EUROPEAN UNION

INTRA

Part I: Description of consignment

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.8. Region of origin Code

I.9. Country of destination ISO 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

Mode	International transport document	Identification

I.16. Transporter
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 ☐ Ambient ☐ Frozen ☐

I.19. Container No / Seal No

I.20. Certified as
Confined establishment ☐

I.21. For transit through a third country ☐
Third country ISO Code
Exit point BCP code
Entry point BCP code

I.22. For transit through Member State(s) ☐
Member State ISO Code

I.23. For export ☐
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

1. 01 LIVE ANIMALS

0106 Other live animals
Mammals:
010611 Primates

Part I: Description of consignment

01061100 Primates

#1.	Commodity	Breed/Category	Sex	Identification system
Species		Identification Number	Age	Quantity

Part II: Certification

II. Health information

II. Health information

I, the undersigned official veterinarian, hereby certify, that:

II.1. The animals (1) of the consignment described in Part I meet the following requirements:

II.1.1. Their confined establishment of dispatch is approved in accordance with Articles 97 and 99 of Regulation (EU) 2016/429 of the European Parliament and of the Council.

II.1.2. They have not shown clinical signs or symptoms of diseases, in particular relevant diseases listed in the Annex to Commission Implementing Regulation (EU) 2018/1882, during the clinical examination, or where this is not possible, a clinical inspection, which was carried out within the last 48 hours prior to the time of departure of the consignment, on (insert date dd/mm/yyyy).

II.2. According to official information, animals of the consignment described in Part I meet the following health requirements:

II.2.1. They come from a confined establishment or a zone:

(2) ○ either [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.]

(2) ○ or [subject to restrictions affecting the species of animals to be moved and established for (3), but derogations from movement restrictions have been granted, and:

(2) ☐ [they comply with the requirements set out in (4);]]

(2) ☐ [and in particular, they are (5).]]

(2) (6) ☐ either [II.2.2. 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) in the last 60 days prior to the date of movement 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) (6) ☐ and/or [II.2.2. 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 they:

(2) ☐ either [II.2.2.1. 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:

(2) ☐ either [II.2.2.1.1. for at least 60 days prior to the date of departure of the consignment;]]

(2) ☐ and/or [II.2.2.1.2. 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) ☐ and/or [II.2.2.1.3. 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) ☐ and/or [II.2.2.2. 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 [II.2.2.2.1. for at least 60 days prior to the date of departure of the consignment]]

(2) ☐ and/or [II.2.2.2.2. 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]]

Part II: Certification

II. Health information			
(2) ☐ and/or	[II.2.2.2.3.	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.2.2.3.	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) ☐ either	[II.2.2.3.1.	have been vaccinated more than 60 days prior to the date of departure of the consignment;]]	
(2) ☐ and/or	[II.2.2.3.2.	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) ☐ and/or	[II.2.2.4.	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) ☐ either	[II.2.2.4.1.	the serological test has been carried out on samples collected at least 60 days prior to the date of departure of the consignment;]]	
(2) ☐ and/or	[II.2.2.4.2.	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) (6) ☐ and/or	[II.2.2.	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 they:	
(2) ☐ either	[II.2.2.1.	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	[II.2.2.1.1.	for at least 60 days prior to the date of departure of the consignment;]]	
(2) ☐ and/or	[II.2.2.1.2.	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 the commencement of the period of protection against attacks by vectors;]]	
(2) ☐ and/or	[II.2.2.1.3.	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 the commencement of the period of protection against attacks by vectors;]]]	
(2) ☐ and/or	[II.2.2.2.	have been kept at least 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 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:	
(2) ☐ either	[II.2.2.2.1.	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 departure of the consignment in an area of at least 150 km radius centred on the place where the animals were kept and are within the immunity period guaranteed in the specifications of the vaccine, and:	
(2) ☐ either	[II.2.2.2.1.1.	have been vaccinated more than 60 days prior to the date of departure of the consignment;]]]	
(2) ☐ and/or	[II.2.2.2.1.2.	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 onset of the immunity set in the specifications of the vaccine;]]]]	

Part II: Certification

II. Health information			
(2) ☐ and/or	[II.2.2.2.2.	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 in an area of at least 150 km radius centred on the place where the animals were kept, and:	
(2) ☐ either	[II.2.2.2.2.1.	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) ☐ or	[II.2.2.2.2.2.	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) (6) ☐ and/or	[II.2.2.	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.2.2.1.	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) ☐ either	[II.2.2.1.1.	Part II, Chapter 2, Section 1, point 5, of Annex V to that Delegated Regulation, and	
(2) ☐ and/or	[II.2.2.1.2.	Part II, Chapter 2, Section 1, point 6, of Annex V to that Delegated Regulation, and	
(2) ☐ and/or	[II.2.2.1.3.	Part II, Chapter 2, Section 1, point 7, of Annex V to that Delegated Regulation, and	
(2) ☐ and/or	[II.2.2.1.4.	Part II, Chapter 2, Section 1, point 8, of Annex V to that 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.2.2.2.	with an approved eradication program 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	[II.2.2.2.1.	Part II, Chapter 2, Section 1, point 5, of Annex V to that Delegated Regulation, and	
(2) ☐ and/or	[II.2.2.2.2.	Part II, Chapter 2, Section 1, point 6, of Annex V to that Delegated Regulation, and	
(2) ☐ and/or	[II.2.2.2.3.	Part II, Chapter 2, Section 1, point 7, of Annex V to that Delegated Regulation, and	
(2) ☐ and/or	[II.2.2.2.4.	Part II, Chapter 2, Section 1, point 8, of Annex V to that 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.2.2.3.	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:	
(2) ☐ either	[II.2.2.3.1.	without any conditions, and:	
(2) ☐ and/or	[II.2.2.3.2.	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) ☐ and/or	[II.2.2.3.3.	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) ☐ and/or	[II.2.2.3.4.	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) ☐ and/or	[II.2.2.3.5.	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	

Part II: Certification

II. Health information		

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.3. To the best of my knowledge and as declared by the operator:

II.3.1. In the confined establishment of departure of the consignment there are no abnormal mortalities with an undetermined cause affecting the animals to be moved.

II.3.2. The animals have not been in contact with animals which are subject to movement restrictions referred to in point II.2.1, or with animals of a lower health status.

II.3.3. Based on the results of the surveillance plan of the confined establishment, the animals do not pose a significant risk at the confined establishment of destination for the spread of diseases for which they are listed.

II.4. Arrangements are made to transport the consignment in accordance with Article 4 of Delegated Regulation (EU) 2020/688.

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 10 days for the 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 I.11: “Place of dispatch”: Indicate a confined establishment approved in accordance with Articles 97 and 99 of Regulation (EU) 2016/429.

Box reference I.12: “Place of destination”: Indicate a confined establishment approved in accordance with Articles 97 and 99 of Regulation (EU) 2016/429.

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), titleand 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) Only in the case of animals belonging to the families Antilocapridae, Bovidae, Camelidae, Cervidae, Giraffidae, Moschidae or Tragulidae.

Certifying Officer/Official veterinarian	
Name (in capital letters)	Qualification and title
Date of declaration	Signature
Stamp	