



**Certificate for the export of other mammals to Northern Ireland**

<b>Part I: Description of consignment</b>	<b>I.1. Consignor</b> Name: Address:  Postal code/region: Country: ISO country code:		<b>I.2. Certificate unique reference number:</b> .....		
			<b>I.3. Central Competent Authority:</b> Department for Environment, Food and Rural Affairs (Defra)		
			<b>I.4. Local Competent Authority:</b> Animal and Plant Health Agency (APHA)		
	<b>I.5. Consignee</b> Name: Address:  Postal code/region: Country: ISO country code:		<b>I.6 Operator responsible for the consignment</b> Name: Address:  Postal Code/region: Country: ISO country code:		
	<b>I.7. Country of origin:</b> UNITED KINGDOM <b>ISO code:</b> GB	<b>I.8. Region of origin:</b>  <b>Code:</b>	<b>I.9. Country of destination:</b> NORTHERN IRELAND <b>ISO code:</b> XI	<b>I.10. Region of destination:</b>  <b>Code:</b>	
	<b>I.11. Place of dispatch</b> Name: Address:  Registration/Approval No: Postal code: Country: ISO code:		<b>I.12. Place of destination</b> Name: Address:  Registration/Approval No: Postal code: Country: ISO code:		

<b>I.13. Place of loading</b> Address:  Postal code:				<b>I.14. Date and time of departure</b>			
<b>I.15. Means of transport</b> Aeroplane <input type="checkbox"/> Ship <input type="checkbox"/> Railway <input type="checkbox"/> Road vehicle <input checked="" type="checkbox"/> Other <input type="checkbox"/> Identification:				<b>I.16 Entry Border Control Post</b>  <b>I.17. Accompanying documents (e.g., CITES, import authorisation)</b> Type Code: Country <span style="float: right;">ISO code:</span> Commercial document reference:			
<b>I.18. Transport conditions</b>		Ambient <input type="checkbox"/>	Chilled <input type="checkbox"/>	Frozen <input type="checkbox"/>			
<b>I.19. Container number/ Seal number</b> Container No <span style="float: right;">Seal No</span>							
<b>I.20. Certified as or for:</b> Laboratory/Research <input type="checkbox"/> Sale <input type="checkbox"/> Breeding/Further keeping <input type="checkbox"/> Confined establishment <input type="checkbox"/>							
<b>I.21 For Transit</b> <input type="checkbox"/> Third country <span style="float: right;">ISO country code</span>			<b>I.22 For Internal market</b> <input type="checkbox"/>				
<del>                     I.21 For Transit <input type="checkbox"/>                      Third country <span style="float: right;">ISO country code</span> </del>			<del>                     I.22 For Internal market <input type="checkbox"/> </del>				
<del> <b>I.23 For re-entry</b> <input type="checkbox"/> </del>			<del> <b>I.23 For re-entry</b> <input type="checkbox"/> </del>				
<b>I.24 Total number of cages/containers</b>		<b>I.25 Total number of animals</b>		<b>I.26 Total net weight/gross weight (kg)</b>			
<b>I.27. Description of consignment:</b>							
CN Code	Species (Scientific name)	Subspecies / Category	Identification system	Identification number/name	Age	Sex	Number
						M/F	
						M/F	
						M/F	

**II.1 Health information**

I, the undersigned official veterinarian hereby certify that the animal(s) of the consignment described in Part I meet the following requirements:

**II.1.1 Country/Territory of origin**

The zone, country or territory described in boxes I.7. and I.8 is a WOA member country where rabies is a notifiable disease.

**II.1.2 Premises of origin as described in box I.11**

- (a) is (approved) or <sup>(1)</sup> (registered) <sup>(1)</sup> by the competent authority of the third country of origin and complies with conditions at least equivalent to Article 102 of Regulation (EU) 2016/429.
- (b) is not subject to national restrictions for animal health reasons.
- (c) where there have been no clinical cases of rabies in the 6 months prior to the date of dispatch.
- (d) has had no reports of Brucella spp. and Tuberculosis complex in the 12 months prior to the date of dispatch
- (e) (in which the animals have remained since birth or were resident for at least 6 months prior to issue of this certificate)  
*or*  
(in which the animals have remained since birth or were resident for at least 3 months prior to issue of this certificate) <sup>(1)(2)</sup>
- (f) have not had any abnormal mortalities with an undetermined cause.
- (g) is under veterinary supervision and is subject to animal health surveillance and biosecurity measures according to the 'Animal Health Regulation' (Reg (EU) 2016/429)

**II.1.3 Animals**

- (a) have been (resident in the country or territory of dispatch since birth or for at least 6 months)<sup>(1)</sup> or, (were imported into the country or territory of dispatch directly from (\_\_\_\_\_, an EU Member State) <sup>(1)</sup> or (\_\_\_\_\_ a Third Country under the conditions at least as strict as those set out in this certificate)) <sup>(1)(3)</sup>
- (b) their movement has been authorised in advance by a written agreement of the Competent Authority of destination <sup>(3)</sup>.
- (c) a clear identification link between the live animals and the health certificate is established or the animals are individually identified. In the case of mammals of the Order Carnivora, they are individually identified by means of an implanted transponder.
- (d) are not Species of Union Concern under Regulation (EU) 1143/2014 unless they have been provided with a permit.
- (e) have not come into contact with other animals not complying with at least the same health requirements as described in this certificate since birth or for the last 60 days and, if applicable, during their transportation from the premises of origin to the place of loading.
- (f) have been examined on (enter date dd/mm/yyyy) \_\_\_\_\_ within the period of 24 hours prior to the time of loading for dispatch to Northern Ireland, and found to be free of clinical signs of infectious diseases including those referred to in Annex I (1) of Delegated Regulation (EU) 2020/692.
- (g) there is no reason to suspect it/they have been exposed to an infectious disease prior to issue of this certificate.
- (h) are not animals to be killed under a national programme for the eradication of disease.
- (i) <sup>(1)</sup> (in the case of bats (Chiroptera) the prevention and control rules in Article 9(1) of Regulation (EU) 2016/429 on the basis of Article 2 and the Annex of Implementing Regulation (EU) 2018/1882, are met.

- (j) <sup>(1)</sup> (in the case of fruit bats of the genus *Pteropus* from Malaysia (Peninsula) or Australia, the animals comply with the following conditions regarding Commission Decision 2006/146/EC
  - a. the animals originate from captive colonies.
  - b. the animals have been isolated in quarantine premises for at least 60 days, and
  - c. the animals have been subjected with negative results to a serum neutralisation or approved ELISA test for antibody against Hendra and Nipah disease viruses, carried out in a laboratory approved for these tests by the competent authorities on samples of blood taken on two occasions with an interval of 21 to 30 days, the second sample to be taken within 10 days of export (copy attached)
- (k) <sup>(1)</sup> (They do not concern the following species from zones or countries whose import is prohibited under Decision 2003/459/EC with regard to monkey pox virus.
  - a. prairie dogs (*Cynomys* sp.) originating in or coming from the United States of America
  - b. rodents of non-domestic species and all squirrels originating in or coming from third countries of the African sub-Saharan region
- (l) <sup>(1)</sup> (in the case of Lagomorphs, the animals come from an establishment in which no clinical signs of myxomatosis, VHD (viral or rabbit haemorrhagic disease) or tularemia have been observed during the 6 months prior to the date of dispatch to Northern Ireland)
- (m) <sup>(1)</sup> (In the case of carnivores<sup>(4)</sup>, other than carnivores moving directly from a confined establishment in the exporting country to a confined establishment in Northern Ireland, the animal(s) were at least 12 weeks old at the time of vaccination against rabies and at least 21 days have elapsed since the completion of the primary anti-rabies vaccination carried out in accordance with the validity requirements set out in Annex III to Regulation (EU) 576/2013 of the European Parliament and of the Council, and any subsequent revaccination was carried out within the period of validity of the preceding vaccination; Please provide vaccination details in the box below <sup>(5)</sup>)

Transponder <sup>(5)</sup>		Date of vaccination	Name and manufacturer of vaccine	Batch number	Validity of vaccination	
Alphanumeric code of the animal	Date of implantation and/or reading (dd/mm/yyyy)				From dd/mm/yyyy	To dd/mm/yyyy

**II.1.4 Parasite treatment <sup>(6)</sup>**

Provide details on testing and, where indicated in line with testing results, details of treatment for endo- and ectoparasites administered within 60 days of the issue of this certificate. (Specify active ingredients and doses of the products used, when applicable). A veterinary statement or certified copy of the test result is attached to the certificate and referenced under I.17. Enter details here

**II.1.5 Animal transport attestation**

- a. Arrangements were made to transport the animals in compliance with Council Regulation (EC) No. 1/2005, IATA guidelines and/or CITES guidelines for transport, where applicable or of other recognised international standards for transport of animals and the animals are fit for the intended transport.
- b. The person in charge of the transport of animals submitted a signed written statement, which demonstrates that suitable arrangements have been made for the feeding, watering, and care of the animal(s) during transport in accordance with the international standards recognised for the transport of animals.

- c. A correctly authorised transporter in compliance with Council Regulation (EC) No. 1/2005 will be used to transport this/these animals to Northern Ireland
- d. Any transport box or container in which animals are loaded into, is used for the first time or it has been cleaned and disinfected with an officially authorised disinfectant before loading.
- e. Arrangements have been made to load the animal(s) for dispatch to Northern Ireland into the transport box/container means of transport described above, which has been so constructed that faeces, urine, litter, or fodder could not flow or fall out of the vehicle or container during transportation.

**Notes:**

- (a) This certificate may only be used for mammal species (including sea mammals) where there is no harmonised EU certificate or no agreed national certificate between the Competent Authority of origin and the Competent Authority of destination. Primates are not eligible under this certificate.
- (b) This certificate shall be completed according to the notes for the completion of certificates provided for in Chapter 4 of Annex I to Commission Implementing Regulation (EU) 2020/2235
- (c) This certificate is valid for 10 days from the date of its issue or until the date of documentary and identity checks carried out at the first Border Control Post of entry to the European Union, whichever occurs earliest. In the case of transport by sea, the validity of the certificate is extended by the additional period corresponding to the duration of the journey by sea.

**Part 1**

**Box I.11:** Approval/registration number must be provided.

**Box I.12:** Approval/registration number must be provided.

**Box I.17:** Accompanying documents reference to the DAERA import authorisation and, including but not limited to, where applicable, the antiparasitic test/treatment under section II.1.4, the CITES documentation, DAERA Invasive species permit or a copy of a licence issued under The Dangerous Wild Animals (Northern Ireland) Order 2004 and a copy of the health certificate for the movement of the animal into the exporting country of dispatch to Northern Ireland.

**Box I.27:**

- **CN Code:** use the appropriate CN codes as declared under Commission Implementing Regulation (EU) 2021/632
- **Species:** This certificate may only be used for mammal species (including sea mammals) where there is no harmonised EU certificate or no agreed national certificate between the Competent Authority of origin and the Competent Authority of destination. Primates or pet mammals under EU Regulation 576/2013 are not eligible under this certificate.
- **Identification system:** Indicate transponder or other means of identification. Individual identification must be used wherever possible, but in the case of small animals, batch identification may be used. A clear link between the live animals and the health certificate must be established. CITES requirements may indicate the type of identification required. In the case of mammals of the Order Carnivora, they are individually identified by means of an implanted transponder.
- **Age and Sex:** only to be completed if appropriate.
- **Identification number:** Indicate the transponder or other identification means alphanumeric code.
- Add as many entries as are needed providing all the entries refer to the same species for which the same attestations apply. A separate schedule may be used in case of a large number of entries. The schedule must be linked and cross referenced across the official documents.

**Notes for completion**

<sup>(1)</sup> Please complete or delete as applicable.

<sup>(2)</sup> This attestation can be selected if the animals were moved to the confined establishment of origin from another approved confined establishment of equal health status located within the same Third Country of origin or an EU or EEA country

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- (3) authorisation must be attached to this certificate and accompany the animals during transit. Authorisation reference must be entered in Box I.17.
- (4) other mammals of the Order Carnivora, other than dogs, cats and ferrets
- (5) A separate schedule may be attached for additional entries which must include cross references to all other relevant documentation.
- (6) must be completed in the case of mammals of the species belonging to the order Carnivora, other than sea mammals. May be deleted for other mammals.

**Official Veterinarian**

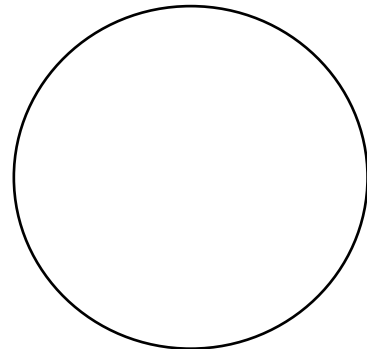
.....  
Name (in capital letters) .....

Date: .....

Qualification and title

Stamp:

Signature:



V.2.9110EHC APPLICATION