	I.1. Consignor				I.2. IMSOC refere	ence	I.2.a. Local refere	nce	
	Name						I.3. Central Comp	etent Authority	
	Address Country ISO Code					I.4. Local Compete	-		
					I.6. Operator conducting assembly operations independently of an				
ent	Name	I.5. Consignee				ducting assembly of	perations independ	iently of an	
Ĩ	Address				Name				
sign	Country ISO Code				Address Approval Numb	Address			
SUD					Country	ISO Code			
ofc	I.7. Country of orig	rin		ISO Code	I.9. Country of de	estination		ISO Code	
on	1.7. Country of of §	5111				estillation			
Part I: Description of consignment	I.8. Region of origi	8. Region of origin Code			I.10. Region of de	estination		Code	
scr	I.11. Place of dispa	itch			I.12. Place of des	tination			
De	Name Address				Name	Name Address			
tI:	Approval Number	r			Approval Number Country ISO Code				
Par	Country		ISO Code						
	I.13. Place of loadi	ng			L14. Date and tir	ne of departure	rture		
	Name	-			I.14. Date and time of departure				
	Address								
	Approval Number	r	ISO Codo						
	Country		ISO Code						
	I.15. Means of Tra Mode		Identification		I.16. Transporter				
	Mode	International transport document	Identification		Name Address				
		document			Activity ID				
					Country ISO Code				
					I.17. Accompany	ing documents			
					Accompanying d reference				
					Date of issue Country				
	I.18. Transport cor	nditions			Place of issue				
	Chilled	hilled Frozen			Ambient 🗆				
		I.19. Container No / Seal No							
	I.20. Certified as				Live aquatic anii	nals for human	Querentine estab	lishment 🗖	
		eelaying Other Other			consumption				
	Ornamental aquae	rnamental aquaculture Breeding 🗆							
	.21. For transit through a third country								
	Third country Exit point			ISO Code BCP code					
	Entry point					BCP code			
	22. For transit through Member State(s)				I.23. For export				
	Member State ISO Code			Third country ISO Code					
				Exit point BCP code					
	24. Estimated journey time				I.25. Journey Log				
	26. Total number of packages I.27. Total quantity			I.28. Total net we	eight	I.28. Total gross w	reight		
	I.30. Description of consignment								
	1. 03 FISH AND CRUSTACEANS, MOLLUSCS AND OTHER AQUATIC INVERTEBRATES 0301 Live fish								
	Ornamental fish 030111 Freshwater fish 03011100 Freshwater fish								
	0301110 #1. Commodity		pecies	Quantity		Package count	Net weigh		

	II. Health infor	rmation						
	T the same day							
	I, the under II.1.	, the undersigned official veterinarian, hereby certify: I.1. According to official information, the aquatic animals in the consignment described in Part I meet the						
	11.1.		nimal health requirements:					
Part II: Certification		II.1.1.	is subject to the movement restrict 191(2), points (b)(i) and (ii), of Regu	ate from (1) \Box [an establishment] (1) \Box [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,				
Cer		II.1.2.	The aquatic animals:					
Part II: ((1) 0 either	[originate from (1) \Box [an establishment] (1) \Box [a habitat] where there are no increased mortalities with an undetermined cause.]					
		(1) ○ or	[originate from a part of (1) \Box [an establishment] (1) \Box [a habitat] which is ind the epidemiological unit where increased mortalities or disease symptoms hav and the Member State of destination (1) \Box [and the Member State (1) \Box [s] of the [has] (1) \Box [have] given consent for the movement to occur.]					
	(1) 🗆 [II.2.	Aquacultur	ture 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 has not indicated any cause for con	/429] (1) [approved in acco /429] where mortality records regularly updated and a docu n a period of 72 hours prior to	rdance with Article 176 or s, movement records and umentary check on those			
		II.2.2.	The aquaculture animals:					
		(1) ○ 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	t (b), of Commission Delegate ours prior to the time of depa	d Regulation (EU) 2020/990			
		(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 t	he derogation laid down in			
	(1)(2)(3) □ [II.3.	necrosis (II Marteilia re	nts for (4)listed species for Viral had HN), infection with HPR-deleted infe efringens, infection with Bonamia e spot syndrome virus	ectious salmon anaemia virus	(ISAV), infection with			
		The aquation	c animals referred to in Part I:					
		(1) ○ either	(1)(2)[originate from a (1) \Box [Mem from (1) \Box [VHS] (1) \Box [IHN] (1) \Box Marteilia refringens] (1) \Box [infection exitiosa] (1) \Box [infection with White Part II of Commission Delegated Reference of the second sec	[infection with HPR-deleted I on with Bonamia ostreae] (1) te spot syndrome virus] in acc	SAV] (1) □ [infection with □ [infection with Bonamia			
		(1) ° or	[originate from a (1) \Box [Member St eradication programme for (1) \Box [(1) \Box [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) □ [infect ingens] (1) □ [infection with 3 1) □ [infection with White sp or compartment which is also	ion 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 Implementin vectors of the relevant listed diseas Commission Delegated Regulation	ng Regulation (EU) 2018/1882 se as they do not fulfil the con	and they are not regarded as			
		(1) ° or	[are aquaculture animals of one of		lumn 4 of the table in the			

iU	ROPEAN	UNION	intended for release into the wild – Model AQUA-INTRA-RELEASE				
	II. Health info	ormation					
			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 15 of Commission Delegated Regulation (EU) 2020/691, and are regarded as disease free.]				
auon		(1) ° or	[are aquaculture animals of one of the vector species listed in column 4 in the table in the Annex to Implementing Regulation (EU) 2018/1882 and are regarded as vectors but they have been kept in isolation in an establishment approved in accordance with Article 16 of Commission Delegated Regulation (EU) 2020/691 and are no longer regarded as vectors.]]				
Part II: Ceruncauon	(1)(5) 🗆 [II.4.	viraemia o necrosis v	rements for (6)species susceptible to Koi herpes virus disease (KHV), infection with Spring nia of carp virus (SVC), Bacterial kidney disease (BKD), infection with Infectious pancreatic sis virus (IPN), infection with Gyrodactylus salaris (GS), infection with Salmonid alphavirus (SAV) ifection with Ostreid herpes virus 1 μ var (OsHV-1 μ var) onsignment originates from a (1) \Box [Member State], (1) \Box [zone] (1) \Box [compartment] which the health guarantees as regards (1) \Box [KHV], (1) \Box [SVC], (1) \Box [BKD], (1) \Box [IPN], (1) \Box [GS], (1) V], (1) \Box [OsHV-1 μ var] which are necessary to comply with the national measures which apply in ember State of destination, and for which the Member State or part thereof, is listed in (1) \Box x I] (1) \Box [Annex II] to Commission Implementing Decision (EU) 2021/260.]				
Par		fulfils the [SAV], (the Memb					
	II.5.						
		(i)	there were no abnormal mortalities with an undetermined cause; and				
		(ii)	the animals have not been in contact with aquatic animals of (4)listed species which did not comply with the requirements referred to in point II.1.				
	II.6.	-	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) \Box [the means of transport] (1) \Box [containers] in accordance with Article 5 of Delegated Regulation (EU) 2020/990, and the consignment is identified by (1) \Box [a legible and visible label on the exterior of the container] (1) \Box [a legible and visible label on the exterior of the container] (1) \Box [a legible and visible label on the exterior of the ship's manifest when transported by well boat], which clearly links the consignment to this animal health certificate.						
	II.8.	Validity of	f 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						
	from the E Protocol o	luropean Ur n Ireland / N	e Agreement on the withdrawal of the United Kingdom of Great Britain and Northern Ireland nion and the European Atomic Energy Community, and in particular Article 5(4) of the Northern Ireland in conjunction with Annex 2 to that Protocol, references to European Union ade 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.						
	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 ap	ppropriate/delete if not applicable.				
 Applies in all cases where the Member State of destination has taken measures in according Article 199 in Regulation (EU)2016/429 and requires that aquatic animals for release in originate from a Member State, zone or compartment which has disease-free status for disease as defined in Article 1, point (3), of Implementing Regulation (EU) 2018/1882. 							
(3) Other than in the cases referred to in Note (2) of this Part, Section II.3 applies only when the							

	II. Health info	rmation						
II: Certification		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.						
	(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.						
	(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	icer/Official veterinarian						
പ്പ	Name (in capital letters)		Qualification and title					
	Date of signature		Signature					
	Stamp							