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.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

Mode	International transport document	Identification

I.16. Transporter
Name
Address
Activity ID
Country ISO Code

I.17. Accompanying documents
Commercial document reference
Date of issue
Country
Place of issue

~~I.18. Transport conditions~~

I.19. Container No / Seal No

I.20. Certified as
Event or activity near borders ☐ Slaughter ☐ Quarantine or similar establishment ☐ Other ☐
Confined establishment ☐ Travelling circus/animal act ☐ Further keeping ☐ Exhibition ☐
Release into the wild ☐

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.25. Journey Log

I.27. Total quantity

I.28. Total gross weight

I.30. Description of consignment

Commodity	Species	Sex	Identification system	Identification Number

Quantity	Age

Part II: Certification

II. Health information

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 the 30 day period prior to the 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 30 day period prior to the departure of the animals.

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 24 hour period prior to 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. They do not come from establishments subject to movement restrictions affecting the species or situated in a restricted zone established for reasons of diseases listed for camelid animals.

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 departure, and the animals in 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 30 day period prior to departure, 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 on the establishments during at least the 12 month period prior to departure, as referred to in Article 23(1)(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 30 day period prior to departure.

(2) ☐ [II.2.5. 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 30 day period prior to departure.]

II.2.6. They come from establishments situated in an area of at least 150 km radius around those establishments in which infection with epizootic haemorrhagic disease virus has not been reported in any establishment during the last 2 years prior to departure.

II.2.7. They come from establishments in which anthrax in ungulates has not been reported during the 15 day period prior to departure.

II.2.8. They come from establishments in which surra (Trypanosoma evansi) has not been reported during the 30 day period prior to departure, and

(2) either ○ [surra has not been reported in the establishments during the last 2 years prior to their departure.]

(2) or ○ [surra has been reported during the last 2 years prior to departure, following the last outbreak the affected establishments have remained under movement restrictions until:

– the infected animals have been removed from the establishments, and

Part II: Certification

II. Health information

– the remaining animals on the establishments have been subjected to a test for surra (Trypanosoma evansi) 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 after the infected animals have been removed from the establishments.]

(2) either ☐ [II.2.9. They originate from a Member State or a zone free from infection with bluetongue virus (serotypes 1-24), where no case of infection with bluetongue virus (serotypes 1-24) has been confirmed during the last 24 months in the targeted animal population and have not been vaccinated with a live vaccine against infection with bluetongue virus (serotypes 1-24) in the 60 day period before the date of movement and the requirements laid down in Article 32(1)(a), (b) or (c) or Article 32(2) of Delegated Regulation (EU) 2020/688 are fulfilled.]

(2) and/or ☐ [II.2.9. They originate from a Member State or a zone covered by the eradication programme for infection with bluetongue virus (serotypes 1-24) and the requirements laid down in Article 32(1)(a), (b) or (c) or Article 32(2) of Delegated Regulation (EU) 2020/688 are fulfilled, and they

(2) either ☐ [II.2.9.1. have been kept in a Member State or zone 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.2.9.1.1. for at least 60 days prior to the date of movement]]

(2) and/or ☐ [II.2.9.1.2. for at least 28 days prior to the date of movement and have been subjected to a serological test, with negative results, carried out on samples collected at least 28 days following the entry date of the animal into the Member State or zone seasonally free from infection with bluetongue virus (serotypes 1-24)]]

(2) and/or ☐ [II.2.9.1.3. for at least 14 days prior to the date of movement and have been subjected to a PCR test, with negative results, carried out on samples collected at least 14 days following the entry date of the animal into the Member State or zone seasonally free from infection with bluetongue virus (serotypes 1-24);]]]

(2) and/or ☐ [II.2.9.2. have been protected against attacks by the 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.2.9.2.1. for at least 60 days prior to the date of movement]]

(2) and/or ☐ [II.2.9.2.2. for at least 28 days prior to the date of movement 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.2.9.2.3. for at least 14 days prior to the date of movement 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.2.9.3. have been vaccinated against those serotypes from 1 to 24 of infection with bluetongue virus which were reported during the past 2 years in that Member State or zone and are within the immunity period guaranteed in the specifications of the vaccine and

(2) either ☐ [II.2.9.3.1. have been vaccinated more than 60 days before the date of movement]]

(2) and/or ☐ [II.2.9.3.2. 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 onset of the immunity set in the specifications of the vaccine;]]]

Part II: Certification

II. Health information

(2) and/or ☐ [II.2.9.4.] 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 during the past 2 years in that Member State or zone and

(2) either ☐ [II.2.9.4.1.] the serological test has been carried out on samples collected at least 60 days before the date of movement]]

(2) and/or ☐ [II.2.9.4.2.] the serological test has been carried out on samples collected at least 30 days before the date of the movement and the animal has been subjected to a PCR test, with negative results, carried out on samples collected not earlier than 14 days before the date of movement;]]]

(2) and/or ☐ [II.2.9.] They originate from a Member State or a zone 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)(a), (b) or (c) or Article 32(2) of Delegated Regulation (EU) 2020/688 are fulfilled, and they

(2) either ☐ [II.2.9.1.] have been protected against attacks by the 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.2.9.1.1.] for at least 60 days prior to the date of movement]]

(2) and/or ☐ [II.2.9.1.2.] for at least 28 days prior to the date of movement 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.2.9.1.3.] for at least 14 days prior to the date of movement 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.2.9.2.] have been kept for the 60 day period prior to departure in an establishment situated in a Member State or in an area of at least 150 km radius centred on the establishment, where surveillance in compliance with the requirements set out in Sections 1 and 2 of Chapter 1 of Part II of Annex V to Delegated Regulation (EU) 2020/689 has been carried out during that period, and

(2) either ☐ [II.2.9.2.1.] the animals have been vaccinated against those serotypes from 1 to 24 of infection with bluetongue virus which were reported during the past 2 years 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.2.9.2.1.1.] have been vaccinated more than 60 days before the date of movement]]]

(2) and/or ☐ [II.2.9.2.1.2.] 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 onset of the immunity set in the specifications of the vaccine;]]]]

(2) and/or ☐ [II.2.9.2.2.] the animals have been immunised against those serotypes from 1 to 24 of infection with bluetongue virus which were reported during the past 2 years in an area of at least 150 km radius centred on the place where the animals were kept, and

(2) either ☐ [II.2.9.2.2.1.] the animals have been subjected with positive results to a serological test carried out on samples collected at least 60 days before the date of movement]]]

Part II: Certification

II. Health information

(2) and/or ☐ [II.2.9.2.2.2.] the animals have been subjected with positive results to a serological test carried out on samples collected at least 30 days before the date of the movement and to a PCR test, with negative results, carried out on samples collected not earlier than 14 days before the date of movement;]]]]

(2) and/or ☐ [II.2.9.] They do not fulfil the requirements laid down in points 1 to 3 of Section 1 of Chapter 2 of Part II 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.2.9.1.] 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)(a), (b) and (c) of Delegated Regulation (EU) 2020/689, and

(2) either ☐ [II.2.9.1.1.] point 5 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and

(2) and/or ☐ [II.2.9.1.2.] point 6 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and

(2) and/or ☐ [II.2.9.1.3.] point 7 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and

(2) and/or ☐ [II.2.9.1.4.] point 8 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and

the requirements laid down in Article 32(1)(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.2.9.2.] 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)(a), (b) and (c) of Delegated Regulation (EU) 2020/689 and

(2) either ☐ [II.2.9.2.1.] point 5 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and

(2) and/or ☐ [II.2.9.2.2.] point 6 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and

(2) and/or ☐ [II.2.9.2.3.] point 7 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and

(2) and/or ☐ [II.2.9.2.4.] point 8 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and

the requirements laid down in Article 32(1)(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.2.9.2.] 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 ☐ [II.2.9.2.1.] without any conditions, and

(2) and/or ☐ [II.2.9.2.2.] subject to the conditions referred to in point 5 of Section 1 of Chapter 2 of Part II of Annex V to Delegated Regulation (EU) 2020/689, and

(2) and/or ☐ [II.2.9.2.3.] subject to under the conditions referred to in point 6 of Section 1 of Chapter 2 of Part II of Annex V to Delegated Regulation (EU) 2020/689, and

(2) and/or ☐ [II.2.9.2.4.] subject to the conditions referred to in point 7 of Section 1 of Chapter 2 of Part II of Annex V to Delegated Regulation (EU) 2020/689, and

Part II: Certification

II. Health information

(2) and/or ☐ subject to the conditions referred to in point 8 of Section 1 of Chapter [II.2.9.2.5. 2 of Part II of Annex V to Delegated Regulation (EU) 2020/689, and

the requirements laid down in Article 32(1)(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, the animals come from establishments where there were no abnormal mortalities with an undetermined cause.

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

II.5. This 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.

(2)(3) ☐ [II.6. Since leaving their establishments of origin and before arriving to this establishment approved for assembly operations, none of the animals of the consignment has undergone more than two assembly operations, and

(2) either ☐ [they come from their establishments of origin.]]

(2) or ☐ [at least one of the animals of the consignment has undergone one assembly operation on an approved establishment.]]

(2) or ☐ [at least one of the animals of the consignment has undergone two assembly operations on approved establishments.]]

Animal welfare attestation

At the time of inspection, the animals covered by this 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).

Part II: Certification

II. Health information

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 European Union in this certificate include the United Kingdom in respect of Northern Ireland.

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.

Part I:

Box reference 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.

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, must 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 can be one or more animals in the consignment.

(2) Delete if not applicable.

(3) Applicable in case the consignment is dispatched from the establishment approved for assembly operations.

Certifying Officer/Official veterinarian

Name (in capital letters)
Date of signature
Stamp

Authority name
Signature