

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		ISO Code		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			ISO Code		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
						Approval Number
						Country
						ISO Code
I.17. Accompanying documents			I.17. Accompanying documents			
			[en] accompanying document number			
					Date of issue	
					Place of issue	
			Country			
I.18. Transport conditions						
Ambient <input type="checkbox"/>		Frozen <input type="checkbox"/>			Chilled <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 germinal product processing establishment(1) described in Box I.11. at which the semen(2)/ oocytes(2)/ in vivo derived embryos(2)/ in vitro produced embryos(2)/ micromanipulated embryos(2) was/were processed and stored:</p> <p>II.1.1. is approved and kept in a register by the competent authority;</p> <p>II.1.2. complies with requirements as regards responsibilities, operational procedures, facilities and equipment set out in Part 4 of Annex I to Delegated Regulation (EU) 2020/686.]</p> <p>II.2. The semen(2)/ oocytes(2)/ in vivo derived embryos(2)/ in vitro produced embryos(2)/ micromanipulated embryos(2) described in Part I is/are intended for artificial reproduction and</p> <p>(2) <input type="checkbox"/> either</p> <p>[II.2.1. has/have been collected or produced, processed and stored in a semen collection centre(2)(3)/ by an embryo collection team(2)(3)/ by an embryo production team(2)(3), and/or processed and stored in a germinal product processing establishment(2)(3), and/or stored in a germinal product storage centre(2)(3) situated in the Member State of its/their collection or production and complying with requirements as regards responsibilities, operational procedures, facilities and equipment set out in Part 1(2)/ Part 2(2)/ Part 3(2)/ Part 4(2)/ Part 5(2) of Annex I to Delegated Regulation (EU) 2020/686, and was/were moved to the germinal product processing establishment indicated in Box I.11. situated in the Member State of its/their collection or production under animal health certification requirements at least as strict as those provided for in:</p> <p>(2) <input type="checkbox"/> either [Model BOV-SEM-A-INTRA(4);]</p> <p>(2) <input type="checkbox"/> and/or [Model BOV-SEM-B-INTRA(4);]</p> <p>(2) <input type="checkbox"/> and/or [Model BOV-SEM-C-INTRA(4);]</p> <p>(2) <input type="checkbox"/> and/or [Model BOV-OOCTYES-EMB-A-INTRA(4);]</p> <p>(2) <input type="checkbox"/> and/or [Model BOV-EMB-B-INTRA(4);]</p> <p>(2) <input type="checkbox"/> and/or [Model BOV-GP-PROCESSING-INTRA(4);]</p> <p>(2) <input type="checkbox"/> and/or [Model BOV-GP-STORAGE-INTRA(4);]</p> <p>(2) <input type="checkbox"/> and/or</p> <p>[II.2.1. has/have been collected or produced, processed and stored in a semen collection centre(2)(3)/ by an embryo collection team(2)(3)/ by an embryo production team(2)(3), and/or processed and stored in a germinal product processing establishment(2)(3), and/or stored in a germinal product storage centre(2)(3) situated in the Member State of its/their collection or production and complying with requirements as regards responsibilities, operational procedures, facilities and equipment set out in Part 1(2)/ Part 2(2)/ Part 3(2)/ Part 4(2)/ Part 5(2) of Annex I to Delegated Regulation (EU) 2020/686, and was/were moved to the germinal product processing establishment indicated in Box I.11. situated in another Member State accompanied by certificate(s) in accordance with:</p> <p>(2) <input type="checkbox"/> either [Model BOV-SEM-A-INTRA(4);]</p> <p>(2) <input type="checkbox"/> and/or [Model BOV-SEM-B-INTRA(4);]</p> <p>(2) <input type="checkbox"/> and/or [Model BOV-SEM-C-INTRA(4);]</p> <p>(2) <input type="checkbox"/> and/or [Model BOV-OOCTYES-EMB-A-INTRA(4);]</p> <p>(2) <input type="checkbox"/> and/or [Model BOV-EMB-B-INTRA(4);]</p> <p>(2) <input type="checkbox"/> and/or [Model BOV-GP-PROCESSING-INTRA(4);]</p> <p>(2) <input type="checkbox"/> and/or [Model BOV-GP-STORAGE-INTRA(4);]</p> <p>(2) <input type="checkbox"/> and/or</p> <p>[II.2.1. has/have been collected or produced, processed and stored in a semen collection centre(2)(3)/ by an embryo collection team(2)(3)/ by an embryo production team(2)(3) , and/or processed and stored in a germinal product processing establishment(2)(3), and/or stored in a germinal product storage centre(2)(3) situated in a third country, territory or zone thereof listed in Annex IX to Commission Implementing Regulation (EU) 2021/404 and complying with requirements as regards responsibilities, operational procedures, facilities and equipment set out in Part 1(2)/ Part 2(2)/ Part 3(2)/ Part 4(2)/ Part 5(2) of Annex I to Delegated Regulation (EU) 2020/686, and entered the Union accompanied by certificate(s) in accordance with:</p> <p>(2) <input type="checkbox"/> either [Model BOV-SEM-A-ENTRY(4);]</p>		

Part II: Certification	II. Health information		
	(2)	<input type="checkbox"/> and/or	[Model BOV-SEM-B-ENTRY (4);]
	(2)	<input type="checkbox"/> and/or	[Model BOV-SEM-C-ENTRY (4);]
	(2)	<input type="checkbox"/> and/or	[Model BOV-OOCYTES-EMB-A-ENTRY (4);]
	(2)	<input type="checkbox"/> and/or	[Model BOV-in-vivo-EMB-B-ENTRY(4);]
	(2)	<input type="checkbox"/> and/or	[Model BOV-in-vitro-EMB-C-ENTRY(4);]
	(2)	<input type="checkbox"/> and/or	[Model BOV-in-vitro-EMB-D-ENTRY(4);]
	(2)	<input type="checkbox"/> and/or	[Model BOV-GP-PROCESSING-ENTRY(4);]
	(2)	<input type="checkbox"/> and/or	[Model BOV-GP-STORAGE-ENTRY(4);]
		II.2.2.	has/have been collected, processed and stored in accordance with animal health requirements set out in Annex III to Delegated Regulation (EU) 2020/686;
	II.2.3.	is/are placed in straws or other packages on which the mark is applied in accordance with requirements provided for in Article 10 of Delegated Regulation (EU) 2020/686 and/or Article 83(a) of Delegated Regulation (EU) 2020/692 and that mark is indicated in Box I.30;	
	II.2.4.	is/are transported in a container which:	
	II.2.4.1.	was sealed and numbered prior to the dispatch from the germinal product processing establishment under responsibility of the centre veterinarian, or by an official veterinarian, and the seal bears the number as indicated in Box I.19;	
	II.2.4.2.	has been cleaned and either disinfected or sterilised before use, or is single-use container;	
(2)(5)	[II.2.4.3.	has been filled in with the cryogenic agent which not have been previously used for other products;]	
(2)(6)	<input type="checkbox"/> [II.2.5.	is/are placed in straws or other packages which are securely and hermetically sealed;	
	II.2.6.	is/are transported in a container where they are separated from each other by physical compartments or by being placed in secondary protective bags.]	

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 unique approval number and the name and address of the germinal product processing establishment of dispatch of the consignment of semen, oocytes, and/or embryos. Only germinal product processing establishments approved by the competent authority and included in the register referred to in Article 101(1)(b) of Regulation (EU) 2016/429 and Article 7 of Delegated Regulation (EU) 2020/686.</p> <p>Box reference I.12: "Place of destination": Indicate the address and unique registration or approval number of the establishment of destination of the consignment of semen, oocytes, and/or embryos.</p> <p>Box reference I.17: "Accompanying documents": Number(s) of related original certificate(s) shall correspond to the serial number of the individual official document(s) or health certificate(s) that accompanied the semen, oocytes and/or embryos described in Part I from the semen collection centre where the semen was collected, and/or the embryo collection and/or production team by which the oocytes and/or embryos were collected or produced, and/or the germinal product processing establishment where the semen, oocytes or embryos were processed and stored, and/or the germinal product storage centre where the semen, oocytes or embryos were stored to the germinal product processing establishment described in Box I.11. The original(s) of those document(s) or those certificate(s) or the officially endorsed copies thereof must be attached to this certificate.</p> <p>Box reference I.19: Seal number shall be indicated.</p> <p>Box reference I.26: Total number of packages shall correspond to the number of containers.</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": Select amongst "Bos taurus", "Bison bison" or "Bubalus bubalis" as appropriate.</p> <p>"Identification number": Indicate identification number of each donor animal.</p> <p>"Identification mark": Indicate mark on the straw or other packages where semen, oocytes and/or embryos of the consignment are placed.</p> <p>"Date of collection/production": Indicate the date on which semen, oocytes and/or embryos of the consignment was/were collected or produced.</p> <p>"Approval or registration number of plant/establishment/centre": Indicate the unique approval number of the semen collection centre where the semen was collected, and/or of the embryo collection and/or production team by which the oocytes or embryos were collected or produced.</p> <p>"Quantity": Indicate number of straws or other packages with the same mark.</p>		

II. Health information		
Part II: Certification	Part II:	
	(1) Only germinal product processing establishments approved by the competent authority and included in the register referred to in Article 101(1)(b) of Regulation (EU) 2016/429 and Article 7 of Delegated Regulation (EU) 2020/686.	
	(2) Delete if not applicable.	
	(3) Only germinal product establishments approved by the competent authority and included in the register referred to in Article 101(1)(b) of Regulation (EU) 2016/429 and Article 7 of Delegated Regulation (EU) 2020/686.	
	(4) The original(s) of the document(s) or the health certificate(s) or the officially endorsed copies of thereof that accompanied the semen, oocytes or embryos described in Part I from the semen collection centre where the semen was collected, and/or the embryo collection or production team by which the oocytes and/or embryos were collected or produced, and/or the germinal product processing establishment where the semen, oocytes or embryos were processed and stored, and/or the germinal product storage centre where the semen, oocytes or embryos were stored to the germinal product processing establishment of the semen, oocytes and/or embryos dispatch described in Box I.11 must be attached to this certificate.	
	(5) Applicable for frozen semen, oocytes or embryos. (6) Applicable for the consignment where in one container semen, oocytes, in vivo derived embryos, in vitro produced embryos and micromanipulated embryos of bovine animals are placed and transported.	
Certifying Officer/Official veterinarian		
Name (in capital letters)	Qualification and title	
Date of signature	Signature	
Stamp		