EUROPEAN UNION **INTRA**

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.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. 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
Breeding ☐ Relaying ☐ Ornamental aquaculture establishment ☐ Other ☐
Quarantine establishment ☐ Live aquatic animals for human consumption ☐

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.26. Total number of packages
I.28. Total net weight

I.27. Total quantity
I.28. Total gross weight

I.30. Description of consignment

Commodity	Species	Quantity	Package count	Net weight

Part II: Certification

II. Health information

I, the undersigned official veterinarian, hereby certify:

II.1. According to official information, the aquatic animals in the consignment described in Part I meet the following animal health requirements:

II.1.1. The aquatic animals do not originate from (1) ☐ [an establishment] (1) ☐ [a habitat] which is subject to the movement restrictions or the emergency measures referred to in Article 191(2)(b)(i) and (ii) of Regulation (EU) 2016/429 which have been established to control listed diseases for which the aquatic animals in the consignment are listed species, or emerging diseases;

II.1.2. The aquatic animals:

(1) ○ either [originate from (1) ☐ [an establishment] (1) ☐ [a habitat] where there are no increased mortalities with an undetermined cause.]

(1) ○ or [originate from a part of (1) ☐ [an establishment] (1) ☐ [a habitat] which is independent of a epidemiological unit where increased mortalities or disease symptoms have occurred, and the Member State of destination (1) ☐ [and the Member State (1) ☐ [s] of transit] (1) ☐ [has] (1) ☐ [have] given consent for the movement to occur.]

(1) ☐ [II.2. Aquaculture animals in the consignment described in Part I meet the following requirements:

II.2.1. They come from an aquaculture establishment which is (1) ☐ [registered in accordance with Article 173 of Regulation (EU) 2016/429] (1) ☐ [approved in accordance with Article 176 or Article 177 of Regulation (EU) 2016/429] where mortality records, movement records and health and production records are regularly updated and a documentary check on those records has been carried out within a period of 72 hours prior to the time of departure and has not indicated any cause for concern;

II.2.2. The aquaculture animals:

(1) ○ either [have undergone a clinical inspection and where relevant, a clinical examination in accordance with Article 15(1)(b) of Commission Delegated Regulation (EU) 2020/990 carried out within a period of 72 hours prior to the time of departure and have not shown symptoms of relevant listed diseases or emerging diseases.]

(1) ○ or [are (1) ☐ [eggs] (1) ☐ [molluscs] which do not require a clinical inspection within a period of 72 hours prior to the time of departure as they are subject to the derogation laid down in Article 15(2) of Commission Delegated Regulation (EU) 2020/990.]]

(1)(2) ☐ [II.3. Requirements for (3)listed species for Viral haemorrhagic septicaemia (VHS), Infectious haematopoietic necrosis (IHN), infection with HPR-deleted infectious salmon anaemia virus (ISAV), infection with Marteilia refringens, infection with Bonamia exitiosa, infection with Bonamia ostreae, and infection with White spot syndrome virus

The aquatic animals referred to in Part I:

(1) ○ either [originate from a (1) ☐ [Member State] (1) ☐ [zone] (1) ☐ [compartment] declared free from (1) ☐ [VHS] (1) ☐ [IHN] (1) ☐ [infection with HPR-deleted ISAV] (1) ☐ [infection with Marteilia refringens] (1) ☐ [infection with Bonamia ostreae] (1) ☐ [infection with Bonamia exitiosa] (1) ☐ [infection with White spot syndrome virus] in accordance with Chapter 4 of Part II of Commission Delegated Regulation (EU) 2020/689.]

(1) ○ or [originate from a (1) ☐ [Member State] (1) ☐ [zone] (1) ☐ [compartment] under an eradication programme for (1) ☐ [VHS] (1) ☐ [IHN] (1) ☐ [infection with HPR-deleted ISAV] (1) ☐ [infection with Marteilia refringens] (1) ☐ [infection with Bonamia ostreae] (1) ☐ [infection with Bonamia exitiosa] (1) ☐ [infection with White spot syndrome virus], and are destined for a Member State, zone or compartment which is also subject to an eradication programme for the same disease, in accordance with the derogation laid down in Article 198 of Regulation (EU) 2016/429.]

(1) ○ or [are one of the vector species listed in column 4 of the table in the Annex to Commission Implementing Regulation (EU) 2018/1882 and they are not regarded as vectors of the category B or category C diseases in question.]]

(1)(4) ☐ [II.4. Requirements for species susceptible to Koi herpes virus disease (KHV), infection with Spring viraemia of carp virus (SVC), Bacterial kidney disease (BKD), infection with Infectious pancreatic necrosis virus (IPN), infection with Gyrodactylus salaris (GS), infection with Salmonid alphavirus (SAV) and infection with Ostreid herpes virus 1 µvar (OsHV-1 µvar)

Part II: Certification

II. Health information

The consignment originates from a (1) ☐ [Member State], (1) ☐ [zone] (1) ☐ [compartment] which fulfils the health guarantees as regards (1) ☐ [KHV], (1) ☐ [SVC], (1) ☐ [BKD], (1) ☐ [IPN], (1) ☐ [GS], (1) ☐ [SAV], (1) ☐ [OsHV-1 µvar] which are necessary to comply with the national measures which apply in the Member State of destination, as provided for in implementing acts adopted by the Commission in accordance with Article 226(3) of Regulation (EU) 2016/429.]

II.5. To the best of my knowledge, and as declared by the operator, the aquatic animals in the consignment show no disease symptoms and come from (1) ☐ [an establishment] (1) ☐ [a habitat] where:

(i) there were no abnormal mortalities with an undetermined cause; and

(ii) the animals have not been in contact with kept animals of (4)listed species which did not comply with the requirements referred to in point II.1.

II.6. Transport requirements

Arrangements have been made to transport the consignment in accordance with the provisions laid down in Articles 3 and 4 of Commission Delegated Regulation (EU) 2020/990.

II.7. Labelling requirements

Arrangements have been made to identify and label (1) ☐ [the means of transport] (1) ☐ [containers] in accordance with Article 5 of Commission Delegated Regulation (EU) 2020/990, and the consignment is identified by (1) ☐ [a legible and visible label on the exterior of the container] (1) ☐ [a legible and visible label on the exterior of the means of transport](1) ☐ [an entry in the ship's manifest when transported by well boat], which clearly links the consignment to this animal health certificate.

II.8. Validity of the animal health certificate

This animal health certificate is valid for a period of 10 days from the date of issuing. In the case of transport by waterway/sea of aquatic animals, this period of 10 days may be extended by the duration of the journey by waterway/sea.

Part II: Certification

II. Health information

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.

'Aquatic animals' are animals as defined in point (3) of Article 4 of Regulation (EU) 2016/429. 'Aquaculture animals' are aquatic animals which are subject to aquaculture as defined in point (7) of Article 4 of Regulation (EU) 2016/429.

Part II of this certificate does not apply to the following aquatic animals:

(a) live molluscs and live crustaceans which are packaged and labelled for human consumption in accordance with the specific requirements for those animals as set out in Sections VII and VIII of Annex III to Regulation (EC) No 853/2004 and which are no longer able to survive as living animals if returned to the aquatic environment;

(b) live molluscs and live crustaceans which are intended for human consumption without further processing, provided they are packaged for retail sale in accordance with the specific requirements for those animals as set out in Sections VII and VIII of Annex III to Regulation (EC) No 853/2004;

(c) molluscs which are packaged and labelled for human consumption in accordance with the specific requirements for those animals as set out in Sections VII and VIII of Annex III to Regulation (EC) No 853/2004 and which are intended for further processing without temporary storage at the place of processing.

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 II:

(1) Keep as appropriate/delete if not applicable.

(2) Only applicable when the Member State/zone/compartment of destination either has disease-free status for a category C disease as defined in point (3) of Article 1 of Commission Implementing Regulation (EU) 2018/1882 or is subject to an optional eradication programme established in accordance with Article 31(2) of Regulation (EU) 2016/429.

(3) Listed species as referred to in columns 3 and 4 of the table in the Annex to Commission Implementing Regulation (EU) 2018/1882.

(4) Only applicable when the Member State of destination has approved national measures for a specific disease in place, which have been approved by the Commission in accordance with Article 226 of Regulation (EU) 2016/429.

Certifying Officer/Official veterinarian

Name (in capital letters) | Qualification and title

Date of signature | Signature

Stamp