

**CHAPTER 38: MODEL ANIMAL HEALTH CERTIFICATE FOR ENTRY INTO THE UNION OF DOGS, CATS AND FERRETS (MODEL ‘CANIS-FELIS-FERRETS’)**

<b>Part I: Description of consignment</b>	<b>COUNTRY UNITED STATES</b>		<b>ANIMAL HEALTH CERTIFICATE TO THE EU</b>		
	<b>I.1 Consignor/Exporter</b> Name		<b>I.2 Certificate Reference</b>	<b>I.2a IMSOC reference</b>	
	Address		<b>I.3 Central Competent Authority</b> USDA APHIS Veterinary Services	<b>QR CODE</b>	
	Country                      ISO country code		<b>I.4 Local Competent Authority</b>		
	<b>I.5 Consignee/Importer</b> Name		<b>I.6 Operator responsible for the consignment</b>		
	Address				
	Country                      ISO country code				
	<b>I.7 Country of Origin</b> ISO Country Code		<b>I.9 Country of destination</b> ISO country code		
	United States                      US				
	<b>I.8 Region of origin</b> Code		<b>I.10 Region of destination</b> Code		
	<b>I.11 Place of dispatch</b> Registration/Approval No		<b>I.12 Place of destination</b> Registration/Approval No		
	Name		Name		
	Address		Address		
Country                      ISO country code		Country                      ISO country code			
United States                      US					
<b>I.13 Place of loading</b>		<b>I.14 Date and time of departure</b>			
<b>I.15 Means of Transport</b>		<b>I.16 Entry Border Control Post</b>			
Aircraft                      Vessel		<b>I.17 Accompanying documents</b>			
Railway                      Road Vehicle					
Identification		Type			
		Country			
		ISO country code			
		Commercial document reference			
		Code			

**Certificate Reference**

Part I: Description of consignment	<b>I.18 Transport Conditions</b>		Ambient	Chilled	Frozen	
	<b>I.19 Container number/Seal number</b>					
	Container No			Seal No		
	<b>I.20 Certified as or for</b>					
	Further keeping	Confined establishment	Quarantine establishment	Other		
	<b>I.21 For transit</b>			<b>I.22 For internal market</b>		
	Third country			ISO country code		
	<b>I.23</b>					
	<b>I.24 Total number of packages</b>		<b>I.25 Total quantity</b>		<b>I.26 Total net weight/gross weight (kg)</b>	
	<b>I.27 Description of consignment</b>					
CN code		Nature of commodity		Test		
010619		Pet animal(s)				
Species	Subspecies/Category	Sex	Identification system	Identification number	Date of Birth	Quantity

**II. Health information**

**II.a Certificate reference**

**II.b IMSOC reference**

I, the undersigned official veterinarian hereby certify that the animals described in Part I:

II.1. come from a country, territory or zone thereof with code: US-0<sup>(1)</sup> which, on the date of issue of this certificate is authorised for the entry into the Union of dogs, cats and ferrets and is listed in Part 1 of Annex VIII to Commission Implementing Regulation (EU) 2021/404;

[II.2.]

[II.3.]

II.4 have been subjected with negative result to a clinical inspection, carried out by an official veterinarian in the third country, territory or zone thereof of origin within 48 hour period prior to loading for dispatch to the Union for the detection of signs indicative of the occurrence of diseases, including the relevant listed diseases referred to in Annex 1 of Delegated Regulation (EU) 2020/692 and emerging diseases.

[II.5. 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<sup>(5)</sup> 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<sup>(6)</sup>, and

[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;]

Part II: Certification

Transponder		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 <sup>(9)</sup> [dd/mm/yyyy]				From [dd/mm/yyyy]	To [dd/mm/yyyy]	
1	2	3	4	5	6	7	8
			&				/
			&				
			&				
			&				
			&				
			&				

[II.6. the dogs have not been treated against infestation with *Echinococcus multilocularis*.]

II.a Certificate reference

II.b IMSOC reference

**Notes:**

This certificate is intended for commercial entries into the Union of dogs, cats and ferrets, including when they are destined to a confined establishment or to an approved quarantine establishment and when the Union is not the final destination of the animals and for entry into the Union of dogs, cats and ferrets moved in accordance with Article 5(4) of Regulation (EU) No 576/2013 of the European Parliament and of the Council.

In accordance with the Agreement on the withdrawal of the United Kingdom of Great Britain and Northern Ireland from the European Union and the European Atomic Energy Community, and in particular Article 5(4) of the Protocol on Ireland / Northern Ireland in conjunction with Annex 2 to that Protocol, references to European Union in this certificate include the United Kingdom in respect of Northern Ireland.

This animal health certificate shall be completed according to the notes for the completion of certificates provided for in Chapter 4 of Annex I to Commission Implementing Regulation (EU) 2020/2235.

**Part I:**

Box I.20: Certified as or for: indicate

- "Further keeping" where dogs, cats or ferrets are moved in accordance with Title V of Part II of Delegated Regulation (EU) 2020/692;
- Confined establishment: as defined in Article 4(48) of Regulation (EU) 2016/429 of the European Parliament and of the Council;
- Approved quarantine establishment: as defined in Article 3(9) of Commission Delegated Regulation (EU) 2020/688
- "others" 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.

**Part II:**

- (1) Code of the zone as it appears in Column 2 of Part 1 of Annex VIII to Implementing Regulation (EU) 2021/404.
- (2) Keep as appropriate.
- (3) Not applicable to the movement of dogs, cats and ferrets other than non-commercial movements kept as pet animals in households that cannot be carried out in accordance with the conditions laid down in Article 245(2) or Article 246(1) and (2) of Regulation (EU) 2016/429.
- (4) Date of loading: it cannot be a date prior to the date of authorisation of the zone for entry into the Union, or a date in a period when restriction measures have been adopted by the Union against entries of these animals from the zone.
- (5) Any revaccination must be considered a primary vaccination if it was not carried out within the period of validity of a previous vaccination.
- (6) A certified copy of the identification and vaccination details of the animals concerned shall be attached to the certificate.
- (7) The rabies antibody titration test referred to in point II.5:
  - 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 an official laboratory;
  - 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 official laboratory on the result of the rabies antibody test referred to in point II.5. 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.5.
- (9) In conjunction with footnote (6), the marking of the animals concerned by the implantation of a transponder 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 infestation with *Echinococcus multilocularis* referred to in point II.6 must:

Part II: Certification

II.a Certificate reference

II.b IMSOC reference

Part II: Certification

- be administered by a veterinarian within a period of not more than 48 hours and ending 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.

(11) The table referred to in point II.6 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)

Signature

Date

Qualification and title

Stamp

**Official veterinarian**

Name (in capital letters)

Signature

Date

Qualification and title

Stamp