Part I: Description of consignment

I.1. Consignor
Name
Address
Country ISO Code

I.2. IMSOC reference

I.2.a. Local reference

I.3. Central Competent Authority

I.4. Local Competent Authority

I.5. Consignee
Name
Address
Country ISO Code

I.6. Operator conducting assembly operations independently of an establishment
Name
Address
Approval Number
Country ISO Code

I.7. Country of origin ISO Code

I.8. Region of origin Code

I.9. Country of destination ISO 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. Transporter
Name
Address
Approval Number
Country ISO Code

I.17. Accompanying documents
[en] accompanying.document.number
Date of issue
Country
Place of issue

I.18. Transport conditions
Ambient ☐ Frozen ☐ Chilled ☐

I.19. Container No / Seal No

I.20. Certified as

Further process ☐	Event or activity near borders ☐	Relaying ☐	Travelling circus/animal act ☐
Dispatch centre ☐	Products for human consumption ☐	Registered equidae ☐	Release into the wild ☐
Germinal products ☐	Organic fertilizers and soil improvers ☐	Slaughter ☐	Further keeping ☐
Technical use ☐	Confined establishment ☐	Ornamental aquaculture establishment ☐	Exhibition ☐
Other ☐	Live aquatic animals for human consumption ☐	Quarantine or similar establishment ☐	

I.21. For transit through a third country ☐
Third country ISO Code
Exit point BCP code
Entry point BCP code

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

I.23. For export ☐
Third country ISO Code
Exit point BCP code

I.24. Estimated journey time

I.25. Journey Log

I.27. Total quantity

I.28. Total gross weight

Part I: Description of consignment

I.30. Description of consignment

Commodity	Species	Sex	Identification system	Identification Number

Quantity	Age

Part II: Certification

II. Health information

I, the undersigned official veterinarian, hereby certify:

II.1. The other carnivores(1)(2) of the consignment described in Part I meet the following requirements:

II.1.1. They are identified:

(3) ☐ either [individually;]

(3) ☐ and/or [as a group of animals of the same species kept together during the movement to destination;]

II.1.2. They have undergone a clinical examination or a clinical inspection on _____ (insert date dd/mm/yyyy) within the the 48 hour period prior to departure and have not shown symptoms or clinical signs of diseases.

II.1.3. They come from a registered or approved establishment in which infection with rabies virus in kept terrestrial animals has not been reported during the 30 day period prior to departure and in which to the best of my knowledge and as declared by the operator, there were no abnormal mortalities with an undetermined cause.

II.2. The other carnivores(1) of the consignment described in Part I meet the following requirements:

(3) ○ either [II.2.1. They have received a complete primary course of anti-rabies vaccination and at least 21 days have elapsed since the completion of the primary anti-rabies vaccination carried out in accordance with the validity requirements set out in Part I of Annex VII to Commision Delegated Regulation (EU) 2020/688, and any subsequent revaccination was carried out within the period of validity of the preceding vaccination.]

(3) ○ or [II.2.1. [They are intended for direct transport in accordance with Article 58(2) of Delegated Regulation (EU) 2020/688 to:

(3) ○ either [the confined establishment indicated in Box I.20 of Part I;]

(3) ○ or [the establishment indicated in Box I.20 of Part I where these animals are kept as fur animals as defined in point 1 of Annex I to Commission Regulation (EU) No 142/2011.]]

☐ [II.2.2. The canidae, other than dogs, due to their scheduled destination(4) indicated in Box I.10, or in Box I.11 where regionalisation is applied:

(3) ○ either [have been treated against Echinococcus multilocularis in accordance with Part 2(2) of Annex VII to Delegated Regulation (EU) 2020/688;]]

(3) ○ or [have not been treated against(5) Echinococcus multilocularis;]]

Identification	Anti-echinococcus treatment		Administering veterinarian
	Name and manufacturer of the product	Date [dd/mm/yyyy] and time [00:00]of treatment	Name in capitals, stamp and signature
_____	_____	_____	_____
_____	_____	_____	_____
_____	_____	_____	_____
_____	_____	_____	_____

Part II: Certification

II. Health information

(3) ○ or [are intended for direct transport in accordance with Article 58(2) of Delegated Regulation (EU) 2020/688 to:

(3) ○ either [the confined establishment indicated in Box I.20 of Part I.]]]

(3) ○ or [the establishment indicated in Box I.20 of Part I.]where these animals are kept as fur animals as defined in point (1) of Annex I to Regulation (EU) No 142/2011.]]]

II.3. Arrangements are made to transport the consignment in accordance with Article 4 of Delegated Regulation (EU) 2020/688.

II.4. This certificate is valid for 10 days from the date of issuing. In the case of transport by waterway/sea of animals, the period of validity of the certificate may be extended by the duration of the journey by waterway/sea.

Animal welfare attestation

At the time of inspection, the animals covered by this health certificate were fit to be transported in accordance with the provisions of Council Regulation (EC) No 1/2005 on the intended journey due to start on _____ (insert date).

Notes:

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 2 of Annex I to Commission Implementing Regulation (EU) 2020/2235.

Part I:

Box reference I.11: “Place of dispatch”: Indicate a registered or an approved establishment of dispatch.

Box reference I.12: “Place of destination”: Indicate a registered or an approved establishment of destination.

Box reference I.30: “Identification number”: Indicate for each animal of the consignment its identification.

Part II:

(1) There can be one or more animals in the consignment.

(2) Other carnivores means animals of the species belonging to the order Carnivora other than dogs, cats and ferrets as defined in Article 3(32) of Delegated Regulation (EU) 2020/688

(3) Delete if not applicable.

(4) Member States or parts thereof listed in the Annex to Commission Implementing Regulation (EU) 2018/878.

(5) The table referred to in point II.2.2. must be used to document the details of the treatment against Echinococcus multilocularis, in accordance with Part 2(2) of Annex VII to Delegated Regulation (EU) 2020/688, if administered after the date the certificate was signed and prior to the scheduled entry into Member States or parts thereof listed in the Annex to Commission Implementing Regulation (EU) 2018/878.

Certifying Officer/Official veterinarian

Name (in capital letters) Qualification and title

Date of signature Signature

Stamp