

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			Country		
	ISO Code			ISO Code		
	I.7. Country of origin			I.9. Country of destination		
ISO Code			ISO Code			
I.8. Region of origin			I.10. Region of destination			
Code			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"/>		Exhibition <input type="checkbox"/>		Release into the wild <input type="checkbox"/>		
Slaughter <input type="checkbox"/>		Further keeping <input type="checkbox"/>		Confined establishment <input type="checkbox"/>		
Other <input type="checkbox"/>		Event or activity near borders <input type="checkbox"/>		Quarantine or similar establishment <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.27. Total quantity			I.28. Total gross weight			
I.30. Description of consignment						
<b>1. 01 LIVE ANIMALS</b>						

<b>Part I: Description of consignment</b>	<b>0106</b> Other live animals			
	Mammals:			
	<b>010613</b> Camels and other camelids (Camelidae)			
<b>01061300</b> Camels and other camelids (Camelidae)				
#1.	Commodity	Sex	Identification system	Identification Number
Species		Quantity	Age	

	II. Health information		
Part II: Certification	I, the undersigned official veterinarian, hereby certify that:		
	II.1.	The camelid animals (1) of the consignment described in Part I meet the following requirements:	
	II.1.1.	They are identified as provided for in Article 73 of Commission Delegated Regulation (EU) 2019/2035.	
	II.1.2.	They, for at least 30 days prior to the date of departure of the consignment, or since birth, if they are younger than 30 days of age,	
	II.1.2.1.	have been continuously resident in the establishment of origin;	
	II.1.2.2.	have not been in contact with kept camelid animals of a lower health status or subject to movement restrictions for animal health reasons;	
	II.1.2.3.	have not been in direct or indirect contact with kept animals that have entered the Union from a third country or territory during the last 30 days prior to the date of departure of the consignment.	
	II.1.3.	They have not shown clinical signs or symptoms of diseases listed for camelid animals during the clinical examination which was carried out, within the last 24 hours prior to the time of departure of the consignment, on (insert date dd/mm/yyyy).	
	II.2.	According to official information, the animals described in Part I meet the following health requirements:	
	II.2.1.	(2) either <input type="checkbox"/> [They come from establishments or zones not subject to movement restrictions affecting camelid animals 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.]	
		(2) or <input type="checkbox"/> [They come from establishments or zones subject to movement restrictions affecting camelid animals and established for (3), but derogations from movement restrictions have been granted, and:	
		(2) <input type="checkbox"/> [they comply with the requirements set out in (4);]	
		(2) <input type="checkbox"/> [and in particular, they are (5).]	
	II.2.2.	They come from establishments in which infection with Brucella abortus, B. melitensis and B. suis in camelid animals has not been reported during the last 42 days prior to the date of departure of the consignment, and the animals of the consignment have been subjected to a test for infection with Brucella abortus, B. melitensis and B. suis with one of the diagnostic methods provided for in Part 1 of Annex I to Commission Delegated Regulation (EU) 2020/688, carried out, with negative results, on a sample taken during the last 30 days prior to the date of departure of the consignment, and in the case of post-parturient females taken at least 30 days after parturition.	
	II.2.3.	They come from establishments in which surveillance for infection with Mycobacterium tuberculosis complex (M. bovis, M. caprae and M. tuberculosis) has been carried out on the camelid animals kept in the establishments during at least 12 months prior to the date of departure of the consignment, as referred to in Article 23(1), point (e) of Delegated Regulation (EU) 2020/688.	
II.2.4.	They come from establishments in which infection with rabies virus in kept terrestrial animals has not been reported during the last 30 days prior to the date of departure of the consignment.		
II.2.5.	They come from establishments situated within an area of at least 150 km radius around those establishments in which infection with epizootic haemorrhagic disease virus:		
	(2) either <input type="checkbox"/> [has not been reported in kept animals of listed species for that disease during the last 2 years prior to the date of departure of the consignment;]		
	(2) and/or <input type="checkbox"/> [has been reported in kept animals of listed species for that disease during the last 2 years prior to the date of departure and the animals have been kept in a zone seasonally free from epizootic haemorrhagic disease in accordance with Parts 1 and 2 of Annex IX to Delegated Regulation (EU) 2020/688:		

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		<p>(2) either <input type="checkbox"/> for at least 60 days prior to the date of departure of the consignment;]</p> <p>(2) and/or <input type="checkbox"/> 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 seasonally disease-free area;]</p> <p>(2) and/or <input type="checkbox"/> 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 animal into the seasonally disease-free area;]</p> <p>(2) and/or <input type="checkbox"/> [has been reported in kept animals of listed species for that disease during the last 2 years prior to the date of departure and the animals have been protected against attacks by vectors during transportation to the place of destination and they have been kept protected against attacks by vectors in a vector protected establishment fulfilling the requirements provided for in Part 3 of Annex IX to Delegated Regulation (EU) 2020/688:</p> <p>(2) either <input type="checkbox"/> for at least 60 days prior to the date of departure of the consignment;]</p> <p>(2) and/or <input type="checkbox"/> 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 the commencement of the period of protection against attacks by vectors;]</p> <p>(2) and/or <input type="checkbox"/> 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 commencement of the period of protection against attacks by vectors;]</p> <p>(2) and/or <input type="checkbox"/> [has been reported in kept animals of listed species for that disease during the last 2 years prior to the date of departure of the consignment and the Member State of destination has informed the Commission and the other Member States that such movement is authorised].</p> <p>II.2.6. They come from establishments 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>II.2.7. They come from establishments 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, and:</p> <p>(2) either <input type="checkbox"/> [surra has not been reported in the establishments during the last 2 years prior to the date of departure of the consignment.]</p> <p>(2) or <input type="checkbox"/> [surra has been reported during the last 2 years prior to the date of departure of the consignment, and following the date of the last outbreak, the affected establishments have remained under movement restrictions until the date on which the infected animals have been removed from the establishments, and the remaining animals in the establishments have been subjected to a test for surra with one of the diagnostic methods provided for in Part 3 of Annex I to Delegated Regulation (EU) 2020/688, carried out, with negative results, on samples taken at least 6 months following the date on which the infected animals have been removed from the establishments.]</p> <p>(2) either <input type="checkbox"/> [II.2.8. They originate from 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 with a live vaccine against infection with bluetongue virus (serotypes 1-24) within 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 Delegated Regulation (EU) 2020/688 are fulfilled.]</p> <p>(2) and/or <input type="checkbox"/> [II.2.8. They originate from 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:</p> <p>(2) either <input type="checkbox"/> [II.2.8.1. have been kept 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:</p>

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<p>(2) either <input type="checkbox"/></p>	<p>[[II.2.8.1.1.</p>	<p>for at least 60 days prior to the date of departure of the consignment;]]</p>
<p>(2) and/or <input type="checkbox"/></p>	<p>[[II.2.8.1.2.</p>	<p>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);]]</p>
<p>(2) and/or <input type="checkbox"/></p>	<p>[[II.2.8.1.3.</p>	<p>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 animal into the Member State or zone thereof seasonally free from infection with bluetongue virus (serotypes 1-24);]]</p>
<p>(2) and/or <input type="checkbox"/></p>	<p>[[II.2.8.2.</p>	<p>have been protected against attacks by vectors during transportation to the place of destination and have been kept protected against attacks by vectors in a vector protected establishment:</p>
<p>(2) either <input type="checkbox"/></p>	<p>[[II.2.8.2.1.</p>	<p>for at least 60 days prior to the date of departure of the consignment;]]</p>
<p>(2) and/or <input type="checkbox"/></p>	<p>[[II.2.8.2.2.</p>	<p>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 the commencement of the period of protection against attacks by vectors;]]</p>
<p>(2) and/or <input type="checkbox"/></p>	<p>[[II.2.8.2.3.</p>	<p>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 the commencement of the period of protection against attacks by vectors;]]</p>
<p>(2) and/or <input type="checkbox"/></p>	<p>[[II.2.8.3.</p>	<p>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 departure of the consignment in that Member State or zone thereof and are within the immunity period guaranteed in the specifications of the vaccine, and:</p>
<p>(2) either <input type="checkbox"/></p>	<p>[[II.2.8.3.1.</p>	<p>have been vaccinated more than 60 days prior to the date of departure of the consignment;]]</p>
<p>(2) and/or <input type="checkbox"/></p>	<p>[[II.2.8.3.2.</p>	<p>have been vaccinated with an inactivated vaccine and subjected to a PCR test, with negative results, carried out on samples collected at least 14 days after the date of the onset of the immunity set in the specifications of the vaccine;]]</p>
<p>(2) and/or <input type="checkbox"/></p>	<p>[[II.2.8.4.</p>	<p>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 during the last 2 years prior to the date of departure of the consignment in that Member State or zone thereof, and:</p>
<p>(2) either <input type="checkbox"/></p>	<p>[[II.2.8.4.1.</p>	<p>the serological test has been carried out on samples collected at least 60 days prior to the date of departure of the consignment;]]</p>
<p>(2) and/or <input type="checkbox"/></p>	<p>[[II.2.8.4.2.</p>	<p>the serological test has been carried out on samples collected at least 30 days prior to the date of the departure of the consignment and the animal has been subjected to a PCR test, with negative results, carried out</p>

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on samples collected not earlier than 14 days prior to the date of departure of the consignment;]] ]

- (2) and/or  [II.2.8. They originate from 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:
  - (2) either  [II.2.8.1. have been protected against attacks by vectors during transportation to the place of destination and have been kept protected against attacks by vectors in a vector protected establishment:
    - (2) either  for at least 60 days prior to the date of departure of the [II.2.8.1.1. consignment;]]
    - (2) and/or  [II.2.8.1.2. 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 the commencement of the period of protection against attacks by vectors;]]
    - (2) and/or  [II.2.8.1.3. 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 the commencement of the period of protection against attacks by vectors;]]
  - (2) and/or  [II.2.8.2. have been kept for the last 60 days prior to the date of departure of the consignment in an establishment situated in a Member State or within an area of at least 150 km radius centred on the establishment, where surveillance in compliance with the requirements set out 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:
    - (2) either  [II.2.8.2.1. 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 the departure of the consignment within an area of at least 150 km radius centred on the place where the animals were kept, and are within the period of immunity guaranteed in the specifications of the vaccine, and:
      - (2) either  [II.2.8.2.1.1. have been vaccinated more than 60 days prior to the date of departure of the consignment;]]
      - (2) and/or  [II.2.8.2.1.2. 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 the onset of the immunity set in the specifications of the vaccine;]]]]
      - (2) and/or  [II.2.8.2.2. 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 within an area of at least 150 km radius centred on the place where the animals were kept, and:
        - (2) either  [II.2.8.2.2.1. the animals have been subjected with positive results to a serological test carried out on samples collected at least 60 days

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prior to the date of departure of the consignment;]]

(2) and/or  [II.2.8.2.2.2. 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) and/or  [II.2.8. 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) either  [II.2.8.1. 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 the conditions referred to in Article 43(2), points (a), (b) and (c), of Delegated Regulation (EU) 2020/689, and in:

(2) either  Part II, Chapter 2, Section 1, point 5, of Annex V to that [II.2.8.1.1. Delegated Regulation, and

(2) and/or  Part II, Chapter 2, Section 1, point 6, of Annex V to that [II.2.8.1.2. Delegated Regulation, and

(2) and/or  Part II, Chapter 2, Section 1, point 7, of Annex V to that [II.2.8.1.3. Delegated Regulation, and

(2) and/or  Part II, Chapter 2, Section 1, point 8, of Annex V to that [II.2.8.1.4. Delegated 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) and/or  [II.2.8.2. 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) either  Part II, Chapter 2, Section 1, point 5, of Annex V to that [II.2.8.2.1. Delegated Regulation, and

(2) and/or  Part II, Chapter 2, Section 1, point 6, of Annex V to that [II.2.8.2.2. Delegated Regulation, and

(2) and/or  Part II, Chapter 2, Section 1, point 7, of Annex V to that [II.2.8.2.3. Delegated Regulation, and

(2) and/or  Part II, Chapter 2, Section 1, point 8, of Annex V to that [II.2.8.2.4. Delegated 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) and/or  [II.2.8.2. 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:

(2) either  without any conditions, and [II.2.8.2.1.

(2) and/or  subject to the conditions referred to in Part II, Chapter 2,

<b>Part II: Certification</b>	<p>II. Health information</p>		
	<p style="text-align: right;">[II.2.8.2.2. Section 1, point 5, of Annex V to Delegated Regulation (EU) 2020/689, and</p> <p>(2) and/or <input type="checkbox"/> subject to under the conditions referred to in Part II, Chapter 2, Section 1, point 6, of Annex V to Delegated Regulation (EU) 2020/689, and</p> <p style="text-align: right;">[II.2.8.2.3.</p> <p>(2) and/or <input type="checkbox"/> 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</p> <p style="text-align: right;">[II.2.8.2.4.</p> <p>(2) and/or <input type="checkbox"/> 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</p> <p style="text-align: right;">[II.2.8.2.5.</p>		
	<p>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.]]</p> <p>(2) <input type="checkbox"/> [II.2.9. They are moved to a Member State or zone thereof with the status free from infectious bovine rhinotracheitis/infectious pustular vulvovaginitis or with an approved eradication programme for infectious bovine rhinotracheitis/infectious pustular vulvovaginitis in bovine animals and they come from an establishment in which infectious bovine rhinotracheitis/infectious pustular vulvovaginitis in camelid animals has not been reported during the last 30 days prior to the date of departure of the consignment.]</p>		
	<p>II.3. To the best of my knowledge and as declared by the operator, the animals come from establishments where there were no abnormal mortalities with an undetermined cause.</p> <p>II.4. Arrangements are made to transport the consignment in accordance with Article 4 of Delegated Regulation (EU) 2020/688.</p> <p>II.5. 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.</p>		
	<p>(2)(6) <input type="checkbox"/> Since the date of departure from their establishments of origin and prior to the date of arrival to this establishment approved for assembly operations, none of the animals of the consignment has undergone more than two assembly operations, and:</p> <p>(2) either <input type="checkbox"/> [they come from their establishments of origin.]]</p> <p>(2) or <input type="checkbox"/> [at least one of the animals of the consignment has undergone one assembly operation in an approved establishment.]]</p> <p>(2) or <input type="checkbox"/> [at least one of the animals of the consignment has undergone two assembly operations in the approved establishments.]]</p>		
	<p>Animal welfare attestation</p> <p>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).</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 the Union in this animal health certificate include the United Kingdom in respect of Northern Ireland.</p> <p>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.</p> <p>Part I:</p> <p>Box I.11: “Place of dispatch”: Indicate an establishment of the origin of the animals in the consignment or an establishment approved for assembly operations in accordance with Articles 97 and 99 of Regulation (EU) 2016/429 of the European Parliament and of the Council.</p>		



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<b>Part II: Certification</b>	Box reference I.12:	“Place of destination”: Indicate an establishment of the final destination of the consignment or an establishment approved for assembly operations in accordance with Articles 97 and 99 of Regulation (EU) 2016/429.	
	Box reference I.17:	“Accompanying documents”: In case the animals are dispatched from an establishment approved for assembly operations in the Member State of origin, the reference number(s) of the official document(s), based on which the animal health certificate for this consignment is issued in this establishment approved for assembly operations, may be indicated.  In case the animals are dispatched from an establishment approved for assembly operations in the Member State of passage, the reference number(s) of the certificate(s), based on which the animal health certificate for this consignment is issued in this establishment approved for assembly operations, shall be indicated.	
	Box reference I.30:	“Identification number”: Indicate identification codes of the animals in the consignment identified in accordance with Article 73 of Delegated Regulation (EU) 2019/2035.	
	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.	
	(6)	Applicable in case the consignment is dispatched from the establishment approved for assembly operations.	
Certifying Officer/Official veterinarian			
Name (in capital letters)		Qualification and title	
Date of declaration		Signature	
Stamp			