



Complete Genome Sequence of the Highly Virulent Aeromonas schubertii Strain WL1483, Isolated from Diseased Snakehead Fish (Channa argus) in China

Lihui Liu,^{a,b,c} Ningqiu Li,^{a,c} Defeng Zhang,^{a,b} Xiaozhe Fu,^a Cunbin Shi,^a Qiang Lin,^{a,c} Guijie Hao^b

Key Laboratory of Fishery Drug Development, Ministry of Agriculture Key Laboratory of Aquatic Animal Immune Technology, Pearl River Fishery Research Institute, Chinese Academy of Fishery Sciences, Guangzhou, Guangdong, China^a; Agriculture Ministry Key Laboratory of Healthy Freshwater Aquaculture, Key Laboratory of Fish Health and Nutrition of Zhejiang Province, Zhejiang Institute of Freshwater Fisheries, Huzhou, Zhejiang, China^b; Freshwater Aquaculture Collaborative Innovation Center of Hubei Province, Wuhan, Hubei, China^c

We sequenced the complete genome of the highly virulent *Aeromonas schubertii* strain WL1483, which was isolated from diseased snakehead fish (*Channa argus*) in China. The full genome sequence of *A. schubertii* WL1483 is 4,400,034 bp, which encodes 4,376 proteins and contains 195 predicted RNA genes.

Received 12 November 2015 Accepted 27 November 2015 Published 21 January 2016

Citation Liu L, Li N, Zhang D, Fu X, Shi C, Lin Q, Hao G. 2016. Complete genome sequence of the highly virulent *Aeromonas schubertii* strain WL1483, isolated from diseased snakehead fish (*Channa argus*) in China. Genome Announc 4(1):e01567-15. doi:10.1128/genomeA.01567-15.

Copyright © 2016 Liu et al. This is an open-access article distributed under the terms of the Creative Commons Attribution 3.0 Unported license.

Correspondence to Defeng Zhang, zhangdefeng08@126.com, or Guijie Hao, 391601350@qq.com.

eromonas schubertii is a Gram-negative, short rod-shaped bacterium, commonly isolated from abscesses, wounds, skin, and pleural fluid (1-3). It can cause septicemia with necrotizing fasciitis and posttraumatic soft tissue infection, especially with trauma associated with freshwater environment or marine animals (4). A. schubertii was first confirmed as the etiological agent of the epizootic in cultured snakehead fish and is becoming a major economic problem in the snakehead farming industry (5, 6). The strain A. schubertii WL1483 was a representative isolate collected from a 2014 disease outbreak among snakeheads in earthen ponds in Foshan, Guangdong Province, China. Virulence studies revealed that A. schubertii WL1483 was highly virulent to snakehead fish (Channa argus). The pathogenesis of A. schubertii is not understood as yet, and fewer sequences of A. schubertii are available. Therefore, the complete genome sequence of A. schubertii WL1483 was determined in this study.

Genomic DNA was extracted from the isolate *A. schubertii* strain WL1483, according to the protocol of the Bacterial DNA kit (OMEGA, USA). The DNA was sequenced using the PacBio RSII and Illumina HiSeq platforms. The hierarchical genome assembly process from the Pacific Biosciences single-molecule real-time analysis toolkit (7) was used to obtain a *de novo* assembly. The PacBio RSII reads were aligned to the trimmed assembly using BLAST to correct the structure error. The original reads of HiSeq were used to fill outer gaps and correct the assembly results. The results showed an assembly with a single contig and a sequence length of 4,400,034 bp, with an average coverage depth of 587.5×. The genome sequence has a mean G+C content of 61.49%.

The assembled sequence was annotated using the GeneMarkS annotation system for prokaryotic genomes (8). tRNA genes were predicted with tRNAscan-SE (9), rRNA genes were predicted with rRNAmmer (10), and sRNAs were predicted by BLAST against the Rfam database (11). A total of 4,376 protein-coding sequences were identified, together with 107 tRNA genes, 81 rRNA genes,

and 7 sRNA genes. Protein annotation using the VFDB database (http://www.mgc.ac.cn/VFs) of virulent factors for bacterial pathogens detected 365 putative virulence factors, and 308 secretory proteins were detected on the genome assembly by SignalP (12).

Nucleotide sequence accession numbers. The sequence and annotation of *A. schubertii* strain WL1483 has been deposited at GenBank under the accession number CP013067, BioSample number SAMN04110141, and BioProject number PRJNA297116.

ACKNOWLEDGMENTS

This work was supported by a grant from Open Foundation of Agriculture Ministry Key Laboratory of Healthy Freshwater Aquaculture (ZJK2014002) and Modern Agro-Industry Technology Research System (nycytx-49-14).

We thank Total Genomics Solution Limited (TGS) and Shanghai OE Biotech Co. Ltd. for bioinformatics analysis support. We report no conflicts of interest and are alone responsible for the content and writing of the paper.

FUNDING INFORMATION

This research received no specific grant from any funding agency in the public, commercial, or not-for-profit sectors.

REFERENCES

- Carnahan AM, Marii MA, Fanning GR, Pass MA, Sam J. 1989. Characterization of *Aeromonas schubertii* strains recently isolated from traumatic wound infections. J Clin Microbiol 27:1826–1830.
- Hickman-Brenner FW, Fanning GR, Arduino MJ, Brenner DJ, Farmer JJ. 1988. Aeromonas schubertii, a new mannitol-negative species found in human clinical specimens III. J Clin Microbiol 26:1561–1564.
- 3. Kokka RP, Lindquist D, Abbot SL, Janda JM. 1992. Structural and pathogenic properties of *Aeromonas schubertii*. Infect Immun 60: 2075–2082.
- Kao TL, Kao ML. 2012. A fatal case of necrotizing Aeromonas schubertii fasciitis after penetrating injury. Am J Emerg Med 30:e3-e5. http:// dx.doi.org/10.1016/j.ajem.2010.10.028.

- Liu JY, Li AH. 2012. First case of Aeromonas schubertii infection in the freshwater cultured snakehead fish, Ophiocephalus argus (Cantor), in China. J Fish Dis 35:335–342. http://dx.doi.org/10.1111/j.1365 -2761.2012.01350.x.
- Chen YF, Liang RS, Zhuo XL, Wu XT, Zou JX. 2012. Isolation and characterization of *Aeromonas schubertii* from diseased snakehead, *Channa maculata* (Lacepède). J Fish Dis 35:421–430. http://dx.doi.org/ 10.1111/j.1365-2761.2012.01362.x.
- Chin C, Alexander DH, Marks P, Klammer AA, Drake J, Heiner C, Clum A, Copeland A, Huddleston J, Eichler EE, Turner SW, Korlach J. 2013. Nonhybrid, finished microbial genome assemblies from long-read SMRT sequencing data. Nat Methods 10:563–569. http://dx.doi.org/ 10.1038/nmeth.2474.
- Besemer J, Lomsadze A, Borodovsky M. 2001. GeneMarkS: a selftraining method for prediction of gene starts in microbial genomes. Im-

plications for finding sequence motifs in regulatory regions. Nucleic Acids Res **29:**2607–2618. http://dx.doi.org/10.1093/nar/29.12.2607.

- Lowe TM, Eddy SR. 1997. tRNAscan-SE: a program for improved detection of transfer RNA genes in genomic sequence. Nucleic Acids Res 25: 955–964. http://dx.doi.org/10.1093/nar/25.5.0955.
- Lagesen K, Hallin P, Rødland EA, Staerfeldt H-H, Rognes T, Ussery DW. 2007. RNAmmer: consistent and rapid annotation of ribosomal RNA genes. Nucleic Acids Res 35:3100–3108. http://dx.doi.org/10.1093/nar/gkm160.
- Gardner PP, Daub J, Tate JG, Nawrocki EP, Kolbe DL, Lindgreen S, Wilkinson AC, Finn RD, Griffiths-Jones S, Eddy SR, Bateman A. 2009. Rfam: updates to the RNA families database. Nucleic Acids Res 37(suppl 1):D136–D140. http://dx.doi.org/10.1093/nar/gkn766.
- 12. Petersen TN, Brunak S, von Heijne G, Nielsen H. 2011. SignalP 4.0: discriminating signal peptides from transmembrane regions. Nat Methods 8:785–786. http://dx.doi.org/10.1038/nmeth.1701.