Part I : Details of consignment

I.1. Consignor
Name
Address
Country ISO Code

I.2. IMSOC Reference
Specimen not to be used for exports from EU

I.2.a. Local Reference

I.5. Consignee
Name
Address
Country ISO Code

I.3. Central competent authority

I.4. Local competent authority

I.7. Country of origin ISO Code

I.9. Country of destination ISO Code

I.8. Region of origin Code

I.10. Region of destination Code

I.11. Place of Dispatch
Name
Address
Approval Number
Country ISO Code

I.12. Place of destination
Name
Address
Approval Number
Country ISO Code

I.13. Place of Loading
Name
Address
Approval Number
Country ISO Code

I.14. Date and time of departure

I.15. Means of Transport

Mode	International transport document	Identification

I.16 Entry Point

I.18. Transport conditions
Ambient ☐

I.17. Accompanying documents
Commercial document reference
Date of issue
Country
Place of issue

I.19. Container No / Seal No

I.20. Certified as
Competition ☐ Pets ☐ Breeding ☐ Circus exhibition ☐

I.21. For transit through a third country ☐
Country ISO Code
EU Exit Authority BCP code
EU Entry Authority BCP code

I.22. For transit through Member State(s) ☐
Country ISO Code

I.25. Total gross weight

I.28. Description of consignment

1. 01 LIVE ANIMALS
0106 Other live animals
Mammals:
010619 Other
01061900 Other

Commodity	Species	Identification system	Identification number	Age

Part II: Certification

II. Health information

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:]

Part II: Certification

II. Health information

Transponder or tattoo					Validity of Vaccination		
Alphanumeric code of the animal	Date of implantation and/or reading (6) [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 blood sampling [dd/mm/yyyy]
1	2	3	4	5	6	7	8

Part II: Certification

II. Health information

(1) ○ either [II.4. the consignment includes dogs destined for a Great Britain 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:

Part II: Certification

II. Health information

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
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Part II: Certification

II. Health information

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

Part II: Certification

II. Health information

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.

(*) Those countries subject to the transitional import arrangements include: an EU member State; Liechtenstein; Norway and Switzerland.

References to European Union legislation within this certificate are references to direct EU legislation which has been retained in Great Britain (retained EU law as defined in the European Union (Withdrawal) Act 2018).

References to Great Britain in this certificate include Channel Islands and Isle of Man.

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.20.: 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.16: Do not use this box until the end of the transitional staging period.

Box I.25.: Identification system: select transponder or tattoo.

Identification number: indicate the transponder or tattoo alphanumeric code.

Part II: Certification

II. Health information

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;
- 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 into Great Britain;
- 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 Great Britain.

Certifying Officer

Name (in capital letters) Qualification and title

Date of signature Signature

Stamp