

Part I: Description of consignment	I.1. Consignor		I.2. IMSOC reference		I.2.a. Local reference	
	Name				I.3. Central Competent Authority	
	Address					
	Country		ISO Code		I.4. Local Competent Authority	
	I.5. Consignee			I.6. Operator conducting assembly operations independently of an establishment		
	Name			Name		
	Address			Address		
	Country			Country		
	ISO Code			ISO Code		
	I.7. Country of origin			I.9. Country of destination		
ISO Code			ISO Code			
I.8. Region of origin			I.10. Region of destination			
Code			Code			
I.11. Place of dispatch			I.12. Place of destination			
Name			Name			
Address			Address			
Approval Number			Approval Number			
Country			Country			
ISO Code			ISO Code			
I.13. Place of loading			I.14. Date and time of departure			
Name						
Address						
Approval Number						
Country						
ISO Code						
I.15. Means of Transport			I.16. Transporter			
Mode	International transport document	Identification	Name			
			Address			
			Approval Number			
			Country			
			ISO Code			
			I.17. Accompanying documents			
			Accompanying document reference			
			Date of issue			
			Country			
			Place of issue			
I.18. Transport conditions						
Ambient <input type="checkbox"/>		Frozen <input type="checkbox"/>		Chilled <input type="checkbox"/>		
I.19. Container No / Seal No						
I.20. Certified as						
Other <input type="checkbox"/>		Release into the wild <input type="checkbox"/>		Breeding <input type="checkbox"/>		
Quarantine establishment <input type="checkbox"/>		Relaying <input type="checkbox"/>		Live aquatic animals for human consumption <input type="checkbox"/>		
Ornamental aquaculture establishment <input type="checkbox"/>						
I.21. For transit through a third country <input type="checkbox"/>						
Third country			ISO Code			
Exit point			BCP code			
Entry point			BCP code			
I.22. For transit through Member State(s) <input type="checkbox"/>			I.23. For export <input type="checkbox"/>			
Member State		ISO Code		Third country		
				ISO Code		
				Exit point		
				BCP code		
I.24. Estimated journey time			I.25. Journey Log			
I.26. Total number of packages		I.27. Total quantity		I.28. Total net weight		
I.30. Description of consignment						
1. 03 FISH AND CRUSTACEANS, MOLLUSCS AND OTHER AQUATIC INVERTEBRATES						
0301 Live fish						
Other live fish						
030193 Carp						

03019300 Carp (Cyprinus spp., Carassius spp., Ctenopharyngodon idellus, Hypophthalmichthys spp., Cirrhinus spp., Mylopharyngodon piceus, Catla catla, Labeo spp., Osteochilus hasselti, Leptobarbus hoeveni, Megalobrama spp.)

#1.	Commodity	Species	Quantity	Package count	Net weight
Part I: Description of consignment					

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) <input type="checkbox"/> [an establishment] (1) <input type="checkbox"/> [a habitat] which is subject to the movement restrictions or the emergency measures referred to in Article 191(2), points (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) <input type="radio"/>	[originate from (1) <input type="checkbox"/> [an establishment] (1) <input type="checkbox"/> [a habitat] where there are no increased mortalities with an undetermined cause.]		
(1) <input type="radio"/> or	[originate from a part of (1) <input type="checkbox"/> [an establishment] (1) <input type="checkbox"/> [a habitat] which is independent of the epidemiological unit where increased mortalities or disease symptoms have occurred, and the Member State of destination (1) <input type="checkbox"/> [and the Member State(1) <input type="checkbox"/> [s] of transit] (1) <input type="checkbox"/> [has] (1) <input type="checkbox"/> [have] given consent for the movement to occur.]		
(1) <input type="checkbox"/>	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) <input type="checkbox"/> [registered in accordance with Article 173 of Regulation (EU) 2016/429] (1) <input type="checkbox"/> [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) <input type="radio"/> either	[have undergone a clinical inspection and where relevant, a clinical examination in accordance with Article 15(1), point (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) <input type="radio"/> or	[are (1) <input type="checkbox"/> [eggs] (1) <input type="checkbox"/> [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 Delegated Regulation (EU) 2020/990.]		
(1) <input type="radio"/> or	[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 16(1) of Delegated Regulation (EU) 2020/990.]		
(1)(2) <input type="checkbox"/>	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		
II.3.	Marteilia refringens, infection with Bonamia exitiosa, infection with Bonamia ostreae, and infection with White spot syndrome virus		
	The aquatic animals described in Part I:		
(1) <input type="radio"/> either	[originate from a (1) <input type="checkbox"/> [Member State] (1) <input type="checkbox"/> [zone] (1) <input type="checkbox"/> [compartment] declared free from (1) <input type="checkbox"/> [VHS] (1) <input type="checkbox"/> [IHN] (1) <input type="checkbox"/> [infection with HPR-deleted ISAV] (1) <input type="checkbox"/> [infection with Marteilia refringens] (1) <input type="checkbox"/> [infection with Bonamia ostreae] (1) <input type="checkbox"/> [infection with Bonamia exitiosa] (1) <input type="checkbox"/> [infection with White spot syndrome virus] in accordance with Part II, Chapter 4, of Commission Delegated Regulation (EU) 2020/689.]		
(1) <input type="radio"/> or	[originate from a (1) <input type="checkbox"/> [Member State] (1) <input type="checkbox"/> [zone] (1) <input type="checkbox"/> [compartment] which is under an eradication programme for (1) <input type="checkbox"/> [VHS] (1) <input type="checkbox"/> [IHN] (1) <input type="checkbox"/> [infection with HPR-deleted ISAV] (1) <input type="checkbox"/> [infection with Marteilia refringens] (1) <input type="checkbox"/> [infection with Bonamia ostreae] (1) <input type="checkbox"/> [Infection with Bonamia exitiosa] (1) <input type="checkbox"/> [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) <input type="radio"/> or	[are wild aquatic animals which have completed quarantine in an establishment approved in accordance with Article 15 of Commission Delegated Regulation (EU) 2020/691 and are regarded as disease-free.]		

Part II: Certification	II. Health information	
	(1) <input type="radio"/> 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 relevant listed disease as they do not fulfil the conditions set out in Annex I to Commission Delegated Regulation (EU) 2020/990.]
	(1) <input type="radio"/> 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 are regarded as vectors but they have been subject to quarantine in an establishment approved in accordance with Article 15 in Commission Delegated Regulation (EU) 2020/691, and are regarded as disease-free.]
	(1) <input type="radio"/> 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 are regarded as vectors but they have been kept in isolation in an establishment approved in accordance with Article 16 of Commission Delegated Regulation (EU) 2020/691 and are no longer regarded as vectors.]
	(1) <input type="radio"/> or	[are aquaculture animals originating from a confined establishment and are destined for a confined establishment in another Member State, both of which are approved in accordance with Article 9 of Commission Delegated Regulation (EU) 2020/691 and which comply with the provisions of Article 9(1) of Commission Delegated Regulation (EU) 2020/990.]
	(1) <input type="radio"/> or	[are aquaculture animals destined for a confined establishment approved in accordance with Article 9 of Commission Delegated Regulation (EU) 2020/691 and which comply with the requirements set out in Article 9(2), point (b) (1) <input type="checkbox"/> [(ii)] (1) <input type="checkbox"/> [(iii)] of Commission Delegated Regulation (EU) 2020/990.]
	(1) <input type="radio"/> or	[are aquaculture animals destined for a confined establishment approved in accordance with Article 9 of Commission Delegated Regulation (EU) 2020/691, for scientific purposes.]
	(1) <input type="radio"/> or	[are destined for a disease control aquatic food establishment approved in accordance with Article 11 of Commission Delegated Regulation (EU) 2020/691.]]
	(1)(4) <input type="checkbox"/>	Requirements for (5)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 μ var (OsHV-1 μ var)
	II.4.	The consignment originates from a (1) <input type="checkbox"/> [Member State], (1) <input type="checkbox"/> [zone] (1) <input type="checkbox"/> [compartment] which fulfils the health guarantees as regards (1) <input type="checkbox"/> [KHV], (1) <input type="checkbox"/> [SVC], (1) <input type="checkbox"/> [BKD], (1) <input type="checkbox"/> [IPN], (1) <input type="checkbox"/> [GS], (1) <input type="checkbox"/> [SAV], (1) <input type="checkbox"/> [OsHV-1 μ var] which are necessary to comply with the national measures which apply in the Member State of destination, and for which the Member State or part thereof is listed in (1) <input type="checkbox"/> [Annex I] (1) <input type="checkbox"/> [Annex II] to Commission Implementing Decision (EU) 2021/260.]
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) <input type="checkbox"/> [an establishment] (1) <input type="checkbox"/> [a habitat] where: (i) there were no abnormal mortalities with an undetermined cause; and (ii) the animals have not been in contact with aquatic animals of (3)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 Delegated Regulation (EU) 2020/990.	
II.7.	Labelling requirements Arrangements have been made to identify and label (1) <input type="checkbox"/> [the means of transport] (1) <input type="checkbox"/> [containers] in accordance with Article 5 of Delegated Regulation (EU) 2020/990, and the consignment is identified by (1) <input type="checkbox"/> [a legible and visible label on the exterior of the container] (1) <input type="checkbox"/> [a legible and visible label on the exterior of the means of transport] (1) <input type="checkbox"/> [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		
	<p>Notes:</p> <p>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.</p> <p>'Aquatic animals' are animals as defined in Article 4, point (3), of Regulation (EU) 2016/429. 'Aquaculture animals' are aquatic animals which are subject to aquaculture as defined in Article 4, point (7), of Regulation (EU) 2016/429.</p> <p>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.</p> <p>Part II:</p> <p>(1) Keep as appropriate/delete if not applicable.</p> <p>(2) Only applicable when the Member State/zone/compartiment of destination either has disease-free status for a Category C disease as defined in Article 1, point (3), of 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.</p> <p>(3) Listed species as referred to in columns 3 and 4 of the table set out in the Annex to Implementing Regulation (EU) 2018/1882.</p> <p>(4) Only applicable when the Member State of destination, or part thereof, has approved national measures for a specific disease as listed in Annex I or Annex II to Implementing Decision (EU) 2021/260, otherwise delete.</p> <p>(5) Susceptible species as referred to in the second column of the table in Annex III to Implementing Decision (EU) 2021/260.</p>		
	<p>Certifying Officer/Official veterinarian</p> <p>Name (in capital letters)</p> <p>Date of signature</p> <p>Stamp</p>	<p>Qualification and title</p> <p>Signature</p>	