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
Accompanying document reference
Date of issue
Country
Place of issue

I.18. Transport conditions
Ambient ☐ Frozen ☐ Chilled ☐

I.19. Container No / Seal No

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

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

0102 Live bovine animals

#1.	Commodity	Subcategory	Sex	Identification system
Species		Identification Number	Age	Quantity

Part II: Certification

II. Health information

I, the undersigned official veterinarian, hereby certify that:

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

II.1.1. They are identified as provided for in Article 38 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 bovine 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 bovine 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 bovine animals.

II.2.2. They come from establishments free from infection with Brucella abortus, B. melitensis and B. suis without vaccination regarding bovine animals, and

(2) either ☐ [the establishments of origin are situated in a Member State or zone thereof with the status free from infection with Brucella abortus, B. melitensis and B. suis regarding the bovine population;]

(2) and/or ☐ [they 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;]

(2) and/or ☐ [they are less than 12 months old;]

(2) and/or ☐ [they are castrated.]

II.2.3. They come from establishments free from infection with Mycobacterium tuberculosis complex (M. bovis, M. caprae and M. tuberculosis), and

(2) either ☐ [the establishments of origin are situated in a Member State or zone thereof with the status free from infection with Mycobacterium tuberculosis complex (M. bovis, M. caprae and M. tuberculosis);]

(2) and/or ☐ [they have been subjected to a test for infection with Mycobacterium tuberculosis complex (M. bovis, M. caprae and M. tuberculosis) with one of the diagnostic methods provided for in Part 2 of Annex I to Delegated Regulation (EU) 2020/688, carried out, with negative results, during the 30 day period prior to departure;]

(2) and/or ☐ [they are less than 6 weeks old.]

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.

II.2.5. 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 kept animals of listed species for that disease during the last 2 years prior to departure.

II.2.6. They come from establishments in which anthrax in ungulates has not been reported during the 15 days period prior to departure.

II.2.7. They come from establishments in which surra (Trypanosoma evansi) has not been reported during the 30 days period prior to departure, and

Part II: Certification

II. Health information			

(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

– 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.2.8. 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.2.8. 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.2.8.1. 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.2.8.1.1. for at least 60 days prior to the date of movement]]

(2) and/or ☐ [II.2.8.1.2. 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.2.8.1.3. 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.2.8.2. 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.2.8.2.1. for at least 60 days prior to the date of movement]]

(2) and/or ☐ [II.2.8.2.2. 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.2.8.2.3. 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.2.8.3. 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

Part II: Certification

II. Health information		

specifications of the vaccine and

(2) either ☐ [II.2.8.3.1.] have been vaccinated more than 60 days before the date of movement]]

(2) and/or ☐ [II.2.8.3.2.] 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;]]]

(2) and/or ☐ [II.2.8.4.] 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.2.8.4.1.] the serological test has been carried out on samples collected at least 60 days before the date of movement.]]]

(2) and/or ☐ [II.2.8.4.2.] 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.2.8.] 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.2.8.1.] 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 ☐ [II.2.8.1.1.] for at least 60 days prior to the date of movement]]

(2) and/or ☐ [II.2.8.1.2.] 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.2.8.1.3.] 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.2.8.2.] 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.2.8.2.1.] 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.2.8.2.1.1.] have been vaccinated more than 60 days before the date of movement]]]

(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 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 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

Part II: Certification

II. Health information			
(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 before the date of movement]]]
(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 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.2.8.	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.2.8.1.	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 to the conditions referred to in Article 43(2)(a), (b) and (c) of Delegated Regulation (EU) 2020/689, and
(2)			either ☐ [II.2.8.1.1. point 5 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and
(2)			and/or ☐ [II.2.8.1.2. point 6 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and
(2)			and/or ☐ [II.2.8.1.3. point 7 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and
(2)			and/or ☐ [II.2.8.1.4. 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.2.8.2.	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.2.8.2.1. point 5 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and
(2)			and/or ☐ [II.2.8.2.2. point 6 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and
(2)			and/or ☐ [II.2.8.2.3. point 7 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and
(2)			and/or ☐ [II.2.8.2.4. 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.2.8.3.	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.2.8.3.1. without any conditions, and
(2)			and/or ☐ [II.2.8.3.2. 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.2.8.3.3. subject to 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

Part II: Certification

II. Health information

(2) and/or ☐ [II.2.8.3.4. 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

(2) and/or ☐ [II.2.8.3.5. subject to the conditions referred to in point 8 of Section 1 of Chapter 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.]]]

(2) ☐ [(2)either ○ [II.2.9. They are moved to a Member State or zone thereof with the status free from enzootic bovine leukosis, and

(2) either ○ [II.2.9.1. they come from establishments free from enzootic bovine leukosis.]]

(2) or ○ [II.2.9.1. they come from establishments not free from enzootic bovine leukosis, and enzootic bovine leukosis has not been reported in those establishments during the 24 month period prior to departure, and

(2) either ☐ [II.2.9.1.1. they are over 24 months of age and they have been subjected to a serological test for enzootic bovine leukosis with one of the diagnostic methods provided for in Part 4 of Annex I to Delegated Regulation (EU) 2020/688, carried out with negative results

(2) either ☐ [II.2.9.1.1.1. on samples taken on two occasions at an interval of at least four months while kept in isolation from the other bovine animals of the establishment]]]

(2) and/or ☐ [II.2.9.1.1.2. on a sample taken during the 30 day period prior to the departure of the consignment, and all bovine animals over 24 months of age kept in the establishment have been subjected to a serological test for enzootic bovine leukosis with one of the diagnostic methods provided for in Part 4 of Annex I to Delegated Regulation (EU) 2020/688, carried out, with negative results, on samples taken on two occasions at an interval of not less than four months during the 12 month period prior to the departure of the consignment;]]]]

(2) and/or ☐ [II.2.9.1.2. they are less than 24 months of age and they were born to dam subjected to a serological test for enzootic bovine leukosis with one of the diagnostic methods provided for in Part 4 of Annex I to Delegated Regulation (EU) 2020/688, carried out, with negative results, on samples taken on two occasions at an interval of not less than four months during the 12 month period prior to the departure of the consignment.]]]

(2) or ○ [II.2.9. They are moved to a Member State or zone thereof with an approved eradication programme for enzootic bovine leukosis, and

(2) either ○ [II.2.9.1. they come from establishments free from enzootic bovine leukosis.]]

(2) or ○ [II.2.9.1. they come from establishments not free from enzootic bovine leukosis, and enzootic bovine leukosis has not been reported in those establishments during the 24 month period prior to departure, and

(2) either ☐ [II.2.9.1.1. they are over 24 months of age and they have been subjected to a serological test for enzootic bovine leukosis with one of the diagnostic methods provided for in Part 4 of Annex I to Delegated Regulation (EU) 2020/688, carried out with negative results

(2) either ☐ [II.2.9.1.1.1. on samples taken on two occasions at an interval of at least four months while kept in isolation from the other bovine animals of the establishment]]]

(2) and/or ☐ [II.2.9.1.1.2. on a sample taken during the 30 day period prior to the departure of the consignment, and all bovine animals

Part II: Certification

II. Health information

over 24 months of age kept in the establishment have been subjected to a serological test for enzootic bovine leukosis with one of the diagnostic methods provided for in Part 4 of Annex I to Delegated Regulation (EU) 2020/688, carried out, with negative results, on samples taken on two occasions at an interval of not less than four months during the 12 month period prior to the departure of the consignment;]]]]

(2) and/or ☐ [II.2.9.1.2. they are less than 24 months of age and they were born to dam, which has been subjected to a serological test for enzootic bovine leukosis with one of the diagnostic methods provided for in Part 4 of Annex I to Delegated Regulation (EU) 2020/688, carried out, with negative results, on samples taken on two occasions at an interval of not less than four months during the 12 month period prior to the departure of the consignment.]]]]

(2) ☐ [(2)either ○ [II.2.10. They are moved to a Member State or zone thereof with the status free from infectious bovine rhinotracheitis/infectious pustular vulvovaginitis and they have not been vaccinated against infectious bovine rhinotracheitis/infectious pustular vulvovaginitis, and

(2) either ○ [II.2.10.1. they come from establishments free from infectious bovine rhinotracheitis/infectious pustular vulvovaginitis, and

(2) either ☐ [II.2.10.1.1. the establishments of origin are situated in a Member State or zone thereof with the status free from infectious bovine rhinotracheitis/infectious pustular vulvovaginitis]]

(2) and/or ☐ [II.2.10.1.2 the animals have been subjected to quarantine for at least 30 days prior to departure and have been subjected to a serological test for the detection of antibodies against whole bovine herpes virus-1 (BoHV-1) with one of the diagnostic methods provided for in Part 5 of Annex I to Delegated Regulation (EU) 2020/688, with a negative result, carried out on a sample taken during the 15 day period prior to the departure of the consignment.]]]

(2) or ○ [II.2.10.1. they come from establishments not free from infectious bovine rhinotracheitis/infectious pustular vulvovaginitis and they have been kept in an approved quarantine establishment for at least 30 days prior to departure and have been subjected to a serological test for the detection of antibodies against whole BoHV-1, with one of the diagnostic methods provided for in Part 5 of Annex I to Delegated Regulation (EU) 2020/688, with a negative result, carried out on a sample taken not less than 21 days after commencement of the quarantine.]]

(2) or ○ [II.2.10. They are moved to a Member State or zone thereof with an approved eradication programme for infectious bovine rhinotracheitis/infectious pustular vulvovaginitis, and

(2) either ○ [II.2.10.1. they come from establishments free from infectious bovine rhinotracheitis/infectious pustular vulvovaginitis, and

(2) either ☐ [II.2.10.1.1. the establishments of origin are situated in a Member State or zone thereof with the status free from infectious bovine rhinotracheitis/infectious pustular vulvovaginitis]]

(2) and/or ☐ [II.2.10.1.2. the establishments of origin are situated in a Member State or zone thereof with an approved eradication programme for infectious bovine rhinotracheitis/infectious pustular vulvovaginitis]]

(2) and/or ☐ [II.2.10.1.3. the animals have been subjected to quarantine for at least 30 days prior to departure and have been subjected to a serological test for the detection of antibodies against whole bovine herpes virus-1 (BoHV-1) with one of the diagnostic methods provided for in Part 5 of Annex I to Delegated Regulation (EU) 2020/688 with a negative result, carried out on a sample taken during the 15 day period prior to the departure of the consignment]]

(2) and/or ☐ the animals are destined for an establishment which keeps bovine

Part II: Certification

II. Health information

[II.2.10.1.4. animals for meat production without contact to bovine animals of other establishments, and from which they are directly moved to the slaughterhouse.]]]

(2) or ○ [II.2.10.1. they come from establishments not free from infectious bovine rhinotracheitis/infectious pustular vulvovaginitis, and

– they have been kept in an approved quarantine establishment for at least 30 days prior to departure, and

– they have been subjected to a serological test for the detection of antibodies against whole BoHV-1, with one of the diagnostic methods provided for in Part 5 of Annex I to Delegated Regulation (EU) 2020/688, with a negative result, carried out on a sample taken not less than 21 days after commencement of the quarantine.]]]

(2) ☐ [(2)either ○ [II.2.11. They are moved to a Member State or zone thereof with the status free from bovine viral diarrhoea and they have not been vaccinated against bovine viral diarrhoea, and

(2) either ○ [II.2.11.1. they come from establishments free from bovine viral diarrhoea, and

(2) either ☐ [II.2.11.1.1. the establishments of origin are situated in a Member State or zone thereof with the status free from bovine viral diarrhoea]]

(2) and/or ☐ [II.2.11.1.2. the establishments of origin have been subjected to a testing regime as referred in point 1(c)(ii) or (iii) of Section 2 of Chapter 1 of Part VI of Annex IV to Delegated Regulation (EU) 2020/689, carried out, with negative results, within the four months period prior to the departure of the consignment]]

(2) and/or ☐ [II.2.11.1.3. the animals have been tested individually to exclude the presence of bovine viral diarrhoea virus prior to the departure of the consignment.]]]

(2) or ○ [II.2.11.1. they come from establishments not free from bovine viral diarrhoea and they have been subjected to a test for bovine viral diarrhoea virus antigen or genome with one of the diagnostic methods provided for in Part 6 of Annex I to Delegated Regulation (EU) 2020/688, carried out with negative results, and

(2) either ☐ [II.2.11.1.1. they have been kept in an approved quarantine establishment for a period of at least 21 days prior to the departure of the consignment

(2) ☐ [and in case of pregnant dams, they have been subjected to a serological test for the detection of antibodies against bovine viral diarrhoea virus with one of the diagnostic methods provided for in Part 6 of Annex I to Delegated Regulation (EU) 2020/688, carried out, with negative results, on samples taken not less than 21 days after commencement of the quarantine]]]

(2) and/or ☐ [II.2.11.1.2. they have been subjected to a serological test for the detection of antibodies against bovine viral diarrhoea virus with one of the diagnostic methods provided for in Part 6 of Annex I to Delegated Regulation (EU) 2020/688, with positive results,

(2) either ☐ [II.2.11.1.2.1. in case of non-pregnant animals, carried out on samples taken prior to departure of the consignment]]]

(2) and/or ☐ [II.2.11.1.2.1. in case of pregnant dams, carried out on samples taken before insemination preceding the current gestation.]]]]

(2) or ○ [II.2.11. They are moved to a Member State or zone thereof with an approved eradication programme for bovine viral diarrhoea, and

(2) either ☐ [II.2.11.1. they come from establishments free from bovine viral diarrhoea, and

(2) either ☐ [II.2.11.1.1. the establishments of origin are situated in a Member State or zone thereof with the status free from bovine viral diarrhoea]]

(2) and/or ☐ the establishments of origin are situated in a Member State or zone

Part II: Certification

II. Health information

[II.2.11.1.2. thereof with an approved eradication programme for bovine viral diarrhoea]]

(2) and/or ☐ [II.2.11.1.3. the establishments of origin have been subjected to a testing regime as referred in point 1(c)(ii) or (iii) of Section 2 of Chapter 1 of Part VI of Annex IV to Delegated Regulation (EU) 2020/689, carried out, with negative results, within the last 4 months prior to the departure of the consignment]]

(2) and/or ☐ [II.2.11.1.4. the animals have been tested individually to exclude the presence of bovine viral diarrhoea virus prior to the departure of the consignment]]

(2) and/or ☐ [II.2.11.1.5. the animals are destined for an establishment which keeps bovine animals for meat production separate from bovine animals of other establishments, and from which they are directly moved to the slaughterhouse]]

(2) and/or ☐ [II.2.11.2. they come from establishments not free from bovine viral diarrhoea and they have been subjected to a test for bovine viral diarrhoea virus antigen or genome with one of the diagnostic methods provided for in Part 6 of Annex I to Delegated Regulation (EU) 2020/688, carried out with negative results, and

(2) either ☐ [II.2.11.2.1. they have been kept in an approved quarantine establishment for a period of at least 21 days prior to the departure of the consignment

(2) ☐ [and in case of pregnant dams, they have been subjected to a serological test for the detection of antibodies against bovine viral diarrhoea virus with one of the diagnostic methods provided for in Part 6 of Annex I to Delegated Regulation (EU) 2020/688, carried out, with negative results, on samples taken not less than 21 days after commencement of the quarantine]]]

(2) and/or ☐ [II.2.11.2.2. they have been subjected to a serological test for the detection of antibodies against bovine viral diarrhoea virus with one of the diagnostic methods provided for in Part 6 of Annex I to Delegated Regulation (EU) 2020/688, with positive results,

(2) either ☐ [II.2.11.2.2.1. in case of non-pregnant animals, carried out on samples taken prior to departure of the consignment]]]

(2) and/or ☐ [II.2.11.2.2.1. in case of pregnant dams, carried out on samples taken before insemination preceding the current gestation.]]]]

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.

(2) ☐ [II.4. According to official information and as declared by the operator, they are semen donor animals, and

II.4.1. they come from a semen collection centre and will be transported directly to another semen collection centre in accordance with Article 19 of Commission Delegated Regulation (EU) 2020/686; and

(2) either ○ [II.4.2. they were continuously resident since the date of their admission at the semen collection centre and were subjected, with negative results, to all compulsory routine tests referred to in point 2 of Chapter I of Part 1 of Annex II to Delegated Regulation (EU) 2020/686 in the period of the preceding 12 months prior to date of that movement; and]

(2) or ○ [II.4.2. they were subjected, with negative results, to all tests referred to in point 1(b) and (c) of Chapter I of Part 1 of Annex II to Delegated Regulation (EU) 2020/686, required before admission to a semen collection centre carried out during the period immediately preceding quarantine and during the quarantine period; and]

II.4.3. the prior consent of the centre veterinarian of the semen collection centre of destination has been obtained by the operator; and

II.4.4. the means of transport used have been cleansed and disinfected before use.]

II.5. Arrangements are made to transport the consignment in accordance with Article 4 of Delegated Regulation (EU) 2020/688.

Part II: Certification

II. Health information		

II.6. 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.7. 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.]]

(2)either ○ [II.8 Bovine animals from vaccination zone I in relation to emergency protective vaccination against lumpy skin disease, in compliance with Article 13(2) of, and Annex IX, Part 3, point (3.1), to Commission Delegated Regulation (EU) 2023/361.]

(2)or ○ [II.8 Bovine animals from vaccination zone II in relation to emergency protective vaccination against lumpy skin disease, in compliance with Article 13(2) of, and Annex IX, Part 3, point (3.2), to Commission Delegated Regulation (EU) 2023/361.]

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) (4)(5).

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 of the European Parliament and of the Council.

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 38 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

Part II: Certification

II. Health information		

operations.

(4) In the case where a consignment is grouped in an establishment approved for assembly operations and comprises animals that were loaded on different dates, the date which the journey commenced for the whole consignment is considered to be the earliest date when any part of the consignment left the establishment of origin.

(5) This statement does not exempt transporters from their obligation in accordance with Union rules in force in particular regarding the fitness to be transported.

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