

Part I: Description of consignment	I.1. Consignor		I.2. IMSOC reference		I.2.a. Local reference	
	Name				I.3. Central Competent Authority	
	Address					
	Country		ISO Code		I.4. Local Competent Authority	
	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			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			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			
			Document Type			
			Accompanying document reference			
			Date of Issue			
			Country			
			Place of issue			
I.18. Transport conditions						
Chilled <input type="checkbox"/>		Ambient <input type="checkbox"/>		Frozen <input type="checkbox"/>		
I.19. Container No / Seal No						
I.20. Certified as						
Travelling circus/animal act <input type="checkbox"/>		Registered equidae <input type="checkbox"/>		Relaying <input type="checkbox"/>		
Technical use <input type="checkbox"/>		Live aquatic animals for human consumption <input type="checkbox"/>		Dispatch centre <input type="checkbox"/>		
Exhibition <input type="checkbox"/>		Release into the wild <input type="checkbox"/>		Germinal products <input type="checkbox"/>		
Ornamental aquaculture establishment <input type="checkbox"/>		Other <input type="checkbox"/>		Further processing <input type="checkbox"/>		
Confined establishment <input type="checkbox"/>		Event or activity near borders <input type="checkbox"/>		Quarantine or similar establishment <input type="checkbox"/>		
Further keeping <input type="checkbox"/>		Products for human consumption <input type="checkbox"/>		Slaughter <input type="checkbox"/>		
				Organic fertilizers and soil improvers <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		
				Exit point		
				ISO Code		
				BCP code		
I.24. Estimated journey time			I.25. Journey Log			

Part I: Description of consignment	I.27. Total quantity		I.28. Total gross weight	
	I.30. Description of consignment			
	1. 01 LIVE ANIMALS 0106 Other live animals Mammals: 010619 Other 01061900 Other			
	#1.	Commodity	Sex	Identification system
	Species	Quantity	Age	

Part II: Certification	II. Health information		
	<p>II. Health information</p> <p>I, the undersigned official veterinarian, hereby certify that:</p> <p>II.1. The animals (1) of the consignment described in Part I are wild terrestrial animals and meet the following requirements:</p> <p>II.1.1. The majority of the animals of the consignment, for at least the 30 days prior to the date of departure of the consignment, or since birth, if they are younger than 30 days of age,</p> <p>II.1.1.1. have been resident in the habitat of origin;</p> <p>II.1.1.2. have not been in contact with kept animals of a lower health status or subject to movement restrictions for animal health reasons;</p> <p>II.1.1.3. have not been in direct or indirect contact with kept animals that have entered the Union from a third country or territory, or zone thereof during the last 30 days prior to the date of departure of the consignment.</p> <p>II.1.2. They have not shown clinical signs or symptoms of listed diseases for animals of the species concerned or emerging diseases during the clinical examination, or where this is not possible, the clinical inspection, which was carried out, within the last 24 hours prior to the time of departure of the consignment, on (insert date dd/mm/yyyy).</p> <p>II.2. According to official information, the animals described in Part I:</p> <p>(2) <input type="radio"/> either [come from a habitat or a zone not subject to movement restrictions affecting the species of animals to be moved and established for reasons of diseases listed for those species or diseases subject to emergency measures relevant for those species, and they have not been in contact with kept animals of a listed species of a lower health status for an adequate period.]</p> <p>(2) <input type="radio"/> or [come from a habitat or a zone subject to movement restrictions affecting the species of animals to be moved and established for (3), but derogations from movement restrictions have been granted, and:</p> <p>(2) <input type="checkbox"/> [they comply with requirements set out in (4);]</p> <p>(2) <input type="checkbox"/> [and in particular, they are (5).]</p> <p>(2) <input type="checkbox"/> [II.3. According to official information, the animals described in Part I are ungulates and meet the following health requirements:</p> <p>(2) <input type="checkbox"/> [II.3.1. They come from a habitat in which infection with <i>Brucella abortus</i>, <i>B. melitensis</i> and <i>B. suis</i> in wild terrestrial animals of listed species for that disease has not been reported during the last 42 days prior to the date of departure of the consignment.]</p> <p>(2) <input type="checkbox"/> [II.3.2. They come from a habitat in which infection with <i>Mycobacterium tuberculosis</i> complex (<i>M. bovis</i>, <i>M. caprae</i> and <i>M. tuberculosis</i>) in wild terrestrial animals of listed species for that disease has not been reported during the last 42 days prior to the date of departure of the consignment.]</p> <p>(2) <input type="checkbox"/> [II.3.3. They come from a habitat in which infection with rabies virus has not been reported during the last 30 days prior to the date of departure of the consignment.]</p> <p>(2) <input type="checkbox"/> [II.3.4. They come from a habitat in which infection with epizootic haemorrhagic disease virus within a radius of 150 km has not been reported in wild terrestrial animals of listed species of that disease during the last 2 years prior to the date of departure of the consignment.]</p> <p>(2) <input type="checkbox"/> [II.3.5. They come from a habitat in which anthrax in ungulates has not been reported during the last 15 days prior to the date of departure of the consignment.]</p> <p>(2) <input type="checkbox"/> [II.3.6. They come from a habitat in which surra (<i>Trypanosoma evansi</i>) has not been reported during the last 30 days prior to the date of departure of the consignment.]]</p> <p>(2) <input type="checkbox"/> [II.4. According to official information the animals described in Part I belong to the families Antilocapridae, Bovidae, Camelidae, Cervidae, Giraffidae, Moschidae or Tragulidae and meet the following health requirements:</p> <p>(2) <input type="checkbox"/> [II.4.1. They originate from a habitat in a Member State or a zone thereof free from infection with bluetongue virus (serotypes 1-24), where no case of infection with bluetongue virus (serotypes 1-24) has been confirmed in the targeted animal population during the last 24 months prior to the date of departure of the consignment and have not been vaccinated</p>		

II. Health information			
Part II: Certification			with a live vaccine against infection with bluetongue virus (serotypes 1-24) in the last 60 days prior to the date of departure of the consignment and the requirements laid down in Article 32(1), point (a), (b) or (c), or Article 32(2) of Commission Delegated Regulation (EU) 2020/688 are fulfilled.]
	(2) <input type="checkbox"/> and/or	[II.4.2.	They originate from a habitat in a Member State or a zone thereof covered by the eradication programme for infection with bluetongue virus (serotypes 1-24) and the requirements laid down in Article 32(1), point (a), (b) or (c), or Article 32(2) of Delegated Regulation (EU) 2020/688 are fulfilled, and they:
	(2) <input type="checkbox"/> either	[II.4.2.1.	have been resident in a Member State or zone thereof seasonally free from infection with bluetongue virus (serotypes 1-24) in accordance with Article 40(3) of Commission Delegated Regulation (EU) 2020/689:
	(2) <input type="checkbox"/> either	[II.4.2.1.1.	for at least 60 days prior to the date of departure of the consignment;]]
	(2) <input type="checkbox"/> and/or	[II.4.2.1.2.	for at least 28 days prior to the date of departure of the consignment and have been subjected to a serological test, with negative results, carried out on samples collected at least 28 days following the date of entry of the animals into the Member State or zone thereof seasonally free from infection with bluetongue virus (serotypes 1-24);]]
	(2) <input type="checkbox"/> and/or	[II.4.2.1.3.	for at least 14 days prior to the date of departure of the consignment and have been subjected to a PCR test, with negative results, carried out on samples collected at least 14 days following the date of entry of the animals into the Member State or zone thereof seasonally free from infection with bluetongue virus (serotypes 1-24);]]
	(2) <input type="checkbox"/> and/or	[II.4.2.2.	have been vaccinated against all serotypes from 1 to 24 of infection with bluetongue virus which were reported in that Member State or zone thereof during the last 2 years prior to the date of departure of the consignment and are within the immunity period guaranteed in the specifications of the vaccine, and:
	(2) <input type="checkbox"/> either	[II.4.2.2.1.	have been vaccinated more than 60 days prior to the date of departure of the consignment;]]
	(2) <input type="checkbox"/> and/or	[II.4.2.2.2.	have been vaccinated with an inactivated vaccine and subjected to a PCR test, with negative results on samples collected at least 14 days after the date of onset of the immunity set in the specifications of the vaccine;]]]
	(2) <input type="checkbox"/> and/or	[II.4.2.3.	have been subjected with positive results to a serological test able to detect specific antibodies against all serotypes 1-24 of infection with bluetongue virus reported in that Member State or zone thereof during the last 2 years prior to the date of departure of the consignment, and:
	(2) <input type="checkbox"/> either	[II.4.2.3.1.	the serological test has been carried out on samples collected at least 60 days prior to the date of departure of the consignment;]]
	(2) <input type="checkbox"/> and/or	[II.4.2.3.2.	the serological test has been carried out on samples collected at least 30 days prior to the date of departure of the consignment and the animals have been subjected to a PCR test, with negative results, carried out on samples collected not earlier than 14 days prior to the date of departure of the consignment;]]]
	(2) <input type="checkbox"/> and/or	[II.4.3.	They originate from a habitat in a Member State or a zone thereof neither free from infection with bluetongue virus (serotypes 1-24) nor covered by the eradication programme for infection with bluetongue virus (serotypes 1-24) and the requirements laid down in Article 32(1), point (a), (b) or (c), or Article 32(2) of Delegated Regulation (EU) 2020/688 are fulfilled, and they:
	(2) <input type="checkbox"/> either	[II.4.3.1.	have been resident at least for the last 60 days prior to the date of departure of the consignment in a habitat situated in a Member State or in an area of at least 150 km radius centred on the habitat, where surveillance in compliance with the requirements laid down in Part II, Chapter 1, Sections 1 and 2, of Annex V to Delegated Regulation (EU) 2020/689 has been carried out during that period, and:
		[II.4.3.1.1.	the animals have been vaccinated against all serotypes from 1 to 24 of infection with bluetongue virus which were reported during the last 2 years prior to the date of departure of the consignment in an area of at least 150 km radius centred on the habitat where the animals were resident and are within the immunity period guaranteed in the specifications

Part II: Certification	II. Health information			
	(2) <input type="checkbox"/> either	[II.4.3.1.1.1. have been vaccinated more than 60 days prior to the date of departure of the consignment;]]]		of the vaccine, and:
	(2) <input type="checkbox"/> and/or	[II.4.3.1.1.2. have been vaccinated with an inactivated vaccine and subjected to a PCR test, with negative results on samples collected at least 14 days after the date of onset of the immunity set in the specifications of the vaccine;]]]		
	(2) <input type="checkbox"/> and/or	[II.4.3.1.2. the animals have been immunised against all serotypes from 1 to 24 of infection with bluetongue virus which were reported during the last 2 years prior to the date of departure of the consignment in an area of at least 150 km radius centred on the habitat where the animals were resident, and:		
	(2) <input type="checkbox"/> either	[II.4.3.1.2.1. the animals have been subjected with positive results to a serological test carried out on samples collected at least 60 days prior to the date of departure of the consignment;]]]		
	(2) <input type="checkbox"/> or	[II.4.3.1.2.2. the animals have been subjected with positive results to a serological test carried out on samples collected at least 30 days prior to the date of departure of the consignment and to a PCR test, with negative results, carried out on samples collected not earlier than 14 days prior to the date of departure of the consignment;]]]		
	(2) <input type="checkbox"/> and/or	[II.4.4. They do not fulfil the requirements laid down in Part II, Chapter 2, Section 1, points 1 to 3, of Annex V to Delegated Regulation (EU) 2020/689 and the competent authority of the Member State of origin authorised movement of those animals to another Member State or zone thereof:		
	(2) <input type="checkbox"/> either	[II.4.4.1. with the status “free from infection with bluetongue virus (serotypes 1-24)” and the Member State of destination has informed the Commission and the other Member States that such movement is authorised subject to the conditions referred to in Article 43(2), points (a), (b) and (c), of Delegated Regulation (EU) 2020/689 and:		
	(2) <input type="checkbox"/> either	[II.4.4.1.1. Part II, Chapter 2, Section 1, point 5, of Annex V to that Regulation, and		
	(2) <input type="checkbox"/> and/or	[II.4.4.1.2. Part II, Chapter 2, Section 1, point 6, of Annex V to that Regulation, and		
	(2) <input type="checkbox"/> and/or	[II.4.4.1.3. Part II, Chapter 2, Section 1, point 7, of Annex V to that Regulation, and		
(2) <input type="checkbox"/> and/or	[II.4.4.1.4. Part II, Chapter 2, Section 1, point 8, of Annex V to that Regulation, and			
	the requirements laid down in Article 32(1), point (a), (b) or (c), or Article 32(2) of Delegated Regulation (EU) 2020/688 and the requirements laid down in Article 33 of that Delegated Regulation are fulfilled]]]			
(2) <input type="checkbox"/> and/or	[II.4.4.2. with an approved eradication programme for infection with bluetongue virus (serotypes 1-24) and the Member State of destination has informed the Commission and the other Member States that such movement is authorised subject to the conditions referred to in Article 43(2), points (a), (b) and (c), of Delegated Regulation (EU) 2020/689 and:			
(2) <input type="checkbox"/> either	[II.4.4.2.1. Part II, Chapter 2, Section 1, point 5, of Annex V to that Regulation, and			
(2) <input type="checkbox"/> and/or	[II.4.4.2.2. Part II, Chapter 2, Section 1, point 6, of Annex V to that Regulation, and			
(2) <input type="checkbox"/> and/or	[II.4.4.2.3. Part II, Chapter 2, Section 1, point 7, of Annex V to that Regulation, and			
(2) <input type="checkbox"/> and/or	[II.4.4.2.4. Part II, Chapter 2, Section 1, point 8, of Annex V to that Regulation, and			
	the requirements laid down in Article 32(1), point (a), (b) or (c), or Article 32(2) of Delegated Regulation (EU) 2020/688 and the requirements laid down in Article 33 of that Delegated Regulation are fulfilled;]]]			
(2) <input type="checkbox"/> and/or	[II.4.4.3. neither free from infection with bluetongue virus (serotypes 1-24) nor covered by the eradication programme for infection with bluetongue virus (serotypes 1-24) and the Member State of destination has informed the Commission and the other Member States that such movement is authorised:			

II. Health information				
Part II: Certification	(2) <input type="checkbox"/> either	[II.4.4.3.1. without any conditions, and:		
	(2) <input type="checkbox"/> and/or	[II.4.4.3.2. subject to the conditions referred to in Part II, Chapter 2, Section 1, point 5, of Annex V to Delegated Regulation (EU) 2020/689, and		
	(2) <input type="checkbox"/> and/or	[II.4.4.3.3. subject to the conditions referred to in Part II, Chapter 2, Section 1, point 6, of Annex V to Delegated Regulation (EU) 2020/689, and		
	(2) <input type="checkbox"/> and/or	[II.4.4.3.4. subject to the conditions referred to in Part II, Chapter 2, Section 1, point 7, of Annex V to Delegated Regulation (EU) 2020/689, and		
	(2) <input type="checkbox"/> and/or	[II.4.4.3.5. subject to the conditions referred to in Part II, Chapter 2, Section 1, point 8, of Annex V to Delegated Regulation (EU) 2020/689, and		
	the requirements laid down in Article 32(1), point (a), (b) or (c), or Article 32(2) of Delegated Regulation (EU) 2020/688 and the requirements laid down in Article 33 of that Delegated Regulation are fulfilled.]]]			
	II.5.	To the best of my knowledge and as declared by the operator, the animals come from a habitat where there were no abnormal mortalities with an undetermined cause.		
	II.6.	Arrangements are made to transport the consignment in accordance with Article 101(1), (2) and (3) of Delegated Regulation (EU) 2020/688.		
	II.7.	This animal health certificate is valid for 10 days from the date of issuing. In the case of transport by waterway/sea of animals, the period of validity of the certificate may be extended by the duration of the journey by waterway/sea.		
	Animal welfare attestation			
At the time of inspection, the animals covered by this animal health certificate were fit to be transported in accordance with the provisions of Council Regulation (EC) No 1/2005 on the intended journey due to start on (insert date).				
Notes:				
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 the Union in this animal health certificate include the United Kingdom in respect of Northern Ireland.				
This animal health certificate shall be completed in accordance with the notes for the completion of certificates provided for in Chapter 2 of Annex I to Commission Implementing Regulation (EU) 2020/2235.				
Part I:				
Box reference	"Place of dispatch": Indicate a habitat of the origin of the animals in the consignment.			
I.11:				
Box reference	"Place of destination": Indicate a habitat or an establishment of the final destination of the consignment.			
I.12:				
Box reference	"Identification number": Indicate identification codes of the animals in the consignment.			
I.30:				
Part II:				
(1)	There may be one or more animals in the consignment.			
(2)	Delete if not applicable.			
(3)	Insert the name of the disease(s).			
(4)	Insert the specific reference to the article(s), title, and number of the relevant legal act(s) adopted by the Commission providing for those requirements.			
(5)	Insert the specific attestation(s) provided for in and required by the relevant legal act(s) adopted by the Commission, as referred to in Article 126(1), points (b)(ii) and (iii), of Regulation (EU) 2016/429 of the European Parliament and of the Council.			
Certifying Officer/Official veterinarian				

Part II: Certification	II. Health information		
	Name (in capital letters)	Qualification and title	
	Date of declaration	Signature	
	Stamp		