PROKARYOTES



Draft Genome Sequence of *Pseudomonas koreensis* CI12, a *Bacillus cereus* "Hitchhiker" from the Soybean Rhizosphere

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ABSTRACT *Pseudomonas koreensis* Cl12 was coisolated with *Bacillus cereus* from a root of a soybean plant grown in a field in Arlington, WI. Here, we report the draft genome sequence of *P. koreensis* Cl12 obtained by Illumina sequencing.

Pseudomonas koreensis was first proposed as a novel Pseudomonas species to classify several isolates from Korean agricultural soils (1). Since then, strains worldwide have been identified with diverse capacities ranging from the production of antibacterial compounds (2) to the suppression of plant diseases caused by oomycete pathogens (3). Recently, whole-genome sequenced-based analyses designated *P. koreensis* as a defined phylogenomic group within the physiologically and genetically heterogeneous *Pseudomonas fluorescens* complex (4, 5). The *P. koreensis* group contains members that have been isolated from *Populus* root systems (6) and is, more generally, enriched with isolates recovered from diverse plants (7). Additionally, a comparative genomic analysis within the *P. fluorescens* complex showed an overrepresentation of traits related to plant-bacterium interactions in genomes from *P. koreensis* isolates (4).

P. koreensis Cl12 was isolated as one of several microbial "hitchhikers" from *Bacillus cereus* cultures purified from field-grown soybean roots (8). These hitchhikers are bacteria that are not visible in colony-purified *B. cereus* cultures until 2 to 4 weeks of incubation at 4°C; although 3 to 5% of *B. cereus* isolates from soybean roots carry hitchhikers, the mechanism underlying the association is unknown. The classification of Cl12 within the *P. fluorescens* complex was determined by independent phylogenetic reconstruction of the *gyrB*, *rpoD*, and *rpoB* genes, as has been described previously (4). *Pseudomonas koreensis* Cl12 was selected as a model for studying bacterial interactions in the rhizosphere. *In vitro* growth of *P. koreensis* Cl12 in root exudate is not significantly affected by the presence of *B. cereus*, but *P. koreensis* Cl12 can impair the growth of other hitchhikers, which is in contrast to the hitchhikers' growth enhancement by *B. cereus* (8).

The *P. koreensis* Cl12 genome was sequenced on the Illumina MiSeq platform. A total of 8,588,279 paired-end reads of 300 bp from a library with an average insert size of 1 kb were generated. Low-quality sequences were trimmed using Trimmomatic (9), and the resulting sequences were then assembled using Velvet (10) and VelvetOptimiser. Contigs were ordered by Mauve (11) using the *P. fluorescens* Pf0-1 genome (12) as a reference, assembled manually by joining with a linker sequence of unknown nucleo-tide character "N," and then gaps were filled with GapFiller (13). The resulting assembly was 6,622,028 bp, consisting of 16 contigs, with an N_{50} contig size of 608,098 bp.

We predict that sequencing new strains of *P. koreensis* will help delineate traits that may mediate its interactions with plant hosts and their associated microbiota. Furthermore, additional genomes belonging to members of the *P. fluorescens* complex may

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Accession number(s). This whole-genome shotgun project has been deposited at DDBJ/ENA/GenBank under the accession no. MPLD00000000. The version described in this paper is the first version.

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REFERENCES

- Kwon SW, Kim JS, Park IC, Yoon SH, Park DH, Lim CK, Go SJ. 2003. Pseudomonas koreensis sp. nov., Pseudomonas umsongensis sp. nov. and Pseudomonas jinjuensis sp. nov., novel species from farm soils in Korea. Int J Syst Evol Microbiol 53:21–27. https://doi.org/10.1099/ijs.0.02326-0.
- Toribio J, Escalante AE, Caballero-Mellado J, González-González A, Zavala S, Souza V, Soberón-Chávez G. 2011. Characterization of a novel biosurfactant producing *Pseudomonas koreensis* lineage that is endemic to Cuatro Ciénegas Basin. Syst Appl Microbiol 34:531–535. https://doi.org/ 10.1016/j.syapm.2011.01.007.
- 3. Hultberg M, Alsberg T, Khalil S, Alsanius B. 2010. Suppression of disease in tomato infected by *Pythium ultimum* with a biosurfactant produced by *Pseudomonas koreensis*. BioControl 55:435–444. https://doi.org/10 .1007/s10526-009-9261-6.
- Garrido-Sanz D, Meier-Kolthoff JP, Göker M, Martín M, Rivilla R, Redondo-Nieto M. 2016. Genomic and genetic diversity within the *Pseudomonas fluorescens* complex. PLoS One 11:e0150183. https://doi.org/10.1371/ journal.pone.0150183.
- Gomila M, Peña A, Mulet M, Lalucat J, García-Valdés E. 2015. Phylogenomics and systematics in *Pseudomonas*. Front Microbiol 6:214. https:// doi.org/10.3389/fmicb.2015.00214.
- Jun SR, Wassenaar TM, Nookaew I, Hauser L, Wanchai V, Land M, Timm CM, Lu TY, Schadt CW, Doktycz MJ, Pelletier DA, Ussery DW. 2015. Diversity of *Pseudomonas* genomes, including *Populus*-associated isolates, as revealed by comparative genome analysis. Appl Environ Microbiol 82:375–383. https://doi.org/10.1128/AEM.02612-15.
- 7. Winsor GL, Griffiths EJ, Lo R, Dhillon BK, Shay JA, Brinkman FSL. 2016.

Enhanced annotations and features for comparing thousands of *Pseudomonas* genomes in the *Pseudomonas* genome database. Nucleic Acids Res 44:D646–D653. https://doi.org/10.1093/nar/qkv1227.

- Peterson SB, Dunn AK, Klimowicz AK, Handelsman J. 2006. Peptidoglycan from *Bacillus cereus* mediates commensalism with rhizosphere bacteria from the *Cytophaga-Flavobacterium* group. Appl Environ Microbiol 72:5421–5427. https://doi.org/10.1128/AEM.02928-05.
- 9. Bolger AM, Lohse M, Usadel B. 2014. Trimmomatic: a flexible trimmer for Illumina sequence data. Bioinformatics 30:2114–2120. https://doi.org/10 .1093/bioinformatics/btu170.
- Zerbino DR, Birney E. 2008. Velvet: algorithms for *de novo* short read assembly using de Bruijn graphs. Genome Res 18:821–829. https://doi .org/10.1101/gr.074492.107.
- Darling ACE, Mau B, Blattner FR, Perna NT. 2004. Mauve: multiple alignment of conserved genomic sequence with rearrangements. Genome Res 14:1394–1403. https://doi.org/10.1101/gr.2289704.
- Silby MW, Cerdeño-Tárraga AM, Vernikos GS, Giddens SR, Jackson RW, Preston GM, Zhang XX, Moon CD, Gehrig SM, Godfrey SAC, Knight CG, Malone JG, Robinson Z, Spiers AJ, Harris S, Challis GL, Yaxley AM, Harris D, Seeger K, Murphy L, Rutter S, Squares R, Quail MA, Saunders E, Mavromatis K, Brettin TS, Bentley SD, 2009. Genomic and genetic analyses of diversity and plant interactions of *Pseudomonas fluorescens*. Genome Biol 10:R51. https://doi.org/10.1186/gb-2009-10-5-r51.
- Boetzer M, Pirovano W. 2012. Toward almost closed genomes with GapFiller. Genome Biol 13:R56. https://doi.org/10.1186/gb-2012-13-6 -r56.