





## Draft Genome Sequence of *Arthrobacter* sp. Strain UCD-GKA (Phylum *Actinobacteria*)

Gregory N. Kincheloe, a Jonathan A. Eisen, a,b David A. Coila

University of California, Davis Genome Center, Davis, California, USA<sup>a</sup>; Department of Evolution and Ecology and Department of Medical Microbiology and Immunology, University of California Davis, Davis, California, USA<sup>b</sup>

**ABSTRACT** Here we present the draft genome of *Arthrobacter* sp. strain UCD-GKA. The assembly contains 4,930,274 bp in 33 contigs. This strain was isolated from the handle of a weight bar in the UC Davis Activities and Recreation Center.

embers of the genus *Arthrobacter* are aerobic, Gram-positive bacteria (1) primarily known for switching between the bacilli and cocci shapes depending on the environmental conditions (2). They are commonly found in soil.

Arthrobacter sp. strain UCD-GKA was isolated from a squatting bar located in the weight room of the UC Davis Activities and Recreation Center. This was part of an undergraduate research project to provide microbial reference genomes of bacteria isolated from the built environment (http://www.microbe.net). A sterile cotton-tipped swab was used to rub the handles of the squat bar and then applied to a lysogeny broth (LB) agar plate. The plate was incubated at 37°C for 4 days. A single colony was dilution streaked, and once isolated, was used to make an overnight culture that was also incubated at 37°C. DNA extraction from this overnight culture utilized the Promega Wizard genomic DNA purification kit. The 16S rRNA gene was amplified using PCR with the 27F and 1391R primers. The PCR product was then purified and used for Sanger sequencing. The resulting consensus sequence was analyzed using NCBI BLAST (3) and phylogenetic tree building using an alignment of related sequences from the Ribosomal Database Project (RDP) (4). A maximum-likelihood phylogenetic 16S rRNA gene tree was inferred using Fast Tree (5), and was visualized in Dendroscope (6). This preliminary tree included other Arthrobacter species, including A. antarcticus, A. psychrophenolicus, A. gangotriensis, and A. sulfureus.

For whole-genome sequencing, a paired-end library was prepared using a KAPA HyperPlus library prep kit (KAPA Biosystems). A portion of an Illumina MiSeq sequencing run generated 383,256 paired-end reads with a read length of 300 bp. After quality trimming and error correction were completed by the A5-miseq assembly pipeline (7, 8), 356,510 quality reads remained in 33 contigs, with 15× coverage and a GC content of 65.7%. Genome completeness was estimated using the Phylosift software (9), which uses a reference list of 37 highly conserved, single copy marker genes (10), 36 of which were found in the assembly in a single copy. The missing marker, RNase H11, was found in 98.5% of bacteria surveyed when these markers were compiled and so it is unclear whether this marker is genuinely absent or the assembly is incomplete. With the use of another metric, which was provided by CheckM (11), completeness was estimated at 99.5% complete.

Annotation was performed using RAST (12). *Arthrobacter* sp. strain UCD-GKA is predicted to contain 4,584 coding sequences and 70 noncoding RNAs. An almost full-length 16S rRNA gene sequence of 1,450 bp was obtained and analyzed as described above. Within the resulting tree, *Arthrobacter* sp. strain UCD-GKA was found

Received 29 November 2016 Accepted 1
December 2016 Published 9 February 2017

**Citation** Kincheloe GN, Eisen JA, Coil DA. 2017. Draft genome sequence of *Arthrobacter* sp. strain UCD-GKA (phylum *Actinobacteria*). Genome Announc 5:e01599-16. https://doi.org/10.1128/genomeA.01599-16.

**Copyright** © 2017 Kincheloe et al. This is an open-access article distributed under the terms of the Creative Commons Attribution 4.0 International license.

Address correspondence to Jonathan A. Eisen, jaeisen@ucdavis.edu.

Kincheloe et al. genameAnnouncements'

in a clade containing mostly *Arthrobacter sulfureus*. However, the tree contained many polyphyletic groups that did not correspond with the 16S rRNA gene phylogeny and therefore we did not assign a species name to this isolate (https://figshare.com/articles/Arthrobacter Tree/4234637).

**Accession number(s).** This whole-genome shotgun project has been deposited at DDBJ/EMBL/GenBank under the accession number MOLR00000000. The version described in this paper is version MOLR01000000.

## **ACKNOWLEDGMENTS**

Sequencing was performed at the DNA Technologies Core, University of California, Davis.

Funding was provided by the Alfred P. Sloan Foundation through a grant to Jonathan A. Eisen and David A. Coil via the "Microbiology of the Built Environment" program.

## **REFERENCES**

- Crocker FH, Fredrickson JK, White DC, Ringelberg DB, Balkwill DL. 2000. Phylogenetic and physiological diversity of *Arthrobacter* strains isolated from unconsolidated subsurface sediments. Microbiology 146: 1295–1310. https://doi.org/10.1099/00221287-146-6-1295.
- Luscombe BM, Gray TRG. 1974. Characteristics of Arthrobacter grown in continuous culture. Microbiology 82:213–222. https://doi.org/10.1099/ 00221287-82-2-213.
- Altschul SF, Gish W, Miller W, Myers EW, Lipman DJ. 1990. Basic local alignment search tool. J Mol Biol 215:403–410. https://doi.org/10.1016/ S0022-2836(05)80360-2.
- Cole JR, Wang Q, Fish JA, Chai B, McGarrell DM, Sun Y, Brown CT, Porras-Alfaro A, Kuske CR, Tiedje JM. 2014. Ribosomal Database Project: data and tools for high-throughput rRNA analysis. Nucleic Acids Res 42:D633–D642. https://doi.org/10.1093/nar/gkt1244.
- Price MN, Dehal PS, Arkin AP. 2010. FastTree 2—approximately maximum-likelihood trees for large alignments. PLoS One 5:e9490. https://doi.org/10.1371/journal.pone.0009490.
- Huson DH, Scornavacca C. 2012. Dendroscope 3: an interactive tool for rooted phylogenetic trees and networks. Syst Biol 61:1061–1067. https:// doi.org/10.1093/sysbio/sys062.
- 7. Coil D, Jospin G, Darling AE. 2015. A5-miseq: an updated pipeline to

- assemble microbial genomes from Illumina MiSeq data. Bioinformatics 31:587–589. https://doi.org/10.1093/bioinformatics/btu661.
- Tritt A, Eisen JA, Facciotti MT, Darling AE. 2012. An integrated pipeline for de novo assembly of microbial genomes. PLoS One 7:e42304. https:// doi.org/10.1371/journal.pone.0042304.
- Darling AE, Jospin G, Lowe E, Matsen FAIV, Bik HM, Eisen JA. 2014. PhyloSift: phylogenetic analysis of genomes and metagenomes. PeerJ 2:e243. https://doi.org/10.7717/peerj.243.
- Wu D, Jospin G, Eisen JA. 2013. Systematic identification of gene families for use as "markers" for phylogenetic and phylogeny-driven ecological studies of bacteria and Archaea and their Major Subgroups. PLoS One 8:e77033. https://doi.org/10.1371/journal.pone.0077033.
- Parks DH, Imelfort M, Skennerton CT, Hugenholtz P, Tyson GW. 2015. CheckM: assessing the quality of microbial genomes recovered from isolates, single cells, and metagenomes. Genome Res 25:1043–1055. https://doi.org/10.1101/gr.186072.114.
- 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. https://doi.org/10.1186/1471-2164-9-75.

Volume 5 Issue 6 e01599-16 genomea.asm.org **2**