



Draft Genome Sequence of *Bacillus salarius* IM0101, Isolated from Hypersaline Soil in Inner Mongolia, China

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ABSTRACT *Bacillus salarius* IM0101 is a halophilic bacterium that was isolated from soil in Inner Mongolia, China. The genome sequence of *B. salarius* IM0101 contains a biomarker gene for polyhydroxyalkanoate (PHA) synthesis. This 6.9-Mb draft genome sequence of *B. salarius* IM0101 will provide new insights into the organism's PHA production machinery.

Bacillus salarius is Gram-positive, rod-shaped, and strictly aerobic bacterium. The optimum growth conditions are 10 to 12% (wt/vol) NaCl, 30°C, and pH 8.0. MK-7 is its major menaquinone. *B. salarius* contains saturated branched fatty acids, including anteiso-C15:0 and anteiso-C17:0. The cell wall of *B. salarius* is A1γ type peptidoglycan with meso-diaminopimelic acid, similar to most of the members of the genus *Bacillus*; also similar to other *Bacillus* species are the components of major fatty acids, lipoquinone, and G+C content (1).

Strain IM0101 was isolated from soil collected in Bayan Nur, Inner Mongolia Autonomous Region, China. The soil sample contained 18% (wt/vol) NaCl. A single colony of *B. salarius* IM0101 was cultured in JCM169 broth (2) for 3 days at 37°C with shaking at 250 rpm. The partial 16S rRNA gene (1,353 bp) of strain IM0101 was identified using the EzBioCloud database (3) and was 99.93% identical to that of *Bacillus salarius*. High-quality genomic DNA (gDNA) of *B. salarius* IM0101 was obtained using the phenol-chloroform method described by Sambrook and Russel (4). The quantity of gDNA was determined by NanoDrop spectrophotometry (Thermo Fisher Scientific, USA). A sequencing library was prepared using the Ion Plus fragment library kit and the Ion PI Hi-Q OT2 200 template kit. The sequencing was performed on the Ion Proton sequencer using the Ion PI Hi-Q sequencing 200 kit and the Ion PI chip (Thermo Fisher Scientific, USA). The average length of the reads was 145 bp. There were 10,229,665 raw reads (208× depth of coverage) generated from the sequencing run.

De novo assembly of the raw reads was performed using SPAdes version 3.12.0 (5). The quality of the genome assemblies was determined using QUAST version 5.0.0 (6). The draft genome sequence of *B. salarius* IM0101 consists of 6,960,367 bp in 326 contigs with an N_{50} value of 112,419 bp and a CG content of 40.1%. Genome annotation and gene prediction were performed using Prokka version 1.13.3 (7). The predicted genome sequence of *B. salarius* IM0101 contains 6,970 protein-coding sequences and 98 tRNA, 5 rRNA, and 2 transfer-messenger RNA (tmRNA) genes.

There is one copy number of the *phaC* gene in the *B. salarius* IM0101 genome. The *phaC* gene encodes the PhaC subunit of polyhydroxyalkanoate (PHA) synthase, the key enzyme of PHA production (8–10).

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Data availability. The whole-genome shotgun sequence of *Bacillus salarius* IM0101 has been deposited at DDBJ/ENA/GenBank under the accession number [RBVX00000000](#) (version RBVX01000000) and SRA accession number [SRR8357421](#).

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REFERENCES

1. Lim J, Jeon C, Lee S, Lee J, Xu L, Jiang C, Kim C. 2006. *Bacillus salarius* sp. nov., a halophilic, spore-forming bacterium isolated from a salt lake in China. *Int J Syst Evol Microbiol* 56:373–377. <https://doi.org/10.1099/ijs.0.63678-0>.
2. Minegishi H, Shimane Y, Echigo A, Ohta Y, Hatada Y, Kamekura M, Maruyama T, Usami R. 2013. Thermophilic and halophilic beta-agarase from a halophilic archaeon *Halococcus* sp. 197A. *Extremophiles* 17: 931–939. <https://doi.org/10.1007/s00792-013-0575-z>.
3. Yoon SH, Ha SM, Kwon S, Lim J, Kim Y, Seo H, Chun J. 2017. Introducing EzBioCloud: a taxonomically united database of 16S rRNA gene sequences and whole-genome assemblies. *Int J Syst Evol Microbiol* 67: 1613–1617. <https://doi.org/10.1099/ijsem.0.001755>.
4. Sambrook J, Russell RW. 2001. Preparation and analysis of eukaryotic genomic DNA, molecular cloning: a laboratory manual, 3rd ed, vol1. Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY.
5. Nurk S, Bankevich A, Antipov D, Gurevich AA, Korobeynikov A, Lapidus A, Prjibelski AD, Pyshkin A, Sirotkin A, Sirotkin Y, Stepanauskas R, Clin-genpeel SR, Woyke T, McLean JS, Lasken R, Tesler G, Alekseyev MA, Pevzner PA. 2013. Assembling single-cell genomes and mini-metagenomes from chimeric MDA products. *J Comput Biol* 20:714–737. <https://doi.org/10.1089/cmb.2013.0084>.
6. Mikheenko A, Prjibelski A, Saveliev V, Antipov D, Gurevich A. 2018. Versatile genome assembly evaluation with QAST-LG. *Bioinformatics* 34:i142–i150. <https://doi.org/10.1093/bioinformatics/bty266>.
7. Seemann T. 2014. Prokka: rapid prokaryotic genome annotation. *Bioinformatics* 30:2068–2069. <https://doi.org/10.1093/bioinformatics/btu153>.
8. Rehm BH. 2003. Polyester synthases: natural catalysts for plastics. *Biochem J* 376:15–33. <https://doi.org/10.1042/BJ20031254>.
9. Schubert P, Steinbüchel A, Schlegel HG. 1988. Cloning of the *Alcaligenes eutrophus* genes for synthesis of poly-beta-hydroxybutyric acid (PHB) and synthesis of PHB in *Escherichia coli*. *J Bacteriol* 170:5837–5847.
10. Peoples OP, Sinskey AJ. 1989. Poly-beta-hydroxybutyrate (PHB) biosynthesis in *Alcaligenes eutrophus* H16. Identification and characterization of the PHB polymerase gene (*phbC*). *J Biol Chem* 264:15298–15303.