EUROPEAN UNION INTRA

	I.1. Consignor			I.2. IMSOC reference	I.2.a. Local refere	nce	
	Name Address				I.3. Central Comp	etent Authority	
	Country ISO Code				I.4. Local Compet	ent Authority	
or consignment	I.S. Consignee Name Address Country ISO Code			I.6. Operator conducting assembly establishment Name Address Approval Number Country	I.6. Operator conducting assembly operations independently of an establishment Name Address Approval Number		
	I.7. Country of origin ISO Code			I.9. Country of destination		ISO Code	
Part I: Description	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 load	ing		I.14. Date and time of departure			
	Name Address Approval Number Country		ISO Code				
	I.15. Means of Tra Mode	insport International	Identification	I.16. Transporter			
	Mode	transport document	identification	Name Address			
				Approval Number Country	ISO Code		
				I.17. Accompanying documents Accompanying document reference Date of issue Country Place of issue			
- 1	I.18. Transport conditions Ambient □ Frozen □			Chilled [Chilled □		
	I.19. Container No	/ Seal No					
	.20. Certified as Other □ Release into the wild □ Ornamental aquaculture Relaying □ establishment □			Breeding \square Live aquatic animals for human consumption \square	Quarantine estab	lishment \square	
	I.21. For transit th	rough a third cour	ıtry				
	Third country Exit point Entry point			ISO Code BCP code BCP code			
	I.22. For transit through Member State(s)			I.23. For export			
	Member State			Third country Exit point	ISO Code BCP code		
١	I.24. Estimated jou		T	I.25. Journey Log			
ı	I.26. Total number		I.27. Total quantity	I.28. Total net weight	I.28. Total gross v	veight	
- 1	I.30. Description of	_					
	1. 03 FISH AND CI 0301 Live fish Other live fis 030193 Ca	sh	LUSCS AND OTHER AQUATIC IN	NVERTEBRATES			

en 1 / 5

EUROPEAN UNION INTRA

	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		
ŗ							
Part I: Description of consignment							
guu							
nsi							
f co							
0 U							
ptio							
Cri							
Des							
τI:							
Par							

en 2 / 5

EUROPEAN UNION

	II. Health information							
	I, the undersigned official veterinarian, hereby certify:							
	II.1.	_	to official information, the aquatic a unimal health requirements:	animals in the consignment d	escribed in Part I meet the			
Part II: Certification	is subject to the movemen			ate from (1) \square [an establishmotions or the emergency measualation (EU) 2016/429 which he aquatic animals in the cons	res referred to in Article ave been established to			
Cer		II.1.2.	The aquatic animals:					
Part II:		(1) o either	[originate from (1) \square [an establish mortalities with an undetermined		there are no increased			
		(1) ∘ or	[originate from a part of (1) \square [an establishment] (1) \square [a habitat] which is independent of the epidemiological unit where increased mortalities or disease symptoms have occurred, and the Member State of destination (1) \square [and the Member State (1) \square [s] of transit] (1) \square [has] (1) \square [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 Article 173 of Regulation (EU) 2016/429] (1) ☐ [approved in accordance with Article Article 177 of Regulation (EU) 2016/429] where mortality records, movement records health and production records are regularly updated and a documentary check on the records has been carried out within a period of 72 hours prior to the time of departments and indicated any cause for concern:						
		II.2.2.	has not indicated any cause for concern; The aquaculture animals:					
		11.2.2. (1) ○ either	[have undergone a clinical inspection and where relevant, a clinical examination in					
		eithei	carried out within a period of 72 hours prior to the time of departure and have not symptoms of relevant listed diseases or emerging diseases.]					
		(1) ○ or	[are (1) ☐ [eggs] (1) ☐ [molluscs] w of 72 hours prior to the time of dep Article 15(2) of Commission Delega	parture as they are subject to	the derogation laid down in			
	(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) \square [Member S (1) \square [VHS] (1) \square [IHN] (1) \square [infection Marteilia refringens] (1) \square [infection with Which Chapter 4, of Commission Delegate	ection with HPR-deleted ISAV on with Bonamia ostreae] (1) te spot syndrome virus] in ac] (1) □ [infection with □ [infection with Bonamia			
		(1) ○ or	[originate from a (1) □ [Member Seradication programme for (1) □ [(1) □ [infection with Marteilia refrese [infection with Bonamia exitiosa] (destined for a Member State, zone programme for the same disease, it of Regulation (EU) 2016/429.]	VHS] (1) [[IHN] (1) [[infectingens] (1) [[infection with (1) [[infection with White sport compartment which is also	tion with HPR-deleted ISAV] Bonamia ostreae] (1) ot syndrome virus], and are o subject to an eradication			
		(1) ○ or	[are one of the vector species listed Implementing Regulation (EU) 201 category B or category C diseases in	8/1882 and they are not regar				
	(1)(4) □ [II.4.	viraemia o	ents for (5)species susceptible to Koi f carp virus (SVC), Bacterial kidney rus (IPN), infection with Gyrodactyl	disease (BKD), infection with	Infectious pancreatic			

en 3/5

EUROPEAN UNION intended for human consumption – Model AQUA-INTRA-H						
	II. Health information					
		and infecti	on with Ostreid herpes virus 1 μναr	(OsHV-1 μvar)		
uo		fulfils the l□ [SAV], (1 the Member	health guarantees as regards (1) □ [: l) □ [OsHV-1 µvar] which are necess	mber State], (1) \square [zone] (1) \square [compartment] which [KHV], (1) \square [SVC], (1) \square [BKD], (1) \square [IPN], (1) \square [GS], (1) sary to comply with the national measures which apply in h the Member State or part thereof, is listed in (1) \square lementing Decision (EU) 2021/260.]		
tificat	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) \square [an establishment] (1) \square [a habitat] where:				
Cer		(i) there were no abnormal mortalities with an undetermined cause; and				
Part II: Certification		(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.				
Pē	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) \square [the means of transport] (1) \square [containers] in accordance with Article 5 of Delegated Regulation (EU) 2020/990, and the consignment is identified by (1) \square [a legible and visible label on the exterior of the container] (1) \square [a legible and visible label on the exterior of the means of transport](1) \square [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.					
	Notes					
	In accordance with the Agreement on the withdrawal of the United Kingdom of Great Britain and Northern Irela 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 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.

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

- live molluscs and live crustaceans which are packaged and labelled for human consumption (a) 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;
- molluscs which are packaged and labelled for human consumption in accordance with the (c) 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 Article 1, point (3), of Implementing Regulation (EU) 2018/1882 or

4/5

EUROPEAN UNION

	II. Health info	rmation					
		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 Implementing Regulation (EU) 2018/1882.					
cation	(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.					
Part II: Certification	(5)	Susceptible species as referred to in the second column of the table in Annex III to Implementing Decision (EU) 2021/260.					
t II:		ng Officer/Official veterinarian					
Par	Name (in capi Date of signat		Qualification and title Signature				
	Stamp						

en 5 / 5