EUROPEAN UNION INTRA

	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 Name Address			I.6. Operator conducting assembly operations independently of an establishment  Name  Address			
ısig	Country ISO Code			Approval			
S				Number Country	ISO Code		
				,			
tion	I.7. Country of origin ISO Code			I.9. Country of destination	ISO Code		
Part I: Description	I.8. Region of origi	in	Code	I.10. Region of destination	Code		
Des	I.11. Place of dispa	atch		I.12. Place of destination			
ij	Name			Name			
art	Annroyal			Address Approval			
щ	Approval Number			Number			
	Country		ISO Code	Country	ISO Code		
	I.13. Place of loadi	ng		I.14. Date and time of departure			
	Name						
	Address Approval						
	Number						
	Country		ISO Code				
	I.15. Means of Tra	T .	- 1	I.16. Transporter			
	Mode	International transport	Identification	Name			
		document		Address Approval			
				Number	100.0-1		
				Country	ISO Code		
				I.17. Accompanying documents			
				Accompanying document reference			
				Date of issue			
				Country Place of issue			
	I.18. Transport conditions						
	Ambient □ Frozen □			Chilled □			
	I.19. Container No	/ Seal No					
	I.20. Certified as						
	Other ☐ Release into the wild ☐ Ornamental aquaculture Releasing ☐			Breeding $\square$ Live aquatic animals for human	Quarantine establishment $\square$		
				consumption $\square$			
	I.21. For transit through a third country						
	Third country			ISO Code			
	Exit point			BCP code			
	Entry point  I.22. For transit through Member State(s)			I.23. For export			
				_			
	Member State			Third country Exit point	ISO Code BCP code		
	I.24. Estimated journey time			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	f consignment					
1. 03 FISH AND CRUSTACEANS, MOLLUSCS AND OTHER AQUATIC INVERTEBRATES							
	<b>0301</b> Live fish Other live fish	h s (Anguilla spp.)					

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EUROPEAN UNION INTRA

		03019230 Of a length of 12   cm or more but less than 20   cm							
	#1.	Commodity	Species	Quantity	Package count	Net weight			
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Part I: Description of consignment									
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## **EUROPEAN UNION**

	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:							
Part II: Certification		II.1.1.	The aquatic animals do not original is subject to the movement restrict 191(2), points (b)(i) and (ii), of Regular control listed diseases for which the or emerging diseases;	ions or the emergency measu alation (EU) 2016/429 which h	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 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		11.2.2. (1) ○ either	o [have undergone a clinical inspection and where relevant, a clinical examination in					
		eithei	carried out within a period of 72 h symptoms of relevant listed diseas	ours prior to the time of depa	•			
		(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) ☐ Requirements for (3)listed species for Viral had necrosis (IHN), infection with HPR-deleted in Marteilia refringens, infection with Bonamia 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 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			

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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	KHV], (1) $\square$ [SVC], (1) $\square$ [BKD sary to comply with the nation the Member State or part the	State], (1) $\square$ [zone] (1) $\square$ [compartment] which $\square$ , (1) $\square$ [SVC], (1) $\square$ [BKD], (1) $\square$ [IPN], (1) $\square$ [GS], (1) to comply with the national measures which apply in e Member State or part thereof, is listed in (1) $\square$ enting Decision (EU) 2021/260.]	
II.5. To the best of my knowledge, and as declared by the operator, the aquatic animals in the consi show no disease symptoms and come from (1) $\square$ [an establishment] (1) $\square$ [a habitat] where:					_	
Cer		(i)	there were no abnormal mortalities	es with an undetermined caus	e; 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 Ire 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 Unin this certificate include the United Kingdom in respect of Northern Ireland.					
	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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## **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:		Certifying Officer/Official veterinarian Name (in capital letters) Qualification and title						
Par	Date of signat		Qualification and title Signature					
	Stamp							

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