ANIMAL HEALTH CERTIFICATE FOR THE MOVEMENT BETWEEN MEMBER STATES OF TERRESTRIAL ANIMALS BETWEEN CONFINED ESTABLISHMENTS (MODEL 'CONFINED-LIVE-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 Address Country	Registration No 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 Address Country	Registration/Approval No ISO country code	**I.12**	**Place of destination** Name Address Country	Registration/Approval No ISO country code
	I.13	**Place of loading**		**I.14**	**Date and time of departure**	
	I.15	**Means of transport** □ Vessel □ Railway Identification Document	□ Aircraft □ Road vehicle □ Other	**I.16**	**Transporter** Name Address Country **I.17 Accompanying documents** Type Country Commercial document reference	Registration/Authorisation No ISO country code Code ISO country code
	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] in the consignment described in Part I meet the following requirements:

II.1.1. Their confined establishment of dispatch is approved in accordance with Articles 97 and 99 of Regulation (EU) 2016/429.

II.1.2. They have not shown clinical signs or symptoms of diseases, in particular relevant diseases listed in Annex of Commission Implementing Regulation (EU) 2018/1882, during the clinical examination, or where this is not possible, a clinical inspection, which was carried out within the 48 hour period prior to departure of the consignment, on ……. (*insert date dd/mm/yyyy*).

II.2. According to official information, animals in the consignment described in Part I meet the following health requirements:

II.2.1. They come from a confined establishment that is not subject to movement restrictions affecting the animals to be moved.

[2][3]*either* [II.2.2. 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 Commission Delegated Regulation (EU) 2020/688 are fulfilled.]

[2][3]*and/or* [II.2.2. 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.2.2.1. 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.2.2.1.1. for at least 60 days prior to the date of movement]]

[2]*and/or* [II.2.2.1.2. 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.2.2.1.3. 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.2.2.2.	have been protected against attacks by the vectors during transportation to the place of destination and have been kept protected against attacks by vectors in a vector protected establishment and
		[(2)*either*	[II.2.2.2.1. for at least 60 days prior to the date of movement]]
		[(2)*and/or*	[II.2.2.2.2. 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.2.2.2.3. 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.2.2.3.	have been vaccinated against those serotypes from 1 to 24 of infection with bluetongue virus which were reported during the past 2 years in that Member State or zone and are within the immunity period guaranteed in the specifications of the vaccine and
		[(2)*either*	[II.2.2.3.1. have been vaccinated more than 60 days before the date of movement]]
		[(2)*and/or*	[II.2.2.3.2. 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.2.2.4.	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.2.2.4.1. the serological test has been carried out on samples collected at least 60 days before the date of movement]]
		[(2)*and/or*	[II.2.2.4.2. 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)(3)*and/or*[II.2.2.	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.2.2.1.	have been protected against attacks by the vectors during transportation to the place of destination and have been kept protected against attacks by vectors in a vector protected establishment and
		[(2)*either*	[II.2.2.1.1. for at least 60 days prior to the date of movement]]

		[(2)*and/or*	[II.2.2.1.2.	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 date of the commencement of the period of protection against attacks by vectors]]
		[(2)*and/or*	[II.2.2.1.3.	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 date of the commencement of the period of protection against attacks by vectors;]]]
	[(2)*and/or*	[II.2.2.2.	have been kept at least 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 laid down 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.2.2.2.1.	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.2.2.2.1.1. have been vaccinated more than 60 days before the date of movement]]]
			[(2)*and/or*	[II.2.2.2.1.2. 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.2.2.2.2.	the animals have been immunised against those serotypes from 1 to 24 of infection with bluetongue virus which were reported during the past 2 years in an area of at least 150 km radius centred on the place where the animals were kept, and
			[(2)*either*	[II.2.2.2.2.1. 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)*or*	[II.2.2.2.2.2. 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)(3)*and/or*[II.2.2.	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 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.2.2.1.	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.2.2.1.1. point 5 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and
		[(2)*and/or*	[II.2.2.1.2. point 6 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and
		[(2)*and/or*	[II.2.2.1.3. point 7 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and
		[(2)*and/or*	[II.2.2.1.4. 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.2.2.2.	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.2.2.2.1. point 5 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and
		[(2)*and/or*	[II.2.2.2.2. point 6 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and
		[(2)*and/or*	[II.2.2.2.3. point 7 of Section 1 of Chapter 2 of Part II of Annex V to that Delegated Regulation, and
		[(2)*and/or*	[II.2.2.2.4. 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.2.2.3.	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.2.2.3.1. without any conditions, and

(2)and/or [II.2.2.3.2. 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.2.2.3.3. 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.2.2.3.4. 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.2.2.3.5. 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.3. To the best of my knowledge and as declared by the operator:

II.3.1. In the confined establishment of dispatch there are no abnormal mortalities with an undetermined cause affecting the animals to be moved.

II.3.2. The animals have not been in contact with animals which are subject to movement restrictions referred to in Point II.2.1., or with animals of a lower health status.

II.3.3. Based on the results of the surveillance plan of the confined establishment, the animals do not pose a significant risk at the confined establishment of destination for the spread of diseases for which they are listed.

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

II.5. This certificate is valid for 10 days from the date of issuing. In the case of transport by waterway/sea of animals, the period of 10 days for the 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 confined establishment approved in accordance with Articles 97 and 99 of Regulation (EU) 2016/429.

Box reference I.12: "*Place of destination*": Indicate a confined establishment approved in accordance with Articles 97 and 99 of Regulation (EU) 2016/429.

Part II:

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

(2) Delete if not applicable.

(3) Only in case of animals belonging to the families Antilocapridae, Bovidae, Camelidae, Cervidae, Giraffidae, Moschidae or Tragulidae.

Official veterinarian

Name (in capital letters) Qualification and title

Local Control Unit name Local Control Unit code

Date

Stamp Signature