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<td>doi:10.1128/genomeA.01087-13</td>
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High-Quality Draft Genome Sequence of Vagococcus lutrae Strain LBD1, Isolated from the Largemouth Bass Micropterus salmoides

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Vagococci are usually isolated from marine hosts and occasionally from endodontic infections. Using 16S rRNA gene comparison, the closest relatives are members of the genera Enterococcus and Carnobacterium. A draft sequence of Vagococcus lutrae was generated to clarify the relationship of Vagococcus to these and other related low-G+C Gram-positive bacteria.

Received 18 November 2013 Accepted 26 November 2013 Published 26 December 2013

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The bacterial genus Vagococcus was proposed in 1989 for Gram-positive, catalase-negative, motile, coccus-shaped bacteria that react with Lancefield group N antiserum (1). Phylogenetic trees based on 16S rRNA gene sequences place Vagococcus adjacent to the genera Enterococcus and Carnobacterium (2). The genus Vagococcus currently consists of eight species (V. fluvialis, V. salmoninarum, V. lutrae, V. fessus, V. carnipilus, V. elongatus, V. penaei, and V. acidifermentans) (1–8).

Most representatives of Vagococcus have been isolated from aquatic environments, suggesting that members of this genus have traits optimized for existence and survival in marine habitats (1–8). V. fluvialis strains have been suggested as a promising candidate probiotic for aquaculture, a critical economic activity practiced worldwide (9). Interestingly, strains of this genus have also been isolated from patients receiving endodontic treatment for periapical lesions (10).

In this study, we sampled the intestine of a largemouth bass (Micropterus salmoides) that was caught in the wild in Maine. Following outgrowth on bile esculin azide agar, we isolated a strain of V. lutrae named LBD1. This strain was subjected to whole-genome sequencing and constitutes the first report of a genome of the species V. lutrae and the genus Vagococcus.

Genomic DNA was isolated with a DNeasy kit (Qiagen, Valencia, CA) and was quantified by a Qubit fluorometric assay (Invitrogen, Carlsbad, CA). The paired-end library (2 × 250 bp) was prepared using a Nextera XT DNA sample preparation kit (Illumina, San Diego, CA). The quality and quantity of the library DNA fragments were measured on an Agilent Technologies 2100 Bioanalyzer (Santa Clara, CA). Sequencing was carried out on the Illumina MiSeq personal sequencer platform at the Massachusetts Eye and Ear Infirmary (MEEI) Ocular Genomics Institute (Boston, MA). CLC Genomics Workbench version 4.9 software (CLC bio, Cambridge, MA) was used for de novo assembly on an i7 Intel dual-core workstation. For V. lutrae LBD1, 7.48 million paired-end reads were collected. The average coverage of the 1.83-Mb LBD1 assembled genome (21 scaffolds; scaffold N50, 81.28 kb) was 720×.

Protein-coding genes were predicted with Prodigal (11) and filtered to remove genes with >70% overlap with tRNAs or rRNAs, which were identified using tRNAscan-SE (12) and RNAmmer (13), respectively. The gene product names were assigned based on top BLAST hits against the SwissProt protein database and a protein family profile search against the TIGRFam equivalogs, followed by top BLAST hits to KEGG protein sequences. For V. lutrae LBD1, we identified 1,736 protein-coding genes (48% with “hypothetical protein” as the gene product name), 3 rRNA genes (one each for 5S, 16S, and 23S), and 49 tRNAs (19 amino acids with RNA-Asn missing, probably in the contig gap regions). Additional analyses performed included Pfam (14), TIGRFam (15), KEGG (16), COG (17), GO (18), and TMHMM (19) analyses.

The availability of this genome sequence begins to illuminate the roles of vagococci as members of the microbiome of fish and other marine animals and may aid future studies related to the aquaculture industry and/or human medicine.

Nucleotide sequence accession numbers. This whole-genome project has been deposited at DDBJ/EMBL/GenBank under accession no. AYSH00000000. The version of V. lutrae LBD1 described in this paper is version AYSH01000000.

ACKNOWLEDGMENTS
This project was funded in part by the National Institute of Allergy and Infectious Diseases, National Institutes of Health, Department of Health and Human Services, under contract no. HHSN272200900018C. Portions of this work were also supported by NIH/NIAID grants no. AI083214 (Harvard-wide Program on Antibiotic Resistance) and no. AI072360.

REFERENCES


