/65/EEC;]	
	(1)	○ or [II.2.	the micromanipulated embryos described in Part I meet the requirements of Chapter III(II)(4) of Annex D to Directive 92/65/EEC;]	
	(1)	□ [II.3.	the consignment consists of embryos of the ovine or caprine species which comply with the following conditions as regards classical scrapie:	
	(1)	○ either	[they were collected from animals which have been kept continuously since birth on a holding or holdings recognised as having a negligible or a controlled risk of classical scrapie in accordance with point 1 of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001;]	
	(1)	○ or	[they were collected from animals which have been kept continuously for the last three years before the collection on a holding or holdings which have complied for the last three years before collection with the requirements laid down in points (a) to (f) of point 1.3. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001;]	
	(1)	○ or	[they were collected from animals which have been kept continuously since birth in a Member State or zone of a Member State with a negligible risk status for classical scrapie approved in accordance with the first subparagraph of point 2.2. of Section A of Chapter A of Annex VIII to Regulation (EC) No 999/2001;]	
	(1)	○ or	[they were collected from ovine animals and	
	(1)	○ either	[are of the ARR/ARR prion protein genotype;]	
	(1)	○ or	[carry at least one ARR allele and were collected after the date of 1 January 2015;]	
(1)	○ either [II.5.	the embryos described in Part I were conceived as a result of artificial insemination of the donor females with semen which was collected, processed, stored and transported under conditions which comply with the requirements of Chapters I(I), II(I) and III(I) of Annex D to Directive 92/65/EEC;]		
(1)	○ or [II.5.	the embryos described in Part I were conceived as a result of in vitro fertilisation of ova complying with the conditions in Chapter III(II)(2) of Annex D to Directive 92/65/EEC with semen which was collected, processed, stored and transported under conditions which comply with the requirements of Chapters I(I), II(I) and III(I) of Annex D to Directive 92/65/EEC;]		
(1)	○ or [II.5.	the ova have not been in contact with semen of the ovine and caprine species;]		
	II.6.	the ova or embryos described in Part I were sent to the place of loading in a sealed container in accordance with point 6 of Chapter III(II) of Annex D to Directive 92/65/EEC and bearing the number detailed in Box I.19.		

II. Health information							
Part II: Certification	<p>Notes</p> <p>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.</p> <p>Part I:</p> <p>Box I.11: Place of dispatch shall correspond to the embryo collection team or embryo production team of embryos collection/production.</p> <p>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.</p> <p>Box I.19: Identification of container and Seal number shall be indicated.</p> <p>Box I.30: "Type": specify if: in vivo derived embryos, in vivo derived oocytes, in vitro produced embryos or micromanipulated embryos.</p> <p>Identification number shall correspond to the official identification of the animal.</p> <p>Date of collection shall be indicated in the following format: dd/mm/yyyy.</p> <p>Approval number of the team shall correspond to the embryo collection team or embryo production team of ova/embryos collection/production.</p> <p>Part II:</p> <p>(1) Delete as appropriate.</p> <p>(2) Only embryo collection or production teams approved by the competent authority and listed in accordance with Article 11(4) of Directive 92/65/EEC.</p>						
	<p>Certifying Officer/Official veterinarian</p> <table border="0"> <tr> <td data-bbox="807 952 807 1093">Name (in capital letters)</td> <td data-bbox="807 952 1487 1093">Authority name</td> </tr> <tr> <td>Date of signature</td> <td>Signature</td> </tr> <tr> <td>Stamp</td> <td></td> </tr> </table>		Name (in capital letters)	Authority name	Date of signature	Signature	Stamp
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Date of signature	Signature						
Stamp							