Changes in the gut microbiome of the sea lamprey during metamorphosis

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Abstract

Vertebrate metamorphosis is often marked by dramatic morphological and physiological changes of the alimentary tract, along with major shifts in diet following development from larva to adult. Little is known about how these developmental changes impact the gut microbiome of the host organism. The metamorphosis of the sea lamprey (*Petromyzon marinus*) from a sedentary filter-feeding larva to a free-swimming sanguivorous parasite is marked by major physiological and morphological changes to all organ systems. The transformation of the alimentary canal includes closure of the larval esophagus and the physical isolation of the pharynx from the remainder of the gut which results in a non-feeding period that can last up to 8 months. To determine how the gut microbiome is affected by metamorphosis, the microbial communities of feeding and non-feeding larval and parasitic sea lamprey were surveyed using both culture-dependent and independent methods. Our results show that the gut of the filter-feeding larva contains a greater diversity of bacteria than that of the blood-feeding parasite, with the parasite gut being dominated by *Aeromonas*, and to a lesser extent *Citrobacter* and *Shewanella*. Phylogenetic analysis of the culturable *Aeromonas* from both the larval and parasitic gut revealed that at least five distinct species were represented. Phenotypic characterization of these isolates revealed that over half were capable of sheep red blood cell hemolysis, but all were capable of trout red blood cell hemolysis. This suggests that the enrichment of *Aeromonas* that accompanies metamorphosis is likely related to the sanguivorous lifestyle of the parasitic sea lamprey.
Introduction

Studies of the mammalian gut have highlighted the importance of the gut microbiome to the metabolism, behaviour, physiology and general health of the host (5, 10, 12, 31). The microbiome can prevent the establishment of bacterial pathogens, aid in digestion, nutrient uptake and even assist in the development of host tissue such as the brain, which in turn affects behaviour (8, 11, 14). In vertebrates that undergo metamorphosis, however, the gut can be subject to major morphological and physiological changes, with post-metamorphic juveniles having a drastically different diet than that of their larval stage. The impact of physiological and dietary changes on the gut microbiome of organisms that undergo metamorphosis remains largely uncharacterized.

Studies of metamorphosing anurans (frogs and toads) have evaluated changes in bacterial species composition and proportion in different life stages of the host. A survey of both tadpoles and adults of northern leopard frogs (Rana pipiens) yielded a prevalence of members of the Enterobacteriaceae, including Escherichia coli, Citrobacter, Klebsiella, Enterobacter, Serratia and Yersinia (9). Citrobacter freundii and Aeromonas hydrophila were the predominant microbes in tadpole and adult intestines, both being found in about 45% of sampled specimens. Another study also observed shifts in the gut microbiome of southern toads (Bufo terrestris) and spring peepers (Pseudacris crucifer) (4). While these studies provide a glimpse into some of the possible changes in the composition of the gut microbiome during anuran metamorphosis, it is unclear how the gut microbiome is affected when metamorphosis is accompanied by a shift to a highly specialized diet.

The sea lamprey (Petromyzon marinus) is a jawless vertebrate having a complex life cycle that involves a shift from a diverse diet of detritus and microbes to a highly specialized diet.
of fish blood. After hatching, newly emerged larvae burrow into the sand, where they feed on detritus and microbes in the water column for 3-7 years (36, 37) (Figure 1). Once metamorphosis has begun, the larvae cannot feed due to major changes in the oral apparatus, which includes blockage of the larval esophagus and physical isolation from the pharynx (35). This non-trophic (non-feeding) period can last up to 8 months (37). As metamorphosis nears completion, a new esophagus is formed and the gut develops longitudinal mucosal folds for greater absorption of nutrients (Figure 1) (37). The digestive system also undergoes several other changes including the degeneration of the bile ducts, bile canaliculi and gall bladder (34). To compensate for the loss of a functional blood bile barrier, parasites use circulating bile-binding proteins to transport toxins through the circulatory system and deposit them in the posterior end of the intestine, resulting in bile accumulation in this region (34). The newly metamorphosed sea lamprey is sanguivorous, having a diet of predominately fish blood and body fluids (37).

Because the sea lamprey undergoes major morphological and physiological changes during metamorphosis, and also goes through a non-trophic stage before acquiring a highly specialized diet, it represents an ideal system to evaluate the impact of changes in host physiology on the gut bacterial community. Only one published study to date has examined the lamprey gut microbiome. This particular work examined the composition of gut bacteria specifically in the larval stage of the pouched lamprey \textit{(Geotria australis)}, although no comparisons were made to that of the parasitic stage (24). In this study, we compared the gut microbiota of the larval and parasitic life stages using 16S rRNA gene sequencing, and also characterized a collection of culturable isolates with respect to hemolysin production, protease secretion and bile tolerance. Our results highlight a significant shift in the relative composition...
of microbes between larval and parasitic sea lamprey stages that are reflective of the shift to a more specialized diet.

**Materials and Methods**

**Animal and tissue collection**

Larval sea lamprey were collected from Oshawa Creek (Ontario, Canada), Harlow creek (Michigan, USA) and Little Garlic River (Michigan, USA) and transported by truck to the University of Regina. When possible, sea lampreys were sampled immediately upon arrival at the University of Regina. To rear parasites, larvae were housed in aquaria with 12-15 cm of sterile, washed and screened sand, filled with dechlorinated city tap water (kept at 18-21 °C), which was aerated and filtered continuously. Larvae were fed a suspension of Baker's yeast (equivalent to 1 g of yeast per animal) twice weekly until the onset of metamorphosis. Lampreys were not fed during the non-trophic phase of metamorphosis. When metamorphosis was complete, post-metamorphic juveniles were transferred to 1700 L aquaria containing aerated and continuously filtered dechlorinated water at 11-15 °C. Rainbow trout (*Oncorhynchus mykiss*) were introduced into the tank as a food source. Prior to tissue collection, animals were anesthetized in 0.05 % tricaine methanesulfonate buffered with 1 % sodium bicarbonate. Following anesthesia, animals were euthanized by decapitation prior to harvesting the gut. Tissue samples were sectioned into anterior, medial and posterior thirds, and placed in 10 mM MgSO₄ solution (100 µl for larval and 400 µl for parasite). Samples from post-metamorphic sea lamprey were obtained from animals that had metamorphosed in lab aquaria. All animal handling and procedures were approved by the President's Committee on Animal Care at the University of Regina and were consistent with the guidelines of the Canadian Council on Animal Care.
DNA extraction and sequencing

Bacteria from larval or parasitic phase sea lamprey were harvested by vortexing a pool of 10 anterior or posterior gut sections from 10 individual lampreys in 10 mM MgSO₄ solution and then physically removing the tissue fragments. This process was repeated 5 times (50 animals total) producing 5 independent samples. Genomic DNA was extracted from the bacterial suspension immediately using the Qiagen Puregene Core Kit Yeast/Bacteria Genomic DNA Kit according to the manufacturer’s instructions. Genomic DNA was used as a template for PCR (polymerase chain reaction) to amplify a 1065 base pair fragment of the 16S rRNA gene using the following primers: forward 16S-335 (5’ ACTCCTACGGGAGGCAGC 3’) and reverse 16S-1400 (5’ ACGGGCGGTGTGTACAA 3’). Twenty-five microliter PCR reactions were prepared using 1x standard Taq buffer (10 mM Tris-HCl, 50 mM KCl, 1.5 mM MgCl₂: NEB, Mississauga ON), 0.2 mM dNTPs, 0.2 µM forward primer, 0.2 µM reverse primer, 0.625 units of standard Taq polymerase (NEB) and 1 µl DNA as template. Cycling parameters were as follow: 94°C for 4 minutes and 32 cycles of 94°C for 30 sec, 55°C for 30 sec and 72°C for 60 sec; and a final 10 minute extension at 72°C. PCR amplicons were visualized by gel electrophoresis using 1 % agarose gels prepared in sodium borate (SB) buffer (5 mM NaOH, 20 mM boric acid, pH 8.5).

PCR reactions were performed on bacterial genomic DNA extracts from intestinal fragments of different life history stages: larval anterior, larval posterior, parasite anterior and parasite posterior. Five independent PCR reactions were performed for each and pools were generated by combining 17 µl from each of the five independent PCR reactions, each of which was generated from a bacterial DNA sample derived from a pool of 10 different tissue samples. The EZ-10 Spin Column PCR Purification Kit (BioBasic, Markham, ON) and the accompanying protocol were used to purify the PCR product pools (BioBasic, Markham, ON). Samples were then quantified...
on a Nanodrop spectrophotometer. Prior to cloning, PCR product pools were dA tailed by combining 1x NEB Standard Buffer, 0.2 mM dATP, 0.5 units NEB Standard Taq polymerase and 3.9 µl pooled PCR product, incubating at 72 °C for 15 minutes followed by snap cooling on ice. The parasite samples were ligated into pGEM-T Vector using protocol from pGEM-T and pGEM-T Easy Vector Systems Technical Manual (Promega, Madison, WI). The vector was then transformed into chemically competent TOP10 cells (Invitrogen, Burlington, ON) according to the manufacturer’s instructions. Transformed cells were plated onto lysogeny broth (LB) plates containing 150 µg/ml ampicillin and top coated with 50 µl of 2 % X-gal. The plates were incubated at 37 °C for 12-16 hours. Larval samples were processed as per parasite samples with the following changes: pooled PCR products were ligated into pCR2.1-TOPO vector (Invitrogen) according to manufacturer’s instructions and then plated onto LB plates containing 50 µg/ml kanamycin. Individual colonies were selected and grown for 16 hours at 37 °C in 150 µl of LB containing 10 % glycerol and either 150 µg/ml ampicillin or 50 µg/ml kanamycin (for parasite and larval samples, respectively). One hundred microliters from each liquid culture was transferred to a 384 well plate, frozen at -20 °C overnight, packaged on dry ice and shipped to Plant Biotechnology Institute in Saskatoon, SK for DNA sequencing. Given preliminary evidence that suggested several predominant microbes in the parasite gut, we sampled more sequences from the parasite to increase the chance of sampling the rare microbes. In total, 451 random 16S rRNA clones were sequenced and analyzed for the parasite anterior and posterior gut samples, and 239 for the larval anterior and posterior samples.

Isolation and phenotypic characterization of culturable isolates
Culturable isolates were obtained by plating washes of gut tissue samples onto LB agar. A total of 62 bacterial isolates were selected from larval, as well as blood-fed and unfed post-metamorphic parasites. Random isolates were selected and identified to genus level by 16S rRNA gene sequencing using the primer set described above. Hemolysin assays were carried out with 48 hour incubations at 30 °C on sheep blood agar (TSA II trypticase soy agar with 10% citrated sheep blood; BD, Mississauga, ON, Canada) and trout blood agar (TSA with 10% citrated trout blood; Fort Qu’Appelle Provincial Trout Hatchery, Fort, Qu’appelle, SK, Canada).

Beta hemolysin activity was expressed as the diameter of lysis (clearing) relative to colony diameter. Protease assays were conducted using skim milk agar (Nutrient Agar with 10% w/v skimmed milk powder). Casein digestion (protease secretion) was scored as positive if clearing was observed following 24 hour incubation. Bile tolerance was evaluated on MacConkey agar (50 g/L Mikrobiolie MacCONKEY™ agar) with varying concentrations of bile salts (1x, 2x, 4x, 6x, 10x; Sigma Aldrich [B8756], St. Louis, MO), with incubation at 30 °C for 12-16 hours. Isolates were scored as bile tolerant if any growth was apparent.

DNA Sequencing of the gyrB gene from culturable Aeromonas isolates

Genomic DNA was extracted from liquid LB cultures grown at 37 °C for 15-20 hours using the Qiagen Puregene Core Kit Yeast/Bacteria genomic DNA Kit according to the manufacturer’s instructions. Genomic DNA was used as template to amplify a portion of the gyrB gene using the following primers: forward 334 (5’ TCCGGCGGTCTGCACGGCGT 3’) and reverse 1464 (5’ TTGTCCGGGTTGTACTCGTC 3’) (32). Twenty-five microliter PCR reactions were performed using 5x KAPA2G Buffer A (1.5 mM MgCl2; Kapabiosystems, Woburn, MA), 0.2 mM dNTPs, 0.2 µM forward primer, 0.2 µM reverse primer, 0.5 units of KAPA2G Robust DNA Polymerase (Kapabiosystems) and 1µl DNA template. Cycling parameters were as follows: 94 °C for 4
minutes and 35 cycles of 94 °C for 30 sec, 60 °C for 30 sec and 72 °C for 60 sec and a final 10 183 
minute extension at 72 °C. PCR amplicons were analyzed using 1 % SB agarose gel 184 
electrophoresis. Following PCR amplification samples were processed for automated DNA 185 
sequencing by adding 1 unit each Calf Intestinal Phosphatase (NEB) and Exonuclease I (NEB) 186 
into each sample, and incubating at 37 °C for 15 minutes, followed by 15 minutes at 80 °C. Eight 187 
microliters of each sample was placed in a 96 well plate and combined with 3.3 μM of the 188 
forward primer (forward 334). Sequencing was performed by Eurofins MWG Operon.

Sequence analysis and phylogenetic reconstruction

Sequences were evaluated for both vector contamination and quality. DNA sequences were 191 
trimmed using the Sequencher 4.8 (GeneCodes Corporation, Michigan) program with the 192 
following parameters: 5' end - trimming no more than 25 %, trim until the first 25 bases contain 193 
less than 3 bases with confidence below 25; 3' end - trim from 3' end until the last 25 bases 194 
contain less than 3 bases with confidences below 25; post fix - remove leading and trailing 195 
ambiguous base, but some sequences had additional sequence kept with lower confidence scores. 196 
The 16S rRNA gene sequences were compared to the Ribosomal Database Project using 198 
Classifier, 16S rRNA training set 9, with a cut-off value of 50% (3), while gyrB sequences were 199 
analyzed using a BLASTN search of Genbank (1). The sequences of the 16S rRNA genes and 200 
gyrB genes have been deposited under GenBank accession numbers JX453764 - JX454446 and 201 
JX453730 - JX453763, respectively. For phylogenetic reconstruction, the 16S rRNA gene and 202 
gyrB sequences were aligned separately using ClustalX, version 2.0 (15) along with known 203 
sequences gathered respectively from the Ribosomal Database and GenBank (2, 3). From these 204 
alignments neighbor-joining phylogenetic trees were constructed with MEGA5 (28).
Comparison of microbial communities between larvae and parasites

A comparison of 16S rRNA gene sequences from both the larval and parasitic sea lamprey revealed a higher bacterial diversity in larvae, with members of the Proteobacteria (36%), Bacteroidetes (30%) and Tenericutes (26%) comprising the majority of diversity sampled. The predominant genera included *Ureaplasma* (15%), *Paludibacter* (11%), *Acinetobacter* (8%) and *Legionella* (5%) (Figure 2). Taxa that were less abundant included *Coxiella* (4%), *Aeromonas* (4%), *Mycoplasma* (3%), *Exiguobacterium* (2%) and *Helicobacter* (2%), along with approximately 20 other genera and unclassified taxa (Supplementary Table 1). An analysis of the distribution of these bacteria along the length of the larval gut revealed some evidence of microbial partitioning, where the larval anterior gut contained predominantly *Acinetobacter* and members of the Bacteriodetes. These groups were underrepresented in the posterior gut, which was dominated by *Ureaplasma* and *Paludibacter*. In contrast, the parasite gut was dominated primarily by *Aeromonas* (84%) with a smaller proportion of *Shewanella* (11%) and *Citrobacter* (2%) species (Figure 2). Other taxa identified in the parasite included *Curvibacter* and *Raoultella*, as well as several others, but these represented only about 3% of the sampled diversity (Supplementary Table 1). The parasite anterior and posterior bacterial populations were quite similar, with most taxa being found throughout; however, *Citrobacter* isolates were more abundant in the posterior than the anterior gut.

Isolation and phenotypic characterization of culturable isolates

Culturable bacteria were isolated from gut tissue samples of the three different stream populations of larval and lab reared parasitic (fed and unfed) sea lampreys, and each isolate was identified by 16S rRNA gene typing. Isolates of *Aeromonas*, *Citrobacter* and *Enterococcus* were cultured from both wild caught larval and lab reared parasite gut tissues. The parasite gut tissues
also yielded isolates of *Shewanella*, *Deinococcus*, *Micrococcus*, *Nitrosomonas* and *Rhodococcus*. With the exception of *Aeromonas*, *Citrobacter* and *Shewanella*, the remaining isolates were not represented in the 16S rRNA metagenomic clone library. Cultured isolates were characterized phenotypically for several environment-specific biochemical capabilities. Firstly, the ability to grow on bile, which is found at high concentrations in the parasite posterior gut (33), was tested using MacConkey agar containing a range of bile salt concentrations. All *Aeromonas* and *Citrobacter* isolates, which were found in anterior and posterior sections of the gut in both larvae and parasites were able to grow on all bile salt concentrations tested, whereas *Enterococcus* isolates, which were also distributed throughout the gut in both lamprey life stages, were not bile tolerant (Table 1). *Shewanella* strains were largely bile tolerant and capable of growing on bile salt concentrations of up to 6x (Table 1).

Isolates were evaluated for their ability to produce enzymes related to digestion. Growth on skim milk agar, an assay developed for evaluating protease secretion, showed that 63% of cultured isolates, including all but two *Aeromonas* isolates (isolates 74 and 91 from the parasite) could hydrolyse casein, whereas all of the *Enterococcus* and *Shewanella* isolates and most of the *Citrobacter* isolates could not (Table 1). Isolates were also evaluated for production of hemolysins using both sheep and trout blood. On sheep blood agar, 31% of the 26 larval isolates tested showed beta hemolysis (39% of the *Aeromonas* isolates) and 54% were gamma/non-hemolytic (61% of the *Aeromonas* isolates). The parasite tended to have more strains that were beta hemolytic on sheep blood agar (40% of all isolates), but a much greater proportion of beta hemolytic *Aeromonas* isolates (68%). On the trout blood agar, approximately 70% of larval and 70% of parasite isolates showed beta hemolysis, with 100% of *Aeromonas* isolates being beta...
hemolytic on trout blood (Table 1). *Citrobacter* and *Shewanella* isolates recovered from the parasite did not exhibit beta hemolytic abilities.

**Phylogeny of Aeromonas**

Given the prevalence of *Aeromonas* species in the parasitic lamprey, the species diversity of the cultured Aeromonads from both larval and parasite samples was evaluated by a phylogenetic approach using the *gyrB* gene (Figure 3). The *gyrB* gene has a mean synonymous (conserved) substitution rate almost four times that of the 16S rRNA gene making it more suitable for establishing species relationships (32). *A. allosaccharophila*, *A. bestiarum*, *A. media*, *A. salmonicida* and *A. veronii* were identified in the larval gut, while *A. media*, *A. salmonicida* and *A. sobria* were found in the parasite gut (Figure 3). There was no apparent correlation between microbial species and either lamprey population (natal stream) or life stage. However, all *Aeromonas* isolates collected from the unfed parasites were *A. media*, with approximately 66% of the cultured isolates from both larva and parasite being identified as *A. media*.

*Aeromonas* strains having similar phenotypic characteristics tend to cluster together in the phylogenetic tree. All *Aeromonas* isolates had beta hemolytic activity on trout blood, whereas sheep blood hemolysis was variable (beta or gamma). The *A. media* group has both beta and gamma hemolytic groups, with the beta hemolytic strains, which were all isolated from the parasite, being monophyletic (Figure 3). In addition, the two isolates that do not produce casein hydrolase, 91 and 74, are also monophyletic within the *A. media* group. The *A. bestiarum* group contains both beta and gamma hemolytic groups, again with beta hemolytic strains being monophyletic. Isolates from the *A. salmonicida*, *A. allosaccharophila* and *A. veronii* were all beta hemolytic, while the *A. sobria* isolates were all gamma hemolytic.
Discussion

The unusually complex lifecycle of the sea lamprey, which includes a non-trophic metamorphic stage and a dramatic shift to a highly specialized diet, provides a unique glimpse into how changes in host physiology impacts the microbial communities in the gut. A comparison of community composition of the larval and parasitic stages using both culture-dependent and culture-independent methods revealed a higher bacterial diversity in the filter-feeding larvae, which is consistent with previous studies that examined microbial diversity in the larval pouched lamprey (24). The predominant bacterial species of the larval pouched lamprey included *Bacillus mycoides* and *Aeromonas hydrophila*, although the genera *Enterobacter* sp., *Pseudomonas* sp., *Clostridium* sp. and *Corynebacterium* were also present (24). Although *Aeromonas*, *Enterobacter*, and *Clostridium* were identified in the larval sea lamprey, the majority of diversity was represented by the Tenericutes (*Ureaplasma* and *Mycoplasma*), as well as members of the Bacteroidetes (*Paludibacter*). It is interesting, however, that *Aeromonas hydrophila* was not recovered from the sea lamprey. Microbial composition of the lamprey, however, has been shown to be correlated with the microbial communities of the river bed and the water in and around the larval burrow (24).

The parasitic stage, which is not a filter feeder and subsists predominantly on a diet of fish blood, has lower microbial diversity than the larval stage, with predominant species of *Aeromonas*, *Citrobacter* and *Shewanella*. Culture-dependent methods recovered additional species that were not present in the 16S rDNA library. This difference may have been due to culturing bias, which favours fast growing species, 16S amplification bias, which could have led to the overrepresentation of the predominant groups like *Aeromonas*, *Citrobacter* and...
Shewanella, and low sampling of clones. The microbiome of parasitic phase lamprey might also be affected in part by the fact they were reared under laboratory conditions, since this environment could potentially expose the lamprey to other microbial consortia. Sampling of wild-caught feeding parasites could shed more light on the influence of lab-rearing on microbial communities, but these animals are extremely difficult to obtain. Nonetheless, there was a clear difference in microbial community structure during the transition from larva to parasite, with a pronounced enrichment of Aeromonas. This community shift may be attributable to the metabolic capabilities of Aeromonas, since many species produce aerolysins that lyse red blood cells and may therefore gain ready access to nutrients in addition to aiding the parasitic host in the digestion of the blood meal and/or nutrient absorption (13, 21). Indeed, our phenotypic tests of culturable isolates revealed that just over half of the culturable Aeromonas isolates tested were capable of lysing sheep blood, while all isolates were capable of lysing trout blood. This suggests that the hemolysins produced by these Aeromonas strains are more specialized to fish blood. Not all isolates have this capability, though. The outgroup (isolate 4), a putative Citrobacter isolate, exhibits alpha hemolytic activity on trout blood. Furthermore, the majority of culturable Aeromonas isolates from both the larval and parasite gut also show protease secretion, indicating possible involvement in the digestion of the protein rich blood diet. Interestingly, the medicinal leech, vampire bat and mosquito, all of which are sanguivorous, have been found to have Aeromonas in their gut (6, 16, 19, 21).

The shift in gut community composition from high bacterial diversity to one predominated by Aeromonas raises several interesting questions about the origin of these isolates and the specific mechanisms underlying their enrichment during metamorphosis. Aeromonads are present in the gut microbiome of many fish (21-23), which is not surprising given the prevalence
of *Aeromonas* in aquatic environments (26). At least five distinct *Aeromonas* species were found to be represented in the larval stage and at least three in the parasite gut; but, the same *Aeromonas* species are found in both life stages. This could suggest that *Aeromonas* may colonize the larval stage from the general environment and persist in the gut during the non-trophic metamorphic period through to adulthood. This is supported by the fact that larvae collected from different localities carry species of *Aeromonas* that are found in the parasite and the fact that there was no evidence of particular phylogenetic groups being represented more in either life stage. Alternatively, the gut microbiome could be purged completely during metamorphosis, allowing for recolonization by free-living bacteria from the general environment. When mosquitoes undergo metamorphosis from larva to adult, they utilize a gut sterilization mechanism, which purges the insect midgut of bacteria (18). Meconial peritrophic membranes that sequester bacteria are formed and surround the larval midgut epithelium and are eventually sloughed off during metamorphosis (18, 25). Such processes may be accompanied by other specificity mechanisms that promote the establishment of specific bacteria from the general environment. In the squid, which forms a mutualistic association with *Vibrio fischeri*, nitric oxide is produced in the light organ tissues that enables colonization of the symbiont through a series of signalling steps (29, 30). For the medicinal leech, the ingested blood meal contains an active complement system capable of killing sensitive bacteria (13, 20, 27). Similar systems could function in the sea lamprey to eliminate some groups of bacteria. It is also possible that the accumulation of toxic bile during metamorphosis could be functioning to impose selection on specific species or groups of bacteria. *Aeromonas* strains tested were all tolerant to high bile concentrations; however, cultured isolates from the parasite were no more bile tolerant than those in the larva suggesting that this alone cannot be responsible for the enrichment observed.
Shewanella and Citrobacter strains that were more prevalent in the parasite tended to be bile tolerant, with the Shewanella strains tolerant to varying degrees. In fact, Citrobacter strains were found to be more prevalent in the parasite posterior gut where bile is known to accumulate. Still, there were several isolates in the parasite that were bile intolerant, suggesting that bile accumulation may not be a primary selective mechanism. If particular host-specificity determinants do function that allow particular strains to persist within the sea lamprey, the colonization by these bacteria may actively exclude other microbes from establishing, possibly through the action of specific anti-microbial compounds. The medicinal leech, for example, has been suggested to possess a single Aeromonas species, Aeromonas veronii bv. sobria (6), which prevents other bacteria from colonizing the digestive tract through the production of antimicrobial compounds (7, 13, 17). Still, it is unclear whether similar, specific mechanisms function in the sea lamprey to promote the persistence and/or colonization of particular strains during sea lamprey metamorphosis. Nonetheless, this study has established that sea lamprey experience a pronounced change in their gut microflora following transition from filter-feeding larvae to sanguivorous parasites. Our results have formed the basis for investigating specific mechanisms that control gut microbiota development, and for establishing the importance of particular species of gut bacteria, such as Aeromonas, in host blood meal digestion and nutrient absorption. Determining the impact of Aeromonas on host fitness may provide a unique target for the development of new lampricidal compounds.

Acknowledgments

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Research Grant to RGM. AT was supported in part by an NSERC Undergraduate Student Research Award.
Figure Legends

Figure 1. Life cycle of the sea lamprey. Adult sea lamprey spawn in the river head-waters. Embryogenesis lasts approximately 17 days, larval lamprey hatch, float downstream and burrow in the sandy river bed where they remain as relatively sedentary filter feeders for 3-7 years. The end of the larval period is marked with a metamorphosis, which involves major morphological and physiological changes to all organ systems. Metamorphosis is a non-trophic period which lasts approximately 6 months. Post-metamorphic juveniles (parasites) migrate downstream to a large body of water (lake or Atlantic Ocean) and commence parasitic feeding on the blood and body fluids of boney fishes. The parasitic feeding phase lasts 12-20 months after which juveniles commence upstream migration, sexually mature into adults, spawn and die. The cross-sections of the larval and parasite intestines depicted in the figure highlight the dramatic morphological changes that occur during metamorphosis.

Figure 2. Distribution of bacterial taxa found in the gut of the lamprey. A. The predominant bacterial groups in the larval stage as indicated. "Tenericutes" comprises Ureaplasma, Mycoplasma, and unclassified Tenericutes; "Other Bacteroidetes" comprises three genera along with related, unclassified taxa; "Other Proteobacteria" comprises ten genera along with unclassified taxa; "Other" comprises seven genera, along with unclassified taxa. B. Prevalent bacterial genera in the parasitic stage. "Other" comprises six taxa, along with unclassified taxa.

Figure 3. Neighbour-joining tree of 34 Aeromonas isolates based on gyrB. Reference taxa include Aeromonas allosaccharophila, AY101777; Aeromonas bestiarum, AY101774; Aeromonas media, AY101782; Aeromonas popoffii, AY101801; Aeromonas salmonicida,
AY101773; Aeromonas sobria, AY101781; Aeromonas veronii bv. sobria, AY101775;
Aeromonas veronii bv. veronii, AY101787; Escherichia coli, HQ660623.
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Table 1. Phenotypic characteristics of culturable isolates from lamprey

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*MacConkey results seen at bile concentrations of 1x, 2x, 4x, 6x, and 10x, where + indicates growth and - indicates no growth. *Blood agar hemolysis results indicated as γ or β. If no hemolysis was observed, the ratio of diameter clearing to diameter of colony. *Isolated from yeast pellet.