



Genome Announcements

Draft genome sequence of a caprolactam degrader bacterium: *Pseudomonas taiwanensis* strain SJ9



Sung-Jun Hong, Gun-Seok Park, Abdur Rahim Khan, Byung Kwon Jung, Jae-Ho Shin*

School of Applied Biosciences, Kyungpook National University, Daegu, Republic of Korea

ARTICLE INFO

Article history:

Received 22 July 2015

Accepted 15 September 2015

Available online 2 March 2016

Associate Editor: John Anthony

McCulloch

Keywords:

Pseudomonas taiwanensis

Bioremediation

Biodegradation

Caprolactam

Nylon

ABSTRACT

Pseudomonas taiwanensis strain SJ9 is a caprolactam degrader, isolated from industrial wastewater in South Korea and considered to have the potential for caprolactam bioremediation. The genome of this strain is approximately 6.2 Mb (G + C content, 61.75%) with 6,010 protein-coding sequences (CDS), of which 46% are assigned to recognized functional genes. This draft genome of strain SJ9 will provide insights into the genetic basis of its caprolactam-degradation ability.

© 2016 Sociedade Brasileira de Microbiologia. Published by Elsevier Editora Ltda. This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

Introduction

Members of the genus *Pseudomonas* that have been isolated and characterized so far have mostly been found as innocuous environmental microorganisms. They have great potential for biotechnological applications owing to their metabolic versatility and adaptability.^{1,2} *Pseudomonas* spp. can thrive in diverse habitats and are known for their ability to colonize soil and participate in soil biochemical processes.^{3,4} The potential of *Pseudomonas* spp. for the degradation and bioremediation of a wide variety of chemicals, including natural and synthetic compounds such as caprolactam,⁵ naphthalene,⁶

and toluene,⁷ has attracted a great research interest. *P. taiwanensis* strain SJ9 was isolated from a wastewater sample collected from a sewage treatment plant in Daegu, South Korea. This work reports the draft genome of *P. taiwanensis* strain SJ9.

The genome of the strain SJ9 was sequenced using an Ion Torrent Personal Genome Machine (PGM) sequencer system.⁸ The sequence reads were assembled using Mimicking Intelligent Read Assembly (MIRA) 3.4.0 and CLC Genomics Workbench (version 6.0), with manual processing using SeqMan software to reduce the contig number. The best assembly results comprised 736 contigs (>400 bp). The draft genome consists of 6,253,055 bp covering almost

* Corresponding author.

E-mail: jhshin@knu.ac.kr (J.-H. Shin).

<http://dx.doi.org/10.1016/j.bjm.2015.09.002>

whole of the predicted average genome, with a G+C content of 61.75%. The assembled contigs were submitted to the RAST annotation server (<http://rast.nmpdr.org>) for subsystem classification and functional annotation.⁹ This analysis predicted 6,010 protein-coding sequences (CDS), of which 46% were assigned to recognized functional genes. Furthermore, 71 tRNA and 12 rRNA genes were also predicted.

The genome also harbored a complete gene cluster coding for caprolactam degrading enzymes such as 2,3-dehydroadipyl-CoA hydratase, acyl-CoA dehydrogenase, aldehyde dehydrogenase, and enoyl-CoA hydratase.^{10,11} This draft genome sequence of *P. taiwanensis* strain SJ9 will help improve the general understanding of the genetic basis of caprolactam degradation by *Pseudomonas* spp.

Nucleotide sequence accession numbers

The draft sequence of *P. taiwanensis* strain SJ9 obtained in this Whole Genome Shotgun project has been deposited at GenBank under the accession no. AXUP00000000. The version described in this paper is the first version, with accession no. AXUP01000000.

Conflicts of interest

The authors declare no conflicts of interest.

Acknowledgments

This study was sponsored by Agricultural Biotechnology Development Program, Ministry of Agriculture, Food and Rural Affairs.

REFERENCES

1. Ridgway HF, Safarik J, Phipps D, Carl P, Clark D. Identification and catabolic activity of well-derived gasoline-degrading bacteria from a contaminated aquifer. *Appl Environ Microbiol.* 1990;56:3565–3575.
2. Loh KC, Cao B. Paradigm in biodegradation using *Pseudomonas putida* – a review of proteomics studies. *Enzyme Microb Technol.* 2008;43:1–12.
3. Raaijmakers JM, Weller DM, Thomashow LS. Frequency of antibiotic-producing *Pseudomonas* spp. in natural environments. *Appl Environ Microbiol.* 1997;63:881–887.
4. Dowling DN, O'Gara F. Metabolites of *Pseudomonas* involved in the biocontrol of plant disease. *Trends Biotechnol.* 1994;12:133–141.
5. Kulkarni RS, Kanekar PP. Bioremediation of epsilon-caprolactam from nylon-6 waste water by use of *Pseudomonas aeruginosa* MCM B-407. *Curr Microbiol.* 1998;37:191–194.
6. Rossello-Mora RA, Lalucat J, Garcia-Valdes E. Comparative biochemical and genetic analysis of naphthalene degradation among *Pseudomonas stutzeri* strains. *Appl Environ Microbiol.* 1994;60:966–972.
7. Zylstra GJ, McCombie WR, Gibson DT, Finette BA. Toluene degradation by *Pseudomonas putida* F1: genetic organization of the tod operon. *Appl Environ Microbiol.* 1988;54:1498–1503.
8. Rothberg JM, Hinz W, Rearick TM, et al. An integrated semiconductor device enabling non-optical genome sequencing. *Nature.* 2011;475:348–352.
9. Aziz RK, Bartels D, Best AA, et al. The RAST Server: rapid annotations using subsystems technology. *BMC Genomics.* 2008;9:75.
10. Buell CR, Joardar V, Lindeberg M, et al. The complete genome sequence of the Arabidopsis and tomato pathogen *Pseudomonas syringae* pv. tomato DC3000. *Proc Natl Acad Sci USA.* 2003;100:10181–10186.
11. Silby MW, Cerdano-Tarraga AM, Vernikos GS, et al. Genomic and genetic analyses of diversity and plant interactions of *Pseudomonas fluorescens*. *Genome Biol.* 2009;10:R51.