## TURKS AND CAICOS ISLANDS GOVERNMENT DEPARTMENT OF AGRICULTURE BUTTERFIELD SQUARE, 16 PARADISE AVENUE PROVIDENCIALES, TURKS AND CAICOS ISLANDS TEL: 1.649.946.5801; FAX: 1.649.941.7587

COUNTRY



## Veterinary certificate to EU

	I.1	Consignor: Name:			I.2. Certificate referen	I.2.a				
		Address:			I.3. Central competent authority					
nent					I.4. Local competent authority					
ısignn		Tel:								
Part 1: details of dispatched consignment	I.5.	Consignee Name:			1.6.					
of dispa		Address:								
etails o										
art 1: de		Postal code:								
Ч	I.7.	Tel: Country of ISO code	I.8		I.9.	I.10.				
		origin TCI   TC								
	I.11				1.12.					
	I.13				I.14.					
	I.15				I.16.					
					I.17. No(s) of CITES					
	I.18	. Description of commodity			I.19. Commodity code (HS code) 010619					
					1.20. Quantity 1.22.					
	I.21									
	I.23				1.24.					
	I.25 Pets	. Commodities certified for:								
	I.26			I.	.27.					
	I.28	. Identification of the commodi	ties				1			
		Species (Scientific name)	Identification system	t	te of application of he microchip or tattoo (dd/mm/yyyy)	Date of birth (dd/mm/yyyy)				
L			ı							

						-					8)	
					II.a. Certificate r	eferenc	e No			II.b		
Part II: Certification	II.	Healt	th in	formation								
		I, the undersigned official veterinarian of The Animal Health Service, Department of Agriculture, TCI Government certify that:										
		II.1. based on the declaration in point II.7. the animals satisfy the definition of 'pet animals' as provided for in point (a) of Article 3 of Regulation (EC) No998/2003;										
		II.2. at least 21 days have elapsed since the completion of the primary vaccination against rabies (1) carried out in accordance with the requirements set out in Annex Ib to Regulation (EC) No 998/2003 and any subsequent revaccination was carried out within the period of validity of the preceding vaccination (2) and details of the current vaccination are provided in the table in point II.4.										
	(3) either	r II.3. the animals come from a third country or territory listed in Section 2 of Part B or in Part or in Part C of Annex II to Regulation (EC) No 998/2003.										
	(3) or	II.3. the animals come, and if transiting another third country or territory, are scheduled to transit through, third country or territory listed in Part 1 of Annex II to Commission Regulation (EU) No 206/2010 and since the dates indicated in the table in point II.4. when blood samples were taken not earlier than 30 days after vaccination from each of the animals by a veterinarian authorised by the competent authority which subsequently proved antibody titres equal to or greater than 0.5 IU/ml in a virus neutralisation test for tables carried out in approved laboratory (4) (5) at least 3 months have elapsed any subsequent revaccination was carried out within the period of validity of preceding vaccination (2).										
		II.4 the details of the current anti-rabies vaccination and the date of sampling are the following:										
			Date of vaccination	Name of manufacturer of vaccine	Bate	h number	Validity (dd/mm/yyyy)		/	Date of the blood sample (dd/mm/yyyy)		
-	anii	mal		(dd/mm/yyyy)				From		То		
	<ul> <li>(3) either (II.5. the dogs have not been treated against <i>Echinococcus multilocularis:</i>)</li> <li>(3) or (II.5. the dogs have been treated against <i>Echinococcus</i> multilocularis and the details of the treatment are documented in the table in point II.6)</li> <li>II.6. the details of the treatment carried out by the administering veterinarian in accordance with Article 7 of Commission Delegated Regulation (EU) No 1152/2011 (6) are the following:</li> </ul>											
					Anti-echino	Anti-echinococcus treatmeat				Administering veterinarian		
	Microch	licrochip or tattoo number Name a of the dog			Name and manufacturer of product			Date (dd/mm/yyyy)and time Na of treatment (00:00)		Name (i	nme (in capital) stamp and signature	
				*				(7)				
									(8) (8)			
									(8)			
									(8)			
	(8) This in	This date must precede the date the certificate was signed. This information may be entered after the date the certificate was signed for the purpose described in point (e) of the Notes and in conjunction with footnote 6.										
	The signat	The signature and the stamp must be in a different colour to that of the printing.										
	Official	Official veterinarian										
Name (in capital letters) : Qua					alification and title:							
	Date	Date:				Signature:						
	Stamp:											

II. Health Information	II.a. Certificate reference No.	II.b.						
(II.7. I have a written declaration signed by the owner or the natural person responsible for animals on behalf of the owner, stating that:								
DECLARATION								
I, the undersigned								
Declare that the animal will accompany me, the owner, or the natural person that I have designated to be responsible of the animals on my behalf and are not intended to be sold or transferred to another owner.								
Place and date:	Sign	ature:						
Notes								
	(a) The original of each certificate shall consist of a single sheet of paper, or, where more text is required it must be in such form that all sheets of paper required are part of an integrated whole and indivisible.							
	(b) The certificate shall be drawn up at least in the language of the Member State of entry and in English. It shall be completed in block letters in the language of the Member State or in English.							
(c) 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.								
(d) When the certificate, including additional sheets referred to in (c), 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 the certificate reference number that has been designated by the competent authority at the top of the pages.								
(e) The certificate is valid for 10 days from the date of issue by the official veterinarian until the date of the checks at the EU travellers' point of entry and for the purpose of further movements within the Union, for a total of 4 months from the date of issue of this certificate or until the date of expiry of the anti-rabies vaccination, whichever date is earlier.								
	of the exporting third country or territory shall ensure t tive 96/93/EC are followed.	that rules and principles of certification equivalent						
Part I								
Box I.11: Place of origin: name and	address of the dispatch establishment. Indicate approv	val or registration number						
Box I.28: Identification system: sele	ect of the following: microchip or tattoo							
Date of application of the	microchip or tattoo: the tattoo must be clearly readabl	le and applied before 3 July 2011						
Identification number: in	dicate the microchip or tattoo number							
Date of birth: indicate only	y if known							
Part II								
(1) Any revaccination must be considered a primary vaccination if it was not carried out within the period validity of a previous vaccination.								
(2) A certified copy of the identification and vaccination details of the animals concerned shall be attached to the certificate.								
	certificate states that certain statements shall be kept as and stamped by the official veterinarian, or completely							

## II. Health Information II.a. Certificate reference No.

- (4) The rabies antibody test referred to in point II.3:
  - 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 3 months before the date of import,
  - must measure a level of neutralizing antibody to rabies virus in serum equal to or greater than 0.5 IU/ml,
  - must be performed by approved in accordance with Article 3 of Council Decision 2000/258/EC designating a specific institute responsible for establishing criteria necessary for standardizing the serological tests to monitor the effectiveness of rabies vaccines (list of approved laboratories available at http://ec.europa.eu/food/animal/veanimals/pets/approval\_en.htm),
  - needs not be renewed on an animal, which following that test with satisfactory results, has been revaccinated against rabies within the period of validity of a previous vaccination.
- (5) A certified copy of the official report from the approved laboratory on the results of the rabies antibody tests referred to in point II.3 shall be attached to the certificate.
- (6) The treatment against Echinococcus multilocularis referred to in point II.5 must:
  - be administered by a veterinary within a period of not more than 120 hours and not less than 24 hours before the time of the scheduled entry of the dogs into one of the Member States or parts thereof listed in Annex I to Regulation (EU) No 1152/2011,
  - 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 *Echinococcus multilocularis* in the host species concerned.
- (7) This date must precede the date the certificate was signed.
- (8) This information may be entered after the date the certificate was signed for the purpose described in point (e) of the Notes and in conjuction with footnote 6.