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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	
n of consignment	I.5. Consignee Name Address Country ISO Code I.7. Country of origin ISO Code			I.6. Operator conducting assembly operations independently of an establishment Name Address Approval Number Country ISO Code		
scription						
	I.8. Region of origin Code			I.10. Region of destination	Code	
Part I: Description	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 loadi	ing		I.14. Date and time of departure		
	Name Address Approval Number Country ISO Code					
	I.15. Means of Tra			I.16. Transporter		
	Mode	International transport document	Identification	Name Address Approval Number Country	ISO Code	
				I.17. Accompanying documents		
				Accompanying documents Accompanying document reference Date of issue Country Place of issue		
	I.18. Transport conditions			Ambient □		
	I.19. Container No	/ Seal No				
	I.20. Certified as Breeding Other Other		Quarantine establishment Live aquatic animals for human consumption	Release into the wild Ornamental aquaculture establishment	Relaying 🗆	
	I.21. For transit th	rough a third cour	try			
	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	Member State ISO Code		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	of consignment				
	1. 03 FISH AND CE 0301 Live fish Other live fis 030199 Oth	h	LUSCS AND OTHER AQUATIC INVER	RTEBRATES		

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		03019917 Other				
	#1.	Commodity	Species	Quantity	Package count	Net weight
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Part I: Description of consignment						
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	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 movement restrict 191(2), points (b)(i) and (ii), of Regu			ate from (1) \square [an establishment] (1) \square [a habitat] which cions or the emergency measures referred to in Article alation (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	r [originate from a part of (1) □ [an establishment] (1) □ [a habitat] which is independer the epidemiological unit where increased mortalities or disease symptoms have occurr and the Member State of destination (1) □ [and the Member State (1) □ [s] of transit] (1 [has] (1) □ [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.	I)(2)(3) Requirements for (4)listed species for Viral had necrosis (IHN), infection with HPR-deleted information 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 Artic 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 in Annex to Implementing Regulation (EU) 2018/1882 and are regarded as vectors but the have been kept in isolation in an establishment approved in accordance with Article Commission Delegated Regulation (EU) 2020/691 and are no longer regarded as vectors				
	(1)(5) □ [II.4.	viraemia o necrosis vi	nents for (6)species susceptible to Koi herpes virus disease (KHV), infection with Spring of carp virus (SVC), Bacterial kidney disease (BKD), infection with Infectious pancreatic virus (IPN), infection with Gyrodactylus salaris (GS), infection with Salmonid alphavirus (SAV) etion with Ostreid herpes virus 1 µvar (OsHV-1 µvar)			
		fulfils the h □ [SAV], (1 the Member	nealth guarantees as regards (1) 🗆 [l) 🗆 [OsHV-1 µvar] which are necess	which State], (1) \square [zone] (1) \square [compartment] which KHV], (1) \square [SVC], (1) \square [BKD], (1) \square [IPN], (1) \square [GS], (1) carry 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 animals in the consignmen 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.					
			animals as defined in Article 4, point hich are subject to aquaculture as d			
	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.					
	Applies in all cases where the Member State of destination has taken measures in accordance with Article 199 in Regulation (EU)2016/429 and requires that aquatic animals for release into the wild originate from a Member State, zone or compartment which has disease-free status for a Category C disease as defined in Article 1, point (3), of 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

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	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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