

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			
	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		I.16. Transporter		I.17. Accompanying documents			
	Mode	International transport document	Identification		Name Address Approval Number Country		ISO Code	
					[en] accompanying document number		Date of issue	
					Country		Place of issue	
I.18. Transport conditions Ambient <input type="checkbox"/> Frozen <input type="checkbox"/> Chilled <input type="checkbox"/>								
I.19. Container No / Seal No								
I.20. Certified as Event or activity near borders <input type="checkbox"/> Slaughter <input type="checkbox"/> Further keeping <input type="checkbox"/> Travelling circus/animal act <input type="checkbox"/> Confined establishment <input type="checkbox"/> Exhibition <input type="checkbox"/> Other <input type="checkbox"/> Release into the wild <input type="checkbox"/> Quarantine or similar establishment <input type="checkbox"/>								
I.21. For transit through a third country <input type="checkbox"/> Third country Exit point Entry point								
I.22. For transit through Member State(s) <input type="checkbox"/> Member State		ISO Code		I.23. For export <input type="checkbox"/> Third country		ISO Code		Exit point
						BCP code		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								
Commodity		Species		Subcategory		Sex		Identification system
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 <i>Brucella abortus</i> , <i>B. melitensis</i> and <i>B. suis</i> without vaccination regarding bovine animals, and		
(2)	either <input type="checkbox"/> [the establishments of origin are situated in a Member State or zone thereof with the status free from infection with <i>Brucella abortus</i> , <i>B. melitensis</i> and <i>B. suis</i> regarding the bovine population;]		
(2)	and/or <input type="checkbox"/> [they have been subjected to a test for infection with <i>Brucella abortus</i> , <i>B. melitensis</i> and <i>B. suis</i> 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 <input type="checkbox"/> [they are less than 12 months old;]		
(2)	and/or <input type="checkbox"/> [they are castrated.]		
II.2.3.	They come from establishments free from infection with <i>Mycobacterium tuberculosis</i> complex (<i>M. bovis</i> , <i>M. caprae</i> and <i>M. tuberculosis</i>), and		
(2)	either <input type="checkbox"/> [the establishments of origin are situated in a Member State or zone thereof with the status free from infection with <i>Mycobacterium tuberculosis</i> complex (<i>M. bovis</i> , <i>M. caprae</i> and <i>M. tuberculosis</i>);]		
(2)	and/or <input type="checkbox"/> [they have been subjected to a test for infection with <i>Mycobacterium tuberculosis</i> complex (<i>M. bovis</i> , <i>M. caprae</i> and <i>M. tuberculosis</i>) 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 <input type="checkbox"/> [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 (<i>Trypanosoma evansi</i>) has not been reported during the 30 days period prior to departure, and		

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	(2)		either <input type="radio"/> [surra has not been reported in the establishments during the last 2 years prior to their departure.]
	(2)		or <input type="radio"/> [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:
			<ul style="list-style-type: none"> - the infected animals have been removed from the establishments, and - the remaining animals on the establishments have been subjected to a test for surra (<i>Trypanosoma evansi</i>) 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 <input type="checkbox"/> [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 <input type="checkbox"/> [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 <input type="checkbox"/> [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 <input type="checkbox"/> for at least 60 days prior to the date of movement]] [II.2.8.1.1.
	(2)		and/or <input type="checkbox"/> 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)]] [II.2.8.1.2.
	(2)		and/or <input type="checkbox"/> 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);]] [II.2.8.1.3.
	(2)	and/or <input type="checkbox"/> [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 <input type="checkbox"/> for at least 60 days prior to the date of movement]] [II.2.8.2.1.
	(2)		and/or <input type="checkbox"/> 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]] [II.2.8.2.2.
	(2)		and/or <input type="checkbox"/> 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;]] [II.2.8.2.3.
(2)	and/or <input type="checkbox"/> [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 specifications of the vaccine and	

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	(2)		either <input type="checkbox"/>	have been vaccinated more than 60 days before the date of movement]]
	(2)		and/or <input type="checkbox"/>	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 <input type="checkbox"/>	[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 <input type="checkbox"/>	the serological test has been carried out on samples collected at least 60 days before the date of movement.]]]
	(2)		and/or <input type="checkbox"/>	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 <input type="checkbox"/>	[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)	[II.2.8.1.	either <input type="checkbox"/>	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 <input type="checkbox"/>	for at least 60 days prior to the date of movement]]
	(2)		and/or <input type="checkbox"/>	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 <input type="checkbox"/>	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)	[II.2.8.2.	and/or <input type="checkbox"/>	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 <input type="checkbox"/>	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 <input type="checkbox"/>	have been vaccinated more than 60 days before the date of movement]]]
	(2)		and/or <input type="checkbox"/>	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 <input type="checkbox"/>	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

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	(2)	either <input type="checkbox"/> [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 <input type="checkbox"/> [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 <input type="checkbox"/> [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 <input type="checkbox"/> [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 <input type="checkbox"/> [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 <input type="checkbox"/> [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 <input type="checkbox"/> [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 <input type="checkbox"/> [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 <input type="checkbox"/> [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 <input type="checkbox"/> [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 <input type="checkbox"/> [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 <input type="checkbox"/> [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 <input type="checkbox"/> [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 <input type="checkbox"/> [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 <input type="checkbox"/> [II.2.8.3.1.	without any conditions, and	
(2)	and/or <input type="checkbox"/> [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 <input type="checkbox"/> [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	

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	(2)		and/or <input type="checkbox"/>	subject to the conditions referred to in point 7 of Section 1 of Chapter [II.2.8.3.4. 2 of Part II of Annex V to Delegated Regulation (EU) 2020/689, and
	(2)		and/or <input type="checkbox"/>	subject to the conditions referred to in point 8 of Section 1 of Chapter [II.2.8.3.5. 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.]]
	(2)	<input type="checkbox"/>	[(2)either	They are moved to a Member State or zone thereof with the status free from enzootic bovine leukosis, and
	(2)	○ [II.2.9.		either ○ they come from establishments free from enzootic bovine leukosis.]]
	(2)		or ○	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)		<input type="checkbox"/>	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)		<input type="checkbox"/>	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)		<input type="checkbox"/>	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)		<input type="checkbox"/>	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 ○		They are moved to a Member State or zone thereof with an approved eradication programme for enzootic bovine leukosis, and
	(2)	[II.2.9.		either ○ they come from establishments free from enzootic bovine leukosis.]]
	(2)		or ○	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)		<input type="checkbox"/>	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)		<input type="checkbox"/>	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]]]	

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	(2)	and/or <input type="checkbox"/> [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 <input type="checkbox"/> [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)	<input type="checkbox"/> [(2)either o [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 o [II.2.10.1.	they come from establishments free from infectious bovine rhinotracheitis/infectious pustular vulvovaginitis, and
	(2)	either <input type="checkbox"/> [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 <input type="checkbox"/> [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 o [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 o [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 o [II.2.10.1.	they come from establishments free from infectious bovine rhinotracheitis/infectious pustular vulvovaginitis, and
(2)	either <input type="checkbox"/> [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 <input type="checkbox"/> [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]]	

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	(2)	and/or <input type="checkbox"/> [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)	[II.2.10.1.4 .	and/or <input type="checkbox"/> the animals are destined for an establishment which keeps bovine animals for meat production without contact to bovine animals of other establishments, and from which they are directly moved to the slaughterhouse.]]]
	(2)	or <input type="checkbox"/> [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)	<input type="checkbox"/> [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 <input type="checkbox"/> [II.2.11.1.	they come from establishments free from bovine viral diarrhoea, and
	(2)	either <input type="checkbox"/> [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)	[II.2.11.1.2 .	and/or <input type="checkbox"/> 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)	[II.2.11.1.3 .	and/or <input type="checkbox"/> the animals have been tested individually to exclude the presence of bovine viral diarrhoea virus prior to the departure of the consignment.]]]
	(2)	[II.2.11.1.	or <input type="checkbox"/> 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)	[II.2.11.1.1	either <input type="checkbox"/> 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)		<input type="checkbox"/> [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)	[II.2.11.1.2 .	and/or <input type="checkbox"/> 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)	[II.2.11.1.2 .1.	either <input type="checkbox"/> in case of non-pregnant animals, carried out on samples taken prior to departure of the consignment]]]

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	(2)		and/or <input type="checkbox"/> in case of pregnant dams, carried out on samples taken [II.2.11.1.2 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 <input type="checkbox"/> they come from establishments free from bovine viral diarrhoea, and [II.2.11.1.	
	(2)		either <input type="checkbox"/> the establishments of origin are situated in a Member State or zone [II.2.11.1.1 thereof with the status free from bovine viral diarrhoea]]	
	(2)		and/or <input type="checkbox"/> the establishments of origin are situated in a Member State or zone [II.2.11.1.2 thereof with an approved eradication programme for bovine viral diarrhoea]]	
	(2)		and/or <input type="checkbox"/> the establishments of origin have been subjected to a testing regime [II.2.11.1.3 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 <input type="checkbox"/> the animals have been tested individually to exclude the presence of [II.2.11.1.4 bovine viral diarrhoea virus prior to the departure of the consignment]]	
	(2)		and/or <input type="checkbox"/> the animals are destined for an establishment which keeps bovine [II.2.11.1.5 animals for meat production separate from bovine animals of other establishments, and from which they are directly moved to the slaughterhouse]]	
	(2)	and/or <input type="checkbox"/> [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 <input type="checkbox"/> they have been kept in an approved quarantine establishment for a [II.2.11.2.1 period of at least 21 days prior to the departure of the consignment	
	(2)		<input type="checkbox"/> [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 <input type="checkbox"/> they have been subjected to a serological test for the detection of [II.2.11.2.2 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 <input type="checkbox"/> in case of non-pregnant animals, carried out on samples [II.2.11.2.2 taken prior to departure of the consignment]]]	
	(2)		and/or <input type="checkbox"/> in case of pregnant dams, carried out on samples taken [II.2.11.2.2 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) <input type="checkbox"/>	[II.4. II.4.1.	According to official information and as declared by the operator, they are semen donor animals, and 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		

Part II: Certification	II. Health information			
	(2) [II.4.2.]	either <input type="radio"/> 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) [II.4.2.]	or <input type="radio"/> 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.		
	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.] <input type="checkbox"/>	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 <input type="radio"/> [they come from their establishments of origin.]]		
	(2)	or <input type="radio"/> [at least one of the animals of the consignment has undergone one assembly operation on an approved establishment.]]		
(2)	or <input type="radio"/> [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) (4)(5).				

Part II: Certification	II. Health information							
	<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 European Union in this certificate include the United Kingdom in respect of Northern Ireland.</p> <p>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.</p> <p>Part I:</p> <p>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.</p> <p>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.</p> <p>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.</p> <p style="text-align: center;">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.</p> <p>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.</p> <p>Part II:</p> <p>(1) There can be one or more animals in the consignment.</p> <p>(2) Delete if not applicable.</p> <p>(3) Applicable in case the consignment is dispatched from the establishment approved for assembly operations.</p> <p>(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.</p> <p>(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.</p>							
<p>Certifying Officer/Official veterinarian</p> <table border="0" style="width: 100%;"> <tr> <td style="width: 50%;">Name (in capital letters)</td> <td style="width: 50%;">Qualification and title</td> </tr> <tr> <td>Date of signature</td> <td>Signature</td> </tr> <tr> <td>Stamp</td> <td></td> </tr> </table>			Name (in capital letters)	Qualification and title	Date of signature	Signature	Stamp	
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