

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			Approval Number		
				Country		
				ISO Code		
I.7. Country of origin			ISO Code		I.9. Country of destination	
					ISO Code	
I.8. Region of origin			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			
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								
	<p>I, the undersigned official veterinarian, hereby certify that the semen described in Part I:</p> <p>II.1. was collected, processed and stored in a semen collection centre(1) approved and supervised by the competent authority in accordance with Chapter I(I) and Chapter I(II) of Annex D to Directive 92/65/EEC;</p> <p>II.2. comes from the donor animals which meet the requirements of Chapter II(II) of Annex D to Directive 92/65/EEC;</p> <p>II.3. was collected, processed, stored and transported under conditions which comply with the requirements of Chapters II(II) and III of Annex D to Directive 92/65/EEC;</p> <p>(2) <input type="radio"/> either</p> <p>[II.4. meets the requirements of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001, as applicable on 31 August 2010;]</p> <p>(2) <input type="radio"/> or</p> <p>[II.4. meets the requirements of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001, as applicable on 31 August 2010, and is destined for a Member State which benefits, for all or part of its territory, from the provisions laid down in point (b) or (c) of Chapter A(I) of Annex VIII to Regulation (EC) No 999/2001, as applicable on 31 August 2010, and the donor animals comply regarding scrapie with the guarantees provided for by the programmes referred to in that point and with the guarantees(3) requested by the Member State of destination.]</p>								
<p>Notes:</p> <p>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 European Union in this certificate include the United Kingdom in respect of Northern Ireland.</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 semen collection centre of origin of the semen.</p> <p>Box I.12: Place of destination shall correspond to the semen collection centre, germinal product processing establishment, germinal product storage centre or to the establishment of semen destination.</p> <p>Box I.19: Identification of container and Seal number shall be indicated.</p> <p>Box I.30: 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 centre shall correspond to the approval number of the semen centre indicated in Box I.11. where the semen was collected.</p> <p>Part II:</p> <p>(1) Only semen collection centres approved by the competent authority and listed in accordance with Article 11(4) of Council Directive 92/65/EEC.</p> <p>(2) Delete as appropriate.</p> <p>(3) Additional guarantees as laid down in Article 2 of Regulation (EC) No 546/2006 [OJ L 94, 1.4.2006, p.28].</p>									
<p>Certifying Officer/Official veterinarian</p> <table border="0" style="width: 100%;"> <tr> <td style="width: 50%;">Name (in capital letters)</td> <td style="width: 50%;">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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