

Genome Sequence of the *Sulfitobacter* sp. Strain 2047-Infecting Lytic Phage Φ CB2047-B

Nana Y. D. Ankrah,^a Charles R. Budinoff,^{a*} William H. Wilson,^b Steven W. Wilhelm,^a Alison Buchan^a

Department of Microbiology, University of Tennessee, Knoxville, Tennessee, USA^a; Bigelow Laboratory for Ocean Sciences, East Boothbay, Maine, USA^b

*Present address: Charles R. Budinoff, Algenol Biofuels, Inc., Fort Myers, Florida, USA.

N.Y.D.A. and C.R.B. contributed equally to this work.

We announce the complete genome sequence of a lytic podovirus, Φ CB2047-B, which infects the bacterium *Sulfitobacter* sp. strain 2047, a member of the *Roseobacter* clade. Genome analysis revealed Φ CB2047-B to be an N4-like phage, with its genome having high nucleotide similarity to other N4-like roseophage genomes.

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Address correspondence to Alison Buchan, abuchan@utk.edu.

Bacteria of the *Roseobacter* lineage, which includes the genus *Sulfitobacter*, are abundant marine heterotrophs that mediate several key biogeochemical processes, including the transformation of organic and inorganic sulfur compounds, the oxidation of carbon monoxide, and the degradation of vascular plant material (1–3). Roseobacters, along with their infecting phages, are excellent models for studying how microbial activities shape biogeochemical cycles (4). Here, we report the genome sequence of phage Φ CB2047-B, which infects *Sulfitobacter* sp. strain 2047.

The phage was isolated from a mesocosm study in Raunefjorden, Norway, using standard virus enrichment and plaque assay techniques (4, 5). Phage DNA was submitted to the Broad Institute and sequenced under the Gordon and Betty Moore Foundation's Marine Phage, Virus, and Virome Sequencing Project. The Broad Institute sequencing data were assembled using the Lasergene SeqMan Pro. The assemblies resulted in the generation of a single contig, which had sequencing coverage of approximately 30 \times . The contig was annotated using RAST and the tRNAscan-SE search server (6, 7). Translated peptides from the phage genome were used as BLASTp queries to the NCBI non-redundant protein sequence database to manually curate possible gene functions and to identify the nearest phage or prophage relatives. The CoreGenesUniqueGenes (CGUG) genome analysis tool (8) was used to identify gene homologues and assign core genes shared with other N4-like phages.

Phage Φ CB2047-B is 74,480 bp (74.5 kb) with a G+C content of 43% and 92 identified open reading frames. The genome sequence indicates this is an N4-like bacteriophage that is highly similar to but genetically distinct from other recently described roseophages (9). Morphological analysis by transmission electron microscopy confirmed that phage Φ CB2047-B belongs to the family *Podoviridae*. The genome content and architecture of Φ CB2047-B are similar to those of other N4 phages. Consistent with most other N4-like phages, the genome possesses 437-bp direct terminal repeat sequences on its distal ends. A CGUG analysis identified 20 highly homologous genes (BLASTp threshold

score, 85) between phage Φ CB2047-B and these previously reported N4-like phages: the enterobacterium phage N4 (accession no. NC_008720), *Pseudomonas* sp. phages LUZ7 (accession no. FN422398) and LIT1 (accession no. FN422399), and N4-like roseophages Φ DSS3P2 (accession no. FJ591093) and Φ EE36P1 (accession no. FJ591094). An analysis focused exclusively on N4-like roseophages (Φ DSS3P2 and Φ EE36P1) identified 41 genes with high homology. Genome-wide nucleotide similarity alignments with the Φ DSS3P2 and Φ EE36P1 genomes showed that phage Φ CB2047-B shares 43.9 and 44.4% nucleotide identity, respectively. Unlike other N4-like phages, Φ CB2047-B contains a deoxycytidine triphosphate (dCTP) deaminase instead of a deoxycytidine monophosphate deaminase, indicating a preference for an alternative route for the generation of dUMP for thymidine biosynthesis. The closest homologue in the NCBI database to the phage dCTP deaminase is from a coliphage, EC1-UPM (accession no. AGC31535), which has 37% identity. The host genome also contains a homologue to this protein that shares 29% identity to the phage gene and suggests genetic divergence.

Nucleotide sequence accession number. The complete sequence of the *Sulfitobacter* phage Φ CB2047-B genome can be accessed under the GenBank accession no. [HQ317387](https://www.ncbi.nlm.nih.gov/nuclseq/HQ317387).

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REFERENCES

- González JM, Kiene RP, Moran MA. 1999. Transformation of sulfur compounds by an abundant lineage of marine bacteria in the α -subclass of the class *Proteobacteria*. *Appl. Environ. Microbiol.* 65:3810–3819.
- Buchan A, González JM, Moran MA. 2005. Overview of the marine *Roseobacter* lineage. *Appl. Environ. Microbiol.* 71:5665–5677. <http://dx.doi.org/10.1128/AEM.71.10.5665-5677.2005>.

3. Gulvik CA, Buchan A. 2013. Simultaneous catabolism of plant-derived aromatic compounds results in enhanced growth for members of the *Roseobacter* lineage. *Appl. Environ. Microbiol.* 79:3716–3723. <http://dx.doi.org/10.1128/AEM.00405-13>.
4. Ankrah NY, May AL, Middleton JL, Jones DR, Hadden MK, Gooding JR, LeCleir GR, Wilhelm SW, Campagna SR, Buchan A. 5 December 2013. Phage infection of an environmentally relevant marine bacterium alters host metabolism and lysate composition. *ISME J.* <http://dx.doi.org/10.1038/ismej.2013.216>.
5. Budinoff CR. 2012. Diversity and activity of roseobacters and roseophage. Ph.D. thesis. University of Tennessee, Knoxville, TN.
6. Lowe TM, Eddy SR. 1997. tRNAscan-SE: a program for improved detection of transfer RNA genes in genomic sequence. *Nucleic Acids Res.* 25: 0955–0964. <http://dx.doi.org/10.1093/nar/25.5.0955>.
7. Aziz RK, Bartels D, Best AA, DeJongh M, Disz T, Edwards RA, Zagnitko O, 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. <http://dx.doi.org/10.1186/1471-2164-9-75>.
8. Mahadevan P, King JF, Seto D. 2009. CGUG: *in silico* proteome and genome parsing tool for the determination of “core” and unique genes in the analysis of genomes up to ca. 1.9 Mb. *BMC Res. Notes* 2:168. <http://www.biomedcentral.com/1756-0500/2/168>.
9. Zhao Y, Wang K, Jiao N, Chen F. 2009. Genome sequences of two novel phages infecting marine roseobacters. *Environ. Microbiol.* 11:2055–2064. <http://dx.doi.org/10.1111/j.1462-2920.2009.01927.x>.