Matvælastofnun
Icelandic Food and Veterinary Authority

# Animal health certificate for imports into the Union of dogs, cats and ferrets

**COUNTRY** Iceland

**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: Ministry of industries and innovation

I.4. Local competent authority: The Icelandic Food and Veterinary Authority

I.5. Consignee
Name
Address

Country
Tel.

I.6.

I.7. Country of origin: Iceland | ISO code: IS

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] |
|---|---|---|---|

**Part II: Certification**

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

I, the undersigned official veterinarian of ..ICELAND..... (*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* [II.3.1. they come from 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 the table];

([1]) *or* [II.3.1. they come from or are scheduled to transit through, a territory or third country listed in Annex I to Commission Decision 2004/211/EC or in Part 1 of Annex II to Commission Regulation (EU) No 206/2010, 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 3 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 details of the current anti-rabies vaccination and the date of sampling for testing the immune response are provided 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] |
|---|---|---|---|---|---|---|---|
| Alpha-numeric code of the animal | Date of implantation and/or reading ([6]) [dd/mm/yyyy] | | | | From [dd/mm/yyyy] | to [dd/mm/yyyy] | |
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([1]) *either* [II.4. are dogs destined for a Member State listed in Annex I to Commission Delegated Regulation (EU) No 1152/2011 and have been treated against *Echinococcus multilocularis*, and the details of the treatment carried out by the administering veterinarian in accordance with Article 7 of Commission Delegated Regulation (EU) No 1152/2011 ([7]) ([8]) are provided in the table below.]

([1]) *or* [II.4. have not been treated against *Echinococcus multilocularis*.]

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

| 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 |
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**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. 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 'others' where the animals are moved in accordance with Article 5(4) of Regulation (EU) No 576/2013 of the European Parliament and of the Council.

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.3.1:

— 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 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 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/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.

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

— 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.3.1 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.3.1.

([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 Annex I to Commission 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.

([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 Annex I to Commission Delegated Regulation (EU) No 1152/2011.

Official veterinarian

Name (in capital letters): Qualification and title:

Date: Signature:'

Stamp:

PART 2

**Explanatory notes for completing the animal health certificates**

(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 Council Directive 96/93/EC are followed.

(g) 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.