

Draft Genome Sequence of *Streptomyces spongiicola* Strain 531S, an Actinobacterium Isolated from Marine Sediment

メタデータ	言語: eng 出版者: 公開日: 2019-04-12 キーワード (Ja): キーワード (En): 作成者: Dohra, Hideo, Kaweewan, Issara, Casareto, Beatriz E., Suzuki, Yoshimi, Kodani, Shinya メールアドレス: 所属:
URL	<a href="http://hdl.handle.net/10297/00026412">http://hdl.handle.net/10297/00026412</a>



# Draft Genome Sequence of *Streptomyces spongiicola* Strain 531S, an Actinobacterium Isolated from Marine Sediment

Hideo Dohra,<sup>a,b</sup> Issara Kaweewan,<sup>c</sup> Beatriz E. Casareto,<sup>a,c</sup> Yoshimi Suzuki,<sup>c</sup> Shinya Kodani<sup>b,c</sup>

<sup>a</sup>Research Institute of Green Science and Technology, Shizuoka University, Suruga-ku, Shizuoka, Japan

<sup>b</sup>Graduate School of Integrated Science and Technology, Shizuoka University, Suruga-ku, Shizuoka, Japan

<sup>c</sup>Graduate School of Science and Technology, Shizuoka University, Suruga-ku, Shizuoka, Japan

**ABSTRACT** *Streptomyces spongiicola* strain 531S (NBRC 113560) was isolated from marine sediment on a beach on Sesoko Island (Okinawa, Japan). We report here the draft genome sequence of *S. spongiicola* 531S, in which 24 potential secondary metabolite gene clusters were predicted with antiSMASH.

*Streptomyces spongiicola* (HNM0071<sup>T</sup>) was previously described as a novel actinomycete species (1). We isolated *S. spongiicola* strain 531S (NBRC 113560) from marine sediment on a beach on Sesoko Island (Okinawa, Japan) using International Streptomyces Project 3 (ISP3) (2) agar medium. To clarify its potential for biosynthesis of secondary metabolites, we determined the draft genome sequence of *S. spongiicola* 531S.

*S. spongiicola* 531S was cultured in ISP2 (2) medium, and the DNA was extracted with a DNeasy blood and tissue kit (Qiagen). A paired-end library of the genome was constructed as previously described (3) and sequenced with the Illumina MiSeq platform (302-bp paired end). The raw read sequences were cleaned with Trimmomatic (4) as described previously (3). The 3,210,816 high-quality reads totaling approximately 730 Mb were assembled with SPAdes v. 3.11.1 (5) with a parameter setting as described previously (3), and contigs less than 500 bp, in which no protein-coding sequence was predicted, were removed. The resulting 84 contigs showed a 105.6-fold coverage, and the  $N_{50}$  value was 395,703 bp.

The draft genome of *S. spongiicola* 531S has a total size of 6,912,677 bp with a G+C content of 72.61%. The average nucleotide identity (ANI) of the genome was calculated using the ani.rb script from the enveomics collection (6). The ANI analysis showed a high ANI value (99.0%) with the complete genome sequence of *S. spongiicola* HNM0071<sup>T</sup> (accession no. CP029254). Gene prediction and annotation were performed using the DFAST-core stand-alone program v. 1.0.7 (7) with a *S. spongiicola* HNM0071<sup>T</sup> protein database constructed with a utility script bundled in the DFAST-core and manually curated. The *S. spongiicola* 531S genome contained 5,874 protein-coding sequences, 81 tRNA genes, and 19 clustered regularly interspaced short palindromic repeat (CRISPR) sequences. Among 5,874 proteins, 5,003 (85.17%) were predicted to be orthologous proteins, with those of *S. spongiicola* HNM0071<sup>T</sup> in reciprocal best matches using a blastp search (6) with the following settings: identity, >50%; E value, <1e-10; and query sequence coverage, >50%.

Secondary metabolite biosynthesis gene clusters of the *S. spongiicola* 531S genome were predicted with the antiSMASH 4.1.0 bacterial version (<https://antismash.secondarymetabolites.org/>) (8). The *S. spongiicola* 531S genome sequence contained 24 putative biosynthetic gene clusters, including 9 clusters involved in the biosynthesis of the following bioactive peptides: 3 bacteriocins, 2 thiopeptides, 1 nonribosomal peptide synthetase (NRPS), and hybrid clusters of a ladderane/arylpolypene/NRPS, a lantipeptide/

**Citation** Dohra H, Kaweewan I, Casareto BE, Suzuki Y, Kodani S. 2019. Draft genome sequence of *Streptomyces spongiicola* strain 531S, an actinobacterium isolated from marine sediment. Microbiol Resour Announc 8:e01198-18. <https://doi.org/10.1128/MRA.01198-18>.

**Editor** Julia A. Maresca, University of Delaware

**Copyright** © 2019 Dohra 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 Shinya Kodani, kodani.shinya@shizuoka.ac.jp.

**Received** 30 August 2018

**Accepted** 10 December 2018

**Published** 17 January 2019

type I polyketide synthase (PKS)/other type of PKS, and a type I PKS/butyrolactone/NRPS. Interestingly, comparative analysis of the biosynthetic gene clusters with *S. spongiicola* HNM0071<sup>T</sup> revealed that *S. spongiicola* 531S lacked two lantipeptide gene clusters, including genes coding for the FxLD family of lantipeptides (accession no. [AWK11671](#) to [AWK11673](#) and [AWK08909](#)). The region of one of the two lantipeptide gene clusters in the *S. spongiicola* HNM0071<sup>T</sup> genome was replaced by two CRISPR sequences and a gene cluster for type I-E CRISPR-associated proteins (accession no. [GBP98722](#) to [GBP98729](#)) in the *S. spongiicola* 531S genome sequence.

The predicted secondary metabolite biosynthesis gene clusters of *S. spongiicola* 531S and their comparative analyses with *S. spongiicola* HNM0071<sup>T</sup> will contribute to study of bioactive compounds and their biosynthetic pathways.

*Streptomyces spongiicola* strain 531S was deposited in the NBRC culture collection (NITE Biological Resource Center, Japan) and designated as NBRC 113560.

**Data availability.** The raw reads of *Streptomyces spongiicola* strain 531S have been deposited in the DDBJ Sequence Read Archive (SRA) under the accession no. [DRA007446](#). This whole-genome shotgun sequencing project has been deposited in DDBJ/ENA/GenBank under the accession no. [BGZL00000000](#). The version described in this paper is the first version, BGZL01000000.

## ACKNOWLEDGMENTS

This study was supported by the Japan Society for the Promotion of Science with grants-in-aids (grant no. 16K01913), the ESPEC Foundation for Global Environment Research and Technology (Charitable Trust), and a 50th Anniversary Grant from the Mitsubishi Corporation for the Global Coral Reef Conservation Project (GCRCP).

We thank the staff at the Sesoko Station, Tropical Biosphere Research Center, University of the Ryukyus, Japan, for providing the necessary facilities during this study and members of Casareto's Marine Biogeochemistry Laboratory, Shizuoka University, Japan, for their help during our survey.

## REFERENCES

- Huang X, Zhou S, Huang D, Chen J, Zhu W. 2016. *Streptomyces spongiicola* sp. nov., an actinomycete derived from marine sponge. *Int J Syst Evol Microbiol* 66:738–743. <https://doi.org/10.1099/ijsem.0.000782>.
- Shirling EB, Gottlieb D. 1966. Methods for characterizing *Streptomyces* species. *Int J Syst Bacteriol* 16:313–340. <https://doi.org/10.1099/00207713-16-3-313>.
- Dohra H, Miyake Y, Kodani S. 2017. Draft genome sequence of *Streptomyces olivochromogenes* NBRC 3561, a bioactive peptide-producing actinobacterium. *Genome Announc* 5:e01048-17. <https://doi.org/10.1128/genomeA.01048-17>.
- 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>.
- Bankevich A, Nurk S, Antipov D, Gurevich AA, Dvorkin M, Kulikov AS, Lesin VM, Nikolenko SI, Pham S, Prjibelski 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>.
- Rodriguez-R LM, Konstantinidis KT. 2016. The enveomics collection: a toolbox for specialized analyses of microbial genomes and metagenomes. *PeerJ* 4:e1900v1. <https://doi.org/10.7287/peerj.preprints.1900v1>.
- Tanizawa Y, Fujisawa T, Nakamura Y. 2018. DFAST: a flexible prokaryotic genome annotation pipeline for faster genome publication. *Bioinformatics* 34:1037–1039. <https://doi.org/10.1093/bioinformatics/btx713>.
- Blin K, Wolf T, Chevrette MG, Lu X, Schwalen CJ, Kautsar SA, Suarez Duran HG, de los Santos ELC, Kim HU, Nave M, Dickschat JS, Mitchell DA, Shelest E, Breitling R, Takano E, Lee SY, Weber T, Medema MH. 2017. antiSMASH 4.0—improvements in chemistry prediction and gene cluster boundary identification. *Nucleic Acids Res* 45:W36–W41. <https://doi.org/10.1093/nar/gkx319>.