



Draft Genome Sequence of Alkalicoccus saliphilus DSM 15402^T, a Haloalkaliphilic Bacterium Isolated from a Mineral Pool

Dacheng Qiu,^a Ziya Liao,^a Weidong Lu,^b Haisheng Wang,^a Jun Li,^c Baisuo Zhao^a

^aGraduate School, Chinese Academy of Agricultural Sciences, Beijing, People's Republic of China ^bSchool of Life Sciences, Qingdao Agricultural University, Qingdao, People's Republic of China ^cInstitute of Agricultural Resources and Regional Planning, Chinese Academy of Agricultural Sciences, Beijing, People's Republic of China

ABSTRACT The haloalkaliphilic bacterium *Alkalicoccus saliphilus* DSM 15402^T was isolated from a mineral pool. It grows aerobically at an optimum of 15% (wt/vol) salinity and pH 9.0. The draft genome consists of approximately 3.52 Mb and contains 3,434 predicted genes. Various genes are potentially involved in the adaptation mechanisms for both osmotic stress and pH homeostasis, providing insight into specific adaptations to this double-extreme environment.

The mesophilic haloalkaliphilic bacterium *Alkalicoccus saliphilus* DSM 15402^T was aerobically isolated from a mineral pool located in Malvizza in the Campania region of Italy (1). This isolate was originally characterized and recommended as *Bacillus saliphilus* and reclassified as *Alkalicoccus saliphilus* by Zhao et al. (2). Its growth occurs at a wide range of 1 to 25% (wt/vol) salinity (optimal salinity, 16%) and at pH 6.5 to 10.0 (optimal pH, 9.0) (2). To comprehend the adaptive strategies of survival under saline-alkaline conditions, the draft genome of *A. saliphilus* DSM 15402^T was sequenced using the Illumina MiSeq platform.

Cells of A. saliphilus grown under optimal conditions were collected (2), and genomic DNA was extracted by using the iTop microbial DNA isolation kit (Beijing, People's Republic of China) according to the manufacturer's instructions. An Illumina sequencing library was constructed using a TruSeq Nano DNA library kit with the whole-genome shotgun (WGS) method. Sequencing was performed at roughly 159imescoverage with a paired-end read length of 2×300 bp. The filtered reads were quality inspected using Quake and the Burrows-Wheeler Aligner (BWA) with the default program parameters and de novo assembled into contigs using A5-miseg version 20150522 (3, 4). A total of 2,438,804 reads were yielded and assembled into 22 contigs. The total length of the draft genome sequence was 3,525,217 bp with a GC content of 45.69% and an N_{50} value of 561,230 bp. Automatic annotation was performed using the NCBI Prokaryotic Genome Annotation Pipeline (PGAP) (https://www.ncbi.nlm.nih.gov/ genome/annotation_prok) and checked by the GenBank curation team for the standard requirements. Then, the genome files in GenBank format (gb file) were uploaded to the Integrated Microbial Genomes Expert Review (IMG ER) tool (https://img.jgi.doe.gov/ cgi-bin/submit/main.cgi) for functional annotation after an analysis project identification (ID) number was registered in the GOLD Database (http://gold.jgi.doe.gov/index). Among the 3,434 predicted genes, 3,357 are potential protein-coding genes (CDSs). A total of 77 RNA genes (6 5S rRNAs, 4 16S rRNAs, 4 23S rRNAs, 59 tRNAs, and 4 other RNA genes) were predicted.

The genome sequence analysis revealed some crucial genes encoding putative proteins potentially associated with the adaptation mechanism of *A. saliphilus* to life under elevated salinity and alkaline pH. The genome harbors 1 gene cluster of *ectA*, *ectB*, and *ectC* genes for ectoine biosynthesis, 1 *glnA* gene for L-glutamine synthesis, 1

Citation Qiu D, Liao Z, Lu W, Wang H, Li J, Zhao B. 2019. Draft genome sequence of *Alkalicoccus saliphilus* DSM 15402^T, a haloalkaliphilic bacterium isolated from a mineral pool. Microbiol Resour Announc 8:e00266-19. https://doi.org/10.1128/MRA.00266-19.

Editor Julia A. Maresca, University of Delaware Copyright © 2019 Qiu et al. This is an openaccess article distributed under the terms of the Creative Commons Attribution 4.0 International license.

Address correspondence to Baisuo Zhao, bszhao@live.com.

Received 6 March 2019 **Accepted** 1 May 2019 **Published** 13 June 2019 *betA* and 2 *betB* genes responsible for glycine betaine synthesis from choline, 3 genes for the glycine betaine-carnitine-choline transporter (BCCT family), and 9 genes for the glycine betaine ABC transporter. These genes show a strategy for coping with high external salinity by accumulating a large amount of compatible solutes, such as ectoine, glutamine, and glycine betaine (5, 6). Furthermore, the presence of 4 genes for the K⁺ uptake protein (Trk family) indicated that *A. saliphilus* also maintains osmotic balance by inducing a massive uptake of K⁺ when coping with a rapid osmotic shock (7). *A. saliphilus* is an obligate haloalkaliphile and therefore must have an adaptive strategy for pH homeostasis, because it has 4 genes of a monovalent cation:proton antiporter (CPA) belonging to the CPA 1 family (1 gene of the K⁺/H⁺ antiporter and 3 genes of the Na⁺/H⁺ antiporter), 7 genes of the multisubunit Na⁺/H⁺ antiporter, 1 gene of the Na⁺/H⁺ antiporter (NhaC family), and 8 genes for F_oF₁ ATP synthase (8–10). The genes on the genome of *A. saliphilus* that are identified in this report might play essential roles in the adaptive mechanisms of this haloalkaliphile.

Data availability. The draft genome sequence of *Alkalicoccus saliphilus* DSM 15402^T has been deposited at GenBank under the accession number PZJJ00000000. The draft genome sequence described in this paper is the first version (PZJJ01000000). The raw sequencing reads have been submitted to the Sequence Read Archive (SRA accession number SRR8449870) and are available in NCBI under BioProject number PRJNA437190 and BioSample number SAMN08640882.

ACKNOWLEDGMENTS

This work was supported by grants 31570110 and 31370158 from the National Science Foundation of China (NSFC) and grant 1610042017001 from the Foundation of Graduate School of the Chinese Academy of Agricultural Sciences (CAAS).

REFERENCES

- Romano I, Lama L, Nicolaus B, Gambacorta A, Giordano A. 2005. *Bacillus saliphilus* sp. nov., isolated from a mineral pool in Campania, Italy. Int J Syst Evol Microbiol 55:159–163. https://doi.org/10.1099/ijs.0.63298-0.
- Zhao B, Lu W, Zhang S, Liu K, Yan Y, Li J. 2017. Reclassification of *Bacillus saliphilus* as *Alkalicoccus saliphilus* gen. nov., comb. nov., and description of *Alkalicoccus halolimnae* sp. nov., a moderately halophilic bacterium isolated from a salt lake. Int J Syst Evol Microbiol 67:1557–1563. https://doi.org/10.1099/ijsem.0.001759.
- Coil D, Jospin G, Darling AE. 2015. A5-miseq: an updated pipeline to assemble microbial genomes from Illumina MiSeq data. Bioinformatics 31:587–589. https://doi.org/10.1093/bioinformatics/btu661.
- Kelley DR, Schatz MC, Salzberg SL. 2010. Quake: quality-aware detection and correction of sequencing errors. Genome Biol 11:R116. https://doi .org/10.1186/gb-2010-11-11-r116.
- Roberts MF. 2005. Organic compatible solutes of halotolerant and halophilic microorganisms. Saline Systems 1:5. https://doi.org/10.1186/1746 -1448-1-5.

- Banciu HL, Muntyan MS. 2015. Adaptive strategies in the doubleextremophilic prokaryotes inhabiting soda lakes. Curr Opin Microbiol 25:73–79. https://doi.org/10.1016/j.mib.2015.05.003.
- Stumpe S, Schlösser A, Schleyer M, Bakker EP. 1996. K+ circulation across the prokaryotic cell membrane: K+-uptake systems, p 473–499. *In* Konings WN, Kaback HR, Lolkema JS (ed), Handbook of biological physics, vol 2. Elsevier, Amsterdam, The Netherlands.
- Padan E, Bibi E, Ito M, Krulwich TA. 2005. Alkaline pH homeostasis in bacteria: new insights. Biochim Biophys Acta 1717:67–88. https://doi .org/10.1016/j.bbamem.2005.09.010.
- Banciu HL, Sorokin DY. 2013. Adaptation in haloalkaliphiles and natronophilic bacteria, p 121–178. *In* Seckbach J, Oren A, Stan-Lotter H (ed), Polyextremophiles. Springer, Dordrecht, The Netherlands.
- Hicks DB, Liu J, Fujisawa M, Krulwich TA. 2010. F1F0-ATP synthases of alkaliphilic bacteria: lessons from their adaptations. Biochim Biophys Acta 1797:1362–1377. https://doi.org/10.1016/j.bbabio.2010.02.028.