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	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   stablishment		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			

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	<b>03019300</b> 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		
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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.	_	to official information, the aquatic a unimal health requirements:	animals in the consignment d	escribed in Part I meet the			
Part II: Certification		II.1.1.	is subject to the movement restrict 191(2), points (b)(i) and (ii), of Regu	ate from (1) $\square$ [an establishment] (1) $\square$ [a habitat] which tions or the emergency measures referred to in Article ulation (EU) 2016/429 which have been established to ne aquatic animals in the consignment are listed species,				
		II.1.2.	The aquatic animals:					
		(1) o either	[originate from (1) $\square$ [an establish mortalities with an undetermined	ment] (1) $\square$ [a habitat] where there are no increased cause.]				
		(1) ○ or	[originate from a part of (1) $\square$ [an establishment] (1) $\square$ [a habitat] which is independent the epidemiological unit where increased mortalities or disease symptoms have occurrand the Member State of destination (1) $\square$ [and the Member State (1) $\square$ [s] of transit] (1) $\square$ [have] given consent for the movement to occur.]					
	(1) □ [II.2.	Aquacultur	ure animals in the consignment described in Part I meet the following requirements:					
		II.2.1.	They come from an aquaculture establishment which is (1) $\square$ [registered in accordance with Article 173 of Regulation (EU) 2016/429] (1) $\square$ [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) o either	[have undergone a clinical inspect accordance with Article 15(1), poin 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	ed 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)(3) □ [II.3.	1)(2)(3)  Requirements for (4)listed species for Viral ha necrosis (IHN), infection with HPR-deleted inf 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	(1)(2)[originate from a (1) $\square$ [Mem from (1) $\square$ [VHS] (1) $\square$ [IHN] (1) $\square$ Marteilia refringens] (1) $\square$ [infecti exitiosa] (1) $\square$ [infection with Whith Part II of Commission Delegated Reference of the commission Dele	[infection with HPR-deleted I on with Bonamia ostreae] (1) te spot syndrome virus] in acc	ISAV] (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, it of Regulation (EU) 2016/429.]	VHS] (1) □ [IHN] (1) □ [infect ringens] (1) □ [infection with (1) □ [infection with White sp or 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 aquaculture animals of one of Annex to Commission Implementing vectors of the relevant listed disease Commission Delegated Regulation	ng Regulation (EU) 2018/1882 se as they do not fulfil the cor	and they are not regarded as			
		(1) ○ or	[are aquaculture animals of one of	the vector species listed in co	olumn 4 of the table in the			

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(3)

	II. Health info	rmation				
Part II: Certification						
	Annex to 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 of Commission Delegated Regulation (EU) 2020/691, and are regarded as disease free.]					
	(1) or [are aquaculture animals of one of the vector species listed in column 4 in the table Annex to Implementing Regulation (EU) 2018/1882 and are regarded as vectors but have been kept in isolation in an establishment approved in accordance with Artic Commission Delegated Regulation (EU) 2020/691 and are no longer regarded as ve					
	(1)(5) □ [II.4.	viraemia o necrosis vi	ements for (6)species susceptible to Koi herpes virus disease (KHV), infection with Spring a of carp virus (SVC), Bacterial kidney disease (BKD), infection with Infectious pancreatic s virus (IPN), infection with Gyrodactylus salaris (GS), infection with Salmonid alphavirus (SAV) ection with Ostreid herpes virus 1 μναr (OsHV-1 μναr)			
Par	fulfils the health guarantees as regards (1) □ [l □ [SAV], (1) □ [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 h the Member State or part thereof, is listed in (1) $\square$ lementing Decision (EU) 2021/260.]		
	II.5.	To the best of my knowledge, and as declared by the operator, the animals in the consignment sh disease symptoms and originate from (1) $\square$ [an establishment] (1) $\square$ [a habitat] where:				
		(i)	there were no abnormal mortalities	es with an undetermined caus	e; and	
		(ii)	the animals have not been in contacomply with the requirements refe	_	listed species which did not	
	II.6. Transport requirements					
	Arrangements have been made to transport the consignment in accordance with the provisions of 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 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 Article 4, point (3), of Regulation (EU) 2010 are aquatic animals which are subject to aquaculture as defined in Article 4, point (7)						
This animal health certificate shall be completed according to the notes for the completion of for in Chapter 2 of Annex I to Commission Implementing Regulation (EU) 2020/2235.				ion of certificates provided		
	Part II:					
	(1) Keep as appropriate/delete if not applicable.					
	(2)	Article 199 originate f	in Regulation (EU)2016/429 and reqrom a Member State, zone or compa	ate of destination has taken measures in accordance with d requires that aquatic animals for release into the wild empartment which has disease-free status for a Category C f Implementing Regulation (EU) 2018/1882.		

Other than in the cases referred to in Note (2) of this Part, Section II.3 applies only when the Member

## **EUROPEAN UNION**

	II. Health info	rmation						
	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 is subject to an optional eradication programme established in accordance with Article 31(2) of Regulation (EU) 2016/429.							
	(4)	Listed species as referred to in columns 3 and 4 of the table in the Annex to Implementing Regulation (EU) 2018/1882.						
Part II: Certification	(5)	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.						
II: Cer	(6)	Susceptible species as referred to in the second column of the table in Annex III to Implementing Decision (EU) 2021/260.						
Part	Certifying Offi Name (in capi Date of signate Stamp		Qualification and title Signature					
	Statitp							

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