

# Draft Genome Sequence of *Pannonibacter phragmitetus* Strain CGMCC9175, a Halotolerant Polycyclic Aromatic Hydrocarbon-Degrading Bacterium

Xinxin Wang,<sup>a,b</sup> Decai Jin,<sup>c</sup> Lisha Zhou,<sup>d</sup> Zhuo Zhang<sup>e</sup>

School of Environmental Science and Engineering, Tianjin University, Tianjin, China<sup>a</sup>; Offshore Environmental Service Co. Ltd., Tianjin, China<sup>b</sup>; Research Center for Eco-Environmental Sciences, Chinese Academy of Sciences, Beijing, China<sup>c</sup>; Shenzhen Key Laboratory of Environmental Microbial Genomics and Application, BGI, Shenzhen, China<sup>d</sup>; Hunan Plant Protection Institute, Changsha, China<sup>e</sup>

X.W. and D.J. contributed equally to this work.

***Pannonibacter phragmitetus* CGMCC9175 is a halotolerant polycyclic aromatic hydrocarbon (PAH)-degrading bacterium isolated from PAH-contaminated intertidal zone sediment. Here, we report the 5.7-Mb draft genome sequence of this strain, which will provide insights into the diversity of *Pannonibacter* and the mechanism of PAH degradation in sediments.**

Received 6 December 2015 Accepted 8 December 2015 Published 28 January 2016

**Citation** Wang X, Jin D, Zhou L, Zhang Z. 2016. Draft genome sequence of *Pannonibacter phragmitetus* strain CGMCC9175, a halotolerant polycyclic aromatic hydrocarbon-degrading bacterium. *Genome Announc* 4(1):e01675-15. doi:10.1128/genomeA.01675-15.

**Copyright** © 2016 Wang et al. This is an open-access article distributed under the terms of the [Creative Commons Attribution 4.0 International license](https://creativecommons.org/licenses/by/4.0/).

Address correspondence to Xinxin Wang, wangxx200899@163.com.

Polycyclic aromatic hydrocarbon (PAH)-degrading bacteria play a key role in the natural remediation of PAH-polluted marine sediments (1), which attract attention from researchers in the field of environmental microbiology. *Pannonibacter phragmitetus* CGMCC9175 was isolated from PAH-contaminated intertidal zone sediment of Bohai Sea in China (2), which can degrade naphthalene, phenanthrene, and anthracene with 5% NaCl. Until recently, only the genome of *P. phragmitetus* DSM 14782 has been available. Here, the draft genome sequence of *P. phragmitetus* CGMCC9175 is presented.

Genomic DNA was extracted and sequenced using the Illumina MiSeq platform. The shotgun sequencing produced 5,361,830 paired-end reads, with an average insert size of 300 bp, yielding approximately 280-fold coverage; the reads were filtered by the NGS QC toolkit version 2.3 (3). The filtered reads were assembled, scaffolded, gap filled, and validated using SOAPdenovo version 2.04 (4), SSPACE version 2.0 (5), GapFiller version 1.10 (6), and bwa version 0.7.4 (7). The final assembly consisted of 40 contigs, with an  $N_{50}$  length of 353,668 bp and a largest length of 653,707 bp. The genome sequence was annotated using the NCBI Prokaryotic Genome Annotation Pipeline (PGAP) ([http://www.ncbi.nlm.nih.gov/genome/annotation\\_prok/](http://www.ncbi.nlm.nih.gov/genome/annotation_prok/)).

The genome consists of 5.7 Mb, with a G+C content of 63.6%. A total of 4,853 coding sequences (CDSs), 167 pseudogenes, 50 tRNAs, 1 noncoding RNA (ncRNA), and 1 rRNA operon were identified. Of the CDSs, 86.1% can be assigned to COGs, with amino acid transport and metabolism as the most abundant class, and 44.1% can be annotated into 2,286 KEGG orthologous groups using KAAS (8), involving 233 metabolic pathways. The ISAs1 family dominated the insertion sequence (IS) elements, as revealed by ISFinder (9). Plasmid-carried genes essential for stabilization and partition were detected, which suggests the presence of

plasmid. A total of 199 tandem repeats, 370 potentially secreted proteins, 3 prophage sequences, and 6 clustered regularly interspaced short palindromic repeat (CRISPR) elements were identified by Tandem Repeats Finder version 4.08 (10), SignalP version 4.1 (11), PFAST (12), and CRISPRFinder (13), respectively. Average nucleotide identity (ANI) analysis (14) revealed that *P. phragmitetus* CGMCC9175 is phylogenetically related to *P. phragmitetus* DSM 14782 (93.3%).

One alkane 1-monooxygenase, 1 catechol 1,2-dioxygenase, 1 homogentisate 1,2-dioxygenase, 2 protocatechuate 3,4-dioxygenase, and 2 biphenyl 2,3-dioxygenase genes were identified, which were responsible for the degradation of alkanes and PAHs. One 2-haloalkanoic acid dehalogenase and 6 haloacid dehalogenase genes were identified, which were responsible for the degradation of halogenated organic compounds. Moreover, 3 genes were identified as being involved in compatible solute synthesis and uptake, including 1 glycine/betaine ABC transporter and 2 betaine-aldehyde dehydrogenase. These genes may be essential to the survival in a hydrocarbon-contaminated saline environment. Information about the genome sequence of *P. phragmitetus* CGMCC9175 offered an opportunity to understand the genetic diversity of *Pannonibacter* and the mechanism of hydrocarbon degradation in intertidal zone sediments.

**Nucleotide sequence accession number.** The draft genome sequence of *P. phragmitetus* CGMCC9175 has been deposited in GenBank under the accession number [LGSQ000000000](https://www.ncbi.nlm.nih.gov/nuccore/LGSQ000000000). The version described in this paper is the first version.

## ACKNOWLEDGMENT

This work was supported by 2013 postdoctoral innovation fund project of Tianjin.

## FUNDING INFORMATION

Postdoctoral Foundation of Tianjin Province, China provided funding to Xinxin Wang under grant number 2013.

The funders had no role in study design, data collection and interpretation, or the decision to submit the work for publication.

## REFERENCES

- Mason OU, Scott NM, Gonzalez A, Robbins-Pianka A, Bælum J, Kimbrel J, Bouskill NJ, Prestat E, Borglin S, Joyner DC, Fortney JL, Jurelevicius D, Stringfellow WT, Alvarez-Cohen L, Hazen TC, Knight R, Gilbert JA, Jansson JK. 2014. Metagenomics reveals sediment microbial community response to deepwater horizon oil spill. *ISME J* 8:1464–1475. <http://dx.doi.org/10.1038/ismej.2013.254>.
- Wang X, Jin D, Zhou L, Wu L, An W, Zhao L. 2014. Draft genome sequence of *Advenella kashmirensis* strain W13003, a polycyclic aromatic hydrocarbon-degrading bacterium. *Genome Announc* 2(1):e00003-14. <http://dx.doi.org/10.1128/genomeA.00003-14>.
- Patel RK, Jain M. 2012. NGS QC toolkit: a toolkit for quality control of next generation sequencing data. *PLoS One* 7:e30619. <http://dx.doi.org/10.1371/journal.pone.0030619>.
- Luo R, Liu B, Xie Y, Li Z, Huang W, Yuan J, He G, Chen Y, Pan Q, Liu Y, Tang J, Wu G, Zhang H, Shi Y, Liu Y, Yu C, Wang B, Lu Y, Han C, Cheung DW. 2012. SOAPdenovo2: an empirically improved memory-efficient short-read *de novo* assembler. *GigaScience* 1:18. <http://dx.doi.org/10.1186/2047-217X-1-18>.
- Boetzer M, Henkel CV, Jansen HJ, Butler D, Pirovano W. 2011. Scaffolding pre-assembled contigs using SSPACE. *Bioinformatics* 27:578–579. <http://dx.doi.org/10.1093/bioinformatics/btq683>.
- Boetzer M, Pirovano W. 2012. Toward almost closed genomes with Gap-Filler. *Genome Biol* 13:R56. <http://dx.doi.org/10.1186/gb-2012-13-6-r56>.
- Li H, Durbin R. 2009. Fast and accurate short read alignment with Burrows-Wheeler transform. *Bioinformatics* 25:1754–1760. <http://dx.doi.org/10.1093/bioinformatics/btp324>.
- Moriya Y, Itoh M, Okuda S, Yoshizawa AC, Kanehisa M. 2007. KAAS: an automatic genome annotation and pathway reconstruction server. *Nucleic Acids Res* 35:W182–W185. <http://dx.doi.org/10.1093/nar/gkm321>.
- Siguiet P, Perochon J, Lestrade L, Mahillon J, Chandler M. 2006. ISfinder: the reference centre for bacterial insertion sequences. *Nucleic Acids Res* 34:D32–D36. <http://dx.doi.org/10.1093/nar/gkj014>.
- Benson G. 1999. Tandem repeats finder: a program to analyze DNA sequences. *Nucleic Acids Res* 27:573–580. <http://dx.doi.org/10.1093/nar/27.2.573>.
- Petersen TN, Brunak S, von Heijne G, Nielsen H. 2011. SignalP 4.0: discriminating signal peptides from transmembrane regions. *Nat Methods* 8:785–786. <http://dx.doi.org/10.1038/nmeth.1701>.
- Zhou Y, Liang Y, Lynch KH, Dennis JJ, Wishart DS. 2011. PHAST: a fast phage search tool. *Nucleic Acids Res* 39:W347–W352. <http://dx.doi.org/10.1093/nar/gkr485>.
- Grissa I, Vergnaud G, Pourcel C. 2007. CRISPRFinder: a Web tool to identify clustered regularly interspaced short palindromic repeats. *Nucleic Acids Res* 35:W52–W57. <http://dx.doi.org/10.1093/nar/gkm360>.
- Richter M, Rosselló-Móra R. 2009. Shifting the genomic gold standard for the prokaryotic species definition. *Proc Natl Acad Sci USA* 106:19126–19131. <http://dx.doi.org/10.1073/pnas.0906412106>.