

<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 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 Name Address Approval Number Country ISO Code		I.12. Place of destination Name Address Approval Number Country ISO Code			
	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					
	I.18. Transport conditions Ambient <input type="checkbox"/> Chilled <input type="checkbox"/> Frozen <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"/>						
I.22. For transit through Member State(s) <input type="checkbox"/>		I.23. For export <input type="checkbox"/>				
I.24. 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:</p> <p>II.1. The <input type="checkbox"/> [semen](1)/ <input type="checkbox"/> [oocytes](1)/ <input type="checkbox"/> [embryos](1) <input type="checkbox"/> [of dogs](1)/ <input type="checkbox"/> [cats](1) described in Part I are intended for artificial reproduction and were obtained from donor animals which</p> <p>II.1.1. have been born and remained since birth in the Union, or have entered the Union in accordance with the requirements for entry into the Union;</p> <p>II.1.2. are</p> <p>(1) <input type="checkbox"/> either <input type="checkbox"/> [ marked by the implantation of a transponder in accordance with Article 17(1) of Regulation (EU) No 576/2013;]</p> <p>(1) <input type="checkbox"/> or <input type="checkbox"/> [ marked by a clearly readable tattoo in accordance with Article 17(1) of Regulation (EU) No 576/2013;]</p> <p>(1) <input type="checkbox"/> or <input type="checkbox"/> [ identified in accordance with Article 70 of Commission Delegated Regulation (EU) 2019/2035;]</p> <p>II.1.3. have received an anti-rabies vaccination that complies with the validity requirements set out in Part 1 of Annex VII to Commission Delegated Regulation (EU) 2020/688.</p> <p>II.2. The <input type="checkbox"/> [semen](1)/ <input type="checkbox"/> [oocytes](1)/ <input type="checkbox"/> [embryos](1) described in Part I comes/come from a registered establishment assigned by the competent authority with a unique registration number as indicated in Box I.11.</p> <p>II.3. According to official information, the <input type="checkbox"/> [semen](1)/ <input type="checkbox"/> [oocytes](1)/ <input type="checkbox"/> [embryos](1) described in Part I was/were obtained from donor animals which</p> <p>II.3.1. come from establishments in which infection with rabies virus has not been confirmed for a period of at least 30 days prior to the date of collection of the <input type="checkbox"/> [semen](1)/ <input type="checkbox"/> [oocytes](1)/ <input type="checkbox"/> [embryos](1);</p> <p>II.3.2. comply with any preventive health measure for diseases or infections other than rabies set out in Part 2 of Annex VII to Delegated Regulation (EU) 2020/688.</p> <p>II.4. To the best of my knowledge and as declared by the operator, the <input type="checkbox"/> [semen](1)/ <input type="checkbox"/> [oocytes](1)/ <input type="checkbox"/> [embryos](1) was/were obtained from donor animals which</p> <p>II.4.1. showed no disease symptoms on the day of collection of the <input type="checkbox"/> [semen](1)/ <input type="checkbox"/> [oocytes](1)/ <input type="checkbox"/> [embryos](1);</p> <p>II.4.2. were not used for natural breeding during a period of at least 30 days prior to the date of collection of the <input type="checkbox"/> [semen](1)/ <input type="checkbox"/> [oocytes](1)/ <input type="checkbox"/> [embryos](1) and during the collection period.</p> <p>II.5. The <input type="checkbox"/> [semen](1)/ <input type="checkbox"/> [oocytes](1)/ <input type="checkbox"/> [embryos](1) described in Part I is/are placed in a sealed transport container and the seal bears the number as indicated in Box I.19.</p> <p>II.6. To the best of my knowledge and based on the documentary check of the data submitted by the operator, the <input type="checkbox"/> [semen](1)/ <input type="checkbox"/> [oocytes](1)/ <input type="checkbox"/> [embryos](1) described in Part I is/are placed in straws or other packages on which the mark is applied in accordance with requirements provided for in Article 11 of Commission Delegated Regulation (EU) 2020/686 and that mark is indicated in Box I.30.</p>		

Part II: Certification	II. Health information								
	<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 reference I.11: “Place of dispatch”: indicate the address and the unique registration number of the establishment of dispatch of the consignment of semen, oocytes or embryos.</p> <p>Box reference I.12: “Place of destination”: Indicate the address and the unique registration number, if assigned by the competent authority, of the establishment of destination of the consignment of semen, oocytes or embryos.</p> <p>Box reference I.30: “Type”: Specify if semen, in vivo derived embryos, in vivo derived oocytes, in vitro produced embryos or micromanipulated embryos.</p> <p>“Species”: Indicate “Canis lupus familiaris” or “Felis silvestris catus” as appropriate.</p> <p>“Identification number”: Indicate individual identification number of each donor animal.</p> <p>“Identification mark”: Indicate mark on the straw or other packages where semen, oocytes or embryos of the consignment are placed.</p> <p>“Date of collection/production”: Indicate the date on which semen, oocytes or embryos of the consignment were collected or produced.</p> <p>“Approval or registration number of plant/establishment/centre”: Indicate the unique registration number of the establishment of the collection or production of semen, oocytes or embryos of the consignment.</p> <p>“Quantity”: Indicate number of straws or other packages with the same mark.</p> <p>Part II:</p> <p>(1) <b>Delete if not applicable.</b></p>								
<p>Certifying Officer/Official veterinarian</p> <table style="width: 100%; border: none;"> <tr> <td style="width: 50%; border: none;">Name (in capital letters)</td> <td style="width: 50%; border: none;">Authority name</td> </tr> <tr> <td style="border: none;">Date of signature</td> <td style="border: none;">Signature</td> </tr> <tr> <td style="border: none;">Stamp</td> <td style="border: none;"></td> </tr> </table>				Name (in capital letters)	Authority name	Date of signature	Signature	Stamp	
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