Fecal Strings Associated with *Streptococcus agalactiae* Infection in Nile Tilapia, *Oreochromis niloticus*

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**Abstract:** This study provides the first description of long fecal strings in *Streptococcus agalactiae*-infected fish. Nile tilapia (*Oreochromis niloticus*) were administered an intraperitoneal injection with 0.1 mL solution containing a 10-fold dilution from $10^1$ to $10^6$ cfu *S. agalactiae*/fish. While infected fish developed clinical signs commonly associated with *S. agalactiae* infection, up to 40% of infected fish in each group also produced considerably longer (up to 20 cm in length) fecal waste strings than unchallenged tilapia. Fecal strings from these infected fish were observed trailing from the anus and often became increasingly opaque and white over time. Though *S. agalactiae* was not isolated from the fecal strings themselves, all *S. agalactiae*-challenged fish sampled were culture-positive. Histologic examination of the intestines of infected fish exhibited the presence of sloughed intestinal mucous cells and enterocytes in the feces and the absence of normal fecal matter. The presence of long brown and/or white, opaque fecal strings may suggest a clinical sign of bacterial disease and the need for further clinical examination.

*Streptococcus agalactiae* affects numerous wild and cultured fish species worldwide and causes disease involving septicemia and colonization of numerous organs [1-4]. Clinical signs of *S. agalactiae* infection include lethargy or excitability, going off feed, ‘C’-shaped body posturing, erratic swimming and whirling, opercular clearing, spinal curvature, and death [2, 3, 5]. Nile tilapia (*Oreochromis niloticus*) are susceptible to *S. agalactiae* infection [3-5]. In a variety of previous *S. agalactiae* challenge studies at the Aquatic Animal Health Research Laboratory, Nile tilapia (*n* > 500; weight ranging from 3 to 100 g) were injected intraperitoneally with doses ranging from $10^1$ to $10^9$ colony-forming units (cfu) *S. agalactiae*/fish and maintained at approximately 30°C. Though these studies were not intended to exclusively examine *S. agalactiae*-related clinical signs, it was nonetheless noted that numerous challenged fish produced long fecal strings. Since fecal strings have not been previously associated with *S. agalactiae* infection, a study was performed to examine the presence of fecal strings after experimental *S. agalactiae* challenge.

Nile tilapia (*O. niloticus*) with a mean weight of 13.9 ± 0.6 g were housed at the USDA/ARS Aquatic Animal Health Laboratory in Chestertown, Maryland, USA. The fish were kept in 57 L glass aquaria supplied with flow-through dechlorinated tap water, maintained at a mean temperature of 30.5 ± 0.6°C, mean dissolved oxygen of 5.3 ± 0.9 mg/L, and mean ammonia concentration of 0.2 ± 0.8 mg/L. The fish were also fed daily to satiation with Aquamax feed (Brentwood, Missouri, USA) and maintained on a 12 h : 12 h light : dark period. Tilapia were challenged with an *S. agalactiae* isolate obtained from a mullet, *Liza kunzingeri*, from Kuwait Bay, Kuwait [3]. The isolate was grown at 30°C for 24 h on 5% sheep blood agar (SBA; Remel, Lenexa, Kansas, USA) before reconstitution in tryptic soy broth (TSB; Remel) to create the challenge solution. Tilapia were administered an intraperitoneal injection with 0.1 mL solution containing a 10-fold dilution from $10^1$ to $10^6$ cfu *S. agalactiae*/fish. Ten fish were used for each dose and sequestered into separate tanks accordingly. Ten control fish were injected with TSB. The intraperitoneal challenge route was utilized because it is a reproducible and reliable method ensuring challenge of all individual fish with a uniform bacterial dose [6, 7]. Fish were monitored for 7 days post-challenge; clinical signs of disease were recorded once daily while mortalities were removed three times daily. Swab samples were aseptically obtained from the nares, brain, anterior kidney, and posterior intestine of 20% of dead fish from each group and cultured overnight on SBA at 30°C to confirm the presence of *S. agalactiae*. In order to confirm isolate identity as *S. agalactiae*, two bacterial isolates obtained from the brains of 2 challenged fish were tested with the BIOLOG MicroLog3™ system according to the manufacturer’s instructions (BIOLOG Inc., Hayward, California, USA) [3]. For histologic examination, intestinal tissues from 10 fish in the challenge groups and 5 fish from the control group were sampled 4 days post-challenge. Samples were fixed in 10% neutral buffered formalin, embedded in paraffin, sectioned at 5 μm, and stained with Harris’ hematoxylin and eosin. Any moribund fish in the study were humanely euthanized with an overdose of tricaine methanesulfonate (MS-222; Argent Chemical Laboratories, Redmond, Washington, USA).

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Within 7 days post-challenge, fish in each group developed overt clinical signs of disease, such as going off feed, lethargy, bilateral exophthalmia, erratic swimming, and death. Mortalities within each challenge group began 1 day post-challenge and most had occurred by 3 days post-challenge. Several of the *S. agalactiae*-infected tilapia produced considerably longer (up to 20 cm in length) fecal waste strings (Fig. 1, Table 1) than unchallenged tilapia, irrespective of weight or challenge dose. Strings were observed trailing from the anus and often became increasingly opaque and white as time increased. Long but brown fecal strings started to appear within 2 days post-challenge and began to appear as white, mucoid fecal strings within 4 days post-challenge. At this latter point, the fecal strings would contain intermittent sections of white, presumptively mucoid material and brown feces or the whole fecal strings would contain entirely white, presumptively mucoid material devoid of brown feces.

Given that fish were not individually identified, the numbers of fish per group showing certain clinical signs was determined as the maximum number observed exhibiting a clinical sign in one of the seven post-challenge days. Note that this may ultimately underestimate the total number of fish that showed a clinical sign if they stop exhibiting a clinical sign from one day to the next. When comparing maximum numbers of fish showing a certain clinical sign in one day with the total overall number of fish challenged (60 fish), the percentage of fish showing a specific clinical sign was: mortalities (53%), confined to tank bottom (43%), lethargy (32%), fecal strings (28%), stationary (25%), slow or no feeding response (18%), darkened coloration (12%), and serpentine swimming (2%). Within each challenge dose group, the maximum percentage of fish to have fecal strings in one day was 20 to 40% and a higher percentage did not appear to be related to dose.

Samples for bacteriology obtained from the nares, brain, anterior kidney, and posterior intestines were all positive for bacterial growth. Bacterial cultures from all sampled organs elicited primarily smooth, white, circular, Gram-positive, oxidase-negative colonies, and isolates were identified as *S. agalactiae* with the BIOLOG system (Probability = 98 to 99%). Control fish injected with TSB did not die or exhibit clinical signs of disease (including no fecal strings) and were negative for *S. agalactiae* growth. Light microscopic examination of intestinal sections from ten challenged fish did not reveal severe or consistent alterations. Two fish had food in the intestinal tract, and two fish exhibited focal areas of moderately increasing mucus cell numbers in the anterior intestine. Three fish showed mild inflammatory cell infiltration in the submucosa, and one fish exhibited mild focal enterocyte necrosis with sloughing of mucosal cells. No significant changes were noted among five control fish when compared to normal tilapia intestinal histologic sections [8].

**Table 1. Percentage of 10 Nile Tilapia (*Oreochromis niloticus*) in Each Dose Group Exhibiting Clinical Signs After Challenge with *Streptococcus agalactiae*¹**

<table>
<thead>
<tr>
<th>Dose (cfu/Fish)</th>
<th>Mortalities</th>
<th>Confined to Tank Bottom</th>
<th>Lethargy</th>
<th>Fecal Strings</th>
<th>Stationary</th>
<th>Slow or No Feeding Response</th>
<th>Darkened Coloration</th>
<th>Serpentine Swimming</th>
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</thead>
<tbody>
<tr>
<td>Control</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
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<td>10⁴</td>
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<td>10</td>
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<tr>
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<tr>
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<tr>
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<td>0</td>
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<tr>
<td>Overall Percent²</td>
<td>53</td>
<td>43</td>
<td>32</td>
<td>28</td>
<td>25</td>
<td>18</td>
<td>12</td>
<td>2</td>
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</tbody>
</table>

¹Tilapia were administered an intraperitoneal injection with 0.1 mL inoculum containing a 10-fold dilution from 10⁴ to 10⁶ colony-forming units (cfu) *S. agalactiae*/fish. Ten fish were used for each dose and sequestered into separate tanks accordingly. Ten control fish were inoculated with TSB. Fish were monitored daily for clinical signs of disease and mortality for 7 d post-challenge.

²Overall percent is determined by the number of fish exhibiting a clinical sign per all 60 fish challenged.

**Fig. (1).** Nile tilapia (*Oreochromis niloticus*) challenged with *Streptococcus agalactiae* exhibiting a long, white mucoid fecal string trailing from the anus.

Long mucoid fecal strings in *S. agalactiae*-infected fish have also not been previously described, but long fecal
strings were commonly observed in individual infected fish in our challenge studies. The feces were not cultured due to the high probability of contamination with water-borne Salmonella. While S. agalactiae was not isolated from the fecal strings themselves, fecal strings were only observed coming from S. agalactiae-challenged fish and not controls, all S. agalactiae-challenged fish sampled were culture-positive (including all posterior intestine samples), and all control fish were culture-negative. This suggests that S. agalactiae-challenged fish developed systemic infections and that the resulting disease included fecal string production. They resembled feces termed “mucoid fecal casts” described in select fish viral diseases, presumably created due to the 1) presence of sloughed intestinal mucous cells and enterocytes in the feces and 2) absence of normal fecal matter, This composition of the fecal strings are suggested by other publications [9-11] and by histologic examination of the intestinal tissue of the challenged fish. Fecal strings are not frequently examined or emphasized in relation to fish disease, though strings presumably containing sloughed intestinal mucosa and infectious organisms have been observed with a number of viral fish pathogens: Infectious Pancreatic Necrosis Virus, Infectious Hematopoietic Necrosis Virus, and Salmonid Herpesvirus Type 1 in salmonids and Spring Viremia of Carp [10, 11]. However, fecal strings are not routinely associated with bacterial diseases in fish. The presence of fecal strings can be cause for concern, suggesting a clinical sign of viral or bacterial disease and the need for further etiologic examination. Fecal analysis is commonly used for diagnosis of bacterial and parasitic infections in mammals [12], but is not so common in aquatic animal medicine. Fish feces have been collected in studies that measure parameters such as: intestinal flora composition [13], feed consumption and digestibility [14], and endocrine activity [15]. However, clinical fecal analysis and observation could add to the battery of non-invasive diagnostic tests for viral and now bacterial diseases available to the aquatic animal clinician. Such clinical tests may be helpful because they can indicate disease presence, though they may not be as useful in acute disease. In this study, acutely affected fish died before the appearance of fecal strings. Fecal strings in the tanks may also promote S. agalactiae fecal-oral transmission, because streptococcal infections are known to occur after ingestion of materials containing streptococcal organisms [16, 17]. Feces-related clinical signs and tests may be more useful for tilapia or other fish species maintained in clearer water (i.e. laboratory or aquarium tanks) than murkier water (i.e. ponds), because the strings can be readily observed as the fish swim. But these fecal strings have also been noted clinging to fish removed from water and may be analyzed even if the fish cannot be directly observed swimming with trailing fecal strings.

ACKNOWLEDGEMENTS

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REFERENCES