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## Draft Genome Sequence of Plant Growth-Promoting and Drought-Tolerant *Bacillus altitudinis* FD48, Isolated from Rice Phylloplane

Sowmya Kumaravel,<sup>a</sup> Sugitha Thankappan,<sup>a</sup> Sridar Raghupathi,<sup>a</sup> DSivakumar Uthandi<sup>a</sup>

<sup>a</sup>Department of Agricultural Microbiology, Tamil Nadu Agricultural University, Coimbatore, Tamil Nadu, India

**ABSTRACT** The genome sequence of a temperature-tolerant strain, *Bacillus altitudinis* FD48, is described here. The reads were assembled into contigs with a total size of 3.7 Mb. The genome information will aid in understanding its role in alleviating stress in crop plants as a potential bioinoculant for agricultural applications.

**B**isolated from rice phylloplane, possessed a 3.7-Mb genome which harbors gene clusters for heat shock proteins (chaperonin), cysteine synthesis, trehalose synthesis, stress response proteins, and siderophore biosynthesis. *B. altitudinis* FD48, a Grampositive bacterium isolated from the leaf surface of rice genotype ADT 43 by a leaf imprinting technique (1), not only survives under induced drought conditions (-0.69 MPa) but also confers various PGP benefits, like solubilization of phosphorus, production of plant growth hormones (indole-3-acetic acid [IAA], cytokinin), exopoly-saccharides, 1-aminocyclopropane-1-carboxylate (ACC) deaminase activity, and saline tolerance up to 7.5%.

Genomic DNA (gDNA) was extracted from an overnight monoculture of *B. altitudinis* FD48 using the standard protocol of the cetyltrimethylammonium bromide (CTAB) method (2), with minor modifications. High-quality gDNA was subjected to TruSeq library preparation and was sequenced using an Illumina HiSeq 4000 sequencer. Quality control (QC) of the raw data was done to qualify reads with a Phred score of >30 for downstream analysis. Reads that passed QC were subjected to *de novo* assembly using Velvet (version 1.2.10), followed by genome finishing using the CONTIGuator tool (3, 4), resulting in a genome size of 3.7 Mb with 15 scaffold sequences and 41.19% G+C content. The  $N_{50}$  contig size was 976,391 bp.

A total of 4,029 predicted genes, including 3,964 protein-coding genes (CDSs) and 65 non-protein-coding genes, were observed, along with 2,882 characterized proteins and 1,080 hypothetical/putative proteins. The 16S rRNA gene sequence of the isolate FD48 showed 99% similarity to that of *B. altitudinis*.

The genome of *B. altitudinis* FD48 comprises several genes related to plant growth promotion mechanisms, such as those for the biogenesis of organic acids involved in inorganic phosphorus solubilization (glucose dehydrogenase, citrate synthase, and lactate dehydrogenase) (5). Additionally, genes responsible for flagellar motility, chemotaxis, and biofilm synthesis, which allow *B. altitudinis* FD48 to move toward plant exudates, thereby facilitating adhesion to plant surfaces, in addition to genes related to growth-stimulating volatile compounds and sporulation, were encountered. In addition, the annotated genome has several genes for components of iron and siderophore uptake systems, nitrogen metabolism, and various antibiotic resistance gene clusters. Interestingly, various stress regulatory and temperature tolerance genes were noted, along with those for osmolyte production (proline and glycine betaine), cold shock proteins, heat shock proteins, and resistance to heavy metals.

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Address correspondence to Sivakumar Uthandi, usiva@tnau.ac.in.

S.K. and S.T. contributed equally to this work. This is a contribution from Tamil Nadu Agricultural University. **Accession number(s).** The complete genome sequence of strain FD48 has been deposited in GenBank under the accession number CP025643.

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## REFERENCES

- 1. Aneja K. 2003. Experiments in microbiology, plant pathology and biotechnology New Age International Ltd. Publishers, New Delhi, India.
- 2. Clark MS. 1997. Plant molecular biology—a laboratory manual, 1st ed. Springer, Berlin, Germany.
- Aziz RK, Bartels D, Best AA, DeJongh M, Disz T, Edwards RA, Formsma K, Gerdes S, Glass EM, Kubal M, Meyer F, Olsen GJ, Olson R, Osterman AL, Overbeek RA, McNeil LK, Paarmann D, Paczian T, Parrello B, Pusch GD, Reich C, Stevens R, Vassieva O, Vonstein V, Wilke A, Zagnitko O. 2008. The RAST server: Rapid Annotations using Subsystems Technology. BMC Genomics 9:75. https://doi.org/10.1186/1471-2164-9-75.
- Overbeek R, Olson R, Pusch GD, Olsen GJ, Davis JJ, Disz T, Edwards RA, Gerdes S, Parrello B, Shukla M. 2013. The SEED and the Rapid Annotation of microbial genomes using Subsystems Technology (RAST). Nucleic Acids Res 42:D206–D214. https://doi.org/10.1093/nar/gkt1226.
- Chen Y, Rekha P, Arun A, Shen F, Lai W-A, Young C. 2006. Phosphate solubilizing bacteria from subtropical soil and their tricalcium phosphate solubilizing abilities. Appl Soil Ecol 34:33–41. https://doi.org/10.1016/j .apsoil.2005.12.002.