Commission Implementing Decision (EU) 2019/294 of 18 February 2019 laying down the list of territories and third countries authorised for imports into the Union of dogs, cats and ferrets and the model animal health certificate for such imports (notified under document C(2019) 1059) (Text with EEA relevance)

ANNEX

PART 1

Model animal health certificate for imports into the Union of dogs, cats and ferrets

COUNTRY: **Veterinary certificate to EU**

Part I: Details of dispatched consignment

I.1. Consignor Name Address Country Tel.	I.2. Certificate reference No / I.2.a. I.3. Central competent authority I.4. Local competent authority
I.5. Consignee Name Address Country Tel.	I.6.
I.7. Country of origin / ISO code / I.8.	I.9. Country of destination / ISO code / I.10. Region of destination / Code
I.11. Place of origin Name / Approval number Address Name / Approval number Address Name / Approval number Address	I.12. Place of destination Name / Approval number Address
I.13. Place of loading	I.14. Date of departure
I.15. Means of transport Aeroplane ☐ Ship ☐ Railway wagon ☐ Road vehicle ☐ Other ☐ Identification Documentary references	I.16. Entry BIP in EU I.17.
I.18. Description of commodity	I.19. Commodity code (HS code) **010619** I.20. Quantity
I.21.	I.22. Number of packages
I.23. Seal/Container No	I.24.

I.25. Commodities certified for: Others ☐ Pets ☐ Approved bodies ☐			
I.26.		I.27. For import or admission into EU ☐	
I.28. Identification of the commodities			
Species (Scientific name)	Identification system	Identification number	Date of birth [dd/mm/yyyy]

COUNTRY **Imports into the Union of dogs, cats, ferrets**

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

Part II: Certification

I, the undersigned official veterinarian of ..... (*insert name of third country*) certify that the animals described in Box I.28:

II.1. come from holdings or businesses described in Box I.11 which are registered by the competent authority and are not subject to any ban on animal health grounds, where the animals are examined regularly and which comply with the requirements ensuring the welfare of the animals held;

II.2. showed no signs of diseases and were fit to be transported for the intended journey at the time of examination by a veterinarian authorised by the competent authority within 48 hours prior to the time of dispatch;

(1) *either* [II.3. are destined for a body, institute or centre described in Box I.12 and approved in accordance with Annex C to Council Directive 92/65/EEC, and come from a territory or third country listed in Annex II to Commission Implementing Regulation (EU) No 577/2013.]

(1) *or* [II.3. 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 (2) carried out in accordance with the validity requirements set out in Annex III to Regulation (EU) No 576/2013 of the European Parliament and of the Council, and any subsequent revaccination was carried out within the period of validity of the preceding vaccination (3), and

(1) *either* [they come from, and in case of transit are scheduled to transit through, a territory or third country listed in Annex II to Commission Implementing Regulation (EU) No 577/2013 and details of the current anti-rabies vaccination are provided in columns 1 to 7 in the table below;]

(1) *or* [they come from or are scheduled to transit through, a territory or third country listed in Part 1 of Annex II to Commission Regulation (EU) No 206/2010 or listed without time limit in Annex I to Commission Implementing Regulation (EU) 2018/659, and

— details of the current anti-rabies vaccination are provided in columns 1 to 7 in the table below, and

— a rabies antibody titration test (4), carried out on a blood sample taken by the veterinarian authorised by the competent authority not less than 30 days after the preceding vaccination and at least three months prior to the date of issue of this certificate, proved an antibody titre equal to or greater than 0,5 IU/ml (5) and any subsequent revaccination was carried out within the period of validity of the preceding vaccination, and the date of sampling for testing the immune response are provided in column 8 in the table below:]

Transponder or tattoo		Date of vaccination [dd/mm/yyyy]	Name and manufacturer of vaccine	Batch number	Validity of vaccination		Date of blood sampling [dd/mm/yyyy]
Alphanumeric code of the animal	Date of implantation and/or reading (6) [dd/mm/yyyy]				From [dd/mm/yyyy]	to [dd/mm/yyyy]	
1	2	3	4	5	6	7	8

(1) *either* [II.4. the consignment includes dogs destined for a Member State listed in the Annex to Commission Implementing Regulation (EU) 2018/878 and those dogs have been treated against *Echinococcus multilocularis*, and the details of the treatment carried out by the administering veterinarian in accordance with Article 6 of Commission Delegated Regulation (EU) 2018/772 (7) (8) are provided in the table below:

COUNTRY **Imports into the Union of dogs, cats, ferrets**

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

Transponder or tattoo alphanumeric code 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
			]

([1]) *or* [II.4. the dogs forming part of the consignment have not been treated against Echinococcus multilocularis.]

Notes

This certificate is valid for 10 days from the date of issue by the official veterinarian. 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.11: Place of origin: name and address of the dispatch establishment. Indicate approval or registration number.

Box I.12: Place of destination: mandatory where the animals are destined for a body, institute or centre approved in accordance with Annex C to Council Directive 92/65/EEC.

Box I.25: Commodities certified for: indicate

— 'Pets' where dogs (Canis lupus familiaris), cats (Felis silvestris catus) or ferrets (Mustela putorius furo) are moved in accordance with Article 5(4) of Regulation (EU) No 576/2013 of the European Parliament and of the Council;

— 'Approved bodies' where dogs, cats or ferrets are moved in accordance with Article 13 of Council Directive 92/65/EEC to an approved body, institute or centre as defined in Article 2(c) of that Directive;

— 'others' where dogs, cats or ferrets are moved in accordance with Article 10 of Council Directive 92/65/EEC.

Box I.28: Identification system: select transponder or tattoo.

Identification number: indicate the transponder or tattoo alphanumeric code.

Part II:

([1]) Keep as appropriate.

([2]) Any revaccination must be considered a primary vaccination if it was not carried out within the period of validity of a previous vaccination.

([3]) A certified copy of the identification and vaccination details of the animals concerned shall be attached to the certificate.

([4]) The rabies antibody titration 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 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;

COUNTRY **Imports into the Union of dogs, cats, ferrets**

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

— 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/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 result of the rabies antibody test referred to in point II.3 shall be attached to the certificate.

([5]) 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.

([6]) In conjunction with footnote (3), 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.

([7]) 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 the Annex to Commission Implementing Regulation (EU) 2018/878;

— 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.

([8]) 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 one of the Member States or parts thereof listed in the Annex to Commission Implementing Regulation (EU) 2018/878.

Official veterinarian

Name (in capital letters): Qualification and title:

Date: Signature:

Stamp:

PART 2

Explanatory notes for completing the animal 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 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(s) of another Member State, and accompanied, if necessary, by an official translation.

(d) If for reasons of identification of the items of the consignment (schedule in point I.28 of the model animal health certificate), additional sheets of paper or supporting documents are attached to the certificate, those sheets of paper or documents 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 or documents 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 the certificate reference number that has been designated by the competent authority at the top of the pages.

(f) The original of the certificate shall be completed and signed by an official veterinarian of the exporting territory or third country. The competent authority of the exporting territory or third country 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 the 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 exporting territory or third country.

Changes to legislation:
There are outstanding changes not yet made to Commission Implementing Decision (EU) 2019/294. Any changes that have already been made to the legislation appear in the content and are referenced with annotations.
View outstanding changes

Changes and effects yet to be applied to :

- Annex omitted by S.I. 2020/1463 reg. 10(5)

Changes and effects yet to be applied to the whole legislation item and associated provisions

- Signature words omitted by S.I. 2020/1463 reg. 10(5)
- Art. 1(2)-(4) substituted for Art. 1(2) by S.I. 2020/1463 reg. 10(2)(c)
- Art. 1a inserted by S.I. 2020/1463 reg. 10(3)
- Art. 2(a) words substituted by S.I. 2020/1463 reg. 10(4)(c)(i)
- Art. 2(a) words substituted by S.I. 2020/1463 reg. 10(4)(c)(ii)
- Art. 2(b) words omitted by S.I. 2020/1463 reg. 10(4)(d)