



Draft Genome Sequences of Plant-Associated *Bacillus* Strains Isolated from the Qinghai-Tibetan Plateau

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ABSTRACT Here, we report the draft genome sequences of 45 plant-associated *Bacillus* strains isolated from the Qinghai-Tibetan plateau. According to their genome sequences, 28 isolates were assigned to 10 *Bacillus* species. Seventeen strains could not be assigned and are subjects of further research.

Bacillus strains isolated from samples taken from different sites of the Qinghai-Tibetan plateau, known as the Third Pole of the world (1, 2), were found to grow significantly between 4°C and 12°C (H. Wu, R. Borriss, P. Xue, F. Liu, and X. Gao, unpublished data), to enhance plant growth, and to suppress plant pathogens (3–5). As a first step to characterize these strains more deeply, 45 of the isolates were genome sequenced and their taxonomy was determined.

Colonies of a fresh culture grown on LB agar plates were selected. Genomic DNA was extracted using the QIAamp DNA minikit (Qiagen, Hilden, Germany), and the sequencing was done in 300-nucleotide (nt) paired-end mode on an Illumina MiSeq version 3 sequencing platform at LGC Genomics (Berlin, Germany). Reads were trimmed and assembled *de novo* using the A5 pipeline (6). Genome coverage of the obtained scaffolds was 45× on average. Scaffolds were submitted to GenBank for gene annotation, which was implemented using the NCBI Prokaryotic Genome Annotation Pipeline (PGAP) (7). The genome-to-genome-distance calculator (GGDC) version 2.1 provided by DSMZ (<http://ggdc.dsmz.de>) was used for genome-based species delineation. Formula 2, which is especially appropriate to analyze draft genomes, was used (Meier-Kolthoff et al., 2013 [8]). In addition, JSpecies WS (<http://jspecies.ribohost.com/jspeciesws/>) was used to determine the average nucleotide identity based on BLAST+ (ANIb) by pairwise genome comparisons (9). The recommended species cutoff was defined as 96%.

According to their draft genome sequences, we have assigned 28 of the isolates as representatives of *Bacillus wiedmannii* (GenBank accession numbers PVRQ00000000 to PVRU00000000, and PYWP00000000), *B. atrophaeus* (PVQM00000000 to PVQO00000000, PVWA00000000, and PVWB00000000), *B. pumilus* (PVQT00000000 to PVQX00000000), *B. halotolerans* (PVWC00000000, PVQP00000000, and PVQQ00000000), *B. subtilis* (PVRJ00000000 and PVRK00000000), *B. thuringiensis* (PVRL00000000 and PVRM00000000), *B. velezensis* (PVRO00000000 and PVRP00000000), *B. paralicheniformis* (PVQR00000000), *B. safensis* (PVQS00000000), and *B. toyonensis* (PVRN00000000). Seventeen strains could not be assigned down to the species level due to their estimated GGDC (<70%) and ANIb (<96%) values. Most of the strains (15 isolates) are related to *B. pumilus* (PVQY00000000, PVQZ00000000, PVRA00000000 to PVRI00000000, PVQK00000000, PVQL00000000, PVWX00000000, and PVWY00000000). The genome sequence of strain RJGP41 (PVQJ00000000) is distantly related to *B. simplex*, while strain

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LLTC93 (PVME00000000) resembles the type strain of *B. xiamenensis*, HYC-10. Further research is in progress in order to clarify the taxonomic position of these cold-adapted strains.

Accession number(s). These whole-genome shotgun projects have been deposited at DDBJ/ENA/GenBank under the accession numbers [PVME00000000](#), [PVQJ00000000](#), [PVQK00000000](#), [PVQL00000000](#), [PVQM00000000](#), [PVQN00000000](#), [PVQO00000000](#), [PVQP00000000](#), [PVQQ00000000](#), [PVQR00000000](#), [PVQS00000000](#), [PVQT00000000](#), [PVQU00000000](#), [PVQV00000000](#), [PVQW00000000](#), [PVQX00000000](#), [PVQY00000000](#), [PVQZ00000000](#), [PVRA00000000](#), [PVRB00000000](#), [PVRC00000000](#), [PVRD00000000](#), [PVRE00000000](#), [PVRF00000000](#), [PVRG00000000](#), [PVRH00000000](#), [PVRI00000000](#), [PVRJ00000000](#), [PVRK00000000](#), [PVRL00000000](#), [PVRM00000000](#), [PVRN00000000](#), [PVRO00000000](#), [PVRP00000000](#), [PVRQ00000000](#), [PVRR00000000](#), [PVRS00000000](#), [PVRT00000000](#), [PVRU00000000](#), [PVWA00000000](#), [PVWB00000000](#), [PVWC00000000](#), [PVWX00000000](#), [PVWY00000000](#), and [PYWP00000000](#). The versions described in this paper are the first versions.

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