



Complete Genome Sequence of a Wild-Type Isolate of *Caulobacter vibrioides* Strain CB1

Derrick C. Scott,^a Kiesha Wilson,^b Keshawn Ross,^a Damyen Ingram,^a Tajah Lewter,^a Jasmine Herring,^a David Duncan,^a Anthea Aikins,^a Bert Ely^b

^aDepartment of Biological Sciences, Delaware State University, Dover, Delaware, USA ^bDepartment of Biological Sciences, University of South Carolina, Columbia, South Carolina, USA

ABSTRACT The complete genome sequence of *Caulobacter vibrioides* strain CB1 consists of a chromosome of 4,137,285 bp, with a GC content of 67.2% and 3,990 coding DNA sequences. This strain contains the typical genome rearrangement that is characteristic of the *Caulobacter* strains that are currently sequenced. However, this strain is so closely related to sequenced strain NA1000 that rearrangements were minimal. This will allow further clarification of the causes of rearrangements in the species.

When the genome sequences of *Caulobacter* isolates NA1000 and K31 were compared, numerous genome rearrangements were observed (1). This phenomenon was dubbed genome scrambling. In contrast, similar comparisons of closely related species of other bacterial genera revealed nominal rearrangements. A phylogenetic analysis of the genomic sequences of additional *Caulobacter* strains has revealed insight into the mechanisms of genome scrambling, but more genomic sequences are needed to gain additional clarity. Here, we report the sequence of the total genomic material of an additional *C. vibrioides* wild-type strain, CB1, which was sampled from tap water in California.

The strain *C. vibrioides* CB1 was isolated from tap water in California (2). The cells were cultivated at 30°C for 48 h in peptone yeast extract medium (3), which contained 2 g Bacto peptone, 1 g yeast extract, 0.5 M MgSO₄, and 0.5 M CaCl₂ per liter. Genomic DNA was isolated using the Qiagen DNeasy tissue kit following the manufacturer's protocol. The primers 16S_533F (GTGCCAGCMGCCGCGGTAA) and 16S_U1492R (GG TTACCTTGTTACGACTT) were used to amplify the 16S rRNA region of the genome, and the amplified DNA was sequenced using Sanger technology on an ABI 3730 sequencer (GENEWIZ, USA). Genomic DNA sequencing was performed by the Delaware Bioinformatics Institute using a PacBio RS II sequencer, as suggested in previous studies for bacterial genomes with high GC contents (4). The resulting genome sequence was assembled using the Hierarchical Genome Assembly Process (HGAP) (5) in SMRT Portal through Amazon Machine Image (AMI) EC2 using the smrtanalysis-2.3.0-ami-20fb4848 image with the default *de novo* parameters. The sequence was annotated using the RAST server (http://rast.nmpdr.org) and the NCBI Prokaryotic Genome Annotation Pipeline (6–8) and then visualized and edited in Artemis (9).

The *C. vibrioides* CB1 complete genome consists of a circular chromosome of 4,137,285 bp, with a GC content of 67.2%. The genome is predicted to contain 3,990 coding sequences (CDSs). The numbers of tRNAs and rRNA operons are 51 and 2, respectively.

C. vibrioides CB1 has no inversions and only one insertion when compared to the closely related NA1000 strain. This is the first instance where two *Caulobacter* strains have revealed so few rearrangements when directly compared. Further studies be-

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Address correspondence to Derrick C. Scott, dcscott@desu.edu.

tween these strains will shed light into the mechanisms of genome rearrangement in *Caulobacter* species.

Data availability. The complete genome sequence of *C. vibrioides* CB1 has been deposited in GenBank under the accession number CP023314. The raw reads are also available under SRA accession number SRP158680.

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REFERENCES

- Scott DC, Ely B. 2016. Conservation of the essential genome among Caulobacter and Brevudimonas species. Curr Microbiol 72:503–510. https://doi.org/10.1007/s00284-015-0964-x.
- 2. Poindexter JS. 1964. Biological properties and classification of the *Caulobacter* group. Bacteriol Rev 28:231–295.
- 3. Johnson RC, Ely B. 1977. Isolation of spontaneously derived mutants of *Caulobacter crescentus*. Genetics 86:25–32.
- Scott DC, Ely B. 2015. Comparison of genome sequencing technology and assembly methods for the analysis of a GC-rich bacterial genome. Curr Microbiol 70:338–344. https://doi.org/10.1007/s00284-014-0721-6.
- Chin C-S, Alexander DH, Marks P, Klammer AA, Drake J, Heiner C, Clum A, Copeland A, Huddleston J, Eichler EE, Turner SW, Korlach J. 2013. Nonhybrid, finished microbial genome assemblies from long-read SMRT sequencing data. Nat Methods 10:563–569. https://doi.org/10.1038/nmeth.2474.
- Angiuoli SV, Gussman A, Klimke W, Cochrane G, Field D, Garrity G, Kodira CD, Kyrpides N, Madupu R, Markowitz V, Tatusova T, Thomson N, White O.

2008. Toward an online repository of standard operating procedures (SOPs) for (meta)genomic annotation. OMICS 12:137–141. https://doi.org/ 10.1089/omi.2008.0017.

- Tatusova T, DiCuccio M, Badretdin A, Chetvernin V, Nawrocki EP, Zaslavsky L, Lomsadze A, Pruitt KD, Borodovsky M, Ostell J. 2016. NCBI Prokaryotic Genome Annotation Pipeline. Nucleic Acids Res 44: 6614–6624. https://doi.org/10.1093/nar/gkw569.
- Haft DH, DiCuccio M, Badretdin A, Brover V, Chetvernin V, O'Neill K, Li W, Chitsaz F, Derbyshire MK, Gonzales NR, Gwadz M, Lu F, Marchler GH, Song JS, Thanki N, Yamashita RA, Zheng C, Thibaud-Nissen F, Geer LY, Marchler-Bauer A, Pruitt KD. 2018. RefSeq: an update on prokaryotic genome annotation and curation. Nucleic Acids Res 46:D851–D860. https://doi .org/10.1093/nar/gkx1068.
- Rutherford K, Parkhill J, Crook J, Horsnell T, Rice P, Rajandream M-A, Barrell B. 2000. Artemis: sequence visualization and annotation. Bioinformatics 16:944–945. https://doi.org/10.1093/bioinformatics/16.10.944.