

**COUNTRY:**

**Veterinary certificate to EU**

<b>Part I : Details of dispatched consignment</b>	I.1. Consignor Name Address  Tel.		I.2. Certificate reference No	I.2.a.			
			I.3. Central competent authority				
			I.4. Local competent authority				
	I.5. Consignee Name Address  Postal code Tel.		I.6.				
	I.7. Country of origin	ISO code	I.8.	I.9. Country destination	of ISO code	I.10. Region of destination	Code
	I.11. Place of origin  Name Address Name Address Name Address Approval number Approval number Approval number		I.12.				
	I.13. Place of loading		I.14. Date of departure				
	I.15. Means of transport  Aeroplane <input type="checkbox"/> Ship <input type="checkbox"/> Railway wagon <input type="checkbox"/> Road vehicle <input type="checkbox"/> Other <input type="checkbox"/> Identification Documentary references		I.16. Entry BIP in EU				
			I.17. No(s) of CITES				
	I.18. Description of commodity			I.19. Commodity code (HS code) <b>010619</b>			
			I.20. Quantity				
I.21.			I.22. Number of packages				
I.23. Seal/Container No			I.24.				
I.25. Commodities certified for: Pets <input type="checkbox"/> Approved bodies <input type="checkbox"/>							
I.26.		I.27. For import or admission into EU <input type="checkbox"/>					
I.28. Identification of the commodities							
Species (Scientific name)	Identification system	Date of application of the microchip or tattoo [dd/mm/yyyy]	Identification number	Date of birth [dd/mm/yyyy]			

**COUNTRY**

**Imports of dogs, cats, ferrets and non-commercial movements into the Union of more than five dogs, cats or ferrets**

**Part II: Certification**

II.	Health information	II.a. Certificate reference No	II.b.																																												
<p>I, the undersigned official veterinarian of ..... (<i>insert name of third country</i>) certify that:</p> <p>II.1. the clinical examination carried out on each of the animals within 24 hours of scheduled dispatch by a veterinarian authorised by the competent authority showed the animals to be fit to be transported on the intended journey at the time of inspection;</p> <p>II.2. at least 21 days have elapsed since the completion of the primary vaccination against rabies<sup>(1)</sup> 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<sup>(2)</sup> and details of the current vaccination are provided in the table in point II.4.</p> <p><sup>(3) either</sup> II.3. the animals come from a third country or territory listed in Section 2 of Part B or in Part C of Annex II to Regulation (EC) No 998/2003;]</p> <p><sup>(3) or</sup> II.3. the animals come from, and if transiting another third country or territory, are scheduled to transit through, a 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 rabies carried out in an approved laboratory<sup>(4)(5)</sup>, at least 3 months have elapsed and any subsequent revaccination was carried out within the period of validity of the preceding vaccination<sup>(2)</sup>.]</p> <p>II.4. the details of the current anti-rabies vaccination and the date of sampling are the following:</p>																																															
<table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th rowspan="2" style="width: 15%;">Microchip or tattoo number of the animal</th> <th rowspan="2" style="width: 15%;">Date of vaccination [dd/mm/yyyy]</th> <th rowspan="2" style="width: 15%;">Name and manufacturer of vaccine</th> <th rowspan="2" style="width: 10%;">Batch number</th> <th colspan="2" style="width: 15%;">Validity [dd/mm/yyyy]</th> <th rowspan="2" style="width: 15%;">Date of the blood sample [dd/mm/yyyy]</th> </tr> <tr> <th style="width: 5%;">From</th> <th style="width: 5%;">To</th> </tr> </thead> <tbody> <tr><td> </td><td> </td><td> </td><td> </td><td> </td><td> </td><td> </td></tr> </tbody> </table>				Microchip or tattoo number of the animal	Date of vaccination [dd/mm/yyyy]	Name and manufacturer of vaccine	Batch number	Validity [dd/mm/yyyy]		Date of the blood sample [dd/mm/yyyy]	From	To																																			
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<p><sup>(3) either</sup> II.5. the dogs have not been treated against <i>Echinococcus multilocularis</i>;</p> <p><sup>(3) or</sup> II.5. the dogs have been treated against <i>Echinococcus multilocularis</i> and the details of the treatment are documented in the table in point II.6.]</p> <p>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<sup>(6)</sup> are the following:</p>																																															
<table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th rowspan="2" style="width: 15%;">Microchip or tattoo number of the dog</th> <th colspan="2" style="width: 35%;">Anti-echinococcus treatment</th> <th style="width: 50%;">Administering veterinarian</th> </tr> <tr> <th style="width: 15%;">Name and manufacturer of the product</th> <th style="width: 20%;">Date [dd/mm/yyyy] and time of treatment [00:00]</th> <th style="width: 35%;">Name (in capital), stamp and signature</th> </tr> </thead> <tbody> <tr><td> </td><td> </td><td style="text-align: center;">(7)</td><td> </td></tr> <tr><td> </td><td> </td><td style="text-align: center;">(8)</td><td> </td></tr> </tbody> </table>				Microchip or tattoo number of the dog	Anti-echinococcus treatment		Administering veterinarian	Name and manufacturer of the product	Date [dd/mm/yyyy] and time of treatment [00:00]	Name (in capital), stamp and signature			(7)				(8)				(8)				(8)				(8)																		
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**COUNTRY****Imports of dogs, cats, ferrets and non-commercial movements into the Union of more than five dogs, cats or ferrets**

II. Health information	II.a. Certificate reference No	II.b.
<p><b>Notes</b></p> <p>(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 a form that all sheets of paper required are part of an integrated whole and indivisible.</p> <p>(b) The certificate shall be drawn up in at least one of the official languages of the Member State of the border inspection post of introduction of the consignment into the Union and of the Member State of destination. However, those Member States may authorise the certificate to be drawn up in the official language of another Member State, and accompanied, if necessary, by an official translation.</p> <p>(c) If for reasons of identification of the items of the consignment (schedule in point I.28), 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.</p> <p>(d) When the certificate, including additional schedules 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.</p> <p>(e) The certificate shall be valid for 10 days from the date of issue by the official veterinarian, except for a non-commercial movement into the Union of more than five dogs, cats and ferrets in which case the certificate is valid 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.</p> <p>(f) The competent authorities of the exporting third country or territory shall ensure that rules and principles of certification equivalent to those laid down in Directive 96/93/EC are followed.</p> <p><b>Part I:</b></p> <p>Box I.11.: Place of origin: name and address of the dispatch establishment. Indicate approval or registration number</p> <p>Box I.28.: <i>Identification system</i> : Select of the following : microchip or tattoo  <i>Date of application of the microchip or tattoo</i> : The tattoo must be clearly readable and applied before 3 July 2011  <i>Identification number</i> : Indicate the microchip or tattoo number  <i>Date of birth</i> : Indicate only if known</p> <p><b>Part II:</b></p> <p>(1) Any revaccination must be considered a primary vaccination if it was not carried out within the period of validity of a previous vaccination.</p> <p>(2) A certified copy of the identification and vaccination details of the animals concerned shall be attached to the certificate.</p> <p>(3) Keep as appropriate. 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.</p> <p>(4) The rabies antibody test referred to in point II.3:</p> <ul style="list-style-type: none"> <li>- 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;</li> <li>- must measure a level of neutralising antibody to rabies virus in serum equal to or greater than 0.5 IU/ml;</li> <li>- must be performed by a laboratory approved in accordance with Article 3 of Council Decision 2000/258/EC designating a specific institute responsible for establishing criteria necessary for standardising the serological tests to monitor the effectiveness of rabies vaccines (list of approved laboratories available at <a href="http://ec.europa.eu/food/animal/liveanimals/pets/approval_en.htm">http://ec.europa.eu/food/animal/liveanimals/pets/approval_en.htm</a>);</li> <li>- 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.</li> </ul> <p>(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.</p> <p>(6) The treatment against <i>Echinococcus multilocularis</i> referred to in point II.5 must:</p> <ul style="list-style-type: none"> <li>- be administered by a veterinarian 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;</li> </ul>		

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<p>- 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.</p> <p>(7) This date must precede the date the certificate was signed.</p> <p>(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 conjunction with footnote (6).</p> <p>The signature and the stamp must be in a different colour to that of the printing.</p>								
<p>Official veterinarian</p> <table><tr><td data-bbox="347 741 616 770">Name (in capital letters):</td><td data-bbox="1075 741 1318 770">Qualification and title:</td></tr><tr><td data-bbox="347 804 408 833">Date:</td><td data-bbox="1075 804 1182 833">Signature:</td></tr><tr><td data-bbox="347 866 424 896">Stamp:</td><td></td></tr></table>			Name (in capital letters):	Qualification and title:	Date:	Signature:	Stamp:	
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Date:	Signature:							
Stamp:								

## FOR REFERENCE ONLY – DO NOT INCLUDE WITH HEALTH CERTIFICATE

### ANNEX Ib

#### Technical requirements for the anti-rabies vaccination (Referred to in Article 5(1)(b)(i))

For the purposes of Article 5(1), an anti-rabies vaccination shall be considered valid provided that the following requirements are complied with:

1. The anti-rabies vaccine must:

- (a) be a vaccine other than a live modified vaccine and fall within one of the following categories:
  - (i) an inactivated vaccine of at least one antigenic unit per dose (WHO standard); or
  - (ii) a recombinant vaccine expressing the immunising glycoprotein of the rabies virus in a live virus vector;
- (b) if administered in a Member State, have been granted a marketing authorisation in accordance with:
  - (i) Directive 2001/82/EC of the European Parliament and of the Council of 6 November 2001 on the Community code relating to veterinary medicinal products ( 1 ); or
  - (ii) Regulation (EC) No 726/2004 of the European Parliament and of the Council of 31 March 2004 laying down Community procedures for the authorisation and supervision of medicinal products for human and veterinary use and establishing a European Medicines Agency ( 2 );
- (c) if administered in a third country, meet at least the requirements laid down in Part C of Chapter 2.1.13 of the Manual of Diagnostic Tests and Vaccines for Terrestrial Animals, 2008 Edition, of the World Organisation for Animal Health.

2. An anti-rabies vaccination may only be considered valid if it meets the following conditions:

- (a) the vaccine was administered on a date indicated in:
  - (i) Section IV of the passport; or
  - (ii) the appropriate section of the accompanying animal health certificate;
- (b) the date referred to in point (a) must not precede the date of microchipping indicated in:
  - (i) Section III(2) of the passport; or
  - (ii) the appropriate section of the accompanying animal health certificate;
- (c) at least 21 days must have elapsed since the completion of the vaccination protocol required by the manufacturer for the primary vaccination in accordance with the technical specification of the marketing authorisation referred to in point 1(b) for the anti-rabies vaccine in the Member State or third country in which the vaccination is administered;
- (d) the period of validity of the vaccination, as prescribed in the technical specification of the marketing authorisation for the anti-rabies vaccine in the Member State or third country where the vaccine is administered, must have been entered by the authorised veterinarian in:
  - (i) Section IV of the passport; or
  - (ii) the appropriate section of the accompanying animal health certificate;
- (e) a revaccination (booster) must be considered a primary vaccination if it was not carried out within the period of validity referred to in point (d) of a previous vaccination.