Molecular Characterization of Novel Cryptosporidium Fish Genotypes in Edible Marine Fish

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Abstract: Current knowledge of Cryptosporidium species/genotypes in marine fish is limited. Following phylogenetic analysis at the 18S rDNA locus, a recent study identified six new genotypes of Cryptosporidium colonizing edible fish found in European seas. Of these, five grouped in a clade together (#Cryptofish 1–5) and one grouped separately (#Cryptofish 7). In the present study, after phylogenetic analyses of #Cryptofish1, #Cryptofish2, #Cryptofish4, #Cryptofish5 and #Cryptofish7 at the actin locus, the presence of two major clades was confirmed. In addition, when possible, longer 18S amplicons were generated. In conclusion, the small genetic distances between these genotypes designated as a novel marine genotype I (#Cryptofish 1-5) suggest that they may be genetic variants of the same species, while the designated novel marine genotype 2 (#Cryptofish 7) is clearly representative of a separate species.

Keywords: piscine Cryptosporidium; edible marine fish; genetic characterization; 18S rDNA gene; actin gene; molecular phylogeny

1. Introduction

The protozoan parasite Cryptosporidium is a waterborne and foodborne pathogen, causing severe diarrhea mainly in young children and immunocompromised persons and is also found in a wide range of vertebrate hosts [1].

To date, Cryptosporidium spp. have also been genetically characterized in more than 25 species of both freshwater and marine fish [2]. Four species and more than 20 piscine genotypes, all with strong host specificity and no reports in humans have been identified, including Cryptosporidium molnari [3,4], C. scophthalmi [5], C. huwi (previously known as piscine genotype (1) [6] and C. bollandii (previously known as piscine genotype (2) [7,8], piscine genotypes 3–8 [9–12], piscine genotype 9 [13], a C. molnari-like genotype [2,10,14–17], five unnamed novel genotypes [15,16], a Koi-Carp genotype [18] and Cryptofish 1–5 and 7 [2]. In addition, other Cryptosporidium spp. that are commonly identified in mammals,
such as C. parvum, C. hominis, C. scrofarum, C. xiaoi [2,9,12,13,19] and Cryptosporidium rat genotype III-like genotype have also been found in fish [11].

Molecular studies previously conducted on fish species revealed an extensive genetic diversity of Cryptosporidium spp. isolates [15,16]. However, the majority of studies on piscine Cryptosporidium have been carried out on ornamental or farmed fish, and scarce data are currently available concerning the molecular identification of Cryptosporidium species and genotypes in wild marine fish.

In a recent work evaluating the prevalence of Cryptosporidium spp. in commercially important edible marine fish in different European seas, sequence and phylogenetic analysis at the 18S rDNA locus identified 6 novel genotypes of Cryptosporidium colonizing these hosts [2]. Of these six genotypes, five grouped together (#Cryptofish 1–5) whereas #Cryptofish 7 emerged separately [2]. Based on the molecular characteristics of these genotypes, the main goal of the present study was thus to perform a comparative genetic characterization at the actin locus between these novel genotypes and with available piscine-derived Cryptosporidium genotypes, to provide further evidence for them as potential new separate species. A more comprehensive Cryptosporidium taxonomy is of major interest for a better understanding of transmission dynamics, public health significance and biology of this parasite.

2. Materials and Methods

2.1. Sampling

Scrapings of the gastrointestinal epithelia from several edible marine fish species (Merlangius merlangus, Pollachius virens, Molva dypterygia and Scomber scombrus) caught in different European seas (English Channel, North Sea, Bay of Biscay, Celtic Sea and Mediterranean Sea) that were previously identified as positive for different Cryptosporidium genotypes at the 18S rDNA locus [2] were selected. Samples were preserved in RCL2® buffer (Alphelys, Plaisir, France) and stored at −20 °C until required. Sections of the stomach and/or intestine were also collected and fixed in 10% buffered formalin for further histological analysis (Table 1).

Table 1. Details of Cryptosporidium piscine isolates typed at the 18S locus in a previous study and re-amplified and sequenced at the 18S rDNA and actin loci for the present study.

| Sample Identification | Fishing Area | Fish Scientific Name | Fish Common Name | Order | Organ Location | Genotype Identified (18S rDNA ) | GenBank Accession Numbers (18S rDNA) | GenBank Accession Numbers (Actin) |
|-----------------------|--------------|----------------------|------------------|-------|----------------|-------------------------------|-----------------------------------|----------------------------------|
| 524 NE Atlantic       | M. merlangus | Whiting              | Gadiformes       | Stomach | Cryptofish2   | MK236539 and MT776545           | MT570031                          |                                  |
| 698 NE Atlantic       | P. virens    | Saithe               | Gadiformes       | Stomach | Cryptofish1   | MK236538                        | NA                                |                                  |
| 710 NE Atlantic       | P. virens    | Saithe               | Gadiformes       | Stomach | Cryptofish1   | MK236538                        | MT570028                          |                                  |
| 716 NE Atlantic       | P. virens    | Saithe               | Gadiformes       | Stomach | Cryptofish1   | MK236538                        | NA                                |                                  |
| 720 NE Atlantic       | P. virens    | Saithe               | Gadiformes       | Stomach | Cryptofish1   | MK236538                        | NA                                |                                  |
| 722 NE Atlantic       | P. virens    | Saithe               | Gadiformes       | Intestine| Cryptofish1  | MK236538                        | MT570027                          |                                  |
| 735 NE Atlantic       | M. dypterygia| Blue ling            | Gadiformes       | Stomach | Cryptofish4   | MK236541                        | MT570032                          |                                  |
| PV-III-4 BO           | NN Sea       | P. virens            | Saithe           | Intestine| Cryptofish1  | MK236538                        | MT570030                          |                                  |
| PV-IV-3 ST            | NN Sea       | P. virens            | Saithe           | Stomach | Cryptofish1   | MK236538                        | MT570029                          |                                  |
| PV-IV-2 BO            | NN Sea       | P. virens            | Saithe           | Intestine| Cryptofish1  | MK236538                        | NA                                |                                  |
| PV-IV-7 ST            | NN Sea       | P. virens            | Saithe           | Stomach | Cryptofish5   | MK236542                        | MT570034                          |                                  |
| SS-VII-4 ST           | English Channel | S. scombrus       | Mackerel         | Scombriformes | Cryptofish7 | MK23654                        | MT570035                          |                                  |

a NE: North East, b NA: Not available c NN: Northern North.

The fish species were collected through research cruises belonging to the French Institut Français de Recherche pour l’Exploitation de la Mer (Institut Français de Recherche pour l’Exploitation de la Mer-IFREMER) in different European seas or through purchases from wholesalers or retailers for commercial catches at Boulogne-sur-Mer (Global Positioning System Coordinates: 50°43’ N–1°37’ E),
the first French fishing port [2,20]. Ifremer research cruises are carried out with the French Oceanographic Fleet under the supervision of the French Ministry of Education and Research. A steering committee evaluates and approves the entire scientific campaign program before implementation. The study was performed in accordance with the EU directive 2010/63/EU and followed all the guidelines of the deontology charter of Ifremer’s research.

2.2. DNA Extraction and PCR Amplification

Genomic DNA (gDNA) was newly extracted from gastric and/or intestinal scrapings of epithelia of fish, using the NucleoSpin™ Kit (Macherey-Nagel, GmbH & Co KG, Düren, Germany) as previously described [19]. DNA was diluted in 100 µL of elution buffer.

DNA from fish samples were reamplified for identification of Cryptosporidium at the 18S rDNA locus (~588 bp) as previously described [19]. In the present study, and when possible, longer 18S amplicons (~825 bp) were also generated using primers described by Xiao et al. [21]. Positive isolates were also analyzed at the actin locus using genus-specific actin primers described by Sulaiman et al. [22] producing a ~1066 bp amplicon. No-template controls (NTCs) were included alongside each PCR. Secondary PCR products were visualized on a 1% agarose gel stained with SYBR Safe (Thermo Fisher Scientific, Perth, Australia) fluoresce under ultraviolet light.

2.3. DNA Sequencing and Analysis

To identify Cryptosporidium species/genotypes, secondary PCR products were purified using the NucleoFast® 96 PCR kit (Macherey Nagel, GmbH & Co KG, Düren, Germany). Purified PCR products were sequenced in both directions, using the secondary PCR primers (Genoscreen, Pasteur Institute of Lille, Lille, France). Obtained nucleotide sequences were aligned using the BioEdit v. 7.0.1 package, and compared with available DNA sequences of Cryptosporidium in GenBank data base using the NCBI BLAST basic local alignment search tool (http://www.ncbi.nlm.nih.gov/BLAST/).

For longer 18S rDNA and actin amplicons, secondary PCR products were purified using a filter tip method [23] and sequenced in both directions using an ABI Prism™ Dye Terminator Cycle Sequencing kit (Applied Biosystems, Foster City, CA, USA) according to the manufacturer’s instructions at 58 °C. Nucleotide sequences identified in this study were deposited in GenBank under the accession numbers: MT570027—MT570035 (actin locus) and MT776545—MT776547 (18S rDNA locus).

2.4. Phylogenetic Analysis at the Actin Locus

Sanger sequencing chromatogram files were imported into Geneious Pro 10.2.6 [24], analysed and aligned with reference sequences from GenBank (C. huvi (AY524772), C. molnari (HM365220), C. molnari (HM365219), C. bollandi (MT160193) and C. scophtalmi (KR340589)) using Clustal W (http://www.clustalw.genome.jp). Distance and Maximum Likelihood (ML) trees were constructed using MEGA v. 7, after first identification of the most appropriate nucleotide substitution model [25]. Bootstrap support based on 1000 replications was included.

2.5. Histopathological Examination

Paraffin-embedded tissues were cut to a thickness of 5 µm and stained with hematoxylin and eosin (H & E). A DMRB microscope (Leica Wetzlar, Germany) equipped with a Leica digital camera connected to an Imaging Research MCID analysis system (MCID Software, Cambridge, UK) was used for observation of the histological sections.

2.6. Detection of Cryptosporidium by Immunofluorescence

A conjugated anti-Cryptosporidium spp. (Sporoglo, Waterborne, New Orleans, LA, USA) was used in direct fluorescent-antibody staining assay according to the manufacturers’ instructions. DAPI (4’,6-diamidino-2-phenylindole) was used for nuclei identification. Slides were examined in a LSM880
18S amplicons (~825bp) were only generated successfully for genotype Cryptofish1 (MT776545), Cryptofish2, Cryptofish4, and Cryptofish5 (designated novel marine genotype 1) may be genetic variants of the same species, while Cryptofish7 (designated novel marine genotype 2) is clearly a separate species. The small genetic distances at the actin locus suggests that Cryptofish1, Cryptofish2, Cryptofish4, and Cryptofish5 and 9.1% from Cryptofish7, while Cryptofish2 exhibited 8.2% genetic distance from Cryptofish7. Finally, Cryptofish7 exhibited 14.3%, 15.5%, 16.2% and 20.2% genetic distance from C. molnari, C. bollandi, C. huwi, and C. scophtalmi, respectively (Figure 1 and Table 2). The small genetic distances at the actin locus suggests that Cryptofish1, Cryptofish2, Cryptofish4 and Cryptofish5 (designated novel marine genotype 1) may be genetic variants of the same species, while Cryptofish7 (designated novel marine genotype 2) is clearly a separate species (Table 2).

3. Results

Nested 18S rDNA PCR and sequencing of DNA from piscine-derived Cryptosporidium isolates using the original shorter amplicons products previously described [2] confirmed their assignment to different genotypes as follows: nine samples belonged to genotype Cryptofish1 (MK236538), one to genotype Cryptofish2 (MK236539), another one to Cryptofish4 (MK236541), two corresponded to genotype Cryptofish5 (MK236542) and one to genotype Cryptofish7 (MK236544). However, longer 18S amplicons (~825bp) were only generated successfully for genotype Cryptofish1 (MT776545, MT776546 and MT776547).

Actin sequences obtained for six samples were representative of Cryptofish1, Cryptofish2, Cryptofish4, Cryptofish5 and Cryptofish7 (Table 1). Phylogenetic analyses of Cryptofish1, Cryptofish2, Cryptofish4, Cryptofish5 and Cryptofish7 at the actin locus identified two major clades: one composed of the sequences of genotypes Cryptofish1, Cryptofish2, Cryptofish4 and Cryptofish5 and the other one composed of the sequence of Cryptofish7 (Figure 1). At the actin locus, the Cryptofish1 genotype exhibited 0.9% genetic distance from Cryptofish2, 0.5–0.6% from Cryptofish4 and Cryptofish5 and 9.1% from Cryptofish7, while Cryptofish2 exhibited 8.2% genetic distance from Cryptofish7. Finally, Cryptofish7 exhibited 14.3%, 15.5%, 16.2% and 20.2% genetic distance from C. molnari, C. bollandi, C. huwi, and C. scophtalmi, respectively (Figure 1 and Table 2). The small genetic distances at the actin locus suggests that Cryptofish1, Cryptofish2, Cryptofish4 and Cryptofish5 (designated novel marine genotype 1) may be genetic variants of the same species, while Cryptofish7 (designated novel marine genotype 2) is clearly a separate species (Table 2).

**Figure 1.** Phylogenetic relationships between Cryptofish1, 2, 4, 5 and 7 genotypes (identified in the present study) and other piscine Cryptosporidium species inferred by ML analysis of actin gene. Percentage support (>50%) from 1000 pseudoreplicates from ML analyses is indicated at the left of the supported node. Scale bars indicate the number of substitutions per nucleotide position.
After examination of histological sections from the digestive tract of fishes, the presence of Cryptosporidium-like bodies in apical position was observed (Figure 2A,B). These structures were 3–4 μm in diameter, spherical and positive in H&E. Immunofluorescence analysis confirmed this with structures similar in size and shape to intracellular stages of Cryptosporidium labelled in red with the Sporoglo antibody (Figure 2C). However, the presence of parasites could not be confirmed in all positive samples due to substantial lysis of tissues.

| Piscine Cryptosporidium | #CryptoFish1 | #CryptoFish2 | #CryptoFish4 | #CryptoFish5 | #CryptoFish7 | C. huwi | C. molnari | C. molnari | C. bollandi |
|-------------------------|--------------|--------------|--------------|--------------|--------------|---------|-----------|-----------|-----------|
| #CryptoFish1 (Intestine, MT570028) | - | - | - | - | - | - | - | - | - |
| #CryptoFish2 (MT570031) | 0.9 | - | - | - | - | - | - | - | - |
| #CryptoFish4 (MT570032) | 0.5 | 0.8 | - | - | - | - | - | - | - |
| #CryptoFish5 (MT570034) | 0.6 | 0.7 | 0.004 | - | - | - | - | - | - |
| #CryptoFish7 (MT570035) | 9.1 | 8.2 | 9.2 | 8.8 | - | - | - | - | - |
| C. huwi (AY524772) | 16.5 | 16.1 | 16.6 | 16.4 | 16.2 | - | - | - | - |
| C. molnari (HM365220) | 15.1 | 14.5 | 15.2 | 15.1 | 14.3 | 17.7 | - | - | - |
| C. molnari (HM365219) | 14.5 | 14.0 | 14.7 | 14.5 | 14.3 | 18.4 | 0.7 | - | - |
| C. bollandi (MT160193) | 17.1 | 16.5 | 17.2 | 17.1 | 15.5 | 18.4 | 15.5 | 15.7 | - |
| C. scophtalmi (KR340589) | 19.0 | 18.6 | 19.5 | 19.3 | 20.2 | 24.3 | 17.6 | 17.4 | 20.2 |

Figure 2. Stained sections of the intestinal tract of one Scomber scombrus (Ss VII 4BO) infected by the genotype Cryptofish7 (A) Presence of round bodies suggestive of the developmental stages of Cryptosporidium spp. observed in the apical position of a (stained with H&E) (B) Detail of A (delimited area). (C) Structure labelled in red with the Sporoglo (Waterborne) antibody and blue with DAPI (4′,6-diamidino-2-phenylindole) suggestive of an intracellular stage of Cryptosporidium spp.

4. Discussion

Molecular characterization of Cryptosporidium has contributed to a better understanding of the diversity and transmission dynamics of this important enteric parasite. In the present study, we analyzed new genotypes of Cryptosporidium previously identified in edible fish [2] at the actin locus in order to provide additional data to support their potential species status.

Previous sequence and phylogenetic analysis at the 18S rDNA gene locus had shown that these genotypes were distributed as follows: 22 (48%) belonged to the #CryptoFish1 genotype that exhibited 7.3–8.5% genetic distance from C. molnari, six (13%) belonged to another genotype #CryptoFish2 that exhibited 8.5–9.5% genetic distance from C. molnari, a single isolate (2%) identified as #CryptoFish4 exhibited 8.2–9.5% genetic distance from C. molnari, 4 (9%) belonged to #CryptoFish5 genotype,
which exhibited 7.6–8.8% genetic distance from *C. molnari*. Finally, the single isolate (2%) #Cryptofish7 exhibited 9.1–10.4% genetic distance from *C. molnari* [2]. #Cryptofish3 genotype which was previously identified [2] and which exhibited 8.9–10.1% genetic distance from *C. molnari* could not be analyzed in the current study.

In the previous analyses at the 18S rDNA locus, the genetic distances between #Cryptofish1, #Cryptofish2, #Cryptofish4 and #Cryptofish5 was 0.3–2.8% and these sequences exhibited 6.1–8.9% genetic distances from #Cryptofish7. Interestingly, in the present study, at the actin locus, the genetic distances between #Cryptofish1, #Cryptofish2, #Cryptofish4 and #Cryptofish5 were smaller (0.6–0.9%) and they exhibited 8.2–9.1% genetic distances from #Cryptofish7. #Cryptofish1, #Cryptofish2, #Cryptofish4 and #Cryptofish5 exhibited 14.0–14.5% (#Cryptofish2 and *C. molnari*) to 19.5% (#Cryptofish4 and *C. scophtalmi*) genetic distance from other fish species. #Cryptofish7 exhibited 14.3%, 15.5%, 16.2% and 20.2% genetic distance from *C. molnari*, *C. bollandi*, *C. huwi*, and *C. scophtalmi*, respectively (Table 2).

The genetic distances between #Cryptofish1, #Cryptofish2, #Cryptofish4 and #Cryptofish5 may be enough to suggest that they are separate species. For example, the genetic distance at both the 18S rDNA and actin loci between *C. erinacei* and *C. parvum* is 0.5% [26] and the genetic distance between *C. muris* and *C. andersoni* at the 18S rDNA and actin loci is 0.9% and 3.5%, respectively. However, further analyses are required to confirm this and they are currently grouped together as novel marine *Cryptosporidium* genotype 1. Additional biological characteristics of the two potential novel marine genotypes are listed in Table 3.

### Table 3. Characteristics of novel piscine *Cryptosporidium* genotypes identified in the present study.

| Novel Marine Genotypes | Genotypes | Scientific Name (Fish Host Common Name) | Order | Fishing Area | Organ Distribution | Overall Prevalence in Fish % (n = 1853) | Distribution According to *Cryptosporidium* Positive Cases % (n = 46) |
|------------------------|-----------|----------------------------------------|-------|--------------|-------------------|------------------------------------------|---------------------------------------------------------------|
| Novel marine genotype 1 | #Crypto1, #Crypto2, #Crypto4, and #Crypto5 | *P. virens* (Saithe), *Molva dypterygia* (Blue ling), *Molva molva* (Ling), *Merlangius merlangus* (Whiting), *Merluccius merluccius* (Hake) | Gadiformes | NE Atlantic | Intestine and/or stomach | 1.78 | 71.7 |
| Novel marine genotype 2 | #Cryptofish7 | *Scomber scombrus* (Mackerel) | Scombriformes | English Channel | Stomach | 0.05 | 2.1 |

* Source [2]. * This fish species was found positive for this novel marine genotype in a previous study [2] but unfortunately, there was insufficient sample to reamplify and sequence again.

This genotype was the most prevalent since it was identified either in the stomach or the intestine of five different fish species belonging to the order gadiformes, all caught in the Atlantic North East (*Pollachius virens*, *Molva dypterygia*, *Molva molva*, *Merlangius merlangus* and *Merluccius merluccius*, with *P. virens* as the most common fish host). The second novel species (#Cryptofish7) (novel marine genotype 2) was identified in the stomach of a single specimen from *Scomber scombrus* (order scombriformes) caught in the English Channel (Table 3) [2]. Interestingly, it has been described that genetically related hosts often harbor related species of *Cryptosporidium* [27]. However, further studies are needed to confirm any specificity related to fish orders, considering that #Cryptofish7 was only found in a single specimen.

In order to verify multiplication of parasites in fish digestive tissues, histological sections from the digestive tract of fishes were analyzed after staining with H&E or by immunofluorescence analysis. Cryptosporidium-like bodies were observed in an apical position suggesting the multiplication of
the parasite and a true infection of the fish rather than just carriage (Figure 2). In addition, as DNA screening was conducted on mucosal scrapings from intestines and stomach tissue, this also suggests actual infections. The presence of parasites could not be studied in all positive fishes due to considerable lysis of tissues, which has been previously described as a difficulty for Cryptosporidium detection in fish hosts [10–12]. In addition, tissues were not available for all specimens. Unfortunately, oocysts could not be obtained to determine the morphological features of the oocyst stage of these piscine genotypes. However, morphological overlap in oocyst size is commonly found among Cryptosporidium spp., and in particular among Cryptosporidium spp. from fish [6]. Moreover, it is widely recognized that morphometrics is not a useful tool for defining most species within this genus [28].

The pathogenesis of the Cryptosporidium species identified in the present study remains unknown. Nevertheless, different studies have reported that piscine species and genotypes can cause pathological effects in fish [3,5] as well as an increase in mortality, particularly in juveniles [8], negatively impacting the fish industry economy.

The high diversity of Cryptosporidium species and genotypes in fish indicates a long-term association of Cryptosporidium with their fish hosts. The understanding of the taxonomy of piscine-derived Cryptosporidium species is of relevance considering that previous studies suggest that they might represent the most primitive Cryptosporidium species 6 [10].

Cryptosporidiosis can be considered important in fish since these animals may act as carriers and may be a source of infection for other hosts including humans. Further studies should be conducted to determine if contact with edible marine fish colonized by these Cryptosporidium species/genotypes pose a risk of zoonotic transmission either through their consumption and/or handling or through the consumption of water contaminated with fully sporulated oocysts shed in fish feces [29]. The four already known piscine species of Cryptosporidium have not yet been found in other hosts [1,30–32] suggesting that these species may not be able to grow in the digestive tract of mammals. However, further research is required to confirm this hypothesis.

5. Conclusions

In the present study, two potential novel Cryptosporidium fish species were further characterized at the actin locus. Further investigations will be performed in order to confirm that these genotypes are different Cryptosporidium species.

Cryptosporidium spp. have a wide host range and this together with the potential for high levels of oocyst shedding, allows a significant level of contamination of the environment. In particular, for fish hosts, the dispersion and transmission of zoonotic parasites would be facilitated by the aquatic habitat of the host, that could potentially release oocysts contributing to Cryptosporidium circulation. Therefore, additional epidemiological studies in wildlife animals are needed to better define the host range and zoonotic potential of the parasite.

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