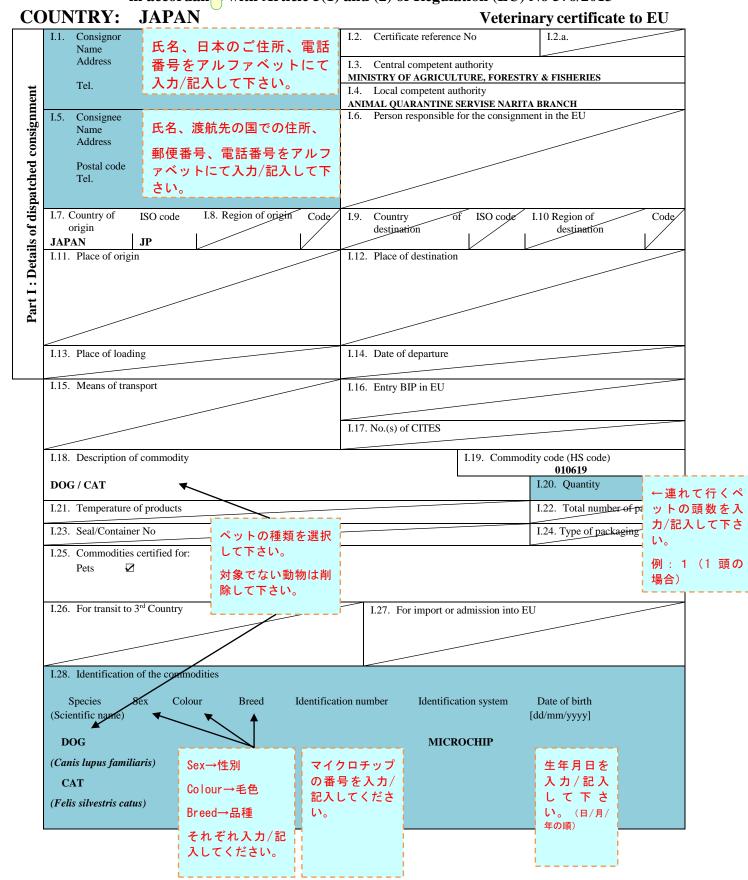
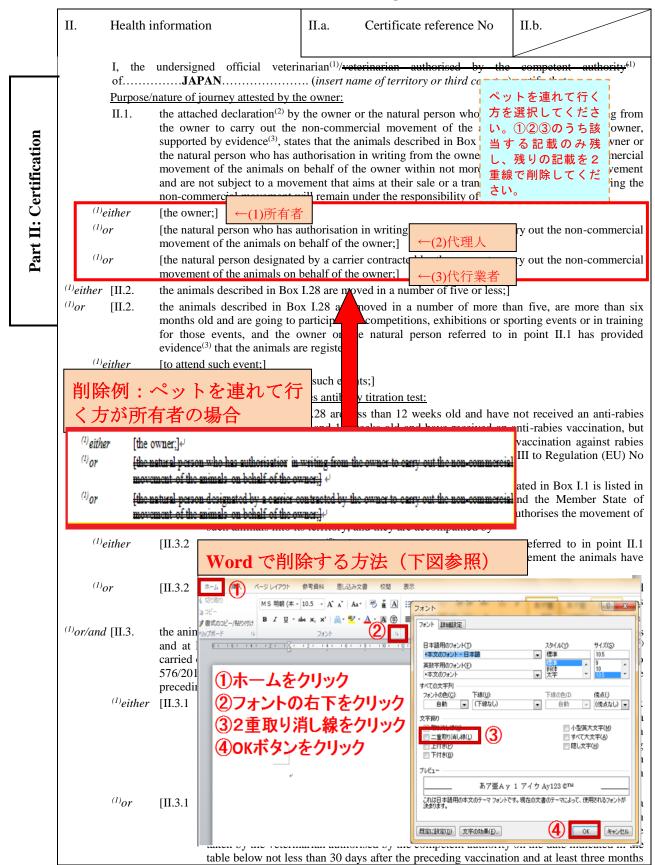
Model animal health fron in accordan

青色もしくは赤字で表示されている箇所について、別添の入力/ 記入用のフォームに入力するか、印刷後に記入して下さい。(こ の記入例に記入しないでください。)



COUNTRY JAPAN

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



※注意:記入用の EU フォーム(ブランク)について、今回の輸出に不要な記載内容をあらかじめ削除したものを送付しております。この記入方法サンプルと記載内容が異なる場合がございますが、赤字の指示内容以外に文章の追加・削除等を行わないで下さい。

COUNTRY TAPAN

Non-commercial movement into a Member State from a territory or third country of dogs, cats or ferrets in accordance

Tra Alphanu code o anin	ımeric	0.5 of t	IU/ml ⁽⁹⁾ and any he preceding va	y subsequent rev	vaccination of the details	was carried ou s of the curre ponse are pro	at within the p nt anti-rabies vided in the ta	o or greater than period of validity vaccination and ble below:
Alphanu code o	ımeric					X7∝1: J:4 €		
code o		Date of	1			Validity of vaccination		
		implantation and/or reading ⁽¹⁰⁾ [dd/mm/yyyy]	Date of vaccination [dd/mm/yyyy]	Name and manufacturer of vaccine	Batch number	From [dd/mm/yyyy]	to [dd/mm/yyyy]	Date of the blood sampling [dd/mm/yyyy]
マイク プ番号 /記入し さい。	を入力	マイクロチップの装着日が不明な場合は読み取り日) (技)を開発を関係を対して、 (日/月/年の順)を入力/記入して下さい。	狂犬病ワク チン接種日 を入力/配入 して下さ い。(日/月/ 年の順)	ワクチンの 製造会名を 英語で入力/ 記入して ださい。	ワの番は番カ/に チッまッを入 力/に さい。	ワクチン の接種日 を入力/記 入してく ださい。 (日/月/年 の順)	ワクチン の有効力/ 配入して くださ い。(日/ 月/年の 順)	日本ので 大手 が 別年間 1 十十十十十十十十十十十十十十十十十十十十十十十十十十十十十十十十十十十
(1)eithe		Delegated R multilocularia accordance w provided in th	eribed in Box I. egulation (EU or, and the deta with Article 7 or the table below.)	No 1152/20	Nant carried Nant carried 必要	ave been tr I out by the Legulation (El	cated against administering U) No 1152/2	to Commission Echinococcus Veterinarian in O11(11)(12)(13) are

国・地域に トを連れて 方は(1)の 線をなく (2)の部 2重線で削 てください。

駆虫が必

Ī	Transponder or		chinococcus catment	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		
	マイクロチップ番号を入力/記入して下さい。	薬品の製品名と製造 会社名をアルファベ ットで入力/記入して ください。	駆除を実施した日付(日/月/年の順)と時刻を入力/記入して下さい。	処置を実施した獣医師の氏名(アルファベットの大文字)、印鑑の押印、署名を記入していただいて下さい。 【印刷後に黒以外のインク(ペン)で直筆してもらうこと】		

†この欄は、エキノコックス駆除が必要な場合のみ、入力/記入してください。

]]

Notes

- (a) This certificate is meant for dogs (Canis lupus familiaris), cats (Felis silvestris catus) and ferrets (Mustela
- (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 Union travellers' point of entry (available at 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 to in point II.3 is not authorised.

COUNTRY JAPAN

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

II.	Health information	II.a.	Certificate reference No	II.b.

You may wish to inquire at 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:

(1) 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 vaccination.
- 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.
- (6) 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 http://ec.europa.eu/food/animal/liveanimals/pets/approval_en.htm);
 - 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.

- 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.
- 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.
- 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 one of the Member States or parts thereof listed in Annex I to Delegated 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.

COUNTRY JAPAN

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

II.	Health information	II.a.	Certificate reference No	II.b.
(12)	The table referred to in point II.4 mu	記入	しないで下さい	mber States or
	ial veterinarian/Authorised veterinarian Aame (in capital letters): Address Telephone:		_	on and title: .RANTINE OFFICER
	Date: Stamp:			Signature:
	rsement by the competent authority (not necessary): Address	cessary whe		official veterinarian) on and title:
	Telephone: Date: Stamp:		Signature:	
	ial at the travellers' point of entry (for the p Name (in capital letters): Address Telephone: E-mail address:	urpose of fi	urther movement into other Mem Title:	aber States)
	Date of completion of the documentary and	d identity cl	necks: Signature:	Stamp:

Part 3

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

