

<b>Part I: Description of consignment</b>	I.1. Consignor Name Address Country		ISO Code	I.2. IMSOC reference	I.2.a. Local reference I.3. Central Competent Authority I.4. Local Competent Authority	
	I.5. Consignee Name Address Country		ISO Code	I.6. Operator conducting assembly operations independently of an establishment Name Address Approval Number Country		
	I.7. Country of origin		ISO Code	I.9. Country of destination		
	I.8. Region of origin		Code	I.10. Region of destination		
	I.11. Place of dispatch Name Address Approval Number Country		ISO Code	I.12. Place of destination Name Address Approval Number Country		
	I.13. Place of loading Name Address Approval Number Country		ISO Code	I.14. Date and time of departure		
	I.15. Means of Transport		I.16. Transporter		I.17. Accompanying documents	
	Mode	International transport document	Identification		Name Address Approval Number Country	ISO Code
					[en] accompanying document number	Date of issue  Place of issue
	I.18. Transport conditions Chilled <input type="checkbox"/>		Frozen <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		ISO Code		Third country		
				ISO Code		
				Exit point		
				BCP code		
I.24. Estimated journey time			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			

Part II: Certification	II. Health information								
	<p>I, the undersigned official veterinarian, hereby certify that:</p> <p>II.1. The ova/embryos(1) described in Part I:</p> <p style="margin-left: 40px;">II.1.1. were collected, processed and stored under conditions which meet the requirements of Directive 92/65/EEC;</p> <p style="margin-left: 40px;">II.1.2. come from donor female swine which meet the requirements of Chapter IV of Annex D to Directive 92/65/EEC;</p> <p style="margin-left: 40px;">II.1.3. meet the requirements of Chapter III of Annex D to Directive 92/65/EEC.</p> <p><input type="checkbox"/> either [II.2. In the case of embryos,</p> <p style="margin-left: 40px;">II.2.1. the semen used for fertilisation meets the requirements of Directive 90/429/EEC;</p> <p style="margin-left: 40px;">II.2.2. the embryos have been washed with trypsin(2).]</p> <p>(1) <input type="checkbox"/> or [II.2. In the case of ova, the ova comes from a donor female swine which meets the conditions of Article 1 of Decision 2008/185/EC(2).]</p>								
<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 destination shall correspond to the embryo collection team of oocytes/embryos collection.</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 oocytes/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 style="margin-left: 40px;">Identification number shall correspond to the official identification of the animal.</p> <p style="margin-left: 40px;">Date of collection shall be indicated in the following format: dd/mm/yyyy.</p> <p style="margin-left: 40px;">Approval number of the team shall correspond to the embryo collection team of oocytes/embryos collection indicated in Box I.11.</p> <p>Part II:</p> <p>(1) Delete as appropriate.</p> <p>(2) This condition applies only to ova and embryos which originate in the Member States or regions thereof not listed in Annexes I and II to Decision 2008/185/EC (OJ L 59, 4.3.2008, p. 19) and destined to the Member States or regions thereof so listed. It shall also apply to movements from Member States or regions thereof listed in Annex II of Decision 2008/185/EC to Member States or regions thereof listed in Annex I of Decision 2008/185/EC.</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%;">Qualification and title</td> </tr> <tr> <td>Date of signature</td> <td>Signature</td> </tr> <tr> <td>Stamp</td> <td></td> </tr> </table>				Name (in capital letters)	Qualification and title	Date of signature	Signature	Stamp	
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