



Draft Genome Sequences of *Sphingomonas mucosissima* DSM 17494 and *Sphingomonas dokdonensis* DSM 21029

 Anja Poehlein,^a Jan Hendrik Wübbeler,^b  Rolf Daniel,^a Alexander Steinbüchel^{b,c}

Department of Genomic and Applied Microbiology and Göttingen Genomics Laboratory, Institut für Mikrobiologie & Genetik, Georg-August-Universität Göttingen, Göttingen, Germany^a; Institut für Molekulare Mikrobiologie & Biotechnologie, Westfälische Wilhelms-Universität Münster, Münster, Germany^b; Environmental Sciences Department, Faculty of Meteorology, Environment and Arid Land Agriculture, King Abdulaziz University, Jeddah, Saudi Arabia^c

ABSTRACT *Sphingomonas mucosissima* and *Sphingomonas dokdonensis* are Gram-negative chemoheterotrophic strictly aerobic rods or cocci. The genomes (3.453 Mb and 3.587 Mb, respectively) contain 3,279 and 3,329 predicted protein-encoding genes, respectively. The genome of *S. dokdonensis* harbors a 90-kb plasmid.

The genus *Sphingomonas* was proposed in 1990 (1), and since then, the scientific interest for sphingomonads, and thus the increasing count of publications for this novel species, are constantly rising (compare at <http://www.bacterio.net/sphingomonas.html>). Members of this genus are Gram-negative chemoheterotrophic strictly aerobic rods or cocci (1). The type species is *Sphingomonas paucimobilis* DSM 1098 (2, 3). The most characteristic feature of this genus is the presence of sphingolipids in their outer cell membrane (1, 4). During a study for the optimization of triacylglycerol synthesis in novel oleaginous bacteria (5), *Sphingomonas jeddahensis* G39^T was enriched and isolated from desert soil and published as a new species of the genus *Sphingomonas* (6). Its closest relative type strains are *Sphingomonas dokdonensis* DSM 21029 and *Sphingomonas mucosissima* DSM 17494. *S. dokdonensis* DSM 21029 was derived from Dokdo, a South Korean island with restricted human access (7). *S. mucosissima* DSM 17494 was isolated from the Colorado Plateau (USA) (8). In this study, both strains were sequenced, and the draft genomes were analyzed and are presented here.

Chromosomal DNA was isolated using the MasterPure complete DNA purification kit (Epicentre, Madison, WI, USA). The extracted DNA was used to generate Illumina shotgun paired-end sequencing libraries, which were sequenced with a MiSeq instrument and the MiSeq reagent kit (version 3), as recommended by the manufacturer (Illumina, San Diego, CA, USA). Quality filtering using Trimmomatic version 0.36 (9) resulted in 2,754,576 and 2,741,920 paired-end reads for *S. dokdonensis* and *S. mucosissima*, respectively. The assembly performed with the SPAdes genome assembler software version 3.10.0 (10) resulted in 16 contigs (>500 bp) for both genomes, with average coverages of 153-fold (*S. dokdonensis*) and 149-fold (*S. mucosissima*). The assembly was validated and the read coverage determined with QualiMap version 2.1 (11). The draft genome of *S. dokdonensis* DSM 21029 (3.453 Mb) exhibited an overall G+C content of 66.46%. The genome of *S. mucosissima* DSM 17494 (3.587 Mb), with an overall G+C content of 65.08%, is slightly larger than that of *S. dokdonensis*. Automatic gene prediction and identification of rRNA and tRNA genes were performed using the software tool Prokka (12). The draft genomes of *S. dokdonensis* and *S. mucosissima* contain 3,279 predicted protein-encoding and 54 RNA genes and 3,329 predicted

Received 17 July 2017 Accepted 19 July 2017 Published 31 August 2017

Citation Poehlein A, Wübbeler JH, Daniel R, Steinbüchel A. 2017. Draft genome sequences of *Sphingomonas mucosissima* DSM 17494 and *Sphingomonas dokdonensis* DSM 21029. *Genome Announc* 5:e00889-17. <https://doi.org/10.1128/genomeA.00889-17>.

Copyright © 2017 Poehlein et al. This is an open-access article distributed under the terms of the [Creative Commons Attribution 4.0 International license](https://creativecommons.org/licenses/by/4.0/).

Address correspondence to Anja Poehlein, apoehle3@gwdg.de.

A.P. and J.H.W. contributed equally to this article.

protein-encoding and 52 RNA genes, respectively. The genome of *S. dokdonensis* putatively harbors a 90-kb plasmid, as genes encoding a RepB family protein, a partitioning protein ParA, and a replication protein B were located in a single contig. Genes encoding a serine palmitoyltransferase and an NADPH-dependent 3-ketodihydrosphingosine reductase, catalyzing the initial steps of *de novo* sphingolipid biosynthesis, have been identified in both genomes (13).

Accession number(s). These whole-genome shotgun projects have been deposited at DDBJ/EMBL/GenBank under the accession numbers [NBBJ00000000](#) and [NBBJ00000000](#) for *S. dokdonensis* DSM 21029 and *S. mucosissima* DSM 17494, respectively. The versions described here are the first versions, NBBJ01000000 and NBBJ01000000, respectively.

ACKNOWLEDGMENT

We thank Melanie Heinemann for technical support.

REFERENCES

1. Yabuuchi E, Yano I, Oyaizu H, Hashimoto Y, Ezaki T, Yamamoto H. 1990. Proposals of *Sphingomonas paucimobilis* gen. nov. and comb. nov., *Sphingomonas parapaucimobilis* sp. nov., *Sphingomonas yanoikuyae* sp. nov., *Sphingomonas adhaesiva* sp. nov., *Sphingomonas capsulata* comb. nov. and two genospecies of the genus *Sphingomonas*. *Microbiol Immunol* 34:99–119. <https://doi.org/10.1111/j.1348-0421.1990.tb00996.x>.
2. Holmes B, Owen RJ, Evans A, Malnick H, Willcox WR. 1977. *Pseudomonas paucimobilis*, a new species isolated from human clinical specimens, the hospital environment, and other sources. *Int J Syst Evol Microbiol* 27:133–146. <https://doi.org/10.1099/00207713-27-2-133>.
3. Takeuchi M, Hamana K, Hiraishi A. 2001. Proposal of the genus *Sphingomonas sensu stricto* and three new genera, *Sphingobium*, *Novosphingobium* and *Sphingopyxis*, on the basis of phylogenetic and chemotaxonomic analyses. *Int J Syst Evol Microbiol* 51:1405–1417. <https://doi.org/10.1099/00207713-51-4-1405>.
4. Chen H, Jogler M, Rohde M, Klenk HP, Busse HJ, Tindall BJ, Spröer C, Overmann J. 2012. Reclassification and emended description of *Caulobacter leidyi* as *Sphingomonas leidyi* comb. nov., and emendation of the genus *Sphingomonas*. *Int J Syst Evol Microbiol* 62:2835–2843. <https://doi.org/10.1099/ijs.0.039636-0>.
5. Röttig A, Hauschild P, Madkour MH, Al-Ansari AM, Almakishah NH, Steinbüchel A. 2016. Analysis and optimization of triacylglycerol synthesis in novel oleaginous *Rhodococcus* and *Streptomyces* strains isolated from desert soil. *J Biotechnol* 225:48–56. <https://doi.org/10.1016/j.jbiotec.2016.03.040>.
6. Wübbeler JH, Oppermann-Sanio FB, Ockenfels A, Röttig A, Osthhaar-Ebker A, Verbarq S, Poehlein A, Madkour MH, Al-Ansari AM, Almakishah NH, Daniel R, Steinbüchel A. 2016. *Sphingomonas jeddahensis* sp. nov., isolated from Saudi-Arabian desert soil. *Int J Syst Evol Microbiol*, in press.
7. Yoon JH, Lee MH, Kang SJ, Lee SY, Oh TK. 2006. *Sphingomonas dokdonensis* sp. nov., isolated from soil. *Int J Syst Evol Microbiol* 56:2165–2169. <https://doi.org/10.1099/ijs.0.64114-0>.
8. Reddy GSN, Garcia-Pichel F. 2007. *Sphingomonas mucosissima* sp. nov. and *Sphingomonas desiccabilis* sp. nov., from biological soil crusts in the Colorado Plateau, USA. *Int J Syst Evol Microbiol* 57:1028–1034. <https://doi.org/10.1099/ijs.0.64331-0>.
9. Bolger AM, Lohse M, Usadel B. 2014. Trimmomatic: a flexible trimmer for Illumina sequence data. *Bioinformatics* 30:2114–2120. <https://doi.org/10.1093/bioinformatics/btu170>.
10. Bankevich A, Nurk S, Antipov D, Gurevich AA, Dvorkin M, Kulikov AS, Lesin VM, Nikolenko SI, Pham S, Pribelski AD, Pyshkin AV, Sirotkin AV, Vyahhi N, Tesler G, Alekseyev MA, Pevzner PA. 2012. SPAdes: a new genome assembly algorithm and its applications to single-cell sequencing. *J Comput Biol* 19:455–477. <https://doi.org/10.1089/cmb.2012.0021>.
11. García-Alcalde F, Okonechnikov K, Carbonell J, Cruz LM, Götz S, Tarazona S, Dopazo J, Meyer TF, Conesa A. 2012. Qualimap: evaluating next-generation sequencing alignment data. *Bioinformatics* 28:2678–2679. <https://doi.org/10.1093/bioinformatics/bts503>.
12. Seemann T. 2014. Prokka: rapid prokaryotic genome annotation. *Bioinformatics* 30:2068–2069. <https://doi.org/10.1093/bioinformatics/btu153>.
13. Bower E, Lowther J, Campopiano D. 2011. Identifying enzymes involved in sphingolipid biosynthesis in the environmentally useful bacterium *Sphingomonas wittichii*. *FASEB J* 25:739.4.