Government of Canada / Gouvernement du Canada
Canadian Food Inspection Agency / Agence canadienne d'inspection des aliments

REFERENCE NUMBER:_____

Veterinary health certificate
For the non-commercial movement of dogs, cats or ferrets in accordance with Regulation (EU) 576/2013

CANADA

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

Part I: Description of consignment

I.1 Consignor/Exporter Name Address Tel.	**I.2 Certificate reference**
	I.3 Consignee Name Address Postal code Tel.
I.4. Central competent authority **Canadian Food Inspection Agency (CFIA)**	**I.5. Country of origin** **CANADA**
I.6. Local competent authority **District of**	**I.7. ISO code** **CA**

I.8 Description of commodity	**I.9 Commodity code (HS code)** **010619**
	I.10. Quantity

I.11 Commodities certified for:
Pets ☐

I.12 Identification of the commodities

Species (Scientific name)	Sex	Colour	Breed	Identification number	Identification System (transponder/tattoo)[10]	Date of birth [dd/mm/yyyy]

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II. Health information	**II.a. Certificate reference number**

Part II: Certification

I, the undersigned official veterinarian(1) /veterinarian authorised by the competent authority(1) of……**CANADA**…….
certify that:

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 authorization 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 (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

(1)either [the owner;]

(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 [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 I.12 are moved in a number of more than five are more than six 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;]

Attestation of rabies vaccination and rabies antibody titration test:

(1)either [II.3. the animals described in Box I.12 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 (4) 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(5); and

(1)either [II.3.1 the animals described in Box I.12 come from a third country listed in Annex II to Implementing Regulation (EU) No 577/2013, either directly, through a third country listed in Annex II to Implementing Regulation (EU) No 577/2013 or through a third country other than those listed in Annex II to Implementing Regulation (EU) No 577/2013 in accordance with point (c) of Article 12(1) of Regulation (EU) No 576/2013(6), and the details of the current anti-rabies vaccination are provided in the table below:]

(1)or [II.3.1 the animals described in Box I.12 come from, or are scheduled to transit through, a third country other than those listed in Annex II to Implementing Regulation (EU) No 577/2013 and a rabies antibody titration test(7), carried out on a blood sample taken by the veterinarian authorised by the competent authority on the date indicated in the table below not less than 30 days after the preceding primary vaccination within a current valid vaccination series and at least three (3) months prior to the date of issue of this certificate, proved an antibody titre equal to or greater than 0,5 IU/ml(8) and any subsequent revaccination was carried out within the period of validity of the preceding vaccination(5), and the details of the current anti-rabies vaccination and the date of sampling for testing the immune response are provided in the table below:]

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Transponder or Tattoo		**Date of vaccination [dd/mm/yyyy**	**Name and manufacturer of vaccine**	**Batch number**	**Validity of vaccination**		**Date of the blood sampling [dd/mm/yyyy]**
Alphanumeric code of the animal	**Date of implantation and/or reading(9) [dd/mm/yyyy]**				**From [dd/mm/yyyy]**	**To [dd/mm/yyyy]**	

Attestation of anti-parasite treatment:

(1)either [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 *Echinococcus multilocularis*, and the details of the treatment carried out by the administering veterinarian in accordance with Article 6 of Delegated Regulation (EU) 2018/772 (10) (11) are provided in the table below:

(1)or [II.4. the dogs described in Box I.12 have not been treated against Echinococcus multilocularis(10).]

Transponder 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 capitals, stamp and signature**

Notes

(a) This certificate is meant for dogs *(Canis lupus familiaris)* cats *(Felis silvestris catus)* and ferrets *(Mustela putorius furo).*

(b) This certificate is valid for 10 days from the date of issue by the official veterinarian until the date of the documentary and identity checks at the designated point of entry into Great Britain, Channel Islands and Isle of Man

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

Part I:

Box I.3: *Consignee:* indicate Great Britain, Channel Islands and Isle of Man as destination.

Box I.12: *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:

(1) Keep as appropriate.

(2) The declaration referred to in point II.1 shall be attached to the certificate and comply with the model and additional

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II. Health information	**II.a. Certificate reference number**

requirements set out in Part 3 of Annex IV to Implementing Regulation (EU) No 577/2013,

(3) 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 attached to the certificate.

(6) 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 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.

(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 authority, at least 30 days after the primary rabies vaccination within a current valid vaccination series and 3 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 EU/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 https://ec.europa.eu/food/animals/pet-movement/approved-labs_en);
- does not have to 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.

A certified copy of the official report from the approved laboratory on the results of the rabies antibody test referred to in point II.3.1 shall be attached to the certificate.

(8) By certifying this result, the official veterinarian confirms that he has verified, to the best of his ability and 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.

(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 entry is made in this certificate and must always precede any vaccination, or where applicable, testing carried out on those animals.

(10) The treatment against *Echinococcus multilocularis* referred to in point II.4 must:

- 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 Great Britain, Channel Islands and Isle of Man;
- 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.

(11) The table referred to in point II.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.

Official veterinarian/Authorised veterinarian (delete as appropriate)

Name (in capital letters): Qualification and title:

Address:

Telephone:

Date: Signature:

Stamp:

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Endorsement 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 body:

Signature:

Stamp:

REFERENCE NUMBER:_____

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II. Health information	**II.a. Certificate reference number**

Section III: Owner/Authorized Person Declaration

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

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.

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.