

GREAT BRITAIN, CHANNEL ISLANDS AND ISLE OF MAN

Pet health certificate for the non-commercial movement to Great Britain, Channel Islands and Isle of Man of dogs, cats or ferrets in accordance with Regulation (EU) No 576/2013

COUNTRY:

Veterinary certificate to Great Britain, Channel Islands and Isle of Man

	I.1. Consignor Name				I.2 Certificate reference number					
	Address				I.3. Consignee Name Address					
	Tel.		Postal code Tel.							
ent	I.4. Central competent authority				I.5. Country of origin					
onsignme	I.6. Local competent authority				I.7. ISO Code of country of origin					
Part I: Details of dispatched consignment	I.8. Description of commodity					I.9. Commodity code (HS code) 010619				
ails of di	I.10. Quantity			.11. Commodities certified for: Pets						
t I: Det	I.12. Identificati	on of the	commo	dities		[
Par	Species (Scientific name)	Sex	C	Colour	Breed		tification umber	Identification system [transponder/ tattoo ⁽¹⁰⁾]	Date of birth [dd/mm/yyyy]	

[11.	Health information II.a. Certificate reference number							
		I, the undersigned official veterinarian (1) / veterinarian authorised by the competent authority (1) of							
		(insert name of country) certify that:							
Part II: Certification		Purpose/nature of journey attested by the owner II.1. the attached declaration (²) by the owner or the natural person who has authorisation in writing from the owner to carry out the non-commercial movement of the animals on behalf of the owner, supported by evidence (³), states that the animals described in Box I.12 will accompany the owner or the natural person who has authorisation in writing from the owner to carry out the non-commercial movement of the animals on behalf of the owner to carry out the non-commercial movement of the animals on behalf of the owner within not more than 5 days of his movement and are not subject to a movement that aims at their sale or a transfer of ownership, and during the non-commercial movement will remain under the responsibility of							
t ⊫:		(¹) <i>either</i> [the owner;]							
Par		(¹) or [the natural person who has authorisation in writing from the owner to carry out the non-commercial movement of the animals on behalf of the owner,]							
		(¹) or [the natural person designated by a carrier contracted by the owner to carry out the non-commercial movement of the animals on behalf of the owner;]							
	(¹) either	r II.2. the animals described in Box I.12 are moved in a number of five or less;							
	(1) or	II.2. the animals described in Box 1.12 are moved in a number of more than five are more than 6 months old and are going to participate in competitions exhibitions or sporting events or in training for those events, and the owner or the natural person referred to in point II.1 has provided evidence (³) that the animals are registered							
		(¹) either [to attend such event;]							
		(¹) or [with an association organising such events;]							

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e animals described hast 21 days have ccordance with the ubsequent revaccin <i>bither</i> [II.3.1 the anim Regulation Implementin Annex II to of Regulatio provided in country other rabies antiboo by the compe preceding prin the date of iss any subseque	d in Box I.12 were a elapsed since the validity requirement ation was carried on nals described in B (EU) No 577/2013, ng Regulation (EU) Implementing Regulation (EU) Implementing Regulation (EU) no (EU) No 576/20 the table below:] mals described in B than those listed in dy titration test (7), tent authority on the mary vaccination we sue of this certifical	at least 12 weeks completion of the ents set out in Ar- nut within the period fox I.12 come from either directly, the No 577/2013 or to ulation (EU) No 57 13 (⁶), and the det Box I.12 come from a Annex II to Imple carried out on a base date indicated i vithin a current val	old at the til e primary au nex III to F od of validity n a third cou rough a thir through a thi 77/2013 in a ails of the cu m, or are sch ementing Re lood sample n the table b id vaccinatio	nti-rabies Regulation of the pre- ntry listed d country l rd country ccordance urrent anti- neduled to gulation (I taken by elow not lo	vaccination (EU) No ceding va in Annex listed in A rother that with poir rabies va transit th EU) No 57 the veteri ess than 3	on (⁴) carried out in 576/2013 and any ccination (⁵); and II to Implementing nnex II to n those listed in it (c) of Article 12(1) ccination are rough a third 77/2013 and a narian authorised
e animals described hast 21 days have ccordance with the ubsequent revaccin <i>bither</i> [II.3.1 the anim Regulation Implementin Annex II to of Regulatio provided in country other rabies antiboo by the compe preceding prin the date of iss any subseque	d in Box I.12 were a elapsed since the validity requirement ation was carried on nals described in B (EU) No 577/2013, ng Regulation (EU) Implementing Regulation (EU) Implementing Regulation (EU) no (EU) No 576/20 the table below:] mals described in B than those listed in dy titration test (7), tent authority on the mary vaccination we sue of this certifical	at least 12 weeks completion of the ents set out in Ar- nut within the period fox I.12 come from either directly, the No 577/2013 or to ulation (EU) No 57 13 (⁶), and the det Box I.12 come from a Annex II to Imple carried out on a base date indicated i vithin a current val	old at the til e primary au nex III to F od of validity n a third cou rough a thir through a thi 77/2013 in a ails of the cu m, or are sch ementing Re lood sample n the table b id vaccinatio	nti-rabies Regulation of the pre- ntry listed d country l rd country ccordance urrent anti- neduled to gulation (I taken by elow not lo	vaccination (EU) No ceding va in Annex listed in A rother that with poir rabies va transit th EU) No 57 the veteri ess than 3	on (⁴) carried out in 576/2013 and any ccination (⁵); and II to Implementing nnex II to n those listed in it (c) of Article 12(1) ccination are rough a third 77/2013 and a narian authorised
	hals described in Box I.12 come from, or are scheduled to transit through a third than those listed in Annex II to Implementing Regulation (EU) No 577/2013 and a y titration test (⁷), carried out on a blood sample taken by the veterinarian authorise ent authority on the date indicated in the table below not less than 30 days after the ary vaccination within a current valid vaccination series and at least 3 months prior ue of this certificate, proved an antibody titre equal to or greater than 0,5 IU/ml (⁸) a nt revaccination was carried out within the period of validity of the preceding and the details of the current anti-rabies vaccination and the date of sampling for					
		Name and	able below:] Batch	Validity of vaccination [dd/mm/yyyy]		Date of the blood sampling
Date of implantation and/or reading (⁹) [dd/mm/yyyy]	vaccination [dd/mm/yyyy]	manufacturer of vaccine	number	From	То	[dd/mm/yyyy]
	vaccination (⁶ testing the im onder or too Date of implantation and/or reading (⁹)	vaccination (⁵), and the details o testing the immune response and onder or too Date of implantation and/or reading (⁹)	vaccination (⁵), and the details of the current anti- testing the immune response are provided in the testing testin	vaccination (⁵), and the details of the current anti-rabies vaccin testing the immune response are provided in the table below:] onder or too Date of implantation and/or reading (⁹) Date of [dd/mm/yyyy]	vaccination (⁵), and the details of the current anti-rabies vaccination and testing the immune response are provided in the table below:] onder or too Date of implantation and/or reading (⁹) Date of implantation and/or	vaccination (⁵), and the details of the current anti-rabies vaccination and the date testing the immune response are provided in the table below:] onder or too Date of implantation and/or reading (⁹) Date of (dd/mm/yyyy) Name and manufacturer of vaccination [dd/mm/yyyy] Batch number

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COUNTRY:

II.	I. Health information			II.a. Certificate reference number				
	Attestatio	n of anti-	parasite treatment					
(') eith				I 12 are destined f	or Great B	Britain Channel Islands and Isle of Man and have		
() 0111		(II.4. the dogs described in Box I.12 are destined for Great Britain, Channel Islands and Isle of Man and have been treated against <i>Echinococcus multilocularis</i> and the details of the treatment carried out by the administering veterinarian in accordance with Article 6 of Delegated Regulation (EU) No 2018/772 (¹⁰) (¹¹) are provided in the table below.]						
(') or) or [II.4. the dogs described in Box I.12 have not been treated against <i>Echinococcus multilocularis</i> (¹⁰).]							
				chinococcus eatment		Administering veterinarian		
	ransponder number of t		Name and manufacturer of the product	Date [dd/mm and time of tre [00:00]	atment	Name in capitals, stamp and signature		
Notes (a) This certificate is meant for dogs (Canis Lupus familiaris) cats (Felis silvestris Cetus) and ferrets (Mustela putorius furo).								
(b)		certificate is valid for 10 days from the date of issue by the official veterinarian until the date of the mentary and identity checks at the designated point of entry into Great Britain, Channel Islands and Isle of						
		he case of transport by sea, that period of 10 days is extended by an additional period corresponding he duration of the journey by sea						
Part I:	:							
Box I.:	3: Cons	<i>signee:</i> in	indicate Great Britain, Channel Islands and Isle of Man as destination.					
Box I.	12: Iden	tification	system: select of the	e following: transp	onder or	tattoo.		
	Iden	tification	number: indicate the	transponder or t	attoo alph	anumeric code.		
	Date of birth/breed as stated by the owner.							

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II.	Health information II.a. Certificate reference number						
Part	set 11.						
Fart							
(1)	Keep as appropriate.						
(²)	The declaration referred to in point II.1 shall be attached to the certificate and comply with the ma additional requirements set out in Part 3 of Annex IV to Implementing Regulation (EU) No 577/201						
(³)	The evidence referred to in point II.1 (e.g. boarding pass, flight ticket) and in point II.2 (e.g. receipt entry to the event, proof of membership) shall be surrendered on request by the competent authority responsible for the checks referred to in point (b) of the Notes.	The evidence referred to in point II.1 (e.g. boarding pass, flight ticket) and in point II.2 (e.g. receipt of entry to the event, proof of membership) shall be surrendered on request by the competent authorities responsible for the checks referred to in point (b) of the Notes.					
(4)	Any revaccination must be considered a primary vaccination if it was not carried out within the period of validity of a previous vaccination.						
(5)	A certified copy of the identification and vaccination details of the animals concerned shall be attac the certificate.	hed to					
(⁶)	The third option is subject to the condition that the owner or the natural person referred to in point I provides, on request by the competent authorities responsible for the checks referred to in point (b declaration stating that the animals have had no contact with animals of species susceptible of rab and remain secure within the means of transport or the perimeter of an international airport during transit through a third country other than those listed in Annex II to Implementing Regulation (EU) No 577/2013. This declaration shall comply with the format, layout and language requirement out in Parts 2 and 3 of Annex I to Implementing Regulation (EU) No 577/2013.), a bies the					
(7)	The rabies antibody titration test referred to in point II.3.1:						
	 must be carried out on a sample collected by a veterinarian authorised by the competent au at least 30 days after the primary rabies vaccination within a current valid vaccination series months before the date of import; 						
	 must measure a level of neutralising antibody to rabies virus in serum equal to or greater th EU/ml; 	an 0,5					
	 must be performed by a laboratory approved in accordance with Article 3 of Council Decision 2000/258/EC (list of approved laboratories available at https://ec.europa.eu/food/animals/permovement/approved-labs_en); 						
	 does not have to be renewed on an animal, which following that test with satisfactory resident period of validity of a previous vaccination. 	ults, has					
	A certified copy of the official report from the approved laboratory on the results of the rabies ant test referred to in point II.3.1 shall be attached to the certificate.	ibody					
(8)	By certifying this result, the official veterinarian confirms that he has verified, to the best of his abilit where necessary with contacts with the laboratory indicated in the report, the authenticity of the laboratory report on the results of the antibody titration test referred to in point II.3,1.	ty and					
(⁹)	In conjunction with footnote (⁶), the marking of the animals concerned by the implantation of a transponder or by a clearly readable tattoo applied before 3 July 2011 must be verified before any is made in this certificate and must always precede any vaccination, or where applicable, testing ca out on those animals.						

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II.	Health information	II.a. Certificate reference number					
(10)	The treatment against Echinococcus multilocular	ris referred to in point II.4 must:					
		period of not more than 120 hours and not less than 24 try of the dogs into Great Britain, Channel Islands and					
	 consist of an approved medicinal product which contains the appropriate dose of praziquantel or pharmacologically active substances, which alone or in combination, have been proven to reduce the burden of mature and immature intestinal forms of <i>Echinococcus multilocularis</i> in the host species concerned. 						
(11)	¹¹) The table referred to in point I1.4 must be used to document the details of a further treatment if administered after the date the certificate was signed and prior to the scheduled entry into Great Britain, Channel Islands and Isle of Man.						
Offici	al veterinarian/Authorised veterinarian (delete as ap	propriate)					
	Name (in capital letters):	Qualification and title:					
	Address						
	Telephone:						
	Date:	Signature:					
	Stamp:						
Endo	rsement by the competent authority (not necessary	when the certificate is signed by an official veterinarian)					
	Name (in capital letters):	Qualification and title:					
	Address						
	Telephone:						
	Date:	Signature:					
	Stamp:						

Official at point of entry in GB					
	Name (in capital letters):	Title:			
	Address				
	Telephone:				
	E-mail address:				
	Date of completion of documentary and identity checks by authorised	l body:			
	Signature:				
	Stamp:				

Explanatory notes for completing the health certificate

- (a) Where the certificate states that certain statements shall be kept as appropriate, statements which are not relevant may be crossed out and initialled and stamped by the official veterinarian, or completely deleted from the certificate.
- (b) The original of each certificate shall consist of a single sheet of paper, or, where more text is required it must be in such a form that all sheets of paper required are part of an integrated whole and indivisible.
- (c) The certificate shall be drawn up in English. It shall be completed in block letters in English.
- (d) If additional sheets of paper or supporting documents are attached to the certificate, those sheets of paper or document shall also be considered as forming part of the original of the certificate by the application of the signature and stamp of the official veterinarian, on each of the pages.
- (e) When the certificate, including additional sheets referred to in point (d), comprises more than one page, each page shall be numbered (page number of total number of pages) at the end of the page and shall bear at the top of each page the certificate reference number that has been designated by the competent authority.
- (f) The original of the certificate shall be issued by an official veterinarian of the country of dispatch or by an authorised veterinarian and subsequently endorsed by the competent authority of the country of dispatch. The competent authority of the country of dispatch shall ensure that rules and principles of certification equivalent to those laid down in Directive 96/93/EC are followed.
- (g) The colour of the signature shall be different from that of printing. This requirement also applies to stamps other than those embossed or watermarked.
- (h) The certificate reference number referred to in boxes I.2 and II.a shall be issued by the competent authority of the country of dispatch.

Written declaration referred to in Article 25(3) of of Regulation (EU) No 576/2013

Section A Model of declaration

I, the undersigned

.....

[owner or the natural person who has authorisation in writing from the owner to carry out the non-commercial movement on behalf of the owner⁽¹⁾]

declare that the following pet animals are not subject to a movement that aims at their sale or a transfer of ownership and will accompany the owner or the natural person who has authorisation in writing from the owner to carry out the non-commercial movement on behalf of the owner⁽¹⁾ within not more than 5 days of his movement.

Transponder/tattoo ⁽¹⁾ alphanumeric code	Animal health certificate number

During the non-commercial movement, the above animals will remain under the responsibility of

⁽¹⁾either [the owner];

- ⁽¹⁾or [the natural person who has authorisation in writing from the owner to carry out the non-commercial movement on behalf of the owner]
- ⁽¹⁾or [the natural person designated by the carrier contracted to carry out the non-

commercial movement on behalf of the owner:

(insert name of the carrier)]

Place and date:

Signature of the owner or natural person who has authorisation in writing from the owner to carry out the non-commercial movement on behalf of the owner⁽¹⁾:

.....

⁽¹⁾ delete as appropriate.