# Microbial communities associated with the parasitic copepod *Lepeophtheirus salmonis.*

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## Abstract

*Lepeophtheirus salmonis* is a naturally occurring marine parasite of salmonid fishes in the Northern hemisphere, and a major problem in salmonid aquaculture. In addition to the direct effects on host fish, *L. salmonis* may act as a vector for diseases. Here, the microbial community of *L. salmonis* recovered from whole genome shotgun sequencing was compared between lice sampled from both the Atlantic and the Pacific, laboratory-reared and wild lice, in addition to lice displaying resistance towards chemical treatments. The analysis shows clear differences in the metagenomic composition between the Atlantic and the Pacific Ocean, whereas the resistance status of the *L*. *salmonis* or the cultivation did not have significant impact.

Keywords: Metagenome, marine bacteria, Atlantic salmon, pathogens, vectors

## 1. Introduction

The salmon louse (*Lepeophtheirus salmonis)* is a marine ectoparasite found on salmonid fish in both the Pacific (subspecies: *L. salmonis oncorhynchi*) and the Atlantic oceans (subspecies: *L. salmonis salmonis*) [1]. High concentrations of salmon lice are found in regions hosting salmonid aquaculture industry [2], and are a potential threat to wild sea trout (*Salmo trutta*) and Atlantic salmon (*Salmo salar*) populations in the North East Atlantic [3, 4]. Management of this pathogen in Norway alone is estimated to cost the aquaculture industry 130-390 US$ million yearly [5]. The salmon lice lifecycle consists of eight life stages separated by molt of the chitinous exoskeleton [6]. Apart from the first two planktonic stages, the salmon louse spends its lifecycle on the surface of the host fish, where it feeds on skin, mucus and blood. Infections with salmon lice can cause morbidity in the host fish due to general and osmotic stress, wounds, and increased susceptibility to secondary infections [7]. The integument of the salmon louse is covered with microorganisms [8] and potential pathogens including *Tenacibaculum*, *Pseudomonas* and *Vibrio* [9] and the microsporidian *Paranucleospora theridion* [10] has been identified. In addition to the direct effects of grazing on the fish epidermis, an increased exposure to secondary infections may also be a result of immunomodulation of the host fish through secretions from parasite exocrine glands [11-13]. Such secretions may reduce the ability of the fish to resist infections with other pathogens. A recent study has also documented that salmon lice infections change the microbial composition of salmon skin [14]. To obtain an overview of organisms associated with the salmon louse, a metagenomic analysis was performed on sequences extracted from lice from multiple locations in the Pacific and the Atlantic Oceans. In addition to identification of bacterial orders, the data was also analyzed for metagenomic patterns caused by origin and resistance properties of the collected salmon lice.

## 2. Data description

*Sampling*

Salmon lice sampled for this dataset belongs to the two subspecies of salmon louse sampled from the Pacific (*Lepeophtheirus salmonis oncorhynchi*) and the Atlantic (*Lepeophtheirus salmonis salmonis*), respectively collected in multiple locations (Figure 1).

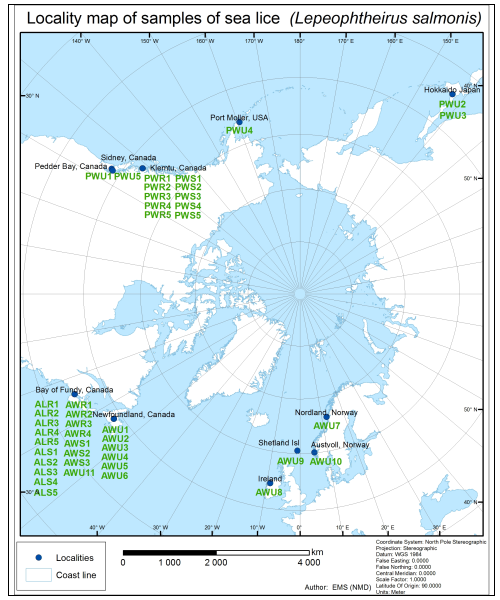


Fig. 1. Geographical distributions of samples. Samples are labelled according to geographical (Atlantic or Pacific), source of lice (laboratory or wild) and resistance status (resistant, sensitive or unknown), see S1.

All samples have previously been described by [15-17]. AWU7-11(S1) contains pooled DNA from multiple individuals in each sample, whereas the remaining samples are from single individuals. Resistance status of the salmon louse has been noted when available. Lice were either collected in the field or cultivated on fish in the laboratory before being submitted to sequencing (S1).

*Bioinformatic analysis*

MetaPhlAn2 was utilized for taxonomic profiling of whole-metagenome shotgun (WMS) samples. Fastq files were first filtered for the presence of salmon lice genomic sequences (http://metazoa.ensembl.org/Lepeophtheirus\_salmonis/Info/Index) using BWA [18]. Libraries contained 10-80 million sequences after filtration. Libraries were then used as input in a MetaPhlAn2 analysis [19] to produce an abundance table of microorganisms present in samples. Sequences assigned to viruses were not included as the sequenced material was based on extraction of DNA only, thus excluding the finding of RNA viruses, resulting therefore in partial data being retrieved. Fastq files were mapped against the MetaPhlAn2 db\_v20/mpa\_v20\_m200.pkl database using bowtie with the “very-sensitive local” setting. The relative abundance of microorganisms was then calculated by Metaphlan2. The resulting abundance file was then used as input for LDA Effect Size (LEfSE) analysis [20] were the alpha value for the factorial Kruskal-Wallis test among classes and the pairwise Wilcoxon tests between subclasses were set to 0.05. Threshold for the logarithmic LDA score for discriminative features were set at the default value of 2. Finally, cladograms were produced setting the root of the tree according to kingdom. Differences in the bacterial metagenome on salmon lice from the Atlantic and Pacific Ocean were assessed using statistical analysis of taxonomic and functional profiles (STAMP) [21]. The Microbiome helper perl script metaphlan\_to\_stamp.pl (<http://msystems.asm.org/content/2/1/e00127-16>) was used to convert the abundance file from MetaPhlAn2 to a STAMP compatible format. Extended error bar plots were produced in STAMP by analyzing differences in the mean proportion of sequences using two-sided Welch’s t-test and Storey FDR multiple test correction. The final data were filtered to remove features with small effect sizes by considering only features were the ratio of the proportions were larger than 2 or the difference in proportions were larger than 0.5.

*Bacterial communities on L. salmonis*

Sequences from Archeae bacteria and eukaryotes were present in the dataset, but very few were recognized by Metaphlan and will not be commented further in this study. The sequences assigned to prokaryotes were initially sorted into 28 phyla, 36 classes and 69 orders. All samples were dominated by two bacterial orders; Flavobacteriales in the Bacteriodetes phylum (30 – 71 %) and Burkholderiales in the Protebacteria phylum (6 – 37 %). The total proportion of Flavobacteriales and Burkholderiales was relatively constant in all samples (average of 58 ± 8 %), however the ratio between the two orders was variable (results not shown). The orders of Flavobacteriales and Burkholderiales contains a great variety of environmental bacteria associated with surface and biofilm in additions to some known pathogenic bacteria that can cause ulcers in fish, e.g. *Flavobacterium psychrophilum.* [22]. Three other bacterial orders Spirochaetales (1 - 16 %), Alphaproteobacteria (3 - 12 %) and Mycoplasmatales (5 - 11 %) were also found in all samples. Other orders included in the Proteobacteria phylum normally associated with marine environment and fish such as Pseudomondales, Pseudoalteromondales, Alteromondales, Aeromondales, Vibrionales and Rickettsiales, were only sporadically represented. These orders are all well-known from the marine environment both as symbionts, and thus part of the normal micro flora, and as pathogens.

All samples were analyzed to search for differences in the composition of the microbial community depending on the geographic origin of the sample. Most samples could be assigned to the Atlantic or the Pacific Ocean based on the microbial composition (fig. 2).

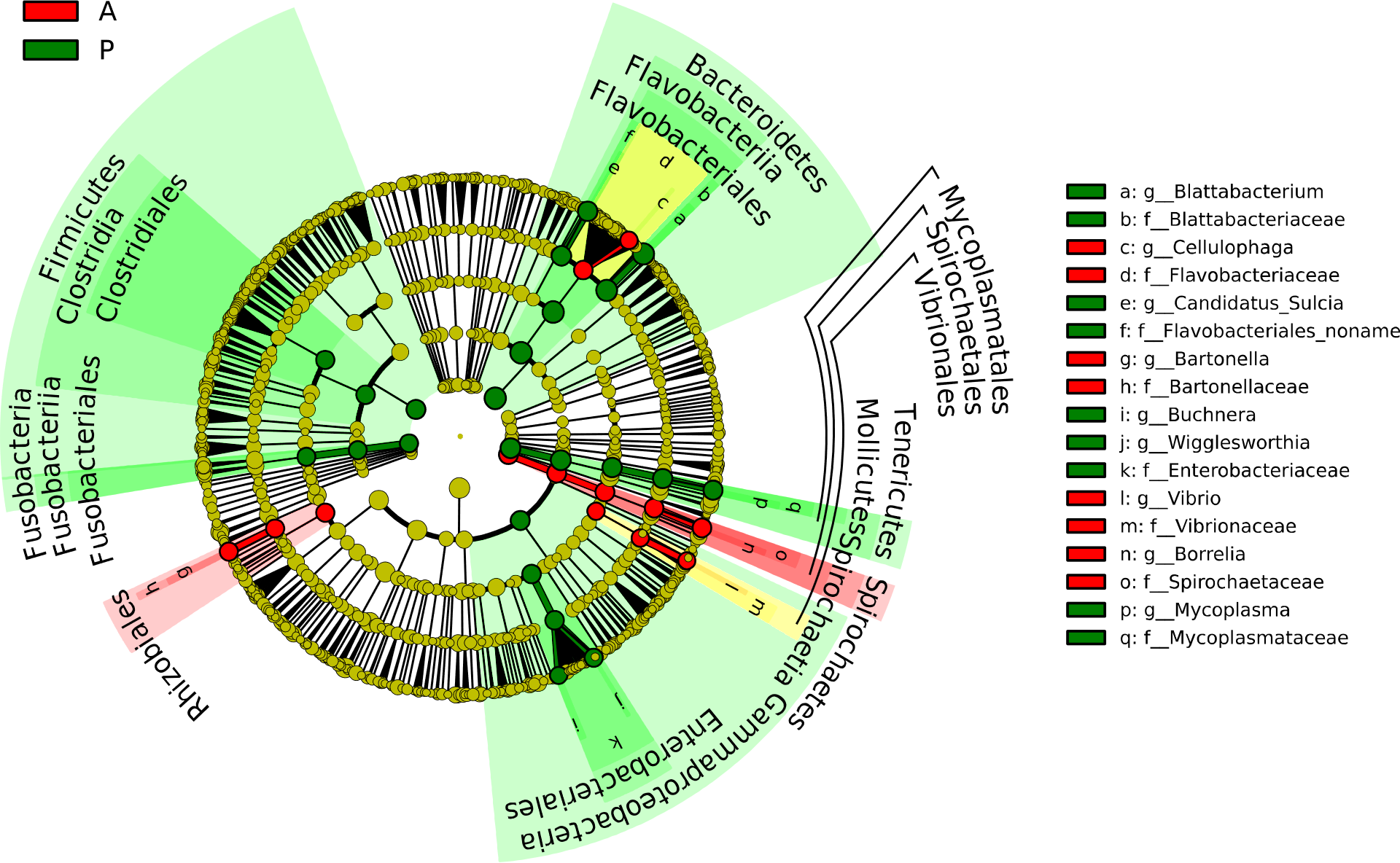


Fig.2. Abundance of bacterial orders. The cladograms report the taxa showing different abundance values (according to LEfSe) in the two geographical areas: Atlantic samples (red) and Pacific samples (green).

It should however be noted that this difference between the two oceans may have been caused by differences in the microbial communities between the Pacific and Atlantic Ocean but could also have been influenced by differences between the two subspecies of *L. salmonis*. Enterobacteriales were present in significantly higher amounts in samples from the Pacific Ocean, while Spirochaetales, Halanaerobiales, Rhizobiales, Rhodobacterales, Sphingobacteriales and Vibrionales were more abundant in the Atlantic samples (Fig 3).

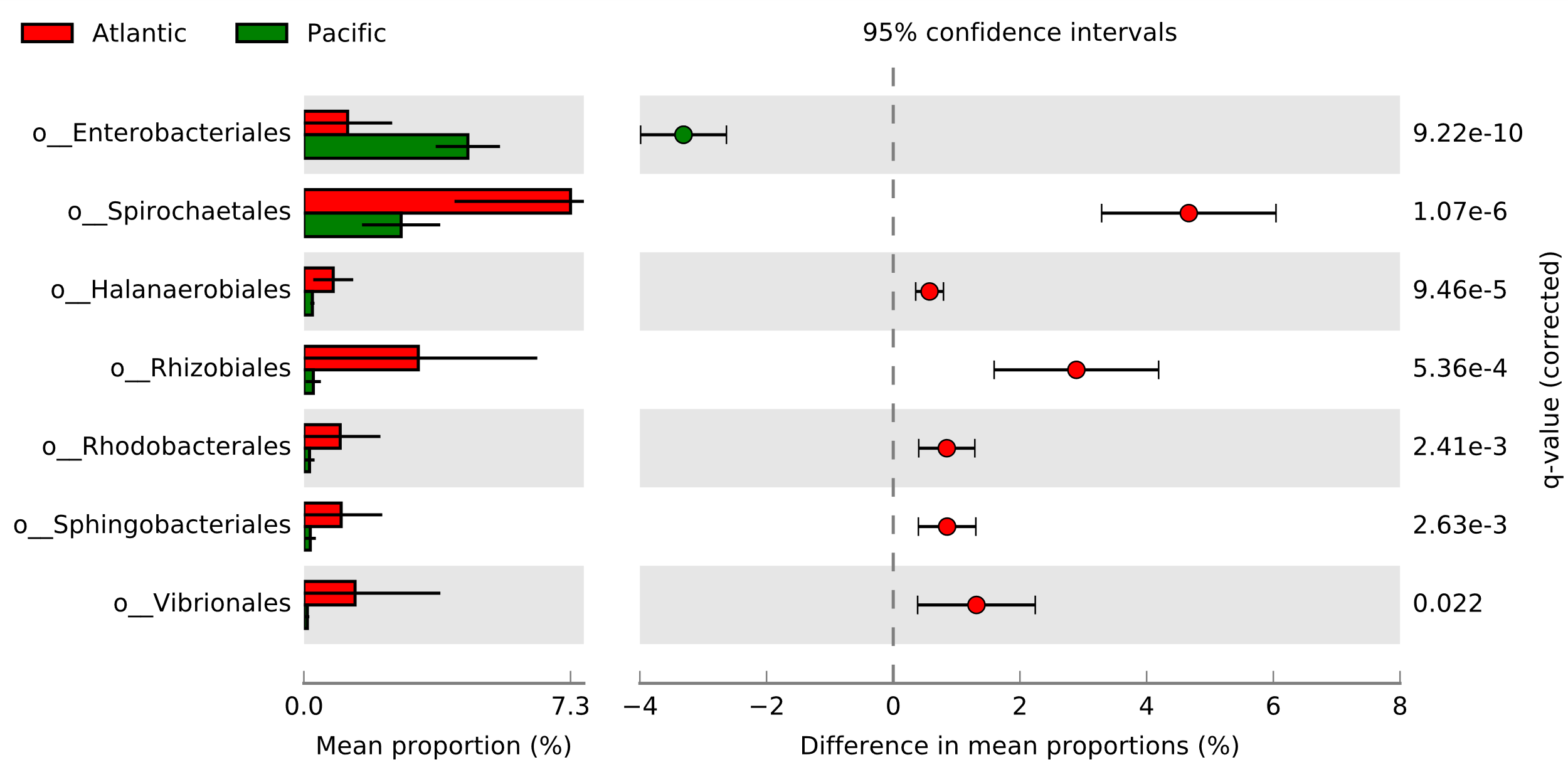


Fig. 3 Extended error bar plot. Bacterial orders with significantly different representation in samples from the Atlantic and Pacific Ocean.

Similar analyses were performed to investigate whether the composition of the microbial community was affected by the resistance status of the salmon louse, or by cultivation in the laboratory. No significant differences were observed. In conclusion, sequences obtained from whole genome sequencing of the parasite *L. salmonis* provides information about associated bacterial orders. Analysis of the data provided metagenomic profiles distinct between the Atlantic and Pacific Ocean but did not demonstrate presence of well described fish pathogens that *L. salmonis* may act as vector for. Future studies should also include analysis of virus and eukaryotes.

## 3. Data Deposition

All sequences utilized in this study can be found in SRA accession PRJNA447894 and PRJNA509381 (ncbi.nlm.nih.gov/bioproject/PRJNA447894 and ncbi.nlm.nih.gov/bioproject/PRJNA509381). For biological information on the samples see supplementary files (S1 and S2).

## 4. Acknowledgements

The analysis of the metagenome data in this study was funded by the Norwegian department for Trade, Industry and Fisheries as part of its funding for the Institute of Marine Research in the internal project 14501. Atlantic sequence data was acquired from the Norwegian Research Council project PrevenT. Pacific sequence data was supported by a Discovery Grant from NSERC Canada.

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