





## Draft Genome Sequence of *Marinobacter* sp. Strain LZ-6, Isolated from the Toxic Dinoflagellate Alexandrium catenella

Qiao Yang,<sup>a</sup> Zhiwei Jiang,<sup>a</sup> Xin Zhou,<sup>a</sup> Xiaoling Zhang<sup>a</sup>

<sup>a</sup>ABI Group, College of Marine Science and Technology, Zhejiang Ocean University, Zhoushan, Zhejiang, China

**ABSTRACT** Here, we report the draft genome sequence of *Marinobacter* sp. strain LZ-6, isolated from the cell culture of a toxic marine dinoflagellate, Alexandrium catenella LZT09. A total of 4,405 predicted protein-coding genes were revealed, including those associated with initial biosynthesis of the key intermediate of paralytic shellfish poisoning toxins (PSTs), namely saxitoxin, and with toxic compound extrusion.

arinobacter species are mesophilic, halotolerant or halophilic, and chemoheterotrophic bacteria most found in saline habitats, and many of them could be of potential interest for biotechnological and bioremediation applications due to their capacity to utilize hydrocarbon contaminants or to degrade plastic or radionuclides (1-4). However, few cultivable Marinobacter species are found to be associated with marine dinoflagellates, although members of Marinobacter are globally distributed in aquatic environments (5). Previously, Marinobacter sp. strain LZ-6 was isolated by serial dilution of the algal cells of toxic Alexandrium catenella and was plated on Difco marine 2216 agar (MA, USA) (6) and cultivated for 2 to 7 days at 28°C during the diversity investigation of the cultivable bacterial community associated with toxic dinoflagellates (7). A. catenella is a typical producer of paralytic shellfish poisoning toxins (PSTs) derived from globally widespread harmful algal blooms (8). For a better understanding of the bacterial roles in alga-bacterium interactions, including host PST biosynthesis, the genome sequence of strain LZ-6 was determined.

Strain LZ-6 was cultured in Difco marine broth 2216 medium at 28°C with shaking (180 rpm) for 2 days. Genomic DNA was extracted with a DNeasy UltraClean microbial DNA isolation kit, according to the instructions from Qiagen (MD, USA). The Illumina  $2 \times 250$ -bp paired-end library was prepared using a TruSeq DNA sample prep kit for Illumina (MA, USA), according to the manufacturer's instructions, and then sequenced using a HiSeq 4000 platform (Illumina, CA, USA). Trimming and quality filtering with Trimmomatic v0.36 (9) yielded  $5 \times 10^6$  Phred Q30 reads. The read quality was assessed with FastQC v0.11.2 (http://www.bioinformatics.babraham.ac.uk/projects/fastqc). Genome assembly was performed with SPAdes v3.5.0 using the standard default settings (10). The assembled 55 contigs have  $340 \times$  coverage with a length of 4,652,078 bp, a GC content of 57.0%, and an  $N_{50}$  value of 312,274 bp. Genome annotation through the NCBI Prokaryotic Genome Annotation Pipeline (PGAP) v1.2.1 (11) predicted 4,465 genes, which comprised 4,405 protein-coding genes, 53 tRNAs, 4 rRNAs, and 3 noncoding RNAs (ncRNAs).

Based on the genomic annotations, multiple homologous genes of four catalytic domains of the key initial sxtA gene involved in saxitoxin biosynthesis (12) were revealed, comprising those encoding methyltransferase (MTF; sxtA1), acetyltransferase (ATF; sxtA2), acyl-carrier-protein (ACP; sxtA3), and aminotransferase (ATF; sxtA4). Furthermore, genes encoding multidrug transporters of the ATP-binding cassette (ABC), multidrug and toxic compound extrusion (MATE) proteins (sxtF and sxtM), and major facilitator superfamily families were also identified (13).

Citation Yang Q, Jiang Z, Zhou X, Zhang X. 2019. Draft genome sequence of Marinobacter sp. strain LZ-6, isolated from the toxic dinoflagellate Alexandrium catenella. Microbiol Resour Announc 8:e00795-19. https://doi.org/ 10.1128/MRA.00795-19.

Editor J. Cameron Thrash, University of Southern California

Copyright © 2019 Yang et al. This is an openaccess article distributed under the terms of the Creative Commons Attribution 4.0 International license.

Address correspondence to Xiaoling Zhang, zhangxiaoling@zjou.edu.cn.

Received 25 July 2019 Accepted 5 August 2019 Published 12 September 2019



**Data availability.** The GenBank accession number for the draft genome sequence of *Marinobacter* sp. strain LZ-6 is SWKL00000000. The BioProject and BioSample accession numbers for this project are PRJNA533500 and SAMN11445130, respectively. The raw sequence reads also have been deposited in the NCBI Sequence Read Archive (SRA) under the accession number SRR9678777.

## **ACKNOWLEDGMENTS**

This work was supported by Natural Science Foundation of Zhejiang Province grant LY18D060007 (to Q. Yang), National Natural Science Foundation of China grant 41876114, and Municipal Science and Technology Project of Zhoushan grant 2017C32083 (to X. Zhang).

## **REFERENCES**

- Boujida N, Palau M, Charfi S, Manresa À, Senhaji NS, Abrini J, Miñana-Galbis D. 2019. Marinobacter maroccanus sp. nov., a moderately halophilic bacterium isolated from a saline soil. Int J Syst Evol Microbiol 69:227–234. https://doi.org/10.1099/ijsem.0.003134.
- Hollensteiner J, Poehlein A, Daniel R. 2019. Complete genome sequence of *Marinobacter* sp. strain JH2, isolated from seawater of the Kiel Fjord. Microbiol Resour Announc 8:e00596-19. https://doi.org/10.1128/MRA.00596-19.
- Palau M, Boujida N, Manresa À, Miñana-Galbis D. 2018. Complete genome sequence of Marinobacter flavimaris LMG 23834<sup>T</sup>, which is potentially useful in bioremediation. Genome Announc 6:e00273-18. https://doi .org/10.1128/genomeA.00273-18.
- Ivanova EP, Ng HJ, Webb HK, Feng G, Oshima K, Hattori M, Ohkuma M, Sergeev AF, Mikhailov VV, Crawford RJ, Sawabe T. 2014. Draft genome sequences of Marinobacter similis A3d10<sup>T</sup> and Marinobacter salarius R9SW1<sup>T</sup>. Genome Announc 2:e00442-14. https://doi.org/10 .1128/genomeA.00442-14.
- Green DH, Bowman JP, Smith EA, Gutierrez T, Bolch CJ. 2006. Marinobacter algicola sp. nov., isolated from laboratory cultures of paralytic shellfish toxin-producing dinoflagellates. Int J Syst Evol Microbiol 56: 523–527. https://doi.org/10.1099/ijs.0.63447-0.
- Fidalgo C, Henriques I, Rocha J, Tacao M, Alves A. 2016. Culturable endophytic bacteria from the salt marsh plant *Halimione portulacoides*: phylogenetic diversity, functional characterization, and influence of metal(loid) contamination. Environ Sci Pollut Res 23:10200–10214. https:// doi.org/10.1007/s11356-016-6208-1.
- 7. Yang Q, Zhang X, Li L, Zhang R, Feng L, Mu J. 2018. *Ponticoccus alexandrii* sp. nov., a novel bacterium isolated from the marine toxigenic dinofla-

- gellate *Alexandrium minutum*. Antonie Van Leeuwenhoek 111:995–1000. https://doi.org/10.1007/s10482-017-0996-2.
- 8. Vandersea MW, Kibler SR, Tester PA, Holderied K, Hondolero DE, Powell K, Baird S, Doroff A, Dugan D, Litaker RW. 2018. Environmental factors influencing the distribution and abundance of *Alexandrium catenella* in Kachemak Bay and Lower Cook Inlet, Alaska. Harmful Algae 77:81–92. https://doi.org/10.1016/j.hal.2018.06.008.
- Bolger AM, Lohse M, Usadel B. 2014. Trimmomatic: a flexible trimmer for Illumina sequence data. Bioinformatics 30:2114–2120. https://doi.org/10 .1093/bioinformatics/btu170.
- Bankevich A, Nurk S, Antipov D, Gurevich AA, Dvorkin M, Kulikov AS, Lesin VM, Nikolenko SI, Pham S, Prjibelski AD, Pyshkin AV, Sirotkin AV, Vyahhi N, Tesler G, Alekseyev MA, Pevzner PA. 2012. SPAdes: a new genome assembly algorithm and its applications to single-cell sequencing. J Comput Biol 19:455–477. https://doi.org/10.1089/cmb.2012.0021.
- Tatusova T, DiCuccio M, Badretdin A, Chetvernin V, Nawrocki EP, Zaslavsky L, Lomsadze A, Pruitt KD, Borodovsky M, Ostell J. 2016. NCBI Prokaryotic Genome Annotation Pipeline. Nucleic Acids Res 44: 6614–6624. https://doi.org/10.1093/nar/gkw569.
- Kellmann R, Mihali TK, Jeon YJ, Pickford R, Pomati F, Neilan BA. 2008. Biosynthetic intermediate analysis and functional homology reveal a saxitoxin gene cluster in cyanobacteria. Appl Environ Microbiol 74: 4044–4053. https://doi.org/10.1128/AEM.00353-08.
- Jagessar KL, Mchaourab HS, Claxton DP. 2019. The N-terminal domain of an archaeal multidrug and toxin extrusion (MATE) transporter mediates proton coupling required for prokaryotic drug resistance. J Biol Chem 294:12807–12814. https://doi.org/10.1074/jbc.RA119.009195.

Volume 8 Issue 37 e00795-19 mra.asm.org **2**