



## Complete Genome Sequence of *Bacillus megaterium* Strain TG1-E1, a Plant Drought Tolerance-Enhancing Bacterium

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**ABSTRACT** Based on a combination of next-generation sequencing and singlemolecule sequencing, we obtained the whole-genome sequence of *Bacillus megaterium* strain TG1-E1, which is a highly salt-tolerant rhizobacterium that enhances plant tolerance to drought stress. The complete genome is estimated to be approximately 5.48 Mb containing a total of 5,858 predicted protein-coding DNA sequences.

Our group characterized *Bacillus megaterium* strain TG1-E1 as a highly salt-tolerant Gram-positive bacterium that is capable of enhancing plant tolerance to drought stress. It was originally isolated from a rhizospheric soil sample of *Spartina anglica* at Zhangpu Yanchang in Fujian Province, China. This rhizobacterium collection is rich in specimens of the *Firmicutes* and *Proteobacteria* phyla, with about 70% belonging to the *Bacillaceae* family. High salinity in the sampling area possibly contributes to the enrichment of *Bacillus* strains in the rhizosphere (1–4). In addition, more than half of the strains isolated in this sampling area can produce phytometabolites, such as auxins and aminocyclopropane-1-carboxylate deaminase (ACCd), displaying the characteristics commonly described in plant tolerance-enhancing strains (5–10). *B. megaterium* TG1-E1 has been deposited in the China General Microbiological Culture Collection Center (CGMCC) with reference number 14422.

DNA samples (at least 100 nM in 10  $\mu$ l) were obtained from bacteria grown in LB medium until an optical density of 1 at 600 nm (OD<sub>600</sub>) was obtained. The sequencing of the B. megaterium TG1-E1 genome was completed by combining next-generation sequencing (NGS) and single-molecule sequencing. NGS was performed with 20  $\mu$ g of DNA with an Illumina HiSeg platform (Core Facility of Genomics, Shanghai Center for Plant Stress Biology, China), and single-molecule sequencing was performed with 20  $\mu$ g of DNA with a PacBio platform (Tianjin Biochip Corporation, China) (11-14). The shotgun sequencing strategy was applied to NGS, and 12,471,203 paired reads (150 bp) were obtained with a sequencing depth of approximately 260-fold. Meanwhile, singlemolecule sequencing produced 98,959 reads with a mean read length of 10,551 bp and an  $N_{50}$  length of 14,471 bp. The total number of sequenced bases was 961,774,920. For de novo assembly, Canu v1.5 was used with default parameters, and the genome correction step was performed using Illumina data with support of Pilon v1.18 (15, 16). The size of the circularized genome was calculated to be about 5.48 Mb. Genes including protein-coding DNA sequences (CDSs) were predicted by a pipeline implemented by Prokka v1.12 (17). On a whole-genome scale, The GC content of this genome

Received 10 June 2018 Accepted 28 August 2018 Published 27 September 2018

Citation Vilchez JI, Tang Q, Kaushal R, Wang W, Lv S, He D, Chu Z, Zhang H, Liu R, Zhang H. 2018. Complete genome sequence of *Bacillus megaterium* strain TG1-E1, a plant drought tolerance-enhancing bacterium. Microbiol Resour Announc 7:e00842-18. https://doi.org/ 10.1128/MRA.00842-18.

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is 38.26%, and 5,858 protein-coding genes, 4 rRNA operons, and 164 tRNA genes were called during annotation.

The whole-genome sequence of *B. megaterium* TG1-E1 reveals information such as the biosynthesis pathways of flagella, spores, and polysaccharides. Concerning characteristics potentially contributing to TG1-E1-induced plant stress tolerance, pathways found within this genome that have potential relevance in aiding plant drought stress include trehalose and antioxidant biosynthesis. In addition, genome annotation also revealed possible mechanisms for plant growth-promoting effects, including bacterial production of acid phosphatases, siderophores, and exopolysaccharides. Further research with this genomic information will help us discover mechanisms through which *B. megaterium* TG1-E1 induces plant drought stress tolerance and will contribute to the subsequent development of biotechnological applications.

**Data availability.** The complete genome sequence of *B. megaterium* TG1-E1 has been deposited in the TBL/EMBL/GenBank databases under the BioProject number PRJNA430758 and the accession number PRKV00000000 (sequences PRKV01000001 to PRKV01000036).

## ACKNOWLEDGMENTS

Huiming Zhang is funded by the Chinese Academy of Sciences (CAS) project number XDPB0404.

We acknowledge the support of the Core Facility of Genomics at PSC.

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