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	I.1. Consignor			I.2. IMSOC reference	I.2.a. Local reference		
	Name Address				I.3. Central Competent Authority		
	Country ISO Code				I.4. Local Competent Authority		
of consignment	I.5. Consignee			I.6. Operator conducting assembly establishment	y operations independently of an		
Ĕ	Name Address			Name			
igi	Country		ISO Code	Approval			
ö				Approval Number			
ot c				Country	ISO Code		
Part I: Description	I.7. Country of origin ISO Code			I.9. Country of destination	ISO Code		
scrip	I.8. Region of origin Code			I.10. Region of destination	Code		
ne	I.11. Place of dispa	atch		I.12. Place of destination			
Τ:	Name Address				Name Address		
Par	Approval			Approval			
į.	Number Country		ISO Code	Nûmber	ISO Code		
			100 code	·	,		
	I.13. Place of loadi	ing		I.14. Date and time of departure	I.14. Date and time of departure		
	Name Address						
	Approval Number						
	Country		ISO Code				
	I.15. Means of Tra	nenort		I.16. Transporter			
	Mode	International	Identification	Name			
		transport document		Address			
				Approval Number			
				Country	ISO Code		
				I.17. Accompanying documents			
				Accompanying document reference	. , .		
				Date of issue	Date of issue Country		
				_			
	I.18. Transport conditions			Place of issue			
	Chilled \square	-		Ambient	Ambient \square		
	I.19. Container No	/ Seal No					
	I.20. Certified as		_	_	_		
	Breeding Live agreetic anim	ala far human	Quarantine establishment	Relaying \square	Other 🗆		
	consumption \square	ive aquatic animals for human Ornamental aquaculture establishment □					
	I.21. For transit through a third country						
	Third country			ISO Code			
	Exit point Entry point			BCP code BCP code			
	I.22. For transit through Member State(s)			I.23. For export			
	Member State	J :	ISO Code	Third country	ISO Code		
		24. Estimated journey time		Exit point	BCP code		
	I.24. Estimated jou			I.25. Journey Log			
	I.26. Total number	r of packages	I.27. Total quantity	I.28. Total net weight	I.28. Total gross weight		
	I.30. Description o	of consignment					
	1. 03 FISH AND CF	1. 03 FISH AND CRUSTACEANS, MOLLUSCS AND OTHER AQUATIC INVERTEBRATES					
	0301 Live fish Ornamental fish 030119 other than. 0301 11 00						

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	03011900 other than. 0301 11 00						
	#1.	Commodity	Species	Quantity	Package count	Net weight	
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Part I: Description of consignment							
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	II. Health information							
	, the undersigned official veterinarian, hereby certify:							
II.1. According to official information, the aquatic animals in the consignment described in Part I m following animal health requirements:					escribed in Part I meet the			
Part II: Certification	II.1.1. The aquatic animals do not origing is subject to the movement restricted (ii), of R control listed diseases for which or emerging diseases;			ions or the emergency measu alation (EU) 2016/429 which h	res referred to in Article ave been established to			
Cert		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 es Article 173 of Regulation (EU) 2016 Article 177 of Regulation (EU) 2016 health and production records are records has been carried out withi	5/429] (1) □ [approved in acco 5/429] where mortality record regularly updated and a docu n a period of 72 hours prior to	rdance with Article 176 or s, movement records and ımentary check on those			
		II.2.2.	has not indicated any cause for concern; The aquaculture animals:					
		(1) o	The aquaculture animals: [have undergone a clinical inspection and where relevant, a clinical examination in					
- 1		either	accordance with Article 15(1), point carried out within a period of 72 h symptoms of relevant listed diseas	nt (b), of Commission Delegate ours prior to the time of depa	d Regulation (EU) 2020/990			
		(1) ○ or	[are (1) \square [eggs] (1) \square [molluscs] w of 72 hours prior to the time of departicle 15(2) of Commission Delega	parture as they are subject to	the derogation laid down in			
	(1)(2) Requirements for (3)listed species for Value (III.3. necrosis (IHN), infection with HPR-deleted Marteilia refringens, infection with Bon with White spot syndrome virus			ectious salmon anaemia virus	(ISAV), infection with			
		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 Whith 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 Steradication programme for (1) □ [(1) □ [infection with Marteilia refr [infection with Bonamia exitiosa] (destined for a Member State, zone programme for the same disease, i 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			

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EUROPEAN UNION intended for human consumption – Model AQUA-INTRA-H						
	II. Health info	rmation				
		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	nber 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 the Member State or part thereof, is listed in (1) \square ementing Decision (EU) 2021/260.]		
II.5. To the best of my knowledge, and as declared by the operator, the aquatic animals in the cons 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 Irefrom 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 this certificate include the United Kingdom in respect of Northern Ireland.					lar Article 5(4) of the	
	Aquatic an	imals' are a	nimals as defined in Article 4, point	(3), of Regulation (EU) 2016/4	29. '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

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	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:	Certifying Officer/Official veterinarian Name (in capital letters) Qualification and title						
Par	Date of signat		Qualification and title Signature				
	Stamp						

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