Model animal health certificate for the non-commercial movement into a Member State from a territory or third country of dogs, cats or ferrets in accordance with Article 5(1) and (2) of Regulation (EU) No 576/2013

**COUNTRY: United States** Veterinary certificate to EU Consignor I.2.a. I.2. Certificate reference No Name Address Central competent authority **USDA APHIS Veterinary Services** of dispatched consignment Tel. Local competent authority I.6. Person responsible for the consignment in the EU Consignee Name Address Postal code Tel. I.7. Country of ISO code I.8. Region of Code Country ISO I.10 Region of Code origin origin destination code destination **United States** Part I: Details I.12. Place of destination I.11. Place of origin I.13. Place of loading I.14. Date of departure I.15. Means of transport I.16. Entry BIP in EU I.17. No.(s) of CITES I.18. Description of commodity I.19. Commodity code (HS code) 010619 Dog Cat Ferret I.20. Quantity I.21. Temperature of products I.22. Total number <del>packages</del> I.23. Seal/Container No I.24. Type of packaging I.25. Commodities certified for: Pets  $\mathbf{X}$ I.26. For transit to 3rd Country I.27. For import or admission into EU 1.28. Identification of the commodities **Species** Identification Date of birth Colour Identification number Sex Breed (Scientific name) system [dd/mm/yyyy]

EN 2019/1293

COUNTRY: United StatesNon-commercial movement into a Member State from a territory or third country of dogs, cats or ferrets in accordance with Article 5(1) and (2) of Regulation (EU) No 576/2013

II.		Health	informat	tion	II.a.	Certificat	e referenc	ce No	II.b.		
<u> </u> 	I, the undersigned official veterinarian <sup>(1)</sup> <b>America</b> (insert name of territory o			territory or third c	ountry) certify that:	the competer	nt authority <sup>(1)</sup> of	the United	States of		
	Purpose/nature of journey attested by the			='		a authorioation in	itin a fram the a				
	II.1.		the attached declaration <sup>(2)</sup> 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 <sup>(3)</sup> , states that the animals described in Box I.28 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 within not more than five 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 [the owner;]								
	<sup>(1)</sup> C		[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;]								
	<sup>(1)</sup> €	<del>)r</del>		[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;]							
	ither	[11.2.	the anima	ıls describ	ed in Box I.28 are	e moved in a numbe	er of five or les	ss;]			
( <del>1)</del> 0	the animals described in Bo going to participate in compe natural person referred to in-		in competitions,	exhibitions or sporti	ng events or i	n training for thos	se events, and the				
	<sup>(1)</sup> €	either	[to attend	such eve	nt;]						
	<sup>(1)</sup> €	er .	<del>[with an a</del>	ssociation	n organising such	events;]					
		Attestation	n of rabies v	<u>accinatio</u>	n and rabies antib	ody titration test:					
<sup>(1)</sup> e.	<sup>(1)</sup> either [II.3.		the animals described in Box I.28 are less than 12 weeks old and have not received an anti-rabies vaccination, or are between 12 and 16 weeks old and have received an anti-rabies vaccination, but 21 days at least have not elapsed since the completion of the primary vaccination against rabies carried out in accordance with the validity requirements set out in Annex III to Regulation (EU) No 576/2013 <sup>(4)</sup> , and  II.3.1 the territory or third country of provenance of the animals indicated in Box I.1 is listed in Annex II to Implementing Regulation (EU) No 577/2013 and the Member State of destination indicated in Box I.5 has informed the public that it authorises the movement of such animals into its territory, and they are								
	<sup>(1)</sup> either		[II.3.2	accompanied by  II.3.2 the attached declaration <sup>(5)</sup> of the owner or the natural person referred to in point II.1 stating that from birth until the time of the non-commercial movement the animals have had no contact with wild animals of species susceptible to rabies;]							
	<sup>(±)</sup> <del>or</del>		<del>[II.3.2</del>	their mo	heir mother, on whom they still depend, and it can be established that the mother received before their birth an anti-rabies vaccination which complied with the validity requirements set out in Annex III to Regulation (EU) No 576/2013;]]						
<sup>(4)</sup> <del>O</del>	<sup>(1)</sup> or/and [II.3.		days have validity re	the animals described in Box I.28 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 <sup>(4)</sup> carried out in accordance with the validity requirements set out in Annex III to Regulation (EU) No 576/2013 and any subsequent revaccination was carried out within the period of validity of the preceding vaccination <sup>(6)</sup> ; and							
	<sup>(4)</sup> either		[II.3.1 the animals described in Box I.28 come from a territory or a third country listed in Annex II to Implementing Regulation (EU) No 577/2013, either directly, through a territory or a third country listed in Annex II to Implementing Regulation (EU) No 577/2013 or through a territory or a third country other than those listed in Annex II to Implementing Regulation (EU) No 577/2013 in accordance with point (e) of Article 12(1) of Regulation (EU) No 576/2013 <sup>(7)</sup> , and the details of the current anti-rabies vaccination are provided in the table below;]								
	country other the antibody titration competent author vaccination and a cqual to or greate validity of the pre			other than those y titration test <sup>(8)</sup> , ent authority on to tion and at least to or greater than of the preceding	Box 1.28 come from the listed in Annex II carried out on a he date indicated it three menths prior 10.5 IU/mil <sup>(6)</sup> and any vaccination (6), and a primmune response	to Implementi blood sample n the table be to the date of subsequent re the details of	ing Regulation (E taken by the value and the the elow not less the issue of this cert evaccination was the current anti-r	EU) No 577/2013 a veterinarian author an 30 days after th ificate, proved an a carried out within t abies vaccination a	and a rabies ised by the e preceding antibody titre the period of		
	Transponder or tattoo						Validity of	vaccination			
A	Alphanumeric code of the animal		Date implant and/ readin	ation or g <sup>(10)</sup>	Date of vaccination [dd/mm/yyyy]	Name and manufacturer of vaccine	Batch number	From [dd/mm/yyyy]	To [dd/mm/yyy]	Date of the blood sampling [dd/mm/ yyyy]	

Alphanumeric code of the animal	Date of implantation and/or reading <sup>(10)</sup> [dd/mm/yyyy]	Date of vaccination [dd/mm/yyyy]	Name and manufacturer of vaccine	Batch number	From [dd/mm/yyyy]	To [dd/mm/yyy]	the blood sampling [dd/mm/

#### COUNTRY: United StatesNon-commercial movement into a Member State from a territory or third country of dogs, cats or ferrets in accordance with Article 5(1) and (2) of Regulation (EU) No 576/2013

II.	Health	information	II.a.	Certificate reference	e No	II.b.
<sup>(1)</sup> either	Regulation (EU) 2018/878 and have been treated against <i>Echinococcus multilocularis</i> , and the deta treatment carried out by the administering veterinarian in accordance with Article 6 of Commission					ris, and the details of the
<sup>(1)</sup> or	[11.4.	• ,		e provided in the table below.] een treated against Echinocod	•	laris <sup>(11)</sup> .]

Transponder or		chinococcus eatment	Administering veterinarian	
tattoo number of the dog	Name and manufacturer of the product	Date [dd/mm/yyyy] and time of treatment [00:00]	Name in capitals, stamp and signature	

#### Notes

- This certificate is meant for dogs (Canis lupus familiaris), cats (Felis silvestris catus) and ferrets (Mustela putorius furo). (a)
- This certificate is valid for 10 days from the date of issue by the official veterinarian until the date of the documentary and (b) checks at the designated Union travellers' point of entry (available http://ec.europa.eu/food/animal/liveanimals/pets/pointsentry\_en.htm).

In the case of transport by sea, that period of 10 days is extended by an additional period corresponding to the duration of the journey by sea.

For the purpose of further movement into other Member States, this certificate is valid from the date of the documentary and identity checks for a total of four months or until the date of expiry of the validity of the anti-rabies vaccination or until the conditions relating to animals less than 16 weeks old referred to in point II.3 cease to apply, whichever date is earlier. Please note that certain Member States have informed that the movement into their territory of animals less than 16 weeks old referred point 11.3 is not authorised. wish to in You may to inquire http://ec.europa.eu/food/animal/liveanimals/pets/index en.htm.

#### Part I:

Box I.5: Consignee: indicate Member State of first destination.

Box I.28: Identification system: select of the following: transponder or tattoo.

Identification number, indicate the transponder or tattoo alphanumeric code.

Date of birth/breed: as stated by the owner.

#### Part II:

(2)

(3)

(4)

(5)

(7)

Keep as appropriate.

The declaration referred to in point II.1 shall be attached to the certificate and comply with the model and additional requirements set out in Part 3 of Annex IV to Implementing Regulation (EU) No 577/2013.

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

Any revaccination must be considered a primary vaccination if it was not carried out within the period of validity of a previous

The declaration referred to in point II.3.2 to be attached to the certificate complies with the format, layout and language requirements laid down in Parts 1 and 3 of Annex I to Implementing Regulation (EU) No 577/2013.

A certified copy of the identification and vaccination details of the animals concerned shall be attached to the certificate.

The third option is subject to the condition that the owner or the natural person referred to in point II.1 provides, on request by the competent authorities responsible for the checks referred to in point (b), a declaration stating that the animals have had no contact with animals of species susceptible of rabies and remain secure within the means of transport or the perimeter of an international airport during the transit through a territory or 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 requirements set out in Parts 2 and 3 of Annex I to Implementing Regulation (EU) No 577/2013.

(8) 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 authority, at least 30 days after the date of vaccination and three months before the date of import;
- must measure a level of neutralising antibody to rabies virus in serum equal to or greater than 0.5 IU/ml;
- must be performed by a laboratory approved in accordance with Article 3 of Council Decision 2000/258/EC (list of approved laboratories available at

 $\underline{http://ec.europa.eu/food/animal/liveanimals/pets/approval en.htm});$ 

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II.	Health information	II.a.	Certificate reference I	No	II.b.		
	<ul> <li>does not have to be renewed on an animal, which following that test with satisfactory results, has been revaccina against rabies within the period of validity of a previous vaccination.</li> </ul>						
	A certified copy of the official report from the approved laboratory on the results of the rabies antibody test referred to in poi II.3.1 shall be attached to the certificate.						
(9)			erinarian confirms that he has verified, to the best of his ability and where necessary with d in the report, the authenticity of the laboratory report on the results of the antibody				
(10)	readable tattoo applied before 3 Jul	marking of the animals concerned by the implantation of a transponder or by a clearly y 2011 must be verified before any entry is made in this certificate and must always plicable, testing carried out on those animals.					
(11)	The treatment against Echinococcus		·				
			iod of not more than 120 hours an ne Member States or parts thereol				
		ombination,	ch contains the appropriate dose of have been proven to reduce the st species concerned.				
(12)	The table referred to in point II.4 mus certificate was signed and prior to the Implementing Regulation (EU) 2018/8	ne schedule					
(13)	The table referred to in point II.4 m certificate was signed for the purpose conjunction with footnote (11).						
Officia	Official veterinarian/Authorised veterinarian						
	Name (in capital letters):		Qu	ualification a	and title:		
	Address						
	Telephone:						
	Date:			S	Signature:		
	Stamp:						
Endor	rsement by the competent authority (not nece	ssary when	the certificate is signed by an offic	ial veterinar	ian)		
	Name (in capital letters):		Qu	ualification a	and title:		
	Address						
	Telephone:						
	Date:		Się	gnature:			
	Stamp:						
Officia	Official at the travellers' point of entry (for the purpose of further movement into other Member States)						
	Name (in capital letters):		Tit	tle:			
	Address						
	Telephone:						
	E-mail address:						
	Date of completion of the documentary and	identity che	cks: Się	gnature:	Stamp:		

## **Declaration**

I, the ur	ndersigned			
[owne	r or the natural person who has authorisation in writing	from the owner to carry out the non-commercial movement on be $\operatorname{owner}^{(I)}$	half of the	
ownersl	hip and will accompany the owner or the carry out the non-commercial movement	abject to a movement that aims at their sale or a e natural person who has authorisation in writing ent on behalf of the owner <sup>(1)</sup> within not more than	g from the	
Tra	nsponder/tattoo <sup>(1)</sup> alphanumeric code	Animal health certificate number		
	·			
Duning	the new commencial maximum the cherr	a animala viill mamain undan tha magnanaihility of		
	the non-commercial movement, the abov • [the owner];	e animals will remain under the responsibility of		
(1)or	2	eation in writing from the owner to carry out owner]	the non-	
<sup>(1)</sup> or	(the natural person designated by the carrier contracted to carry out the non-commercial moven on behalf of the owner:			
	Place and date:			
	Signature of the owner or natural personal out the non-commercial movement on be	on who has authorisation in writing from the ownerhalf of the owner <sup>(<math>I</math>)</sup> :	er to carry	
(1)	delete as appropriate.			

## ANNEX I

### Part 1

## Format and layout of the declaration referred to in point (a) of Article 7(2) and of Article 11(2) of Regulation (EU) No 576/2013

## **DECLARATION**

I, the undersigned	(1)
	riting from the owner to carry out the non-commercial movement of als on behalf of the owner $^{(2)}$ ]
declare that from birth until the time of the r have had no contact with wild animals of spe	non-commercial movement the following pet animals ecies susceptible to rabies:
Transponder/tattoo <sup>(2)</sup> alphanumeric code <sup>(2)</sup>	Passport/Animal health certificate <sup>(2)</sup> number <sup>(2)</sup>
Place and date:	
Signature:	
<ul><li>(1) to be completed in block letters.</li><li>(2) delete as appropriate.</li></ul>	

Page \_\_\_\_ of \_\_\_\_