INTRA

	I.1. Consignor			I.2. IMSOC reference	I.2.a. Local reference		
	Name Address				I.3. Central Competent Authority		
	Country		ISO Code		I.4. Local Competent Authority		
int	I.5. Consignee			I.6. Operator conducting assembly of establishment	l perations independently of an		
ıme	Name			Name			
igr	Address Country		ISO Code	Address			
ons	,			Approval Number			
of consignment				Country	ISO Code		
Part I: Description	I.7. Country of orig	gin	ISO Code	I.9. Country of destination	ISO Code		
rip	I.8. Region of origi	in	Code	I.10. Region of destination	Code		
)esc	I.11. Place of dispa	ntch		I.12. Place of destination			
I: I	Name			Name			
ırt	Address Approval			Address			
Pē	Number			Approval Number			
	Country		ISO Code	Country	ISO Code		
	I.13. Place of loadi	ing		I.14. Date and time of departure			
	Name						
	Address Approval						
	Number		700 O 1				
	Country		ISO Code				
	I.15. Means of Tra	T .	7.3	I.16. Transporter			
	Mode	International transport	Identification	Name Address			
	document			Approval Number			
					ISO Code		
				Country	150 Code		
				I.17. Accompanying documents Document Type			
				Accompanying document			
				reference Date of Issue			
				Date of Issue Country			
				Place of issue			
	I.18. Transport cor	nditions					
	Ambient \square		Chilled \square	Frozen \square	Frozen Ll		
	I.19. Container Nu	mber / Seal Numbe	er				
	I.20. Certified as						
	Other \square		Further keeping \square	Exhibition	Confined establishment \square		
	Release into the wild \square Travelling circus/animal act \square			Breeding	Quarantine or similar establishment \square		
	Event or activity n	near borders \square					
	L21. For transit through a third country						
	Third country			ISO Code			
	Exit point Entry point			BCP code BCP code			
	I.22. For transit through Member State(s)			I.23. For export			
	Member State	0	ISO Code	Third country	ISO Code		
	MICHINEL STATE		150 code	Third country ISO Code Exit point BCP code			
	I.24. Estimated jou	ırney time		I.25. Journey Log			
	I.27. Total quantity	у		I.28. Total gross weight			
	I.30. Description o	f consignment					
	1. 01 LIVE ANIMA	LS					

en 1/11

	0102 Live bovine animals		-	
	#1. Commodity	Subcategory	Sex	Identification system
	Species	Identification Number	Age	Quantity
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Part I: Description of consignment				
Paı				

en 2 / 11

	II. Health infor	rmation								
	I, the under	rsigned offic	cial veterina	arian, hereby certify tha	L t:					
	II.1.	•		of the consignment des		in Part I meet	the follo	owing 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 30 days prior to the date of departure of the consignment, or since birth, if they are younger than 30 days of age,								
cati		have been continuously resident in the establishment of origin;								
Part II: Certification		have not been in contact with kept bovine animals of a lower health status or subject to movement restrictions for animal health reasons;								
rt II: (II.1.2.3.	have not been in direct or indirect contact with kept animals that have entered the Union from a third country or territory during the last 30 days prior to the date of departure of the animals.								
Paı	II.1.3.	-	on which wa	clinical signs or symptor as carried out, within the (insert date dd)	e last 24	hours prior to		e animals during the clinical ne of departure of the		
	II.2.	According requireme		formation, the animals	describ	ed in Part I me	eet the f	ollowing health		
	II.2.1.	(2)	(2) • either [They come from establishments or zones not subject to movement restrictions affecting bovine animals and established for reasons of diseases listed for those species or diseases subject to emergency measures relevant for those species, and they have not been in contact with kept animals of a listed species of a lower health status for an adequate period.]							
	(2)	o or	animals an	e from establishments o d established for granted, and:	r zones	•		restrictions affecting bovine rom movement restrictions		
		□ (2)	[they comp	oly with the requiremen	ts set o	ut in	(4)]]		
		□ (2)	[and in par	ticular, they are		(5).]]				
	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 last 30 days prior to the date of departure, and in the case of post-parturient females taken at least 30 days after parturition;]								
	(2) □ and/or	[they are le	ess than 12 r	nonths old;]						
	(2) □ and/or	[they are ca	astrated.]							
	II.2.3.	-		lishments free from infe f. tuberculosis), and:	ection w	vith Mycobacte	rium tu	berculosis complex (M.		
	(2) □ either			origin are situated in a l cterium tuberculosis co				f with the status free from nd M. tuberculosis);]		
	(2) □ and/or	caprae and Delegated	l M. tubercu Regulation (losis) with one of the dia	agnosti	c methods pro	vided fo	culosis complex (M. bovis, M. or in Part 2 of Annex I to ng the last 30 days prior to		
	(2) □ and/or	[they are le	ess than 6 w	eeks old.]						
	II.2.4.	-					_	t terrestrial animals has not onsignment.		
	II.2.5.	been reported during the last 30 days prior to the date of departure of the consignment. I.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:								

en 3/11

	II. Health info	ormation							
	(2)		een reported in kept animals of liste	d species for that disease dur	 ing the last 2 years prior to				
	either		departure of the consignment;]	anion for that diagona during	the last 2 years prior to the				
ion	(2) □ and/or	date of dep epizootic h	[has been reported in kept animals of listed species for that disease during the last 2 years prior to the date of departure of the consignment and the animals have been kept in a zone seasonally free from epizootic haemorrhagic disease in accordance with Parts 1 and 2 of Annex IX to Delegated Regulation (EU) 2020/688:						
Part II: Certification		(2) □ either							
		(2) □ and/or							
Par		(2) □ and/or							
	(2) □ and/or	date of dep	the last 2 years prior to the against attacks by vectors rotected against attacks by rided for in Part 3 of Annex						
	-	(2) □ either	for at least 60 days prior to the dat	e of departure of the consigni	ment;]				
		for at least 28 days prior to the date of departure of the consignment and have been subjected to a serological test, with negative results, carried out on samples collected at le 28 days following the date of the commencement of the period of protection against attact by vectors;]							
		(2) □ and/or	for at least 14 days prior to the dat subjected to a PCR test, with negati days following the date of commer vectors;]	ive results, carried out on sam	nples collected at least 14				
	(2) □ and/or	date of dep		species for that disease during the last 2 years prior to the e Member State of destination has informed the at such movement is authorised].					
	II.2.6.	•	e from establishments in which anth to the date of departure of the cons	hrax in ungulates has not been reported during the last 15 signment.					
	II.2.7.		e from establishments in which surr s prior to the date of departure of th	a (Trypanosoma evansi) has not been reported during the ne consignment, and:					
	(2) o either	[surra has departure.	not been reported in the establishm]	nents during the last 2 years p	rior to the date of their				
	(2) or	following to restriction and the rediagnostic with negative	been reported during the last 2 year the date of the last outbreak, the affe s until the date on which the infecte maining animals in the establishme methods provided for in Part 3 of A rive results, on samples taken at leas ave been removed from the establish	ected establishments have rer od animals have been remove nts have been subjected to a t nnex I to Delegated Regulatio of 6 months following the date	mained under movement d from the establishments, est for surra with one of the in (EU) 2020/688, carried out,				
	(2) □ either	[II.2.8.	They originate from a Member Stavirus (serotypes 1-24), where no cabeen confirmed in the targeted and date of departure of the consignment against infection with bluetongue of departure of the consignment at (b) or (c), or Article 32(2) of Delega	use of infection with bluetong imal population during the la ent, and have not been vaccin virus (serotypes 1-24) in the la nd the requirements laid dow	ue virus (serotypes 1-24) has st 24 months prior to the lated with a live vaccine ast 60 days prior to the date on in Article 32(1), points (a),				
	(2) □ and/or	[II.2.8.	They originate from a Member Sta programme for infection with blue						

en 4/11

					2021/1011 (2021/103) Model Bo V MVIRTA					
		II. Health info	rmation							
					down in Article 32(1), points (a), (b) or (c), or Article 32(2) of Delegated Regulation (EU) 2020/688 are fulfilled, and they:					
	u		(2) □ either	[II.2.8.1.	have been kept in a Member State or zone thereof seasonally free from infection with bluetongue virus (serotypes 1-24) in accordance with Article 40(3) of Commission Delegated Regulation (EU) 2020/689:					
Part II: Certification	catio	(2) □ either	[II.2.8.1.1.	for at least	60 days prior to the date of depature of the consignment;]]					
	t II: Certifi	(2) □ and/or	[II.2.8.1.2.	subjected to 28 days fol	28 days prior to the date of departure of the consignment and have been to a serological test, with negative results, carried out on samples collected at least llowing the date of entry of the animals into the Member State or zone thereof free from infection with bluetongue virus (serotypes 1-24);]]					
	Par	(2) □ and/or	[II.2.8.1.3.	for at least 14 days prior to the date of departure of the consignment and have been subjected to a PCR test, with negative results, carried out on samples collected at least 14 days following the date of entry of the animals into the Member State or zone thereof seasonally free from infection with bluetongue virus (serotypes 1-24);]]]						
			(2) □ and/or	[II.2.8.2.	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:					
L		(2) □ either	[II.2.8.2.1.	for at least	60 days prior to the date of departure of the consignment;]]					
		(2) □ and/or	[II.2.8.2.2.	subjected	28 days prior to the date of departure of the consignment and have been to a serological test, with negative results, carried out on samples collected at least llowing the date of the commencement of the period of protection against attacks [3]					
		(2) □ and/or	[II.2.8.2.3.	for at least 14 days prior to the date of departure of the consignment and have been subjected to a PCR test, with negative results, carried out on samples collected at least 14 days following the date of the commencement of the period of protection against attacks l vectors;]]]						
			(2) □ and/or	[II.2.8.3.	have been vaccinated against all serotypes from 1 to 24 of infection with bluetongue virus which were reported in that Member State or zone thereof during the last 2 years prior to the date of departure of the consignment and are within the immunity period guaranteed in the specifications of the vaccine and:					
		(2) □ either	[II.2.8.3.1.	have been	vaccinated more than 60 days prior to the date of departure of the consignment;]]					
		(2) □ and/or	[II.2.8.3.2.	results, car	vaccinated with an inactivated vaccine and subjected to a PCR test, with negative rried out on samples collected at least 14 days after the date of the onset of the 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 in that Member State or zone thereof during the last 2 years prior to the date of departure of the consignment, and:					
		(2) □ either	[II.2.8.4.1.		gical test has been carried out on samples collected at least 60 days prior to the parture of the consignment.]]]					
		and/or date of departure of the consigni		date of dep	gical test has been carried out on samples collected at least 30 days prior to the parture of the consignment and the animals have been subjected to a PCR test, rive results, carried out on samples collected not earlier than 14 days prior to the parture of the consignment.]]]					
		(2) □ and/or [II.2.8.	(serotypes (serotypes	1-24) nor co 1-24) and th	Member State or a zone thereof neither free from infection with bluetongue virus overed by the eradication programme for infection with bluetongue virus ne requirements laid down in Article 32(1), points (a), (b) or (c), or Article 32(2) of (EU) 2020/688 are fulfilled, and they:					
		(2) [II.2.8.1. have been protected a		[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					

en 5 / 11

(2) [eithe (2) [and/ (2) [eithe (2) [eithe	er □ /or	[II.2.8.1.1. [II.2.8.1.2.	for at least subjected t 28 days fol	28 days prior to the date	e of departure of the consigni	ment;]]				
eithe (2) [and/ (2) [and/ (2) [(2) [er □ /or	[II.2.8.1.2.	for at least subjected t 28 days fol	60 days prior to the date 28 days prior to the date	e of departure of the consigni	ment;]]				
eithe (2) [and/ (2) [and/ (2) [(2) [er □ /or	[II.2.8.1.2.	for at least subjected t 28 days fol	60 days prior to the date 28 days prior to the date	e of departure of the consigni	ment;]]				
hand/ (2) [and/ and/ (2) [(2) [/or □		subjected t 28 days fol							
(2) [[II.2.8.1.3.	by vectors;	lowing the date of the co	for at least 28 days prior to the date of departure of the consignment and have been subjected to a serological test, with negative results, carried out on samples collected at least 28 days following the date of the commencement of the period of protection against attacks by vectors;]]					
(2) [subjected t	o a PCR test, with negati	e of departure of the consignive results, carried out on samencement of the period of p	nples collected at least 14				
		(2) □ and/or	[II.2.8.2.	i.8.2. have been kept for the last 60 days prior to the date of departure of the consignment in an establishment situated in a Member State or within an a of at least 150 km radius centred on the establishment, where surveillance compliance with the requirements set out in Part II, Chapter 1, Sections 1 at 2,of Annex V to Delegated Regulation (EU) 2020/689 has been carried out du that period, and:						
		[II.2.8.2.1.	against all serotypes from 1 to rted during the last 2 years po of at least 150km radius cent the immunity period guarant	rior to the date of departure tred on the place where the						
		(2) □ either	[II.2.8.2.1. 1.	have been vaccinated n consignment;]]]	more than 60 days prior to the	e date of departure of the				
		(2) □ and/or	[II.2.8.2.1. 2.	with negative results or	with an inactivated vaccine an n samples collected at least 14 the specifications of the vacci	4 days after the date of onset				
(2) [and/	☐ [II.2.8.2.2. the animals have been immunised			e virus which were repor	rted during the last 2 years p	rior to the date of departure				
		(2) □ either	[II.2.8.2.2. 1.		subjected with positive resuled at least 60 days prior to the	ts to a serological test carried e date of departure of the				
		(2) □ and/or	[II.2.8.2.2. 2.	out on samples collecte consignment and to a P	subjected with positive resulted at least 30 days prior to the PCR test, with negative results an 14 days prior to the date of	s, carried out on samples				
(2) [or[II		Delegated I	Regulation ((EU) 2020/689 and the co	in Part II, Chapter 2, Section 1 mpetent authority of the Mer ther Member State or zone th	mber State of origin				
		(2) □ either	[II.2.8.1.	the Member State of de Member States that suc	m infection with bluetongue estination has informed the Coch movement is authorised su 3(2), points (a), (b) and (c), of 1	ommission and the other abject to the conditions				
(2) [eithe		[II.2.8.1.1.	Part II, Cha	pter 2, Section 1, point 5	5, of Annex V to that Delegate	d Regulation, and				
(2) [and/		[II.2.8.1.2.	Part II, Cha	pter 2, Section 1, point 6	6, of Annex V to that Delegate	d Regulation, and				
(2) [and/										

en 6/11

		_			2021/1011 (2021/100) Model BOV 1111141 A			
		II. Health info	rmation					
		(2) □ and/or	[II.2.8.1.4.	Part II, Cha	apter 2, Section 1, point 8, of Annex V to that Delegated Regulation, and			
					ticle 32(1), points (a), (b) or (c), or Article 32(2) of Delegated Regulation (EU) id down in Article 33 of that Delegated Regulation are fulfilled;]]			
	Part II: Certification		(2) □ and/or	[II.2.8.2.	with an approved eradication programme for infection with bluetongue virus (serotypes 1-24) and the Member State of destination has informed the Commission and the other Member States that such movement is authorised subject to the conditions referred to in Article 43(2), points (a), (b) and (c), of Delegated Regulation (EU) 2020/689, and:			
	t II: C	(2) □ either	[II.2.8.2.1.	Part II, Cha	apter 2, Section 1, point 5, of Annex V to that Delegated Regulation, and			
	Par	(2) □ and/or	[II.2.8.2.2.	Part II, Cha	apter 2, Section 1, point 6, of Annex V to that Delegated Regulation, and			
		(2) □ and/or	[II.2.8.2.3.	Part II, Cha	apter 2, Section 1, point 7, of Annex V to that Delegated Regulation, and			
		(2) □ and/or	[II.2.8.2.4.	Part II, Cha	apter 2, Section 1, point 8, of Annex V to that Delegated Regulation, and			
					ticle 32(1), point (a), (b) or (c), or Article 32(2) of Delegated Regulation (EU) id 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 ar	without any conditions, and:			
	(2) [II.2.8.3.2. subject to the conditions referred and/or Delegated Regulation (EU) 2020/6			the conditions referred to in Part II, Chapter 2, Section 1, point 5, of Annex V to Regulation (EU) 2020/689, and				
		(2) □ and/or	[II.2.8.3.3.		the conditions referred to in Part II, Chapter 2, Section 1, point 6, of Annex V to Regulation (EU) 2020/689, and			
		(2) □ and/or	[II.2.8.3.4.	•	the conditions referred to in Part II, Chapter 2, Section 1, point 7, of Annex V to Regulation (EU) 2020/689, and			
		(2) □ and/or	[II.2.8.3.5.		the conditions referred to in Part II, Chapter 2, Section 1, point 8, of Annex V to Regulation (EU) 2020/689, and			
		_			ticle 32(1), points (a), (b) or (c), or Article 32(2) of Delegated Regulation (EU) id down in Article 33 of that Delegated Regulation are fulfilled.]]]			
		(2) [(2) o either	[II.2.9.	They are n leukosis, a	noved to a Member State or zone thereof with the status free from enzootic bovine nd:			
			(2) o 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 last 24 months prior to the date of departure of the consignment, and:			
		either enzootic bovine leukosis with one			ver 24 months of age and they have been subjected to a serological test for ovine leukosis with one of the diagnostic methods provided for in Part 4 of Annex ted 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 4 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 last 30 days prior to the date of 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			

en 7/11

01	NON				2024/1044 (2021/	403) Model BOV-INTRA-A
	II. Health info	rmation				
				-	occasions at an interval of not r to the date of departure of t	•
Part II: Certification	(2) □ and/or	[II.2.9.1.2.	they are less than 24 months of age and they were born to dams 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 4 months during the last 12 months prior to the date of departure of the consignment.]]]			
ertific	(2) or	[II.2.9.	-	noved to a Member State le for enzootic bovine le	or zone thereof with an apparate or zone thereof with an apparate or zone.	roved eradication
t II: C		(2) o either	[II.2.9.1.	they come from establi	shments free from enzootic b	oovine leukosis.]]
		(2) or	[II.2.9.1.	enzootic bovine leukos	shments not free from enzoo is has not been reported in th r to the date of departure the	ose establishments during
	(2) 🗆 either	[II.2.9.1.1.	enzootic b	ovine leukosis with one	d they have been subjected to of the diagnostic methods pro 0/688, carried out with negativ	ovided for in Part 4 of Annex
		(2) □ either	[II.2.9.1.1. 1.		vo occasions at an interval of her bovine animals of the est	
		(2) □ and/or	[II.2.9.1.1. 2.	consignment, and all be establishment have bee leukosis with one of the Delegated Regulation (I samples taken on two of	ng the last 30 days prior to the ovine animals over 24 monthen subjected to a serological to diagnostic methods provide EU) 2020/688, carried out, with occasions at an interval of not reached.	s of age kept in the est for enzootic bovine d for in Part 4 of Annex I to h negative results, on t less than 4 months during
	(2) □ and/or	[II.2.9.1.2.	test for ena Annex I to samples ta	zootic bovine leukosis w Delegated Regulation (E ken on two occasions at	e and they were born to dams ith one of the diagnostic meth U) 2020/688, carried out, with an interval of not less than 4 are of the consignment.]]]]	nods provided for in Part 4 of n negative results, on
	(2) [(2) o either	[II.2.10.	bovine rhi	notracheitis/infectious p	or zone thereof with the stat ustular vulvovaginitis and th cheitis/infectious pustular vul	ey have not been vaccinated
		(2) o either	[II.2.10.1.		shments free from infectious ous pustular vulvovaginitis, a	
	(2) □ either	[II.2.10.1.1			uated in a Member State or z racheitis/infectious pustular v	
	(2) □ and/or	[II.2.10.1.2	departure of antibod methods p negative re	of the consignment and ies against whole bovine rovided for in Part 5 of A	o quarantine for at least 30 da have been subjected to a serce herpes virus-1 (BoHV-1) with Annex I to Delegated Regulati mple taken during the last 15	ological test for the detection n one of the diagnostic on (EU) 2020/688, with a
		(2) or	[II.2.10.1.	rhinotracheitis/infection approved quarantine edeparture of the consignation of antibour methods provided for it with a negative result,	shments not free from infections pustular vulvovaginitis ar stablishment for at least 30 d gament and have been subjectives against whole BoHV-1, we need to Delegate carried out on a sample taken nent of the quarantine.]]	nd they have been kept in an ays prior to the date of ted to a serological test for ith one of the diagnostic ted Regulation (EU) 2020/688,
	(2) ∘ or	[II.2.10.			or zone thereof with an apparation of the or zone thereof with an apparation of the or zone thereof with an apparation of zone control of the or zone thereof with an apparation of zone the zone zone the zone zone the zone zone zone zone zone zone zone zon	

en 8/11

_) I N	ION		2024/1044 (2021/403) Woder DOV-INTRA-2					
		II. Health info	rmation						
			(2) o 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]]]]					
	cation	(2) □ and/or	[II.2.10.1.2	e establishments of origin are situated in a Member State or zone thereof with an proved eradication programme for infectious bovine rhinotracheitis/infectious pustula lvovaginitis]]]]					
art II. Cartif		(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 last 15 days prior to the date of departure of the consignment]]]					
•		(2) □ and/or	[II.2.10.1.4	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	II.2.10.1. they come from establishments not free from infectious bovine rhinotracheitis/infectious pustular vulvovaginitis, and:					
		_	-	een kept in an approved quarantine establishment for at least 30 days prior to the date of f the consignment, 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 the date of commencement of the quarantine.]]]						
		(2) [(2) 0 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) o [II.2.11.1. they come from estate either		II.2.11.1. they come from establishments free from bovine viral diarrhoea, and:					
	- 1	(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 have been subjected to a testing regime as referred in Part VI, Chapter 1, Section 2, point 1(c)(ii) or (iii), of Annex IV to Delegated Regulation (EU) 2020/689, carried out, with negative results, within the 4 months period prior to the date of 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 date of 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 date of departure of the consignment, [and in the 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 ess than 21 days after the date of commencement of the quarantine;] (2)]]					
		(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 in the case of non-pregnant dams, carried out on samples taken prior to the date of departure of the consignment]]]]					
			(2) □ and/or	II.2.11.1.2 in the case of pregnant dams, carried out on samples taken before the date of insemination preceding the current gestation.]]]]					
		(2) or	[II.2.11.	They are moved to a Member State or zone thereof with an approved eradication					

en 9 / 11

	II. Health info	ormation			2021/1011 (2021/				
			programme	e for bovine viral diarrh	noea, and:				
		(2) □ either	[II.2.11.1.	they come from establi	shments free from bovine vir	ral diarrhoea, and:			
l ä	(2) \square either	[II.2.11.1.1 ·		hments of origin are sit ovine viral diarrhoea;]]	tuated in a Member State or z]	one thereof with the status			
fication	(2) □ and/or	[II.2.11.1.2		the establishments of origin are situated in a Member State or zone thereof with an approved eradication programme for bovine viral diarrhoea;]]]					
Part II: Certification	(2) □ and/or	[II.2.11.1.3	Chapter 1, S carried out,	ne establishments of origin have been subjected to a testing regime as referred in Part VI, hapter 1, Section 2, point 1(c)(ii) or (iii), of Annex IV to Delegated Regulation (EU) 2020/689 arried out, with negative results, within the last 4 months prior to the date of departure of the consignment;]]]					
Pa	(2) □ and/or	[II.2.11.1.4		he animals have been tested individually to exclude the presence of bovine viral diarrhoea rirus prior to the date of departure of the consignment;]]]					
	(2) □ and/or	[II.2.11.1.5 ·	production	he animals are destined for an establishment which keeps bovine animals for meat roduction separate from bovine animals of other establishments, and from which they are irectly moved to the slaughterhouse;]]]					
		(2) □ and/or		have been subjected to with one of the diagnos	shments not free from boving a test for bovine viral diarrh stic methods provided for in I EU) 2020/688, carried out with	oea virus antigen or genome Part 6 of Annex I to			
	(2) □ either	[II.2.11.2.1	days prior t have been s diarrhoea v Delegated R	they have been kept in an approved quarantine establishment for a period of at least 21 days prior to the date of departure of the consignment, [and in case of pregnant dams, the have been subjected to a serological test for the detection of antibodies against bovine vira 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 n less than 21 days after the date of commencement of the quarantine;] (2)]]					
	(2) 🗆 and/or	[II.2.11.2.2	viral diarrh	-	logical test for the detection o he diagnostic methods provid 8, with positive results,	_			
		(2) □ either		in the case of non-preg of departure of the con	nant dams, carried out on sansignment]]]]]	mples taken prior to the date			
		(2) □ and/or		1 0	dams, carried out on sample g the current gestation.]]]]]	s taken before the date of			
	II.3.				by the operator, the animals c th an undetermined cause.	ome from establishments			
	II.4.		ents are mad (EU) 2020/68		gnment in accordance with A	article 4 of Delegated			
 II.5. This animal health certificate is valid for 10 days from the date of issuing. In the case of tran waterway/sea of animals, the period of validity of the certificate may be extended by the durgiourney by waterway/sea. (2)(6) Since the date of depature from their establishments of origin and prior to the date of arrivations. [II.6. establishment approved for assembly operations, none of the animals of the consignment has undergone more than two assembly operations, and: (2) o [they come from their establishments of origin.]] either 									
					ons, none of the animals of the				
(2) ○ or [at least one of the anir establishment.]]				nals of the consignmen	t has undergone one assembl	y operation in an approved			
	(2) ∘ or	[at least on establishm		nals of the consignmen	t has undergone two assembl	y operations in the approved			
	Animal we	elfare attesta							
	At the time of inspection, the animals covered by this animal health certificate were fit to be transported in accordance with the provisions of Council Regulation (EC) No 1/2005 on the intended journey due to start on								

en 10/11

	IROPEAN NION		2024/1044 (2021/	403) Model BOV-INTRA-X					
	II. Health info	rmation							
		(incent data) (7) (0)							
	Notes:	(insert date) (7) (8).							
		nce with the Agreement on the withdrawal of th	ne United Kingdom of Great B	ritain and Northern Ireland					
		uropean Union and the European Atomic Energ	_						
ation	Protocol on Ireland/Northern Ireland in conjunction with Annex 2 to that Protocol, references to the Union in this animal health certificate include the United Kingdom in respect of Northern Ireland.								
Certification	provided fo	This animal health certificate shall be completed in accordance with the notes for the completion of certificates provided for in Chapter 2 of Annex I to Commission Implementing Regulation (EU) 2020/2235.							
<u>.</u>	Part I:	Part I:							
Part II	Box reference I.11:	"Place of dispatch": Indicate an establishment establishment approved for assembly operatio (EU) 2016/429 of the European Parliament and	ns in accordance with Article	_					
	Box reference I.12:	"Place of destination": Indicate an establishment approved for assembly operation (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),							
		In case the animals are dispatched in the Member State of passage, the which the animal health certificate approved for assembly operations,	e reference number(s) of the e e for this consignment is issue	certificate(s), based on					
	Box reference I.30:	"Identification number": Indicate identification accordance with Article 38 of Delegated Regula		consignment identified in					
	Part II:								
	(1)	There may be one or more animals in the cons	ignment.						
	(2)	Delete if not applicable.							
	(3)	Insert the name of the disease(s).							
	(4)	Commission providing for those requirements	title, and number of the relevant legal act(s) adopted by the ats.						
	(5)	Insert the specific attestation(s) provided for in Commission, as referred to in Article 126(1), po							
	(6)	Applicable in case the consignment is dispatch operations.	ed from the establishment ap	pproved for assembly					
	(7)	In the case where a consignment is grouped in comprises animals that were loaded on differe whole consignment is considered to be the ear establishment of origin.	nt dates, the date which the j	ourney commenced for the					
	(8)	This statement does not exempt transporters force in particular regarding the fitness to be t		dance with Union rules in					
		cer/Official veterinarian	Qualification and title						
	Name (in capi Date	tal letters)	Qualification and title Signature						
	Stamp								

en **11** / 11