



Draft Genome Sequence of *Dermaococcus nishinomiyaensis* TSA37, Isolated from Wood Ash

 Alexander N. Williams,^a  Kyle S. MacLea^{a,b,c}

^aBiotechnology Program, University of New Hampshire, Manchester, New Hampshire, USA

^bBiology Program, University of New Hampshire, Manchester, New Hampshire, USA

^cDepartment of Life Sciences, University of New Hampshire, Manchester, New Hampshire, USA

ABSTRACT *Dermaococcus nishinomiyaensis* is a common bacterial resident of the human skin microbiome, among other environments. *D. nishinomiyaensis* strain TSA37 was isolated from the ash pan of a residential wood pellet stove. A genome assembly of 3,130,592 bp was generated, with an N_{50} value of 197,547 bp and a calculated G+C content of 69.01%.

Dermaococcus nishinomiyaensis is a species of aerobic Gram-positive cocci of the family *Dermaococcaceae* (1, 2). It is a known skin commensal (3) and is largely nonpathogenic to humans, though recent studies have implicated *D. nishinomiyaensis* in peritoneal dialysis-related peritonitis (4), catheter-related bacteremia (5), and polymicrobial infections of the skin (6) and urinary tract (7, 8). Beyond the human skin microbiome, *D. nishinomiyaensis* has also been isolated from an indoor track facility (9), cured meat (10), well water (11), and the gut of adult *Sarcophaga* flesh flies (12). Given a growing awareness of *D. nishinomiyaensis* beyond the human skin in other areas of the human-built environment, comparisons of skin commensal strains with strains found in other human-adjacent spaces may shed light on conditions under which the strains may exhibit pathogenicity.

D. nishinomiyaensis strain TSA37 was isolated from the ash pan of a residential wood pellet stove. A 30-ml sample of wood ash was collected, and a small portion was spread using a sterile cotton-tipped applicator onto a tryptic soy agar (TSA) plate and incubated at 37°C for 48 hours. After it was subcultured on a TSA plate formulated with 50.0 mg/liter cycloheximide for 24 hours, a small circular yellow colony (Gram-positive cocci) was isolated. Genomic DNA was purified from 5 ml inoculated tryptic soy broth (grown at 37°C for 24 hours) using the QIAamp DNA minikit (Qiagen, Valencia, CA, USA). The purified genomic DNA was fragmented and tagged with sequence adapters using the HyperPlus kit v.3.16 (catalog number KR1145; KAPA, Wilmington, MA, USA) and then sequenced with an Illumina HiSeq 2500 instrument at the University of New Hampshire (UNH) Hubbard Center for Genome Studies (Durham, NH, USA). The resulting 250-bp paired-end reads were bioinformatically paired and trimmed using Trimmomatic v.0.38 (with the following settings: paired-end mode with a window size of 4, quality requirement of 15, and minimum read length of 36), and 1,927,890 trimmed reads covering 3,188,889 bp were then assembled into a draft genome sequence using SPAdes v.3.13.0 with default settings (13, 14) and assessed for quality measures using QUAST v.5.0.2 (15). Contaminants and contigs of <500 bp were removed, and genes and features on the remaining contigs were identified and annotated using the NCBI Prokaryotic Genome Annotation Pipeline (PGAP) v.4.8 (16).

Assignment of the strain to the species *D. nishinomiyaensis* was verified using BLAST (17), average nucleotide identity (98.05% using EzBioCloud) (18), and the NCBI SRA Taxonomy Analysis Tool (STAT) (not yet published but available on the NCBI Sequence

Citation Williams AN, MacLea KS. 2019. Draft genome sequence of *Dermaococcus nishinomiyaensis* TSA37, isolated from wood ash. *Microbiol Resour Announc* 8:e01370-19. <https://doi.org/10.1128/MRA.01370-19>.

Editor David A. Baltus, University of Arizona

Copyright © 2019 Williams and MacLea. 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 Kyle S. MacLea, Kyle.MacLea@UNH.edu.

K.S.M. dedicates this work to Normand A. B. Clement (1908 to 1977).

Received 31 October 2019

Accepted 11 November 2019

Published 12 December 2019

Read Archive “Analysis” tab). *D. nishinomiyaensis* strain TSA37 was found to have a complete genome size of 3,130,592 bp across 37 contigs, with an average sequence coverage of 154×, an N_{50} value of 197,547 bp, and a G+C content of 69.01%. A total of 2,864 genes, 2,801 coding sequences, 93 pseudogenes, and 3 noncoding RNAs (ncRNAs) were identified using PGAP analysis. One CRISPR array was also detected using PGAP analysis. A BLASTn Megablast search of the 29-bp repeat sequence for the array returned *Pseudomonas* phage JG054 as the best hit, with marginal scoring (E value, 0.70; query coverage, 62%; identity, 100%).

Data availability. This *Dermaococcus nishinomiyaensis* whole-genome shotgun project has been deposited in DDBJ/ENA/GenBank under the accession number SUPQ00000000. The version described in this paper is the first version, SUPQ01000000. The raw Illumina data from BioProject PRJNA534294 were submitted to the NCBI Sequence Read Archive (SRA) under experiment accession number SRX6871076.

ACKNOWLEDGMENTS

Sequencing was undertaken at the Hubbard Center for Genome Studies at UNH, supported by NH-INBRE, with the assistance of Kelley Thomas and Stephen Simpson. This work was a project of the Microbiology Education through Genome Annotation-New Hampshire (MEGA-NH) program.

Bacterial and DNA isolation costs were funded via student laboratory fees of the BSCI 737 Microbial Genomics course at the University of New Hampshire (spring 2019 semester). Sequencing costs were supported by New Hampshire-INBRE through an institutional development award (IDeA), P20GM103506, from the National Institute of General Medical Sciences of the NIH. The funders had no role in study design, data collection and interpretation, or the decision to submit the work for publication.

REFERENCES

- Stackebrandt E, Koch C, Gvozdiak O, Schumann P. 1995. Taxonomic dissection of the genus *Micrococcus*: *Kocuria* gen. nov., *Nesterenkonia* gen. nov., *Kytococcus* gen. nov., *Dermaococcus* gen. nov., and *Micrococcus* Cohn 1872 gen. emend. *Int J Syst Bacteriol* 45:682–692. <https://doi.org/10.1099/00207713-45-4-682>.
- Stackebrandt E, Schumann P. 2000. Description of *Bogoriellaceae* fam. nov., *Dermaococcaceae* fam. nov., *Rarobacteraceae* fam. nov. and *Sanguibacteraceae* fam. nov. and emendation of some families of the suborder *Micrococcineae*. *Int J Syst Evol Microbiol* 50:1279–1285. <https://doi.org/10.1099/00207713-50-3-1279>.
- Kocur M, Schleifer KH, Kloos WE. 1975. Taxonomic status of *Micrococcus nishinomiyaensis* Oda 1935. *Int J Syst Bacteriol* 25:290–293. <https://doi.org/10.1099/00207713-25-3-290>.
- Tanaka A, Watanabe Y, Ito C, Murata M, Shinjo H, Otsuka Y, Takeda A. 2019. Successful treatment of peritoneal dialysis-related peritonitis caused by *Dermaococcus nishinomiyaensis*. *CEN Case Rep* 8:183–187. <https://doi.org/10.1007/s13730-019-00388-2>.
- Joron C, Roméo B, Le Flèche-Matéos A, Rames C, El Samad Y, Hamdad F. 2019. *Dermaococcus nishinomiyaensis* as a cause of persistent paediatric catheter-related bacteraemia. *Clin Microbiol Infect* 25:1054–1055. <https://doi.org/10.1016/j.cmi.2019.02.023>.
- Katoulis AC, Koumaki D, Liakou AI, Vrioni G, Koumaki V, Kontogiorgi D, Tzima K, Tsakris A, Rigopoulos D. 2015. Aerobic and anaerobic bacteriology of hidradenitis suppurativa: a study of 22 cases. *Skin Appendage Disord* 1:55–59. <https://doi.org/10.1159/000381959>.
- Seifu WD, Gebissa AD. 2018. Prevalence and antibiotic susceptibility of uropathogens from cases of urinary tract infections (UTI) in Shashemene referral hospital, Ethiopia. *BMC Infect Dis* 18:30. <https://doi.org/10.1186/s12879-017-2911-x>.
- Bonkat G, Rieken M, Rentsch CA, Wyler S, Feike A, Schäfer J, Gasser T, Trampuz A, Bachmann A, Widmer AF. 2011. Improved detection of microbial ureteral stent colonisation by sonication. *World J Urol* 29:133–138. <https://doi.org/10.1007/s00345-010-0535-5>.
- Klein BA, Lemon KP, Gajare P, Jospin G, Eisen JA, Coil DA. 2017. Draft genome sequences of *Dermaococcus nishinomiyaensis* strains UCD-KPL2534 and UCD-KPL2528 isolated from an indoor track facility. *Genome Announc* 5:e01652-16. <https://doi.org/10.1128/genomeA.01652-16>.
- Cordero MR, Zumalacarregui JM. 2000. Characterization of *Micrococcaceae* isolated from salt used for Spanish dry-cured ham. *Lett Appl Microbiol* 31:303–306. <https://doi.org/10.1046/j.1472-765x.2000.00818.x>.
- Oda M. 1935. Bacteriological studies on water used for brewing sake (part 6). I. Bacteriological studies on “miyamizu” (8) and (9). *Micrococcus* and *Actinomyces* isolated from “miyamizu.” *Jozogaku Zasshi* 13:1202–1228. (In Japanese.)
- Gupta AK, Rastogi G, Nayduch D, Sawant SS, Bhonde RR, Shouche YS. 2014. Molecular phylogenetic profiling of gut-associated bacteria in larvae and adults of flesh flies. *Med Vet Entomol* 28:345–354. <https://doi.org/10.1111/mve.12054>.
- 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>.
- Gurevich A, Saveliev V, Vyahhi N, Tesler G. 2013. QUAST: quality assessment tool for genome assemblies. *Bioinformatics* 29:1072–1075. <https://doi.org/10.1093/bioinformatics/btt086>.
- Tatusova T, DiCuccio M, Badretdin A, Chetvertin 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>.
- 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](https://doi.org/10.1016/S0022-2836(05)80360-2).
- Yoon S-H, Ha S-M, Kwon S, Lim J, Kim Y, Seo H, Chun J. 2017. Introducing EzBioCloud: a taxonomically united database of 16S rRNA gene sequences and whole-genome assemblies. *Int J Syst Evol Microbiol* 67:1613–1617. <https://doi.org/10.1099/ijsem.0.001755>.