

Draft Genome Sequence of the 2-Chloro-4-Nitrophenol-Degrading Bacterium Arthrobacter sp. Strain SJCon

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We report the 4.39-Mb draft genome sequence of the 2-chloro-4-nitrophenol-degrading bacterium *Arthrobacter* sp. strain SJ-Con, isolated from a pesticide-contaminated site. The draft genome sequence of strain SJCon will be helpful in studying the genetic pathways involved in the degradation of several aromatic compounds.

Received 24 January 2013 Accepted 6 February 2013 Published 7 March 2013

Citation Vikram S, Kumar S, Vaidya B, Pinnaka AK, Raghava GPS. 2013. Draft genome sequence of the 2-chloro-4-nitrophenol-degrading bacterium Arthrobacter sp. strain SJCon. Genome Announc. 1(2):e00058-13. doi:10.1128/genomeA.00058-13.

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rthrobacter spp. are widely distributed in the environment, and members of this genus have been found to be involved in various metabolic activities, like chromium reduction (1), biodegradation of 4-nitrophenol, 4-chlorophenol (2), and 4-fluorocinnamic acid (3), and biotransformation of benzene and toluene (4). Arthrobacter sp. strain SJCon was isolated from a pesticidecontaminated site in Punjab, India, by an enrichment method. The strain SJCon can degrade nitroaromatic compounds, such as p-nitrophenol and 2-chloro-4-nitrophenol, as sole sources of carbon and energy (5). Whole-genome sequencing of Arthrobacter sp. SJCon was done using Ion torrent technology at CSIR-IMTECH and produced a total of 2,234,522 reads of average length 147.6 nucleotides (nt), with a coverage of $73.3 \times$ of a 4.5-Mb genome (expected size). All reads have been assembled by using Newbler version 2.5.3 (Roche) at the default parameters. A total of 142 contigs of >500 bp length were constructed, with an N₅₀ of 63.6 kb; the largest contig assembled measured 143.0 kb. The final genome draft, which consists of 4,389,620 bp, has been used for genome annotation by the RAST (Rapid Annotations using Subsystems Technology) system (6), RNAmmer 1.2 (7), and ARAGORN software (8). A total of 4,242 coding sequences (CDSs), 3 rRNAs, and 54 tRNAs were predicted. We have also reconfirmed the 16S rRNA gene sequence by Sanger's sequencing and found 100% identity with the RNAmmer-predicted 16S rRNA sequence. However, the 16S rRNA sequence of strain SJCon, available at the GenBank database (accession no. GQ927310.2), showed only 98.94% identity with the 16S rRNA gene sequence from this annotated genome, which might be due to a discrepancy in the previous gene sequence submission. RAST annotation shows that Arthrobacter chlorophenolicus A6 (score 542), Arthrobacter sp. FB24 (score 537), and Arthrobacter aurescens TC1 (score 477) are the closest neighbors of the strain SJCon. This annotation indicates that the genes of glycolysis and gluconeogenesis, galactose metabolism, tricarboxylic acid (TCA) cycle, propanoate metabolism, and the pentose phosphate pathway are present in the genome of strain SJCon. In the annotation, we found the genes of

flavin adenine dinucleotide (FAD)-binding monooxygenase, nitrilotriacetate monooxygenase component B (EC 1.14.13.-), 4-hydroxyphenylacetate 3-monooxygenase (EC 1.14.13.3), cyclohexanone monooxygenase (EC 1.14.13.22), catechol 2,3dioxygenase (EC 1.13.11.2), protocatechuate 3,4-dioxygenase alpha and beta chain (EC 1.13.11.3) and catechol 1,2-dioxygenase (EC 1.13.11.1), 2-keto-4-pentenoate hydratase (EC 4.2.1.80), 3-oxoadipate coenzyme A (CoA)-transferase subunit A and B (EC 2.8.3.6), and 3-ketoacyl-CoA thiolase (EC 2.3.1.16) to be involved in the catabolism of aromatic hydrocarbons.

Nucleotide sequence accession numbers. This Whole Genome Shotgun project has been deposited at DDBJ/EMBL/GenBank under the accession no. AOFD00000000. The version described in this paper is the first version, AOFD01000000.

ACKNOWLEDGMENTS

This work was funded by IMTECH-CSIR. We acknowledge the late R.K. Jain for isolating strain SJCon.

S.K. and S.V. are supported by a research fellowship from the Council of Scientific and Industrial Research (CSIR). Whole-genome assembly and annotation data are available at our genomics web portal (http://crdd .osdd.net/raghava/genomesrs/).

REFERENCES

- 1. Megharaj M, Avudainayagam S, Naidu R. 2003. Toxicity of hexavalent chromium and its reduction by bacteria isolated from soil contaminated with tannery waste. Curr. Microbiol. 47:51–54.
- Sahoo NK, Pakshirajan K, Ghosh PK, Ghosh A. 2011. Biodegradation of 4-chlorophenol by *Arthrobacter chlorophenolicus* A6: effect of culture conditions and degradation kinetics. Biodegradation 22:275–286.
- Hasan SA, Wietzes P, Janssen DB. 2012. Biodegradation kinetics of 4-fluorocinnamic acid by a consortium of *Arthrobacter* and *Ralstonia* strains. Biodegradation 23:117–125.
- Unell M, Nordin K, Jernberg C, Stenström J, Jansson JK. 2008. Degradation of mixtures of phenolic compounds by *Arthrobacter chlorophenolicus* A6. Biodegradation 19:495–505.
- Arora PK, Jain RK. 2011. Pathway for degradation of 2-chloro-4nitrophenol in *Arthrobacter* sp. SJCon. Curr. Microbiol. 63:568–573.

- 6. 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.
- Lagesen K, Hallin P, Rødland EA, Staerfeldt HH, Rognes T, Ussery DW. 2007. RNAmmer: consistent and rapid annotation of ribosomal RNA genes. Nucleic Acids Res. 35:3100–3108.
- 8. Laslett D, Canback B. 2004. ARAGORN, a program to detect tRNA genes and tmRNA genes in nucleotide sequences. Nucleic Acids Res. 32:11–16.