ANIMAL HEALTH CERTIFICATE FOR THE MOVEMENT BETWEEN MEMBER STATES OF WILD TERRESTRIAL ANIMALS (MODEL 'WILD-ANIMALS-INTRA')

EUROPEAN UNION					INTRA
Part I: Description of consignment	I.1 **Consignor** Name Address Country ISO country code		I.2 **IMSOC reference** I.2a **Local reference** I.3 **Central Competent Authority** I.4 **Local Competent Authority**		**QR CODE**
	I.5 **Consignee** Name Address Country ISO country code		I.6 **Operator conducting assembly operations independently of an establishment** Name Registration No Address Country ISO country code		
	I.7 **Country of origin** ISO country code		I.9 **Country of destination** ISO country code		
	I.8 **Region of origin** Code		I.10 **Region of destination** Code		
	I.11 **Place of dispatch** Name Registration/Approval No Address Country ISO country code		I.12 **Place of destination** Name Registration/Approval No Address Country ISO country code		
	I.13 **Place of loading**		I.14 **Date and time of departure**		
	I.15 **Means of transport** □ Vessel □ Aircraft □ Railway □ Road vehicle Identification □ Other Document		I.16 **Transporter** Name Registration/Authorisation No Address Country ISO country code I.17 **Accompanying documents** Type Code Country ISO country code Commercial document reference		
	I.18 **Transport conditions** □ Ambient		□ Chilled	□ Frozen	
	I.19 **Container number/Seal number** Container No Seal No				

I.20	**Certified as or for**		
□ Further keeping	□ Slaughter	□ Confined establishment	□ Germinal products
□ Registered equine animal	□ Travelling circus/animal act	□ Exhibition	□ Event or activity near borders
□ Release into the wild	□ Dispatch centre	□ Relaying area/purification centre	□ Ornamental aquaculture establishment
□ Further processing	□ Organic fertilizers and soil improvers	□ Technical use	□ Quarantine or similar establishment
□ Products for human consumption	□ Pollination	□ Live aquatic animals for human consumption	□ Other

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

I.22	**□ For transit through Member State(s)**		**I.23**	**□ For export**	
	Member State	ISO country code		Third country	ISO country code
	Member State	ISO country code		Exit point	BCP code
	Member State	ISO country code			

I.24	**Estimated journey time**	**I.25**	**Journey log**	□ yes	□ no
I.26	**Total number of packages**	**I.27**	**Total quantity**		
I.28	**Total net weight/gross weight (kg)**	**I.29**	**Total space foreseen for the consignment**		

I.30 Description of consignment

CN code	Species	Subspecies/Category	Sex	Identification system	Identification number	Age	Quantity Type
Region of origin		Cold store		Identification mark	Type of packaging		Net weight
Slaughterhouse		Treatment type		Nature of commodity	Number of packages		Batch No
		Date of collection/production		Manufacturing plant	Approval or registration number of plant/establishment/centre	Test	

Part II: Certification

II. Health information

II.a Certificate reference	II.b IMSOC reference

I, the undersigned official veterinarian, hereby certify that:

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

II.1.1. The majority of the animals of the consignment, 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.1.1.1. have been resident in the habitat of origin;

II.1.1.2. have not been in contact with kept animals of a lower health status or subject to movement restrictions for animal health reasons;

II.1.1.3. 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 animal.

II.1.2. They have not shown clinical signs or symptoms of listed diseases for animals of the species concerned or emerging diseases during the clinical examination, or where this is not possible, the clinical inspection, 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 wild terrestrial animals described in Part I do not come from a habitat subject to movement restrictions or situated in a restricted zone established for reasons of listed diseases for animals of the species concerned.

(2)[II.3. According to official information, the wild terrestrial animals described in Part I are ungulates and meet the following health requirements:

(2)[II.3.1. They come from a habitat in which infection with *Brucella abortus, B. melitensis* and *B. suis* in wild terrestrial animals of listed species for that disease has not been reported during the last 42 days prior to departure.]

(2)[II.3.2. They come from a habitat in which infection with *Mycobacterium tuberculosis* complex *(M. bovis, M. caprae* and *M. tuberculosis)* in wild terrestrial animals of listed species for that disease has not been reported during the last 42 days prior to departure.]

(2)[II.3.3. They come from a habitat in which infection with rabies virus has not been reported during the 30 day period prior to departure.]

(2)[II.3.4. They come from a habitat in which infection with epizootic haemorrhagic disease virus within a radius of 150 km has not been reported in wild terrestrial animals of listed species of that disease during the last 2 years prior to departure.]

(2)[II.3.5. They come from a habitat in which anthrax in ungulates has not been reported during the 15 day period prior to departure.]

(2)[II.3.6. They come from a habitat in which surra (*Trypanosoma evansi*) has not been reported during the last 30 days prior to departure.]]

[(2)][II.4. According to official information the wild terrestrial animals described in Part I belong to the families Antilocapridae, Bovidae, Camelidae, Cervidae, Giraffidae, Moschidae or Tragulidae and meet the following health requirements:

(2) *either*[II.4.1. They originate from a habitat in 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 Commission Delegated Regulation (EU) 2020/688 are fulfilled.]

(2)*and/or*[II.4.2. They originate from a habitat in 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.4.2.1. have been resident 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.4.2.1.1. for at least 60 days prior to the date of movement]]

(2)*and/or* [II.4.2.1.2. 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.4.2.1.3. 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.4.2.2. have been vaccinated against those serotypes from 1 to 24 of infection with bluetongue virus which were reported during the past 2 years in that Member State or zone and are within the immunity period guaranteed in the specifications of the vaccine and

(2)*either* [II.4.2.2.1. have been vaccinated more than 60 days before the date of movement]]

(2)*and/or* [II.4.2.2.2. 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.4.2.3.		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.4.2.3.1. the serological test has been carried out on samples collected at least 60 days before the date of movement]]
			(2)*and/or*	[II.4.2.3.2. 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.4.3.	They originate from a habitat in 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		
			[II.4.3.1.	have been resident at least for the 60 day period prior to departure in a habitat situated in a Member State or in an area of at least 150 km radius centred on the habitat, where surveillance in compliance with the requirements laid down in Sections 1 and 2 of Chapter 1 of Part II of Annex V to Regulation (EU) 2020/689 has been carried out during that period and
			(2)*either*	[II.4.3.1.1. 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 habitat where the animals were resident and are within the immunity period guaranteed in the specifications of the vaccine and
				(2)*either* [II.4.3.1.1.1. have been vaccinated more than 60 days before the date of movement]]]
				(2)*and/or* [II.4.3.1.1.2. 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.4.3.1.2. 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 habitat where the animals were resident, and
				(2)*either* [II.4.3.1.2.1. 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)*or*	[II.4.3.1.2.2.	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.4.4.	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.4.4.1.	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.4.4.1.1.	point 5 of Section 1 of Chapter 2 of Part II of Annex V to that Regulation, and
		(2)*and/or*	[II.4.4.1.2.	point 6 of Section 1 of Chapter 2 of Part II of Annex V to that Regulation, and
		(2)*and/or*	[II.4.4.1.3.	point 7 of Section 1 of Chapter 2 of Part II of Annex V to that Regulation, and
		(2)*and/or*	[II.4.4.1.4.	point 8 of Section 1 of Chapter 2 of Part II of Annex V to that 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.4.4.2.	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)(a), (b) and (c) of Delegated Regulation (EU) 2020/689 and	
		(2)*either*	[II.4.4.2.1.	point 5 of Section 1 of Chapter 2 of Part II of Annex V to that Regulation, and
		(2)*and/or*	[II.4.4.2.2.	point 6 of Section 1 of Chapter 2 of Part II of Annex V to that Regulation, and
		(2)*and/or*	[II.4.4.2.3.	point 7 of Section 1 of Chapter 2 of Part II of Annex V to that Regulation, and
		(2)*and/or*	[II.4.4.2.4.	point 8 of Section 1 of Chapter 2 of Part II of Annex V to that 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.4.4.3. 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.4.4.3.1. without any conditions, and (2)*and/or* [II.4.4.3.2. 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.4.4.3.3. 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 (2)*and/or* [II.4.4.3.4. 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.4.4.3.5. 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.]]] II.5. To the best of my knowledge and as declared by the operator, the wild terrestrial animals come from a habitat where there were no abnormal mortalities with an undetermined cause. II.6. Arrangements are made to transport the consignment in accordance with Article 101(1), (2) and (3) of Delegated Regulation (EU) 2020/688. II.7. 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. **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)*. **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 a habitat of the origin of the animals in the consignment.

Box reference I.12: "*Place of destination":* Indicate a habitat or an establishment of the final destination of the consignment.

Box reference I.30: "*Identification number*": Indicate identification codes of the animals in the consignment.

Part II:

(1) There can be one or more animals in the consignment.

(2) Delete if not applicable.

Official veterinarian

Name (in capital letters) Qualification and title

Local Control Unit name Local Control Unit code

Date

Stamp Signature