

Part I : Details of consignment	I.1. Consignor Name Address Country		I.2. IMSOC Reference Specimen not to be used for exports from EU I.2.a. Local Reference	
	I.5. Consignee Name Address Country		I.3. Central competent authority I.4. Local competent authority	
	I.7. Country of origin		I.9. Country of destination	
	I.8. Region of origin		I.10. Region of destination	
	I.11. Place of Dispatch Name Address Approval Number Country		I.12. Place of destination Name Address Approval Number Country	
	I.13. Place of Loading Name Address Approval Number Country		I.14. Date and time of departure	
	I.15. Means of Transport		I.16 Entry Point	
	I.18. Transport conditions Chilled <input type="checkbox"/> Ambient <input type="checkbox"/> Frozen <input type="checkbox"/> Controlled temperature <input type="checkbox"/>		I.17. Accompanying documents Commercial document reference Date of issue Country Place of issue	
	I.19. Container No / Seal No			
	I.20. Certified as Competition <input type="checkbox"/> Sales <input type="checkbox"/> Registered equidae <input type="checkbox"/> Slaughter <input type="checkbox"/> Production <input type="checkbox"/> Ornamental bird food <input type="checkbox"/> Fattening <input type="checkbox"/> Racing <input type="checkbox"/> Quarantine <input type="checkbox"/> Transhumance <input type="checkbox"/> Rodent food <input type="checkbox"/> Pets <input type="checkbox"/> Circus exhibition <input type="checkbox"/> Artificial reproduction <input type="checkbox"/> Game Restocking <input type="checkbox"/> Other <input type="checkbox"/> Storage <input type="checkbox"/> Training <input type="checkbox"/> Technical use <input type="checkbox"/> Animal Feedingstuff <input type="checkbox"/> Laboratory <input type="checkbox"/> Unregistered equidae <input type="checkbox"/> Pharmaceutical use <input type="checkbox"/> Pollination <input type="checkbox"/> Consignments according to Regulation No 999/2001 <input type="checkbox"/> Pet food <input type="checkbox"/> Breeding and production <input type="checkbox"/> Organic fertilizers <input type="checkbox"/> Category 3 fish oil/fish meal for detoxification according to Regulation 2015/786 <input type="checkbox"/> Breeding <input type="checkbox"/> Human consumption <input type="checkbox"/> Approved Bodies <input type="checkbox"/> Further process <input type="checkbox"/> Relaying <input type="checkbox"/> Production of petfood <input type="checkbox"/> Ornamental use/research <input type="checkbox"/>			
	I.21. For transit through a third country <input type="checkbox"/> Country _____ ISO Code _____ EU Exit Authority _____ BCP code _____ EU Entry Authority _____ BCP code _____		I.22. For transit through Member State(s) <input type="checkbox"/> Country _____ ISO Code _____	
	I.23. Total number of packages	I.24. Total quantity	I.25. Total net weight	I.25. Total gross weight

Part I : Details of consignment	I.28. Description of consignment				
	1. 03 FISH AND CRUSTACEANS, MOLLUSCS AND OTHER AQUATIC INVERTEBRATES				
	0304 Fish fillets and other fish meat (whether or not minced), fresh, chilled or frozen				
	Commodity	Species	Quantity	Batch number	Manufacturing plant
Cold store	Cutting plant	Date of freezing	Date of production	Date of slaughter	
Net weight	Product Description	Package count	Identification mark		

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Part II: Certification	II. Health information		
	<p>II.1. (1)Public health attestation</p> <p>I, the undersigned, declare that I am aware of the relevant provisions of Regulation (EC) No 178/2002 of the European Parliament and of the Council of 28 January 2002 laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety, Regulation (EC) No 852/2004 of the European Parliament and of the Council of 29 April 2004 on the hygiene of foodstuffs and Regulation (EC) No 853/2004 of the European Parliament and of the Council of 29 April 2004 laying down specific hygiene rules for food of animal origin and Regulation (EU) 2017/625 of the European Parliament and of the Council of 15 March 2017 on official controls and other official activities performed to ensure the application of food and feed law, rules on animal health and welfare, plant health and plant protection products, amending Regulations (EC) No 999/2001, (EC) No 396/2005, (EC) No 1069/2009, (EC) No 1107/2009, (EU) No 1151/2012, (EU) No 652/2014, (EU) 2016/429 and (EU) 2016/2031 of the European Parliament and of the Council, Council Regulations (EC) No 1/2005 and (EC) No 1099/2009 and Council Directives 98/58/EC, 1999/74/EC, 2007/43/EC, 2008/119/EC and 2008/120/EC and repealing Regulations (EC) No 854/2004 and (EC) No 882/2004 of the European Parliament and of the Council, Council Directives 89/608/EEC, 89/662/EEC, 90/425/EEC, 91/496/EEC, 96/23/EC, 96/93/EC and 97/78/EC and Council Decision 92/438/EEC (Official Controls Regulation) and certify that the fishery products described above were produced in accordance with those requirements, in particular that they:</p> <ul style="list-style-type: none"> - come from (an) establishment(s) implementing a programme based on the hazard analysis and critical control points (HACCP) principles in accordance with Article 5 of Regulation (EC) No 852/2004, - have been caught and handled on board vessels, landed, handled and where appropriate prepared, processed, frozen and thawed hygienically in compliance with the requirements laid down in Section VIII, Chapters I to IV of Annex III to Regulation (EC) No 853/2004, - satisfy the health standards laid down in Section VIII, Chapter V of Annex III to Regulation (EC) No 853/2004 and the criteria laid down in Commission Regulation (EC) No 2073/2005 of 15 November 2005 on microbiological criteria for foodstuffs; - have been packaged, stored and transported in compliance with Section VIII, Chapters VI to VIII of Annex III to Regulation (EC) No 853/2004; - have been marked in accordance with Section I of Annex II to Regulation (EC) No 853/2004, - fulfil the guarantees covering live animals and products thereof, if of aquaculture origin, provided by the residue plans approved by the appropriate authority where applicable. and - have satisfactorily undergone the official controls laid down in Articles 69 to 71 of Commission Implementing Regulation (EU) 2019/627 of 15 March 2019 laying down uniform practical arrangements for the performance of official controls on products of animal origin intended for human consumption in accordance with Regulation (EU) 2017/625 of the European Parliament and of the Council and amending Commission Regulation (EC) No 2074/2005 as regards official controls. 		
<p>II.2 (2)(4) Animal health attestation for fish and crustaceans of aquaculture origin</p> <p>II.2.1 (3)(4) <input type="checkbox"/> [Requirements for species susceptible to epizootic haematopoietic necrosis (EHN), taura syndrome and yellowhead disease]</p> <p>I, the undersigned official inspector, hereby certify that the aquaculture animals or products thereof referred to in Part I of this certificate:</p> <p>(5) originate from a country/territory, zone or compartment declared free from (4) <input type="checkbox"/> [EHN] (4) <input type="checkbox"/> [taura syndrome] (4) <input type="checkbox"/> [yellowhead disease] in accordance with the relevant OIE Standard by the competent authority of my country, or via a process equivalent to Decision 2009/177</p> <p>(i) where the relevant diseases are notifiable to the competent authority and reports of suspicion of infection of the relevant disease must be immediately investigated by the competent authority,</p> <p>(ii) all introduction of species susceptible to the relevant diseases come from an area declared free of the disease, and</p>			

II. Health information

(iii) species susceptible to the relevant diseases are not vaccinated against the relevant diseases.]

II.2.2 (3)(4) [Requirements for species susceptible to viral haemorrhagic septicaemia (VHS), infectious haematopoietic necrosis (IHN), infectious salmon anaemia (ISA), koi herpes virus (KHV) and white spot disease intended for Great Britain, or a zone or compartment therein declared disease free or subject to a surveillance or eradication programme for the relevant disease

I, the undersigned official inspector, hereby certify that the aquaculture animals or products thereof referred to in Part I of this certificate:

(6) originate from a country/territory, zone or compartment declared free from (4) [VHS] (4) [IHN] (4) [ISA] (4) [KHV] (4) [White spot disease] in accordance with the relevant OIE Standard by the competent authority of my country, or via a process equivalent to Decision 2009/177

(i) where the relevant diseases are notifiable to the competent authority and reports of suspicion of infection of the relevant disease must be immediately investigated by the competent authority,

(ii) all introduction of species susceptible to the relevant diseases come from an area declared free of the disease, and

(iii) species susceptible to the relevant diseases are not vaccinated against the relevant diseases.]

II.2.3 (4)(7) [Requirements for species susceptible to Spring viraemia of carp (SVC), Bacterial kidney disease (BKD), Infectious pancreatic necrosis virus (IPN) and Infection with Gyrodactylus salaris (GS)

I, the undersigned official inspector, hereby certify that the aquaculture animals referred to above,

either (4) [originate from a country/territory or part thereof:

(a) where (4) [SVC] (4) [GS] (4) [BKD] (4) [IPN] is (are) are notifiable to the competent authority and reports of suspicion of infection of the relevant disease(s) must be immediately investigated by the competent authority,

(b) where all aquaculture animals of species susceptible to the relevant disease(s) introduced into that country/territory or part thereof comply with the requirements set out in II.2.3 of this certificate,

(c) where species susceptible to the relevant disease(s) are not vaccinated against the relevant disease(s), and

(d) either (4) [which, in the case of (4) [IPN] (4) [BKD], complies with requirements for disease freedom equivalent to those laid down in Decision 2009/177.]

and/or (4) [which, in the case of (4) [SVC] (4) [GS], complies with requirements for disease freedom laid down in the relevant OIE Standard.]

and/or (4) [which, in the case of (4) [SVC] (4) [IPN] (4) [BKD], comprises one individual farm which under the supervision of the competent authority:

(i) has been emptied, cleansed and disinfected, and fallowed in at least 6 weeks,

(ii) has been restocked with animals from areas certified free from the relevant disease by the competent authority.]]

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and/or (4) [in the case of wild aquatic animals susceptible to (4) [SVC] (4) [IPN] (4) [BKD], have been subject to quarantine under conditions at least equivalent to those laid down in Decision 2008/946/EC.]

and/or (4) [in the case consignments for which GS requirements apply, have been held, immediately prior to export, in water with a salinity of at least 25 parts per thousand for a continuous period of at least 44 days and no other live aquatic animals of the species susceptible to GS have been introduced during that period.]

and/or (4) [in the case of eyed fish eggs for which GS requirements apply, have been disinfected by a method demonstrated to be effective against GS.]]

II.2.4 Transport and labelling requirements

I, the undersigned official inspector, hereby certify that:

- II.2.4.1 the aquaculture animals referred to above are placed under conditions in which the water quality does not alter their health status,
- II.2.4.2 prior to loading the transport container or well boat is clean and disinfected or previously unused; and
- II.2.4.3 the consignment is identified by a legible label on the exterior of the container, or when transported by well boat, in the ship's manifest, with the relevant information referred to in boxes 1.7 to 1.11 of Part I of this certificate, and the following statement:

'(4) [Fish] (4) [Crustaceans] intended for human consumption in Great Britain'.

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Part II: Certification	II. Health information		
	<p>Notes</p> <p>See notes in Annex II of Commission Implementing Regulation (EU) 2019/628 of 8 April 2019 concerning model official certificates for certain animals and goods and amending Regulation (EC) No 2074/2005 and Implementing Regulation (EU) 2016/759 as regards these model certificates</p> <p>(*) Those countries subject to the transitional import arrangements include: an EU member State; Liechtenstein; Norway, Switzerland, Iceland and the Faroe Islands.</p> <p>References to Great Britain in this certificate include Channel Islands and Isle of Man.</p> <p>References to European Union legislation within this certificate are references to direct EU legislation which has been retained in Great Britain (retained EU law as defined in the European Union (Withdrawal) Act 2018).</p> <p>Part I:</p> <ul style="list-style-type: none"> - Box reference 1.8: Region of origin: For frozen or processed bivalve molluscs, indicate the production area. - Box reference 1.20: Tick 'Canning industry' for whole fish initially frozen in brine at — 9 °C or at a temperature higher than - 18 °C and intended for canning in accordance with the requirements of Section VIII, Chapter I; point II(7) of annex III to Regulation (EC) No 853/2004. Tick 'Human consumption' for the other cases. - Box reference 1.25: Insert the appropriate Harmonised System (HS) code(s) using headings such as: 0301, 0302, 0303, 0304, 0305, 0306, 0307, 0308, 0511, 1504, 1516, 1518, 1603, 1604, 1605 or 2106. - Box reference 1.25: Nature of commodity: specify whether aquaculture or wild origin. <p style="margin-left: 100px;">Treatment type: specify whether live, chilled, frozen or processed.</p> <p style="margin-left: 100px;">Manufacturing plant: includes factory vessel, freezer vessel, reefer vessels, cold store and processing plant.</p> <p>Part II:</p> <p>(1) Part II.1 of this certificate does not apply to countries with special public health certification requirements laid down in equivalence agreements or other GB legislation.</p> <p>(2) Part II.2 of this certificate does not apply to:</p> <ul style="list-style-type: none"> (a) non-viable crustaceans, meaning crustaceans that cannot survive as living animals if returned to the environment from which they were obtained, (b) fish which are slaughtered and eviscerated before dispatch, (c) aquaculture animals and products thereof, which are placed on the market for human consumption without further processing, provided that they are packed in retail-sale packages which comply with the provisions for such packages in Regulation (EC) No 853/2004, (d) crustaceans destined for processing establishments authorised in accordance with the Aquatic Animal Health (England and Wales) Regulations 2009 or the Aquatic Animal Health (Scotland) Regulation 2009, or for dispatch centres, purification centres or similar businesses which are equipped with an effluent treatment system that inactivates the pathogens in question, or where the effluent undergoes other types of treatment reducing the risk of transmitting diseases to the natural waters to an acceptable level, and (e) crustaceans which are intended for further processing before human consumption without temporary storage at the place of processing and packed and labelled for that purpose in accordance with Regulation (EC) No 853/2004. 		

Part II: Certification	II. Health information		
	<p>(3) Parts II.2.1 and II.2.2 of this certificate only apply to species susceptible to one or more of the diseases referred to in the heading of the point concerned. Susceptible species are listed in Annex 1A to Regulation (EC) No 1251/2008.</p> <p>(4) Keep as appropriate.</p> <p>(5) For consignments of species susceptible to EHN, taura syndrome and/or yellowhead disease this statement must be kept for the consignment to be authorised into Great Britain.</p> <p>(6) In order to be authorised into Great Britain, zone or compartment (boxes I.9 and I.10 of Part I of the certificate) declared free from VHS, IHN, ISA, KHV or white spot disease or with a surveillance or eradication programme, one of these statements must be kept if the consignment contain species susceptible to the disease(s) for which disease freedom or programme(s) apply(ies).</p> <p style="padding-left: 40px;">Data on the disease status of each farm and mollusc farming area in Great Britain are accessible at https://www.gov.uk/government/groups/fish-health-inspectorate#disease-status-of-fish-shellfish-and-crustacea-in-england-and-wales</p> <p style="padding-left: 40px;">https://www.gov.scot/publications/registers-of-authorised-aquaculture-production-businesses-and-authorised-processing-establishments/</p> <p style="padding-left: 40px;">https://www.gov.scot/publications/health-status-of-fish-and-shellfish-diseases-in-scotland/</p> <p>(7) Part II.2.3 of this certificate only applies to consignments intended for GB or part thereof (boxes I.9 and I.10 of Part I of the certificate), which is regarded as disease-free, or subject to a surveillance or eradication programme for SVC, BKD, IPN or GS, and the consignment comprises species listed in Schedule 1 of The Aquatic Animal Health (England and Wales) Regulations 2009 or The Aquatic Animal Health (Scotland) Regulations 2009 as susceptible to the disease(s) for which the disease-free status or programme(s) apply(ies).</p> <p style="padding-left: 40px;">Part II.2.3 also applies to consignments of fish of any species originating from waters where species listed in Schedule 1 of The Aquatic Animal Health (England and Wales) Regulations 2009 or The Aquatic Animal Health (Scotland) Regulations 2009 as species susceptible to Infection with GS, are present, where those consignments are intended for Great Britain or part thereof listed in Annex I to Decision 2010/221/EU as free of GS.</p> <p style="padding-left: 40px;">Consignments of wild aquatic animals for which SVC, IPN and/or BKD related requirements are applicable, may be imported regardless of the requirements in Part II.2.3 of this certificate if they are intended for a quarantine facility complying with the requirements laid down in Decision 2008/946/EC.</p> <p>- The colour of the stamp and signature must be different to that of the other particulars in the certificate.</p>		
	Certifying Officer Name (in capital letters) Date of signature Stamp	Qualification and title Signature	