



# Draft Genome Sequence of *Staphylococcus succinus* Strain GN1, Isolated from a Basement Floor in Milwaukee, WI

Grant P. Nickolson,<sup>a</sup> Nasim Maghbolí Balasjín,<sup>a</sup>  Christopher W. Marshall<sup>a</sup>

<sup>a</sup>Department of Biological Sciences, Marquette University, Milwaukee, Wisconsin, USA

**ABSTRACT** A strain of *Staphylococcus succinus* was sampled from the floor of the basement of a house and isolated in an undergraduate classroom in Milwaukee, WI. Here, we report the draft genome sequence of this strain.

A sterile swab was used to sample the basement floor of a 1916 home in Milwaukee, WI, to see if environmental isolates from a home environment had any measurable antibiotic resistance. A sterile cotton swab was prewetted with sterile water, and then the bare concrete floor of the basement was swabbed. The swab was then immediately placed in the freezer. The next day, the swab was streaked onto brain heart infusion (BHI) agar and incubated at 30°C for 24 h in order to isolate colonies for further experiments and DNA extraction. A single colony, named isolate GN1, was selected at random and streaked twice for isolation. A single colony was then grown in BHI liquid medium for physiological tests and DNA extraction. DNA was extracted using the DNeasy blood and tissue kit (Qiagen). The sample was then sent to the Microbial Genome Sequencing Center for whole-genome sequencing. Libraries were prepared using a modified version of the Illumina Nextera kit as previously described (1) and sequenced on an Illumina NextSeq 550 instrument. The adapters were removed from the 151-bp reads prior to analysis using *bcl2fastq*. A total of 3,832,226 paired-end reads were recovered. Sequence analyses were done using default settings in the KBase.us narrative interface (2), and the publicly available narrative containing all analyses and data can be found at <https://kbase.us/n/51640/53/>.

After the raw reads were uploaded to KBase, SPAdes v3.13.0 was used to assemble the genome sequence (3). The assembly stats compiled by QUAST (4) demonstrate that the assembly contained contigs ranging from 593 bp to 1.4 million bp and had a genome size of 2,838,015 bp and a G+C content of 33%. The assembly consisted of 11 contigs with an  $N_{50}$  value of 782,976 bp (Table 1). The assembled genome sequence was annotated using the Rapid Annotations using Subsystems Technology (RAST) v0.1.1 toolkit to detect gene locations and functions (5, 6). We also uploaded the assembly to NCBI with the associated BioSample accession number, where gene predictions and annotations were completed using NCBI Prokaryotic Genome Annotation Pipeline (PGAP) (7). The Genome Taxonomy Database (GTDB) v1.1.0 was used for taxonomic identification (8) and FastANI (9) for relatedness. Based on the average nucleotide identity, our isolate was 98.5% identical to *Staphylococcus succinus* (GenBank accession number [GCF\\_001006765.1](https://www.ncbi.nlm.nih.gov/nuccore/GCF_001006765.1)). A phylogenetic tree of our isolate, GN1, and the available *S. succinus* genome assemblies (Fig. 1) was created using the GToTree v1.5.46 phylogenomics workflow (10). Single-copy marker genes were identified using HMMER v3.2.1 (11), aligned using Muscle v3.8 (12), and trimmed using trimAl v1.4 (13), and the phylogenetic tree was generated using IQ-TREE v2.0.3 (14). The pipeline was run using GNU Parallel v20201122 (15). Because of the high sequence similarity and placement on the phylogenetic tree, our isolate is likely a strain of *S. succinus* that we will refer to as *S. succinus* strain GN1.

*Staphylococcus succinus* strains are typically aerobic, Gram-positive bacteria that are commonly isolated from ripened cheese (16, 17). *Staphylococcus* strains are spherical

**Citation** Nickolson GP, Maghbolí Balasjín N, Marshall CW. 2021. Draft genome sequence of *Staphylococcus succinus* strain GN1, isolated from a basement floor in Milwaukee, WI. *Microbiol Resour Announc* 10:e00580-21. <https://doi.org/10.1128/MRA.00580-21>.

**Editor** Steven R. Gill, University of Rochester School of Medicine and Dentistry

**Copyright** © 2021 Nickolson 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 Christopher W. Marshall, [christopher.marshall@marquette.edu](mailto:christopher.marshall@marquette.edu).

**Received** 9 June 2021

**Accepted** 14 June 2021

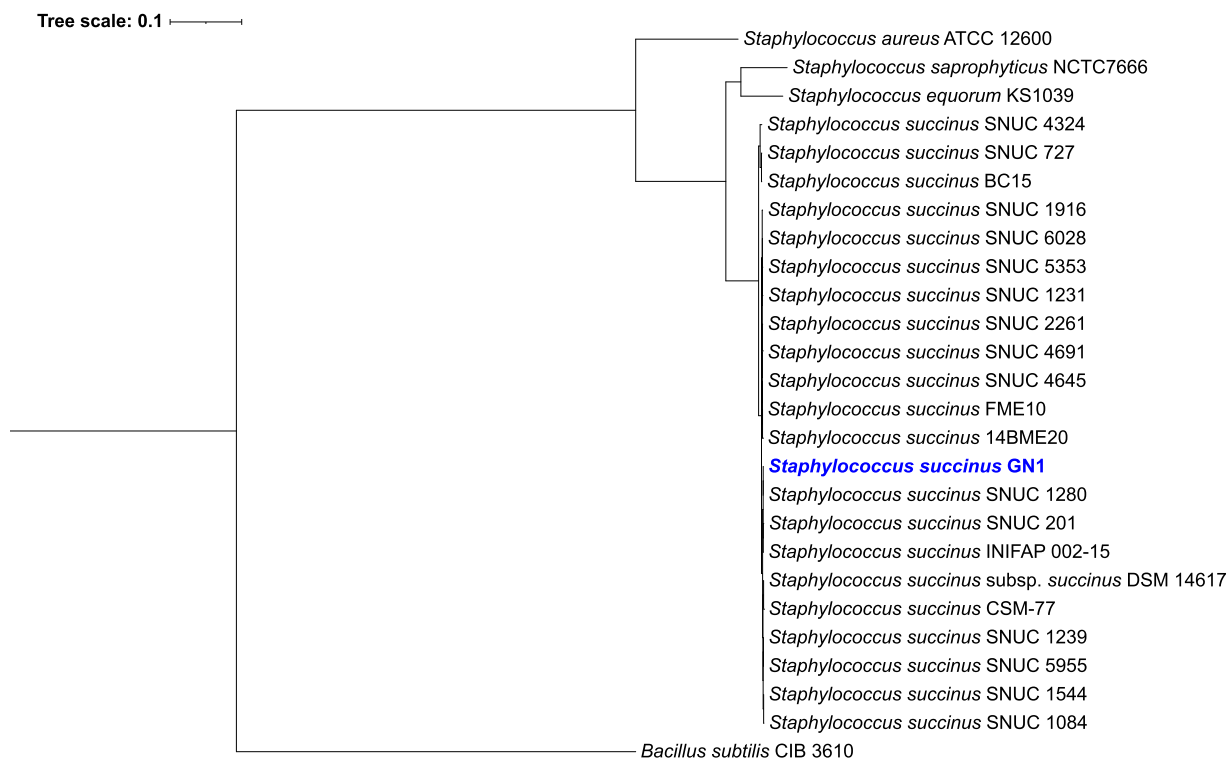
**Published** 8 July 2021

**TABLE 1** Genome assembly statistics

Statistic	Value
GC%	32.85
Genome size (bp)	2,837,157
No. of contigs	11
$N_{50}$ (bp)	1,426,624
No. of predicted coding genes	2,758

bacteria that aggregate in grapelike clusters (18), consistent with the morphology of our isolate as determined by Gram stain. We tested susceptibilities by disk diffusion to eight different antibiotics (neomycin, kanamycin, tetracycline, ciprofloxacin, bacitracin, streptomycin, chloramphenicol, and ampicillin) and found that isolate GN1 was susceptible to all antibiotics. We also tested *Staphylococcus succinus* strain GN1 on a limited number of carbon sources and found that it was able to utilize a pyruvate, lactose, or citrate source on either M9 minimal medium (pyruvate, lactose) or Simmons citrate agar (citrate).

**Data availability.** This genome is part of a larger collection of genomes deposited at the National Center for Biotechnology Information (NCBI) under the BioProject accession number [PRJNA665534](https://www.ncbi.nlm.nih.gov/bioproject/PRJNA665534). This is the repository for an undergraduate microbiology laboratory course. Students isolate a single colony, conduct physiological assays, sequence the genome, and conduct bioinformatics analyses. The sequences for isolate GN1 were deposited at the NCBI Sequence Read Archive under BioProject accession number [PRJNA665534](https://www.ncbi.nlm.nih.gov/bioproject/PRJNA665534), BioSample accession number [SAMN16337256](https://www.ncbi.nlm.nih.gov/biosample/SAMN16337256), and SRA accession number [SRR12762898](https://www.ncbi.nlm.nih.gov/sra/SRR12762898). The assembly accession number is [JAHKBG0000000001](https://www.ncbi.nlm.nih.gov/assembly/JAHKBG0000000001). A public narrative was created in KBase that assembled the genome, generated trees, and computed pangenome analyses (2). A static narrative can be found at <https://kbase.us/n/51640/53/>, which links to the full manipulatable data set.



**FIG 1** Phylogenetic tree generated using GToTree and IQ-TREE of closely related *Staphylococcus succinus* assemblies and related *Staphylococcus* species with *Bacillus subtilis* as the outgroup. Our isolate, GN1, is indicated in blue. The scale bar indicates substitutions per site.

## ACKNOWLEDGMENTS

We thank Stacia Peiffer for technical support.

This work was supported by the Marquette University Department of Biological Sciences educational funding.

## REFERENCES

1. Baym M, Kryazhinskiy S, Lieberman TD, Chung H, Desai MM, Kishony R. 2015. Inexpensive multiplexed library preparation for megabase-sized genomes. *PLoS One* 10:e0128036. <https://doi.org/10.1371/journal.pone.0128036>.
2. Arkin AP, Cottingham RW, Henry CS, Harris NL, Stevens RL, Maslov S, Dehal P, Ware D, Perez F, Canon S, Sneddon MW, Henderson ML, Riehl WJ, Murphy-Olson D, Chan SY, Kamimura RT, Kumari S, Drake MM, Brettin TS, Glass EM, Chivian D, Gunter D, Weston DJ, Allen BH, Baumohl J, Best AA, Bowen B, Brenner SE, Bun CC, Chandonia J-M, Chia J-M, Colasanti R, Conrad N, Davis JJ, Davison BH, DeJongh M, Devoid S, Dietrich E, Dubchak I, Edirisinghe JN, Fang G, Faria JP, Frybarger PM, Gerlach W, Gerstein M, Greiner A, Gurtowski J, Haun HL, He F, Jain R, Joachimiak MP, Keegan KP, Kondo S, et al. 2018. KBase: the United States Department of Energy Systems Biology Knowledgebase. *Nat Biotechnol* 36:566–569. <https://doi.org/10.1038/nbt.4163>.
3. 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>.
4. 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>.
5. Brettin T, Davis JJ, Disz T, Edwards RA, Gerdes S, Olsen GJ, Olson R, Overbeek R, Parrello B, Pusch GD, Shukla M, Thomason JA, III, Stevens R, Vonstein V, Wattam AR, Xia F. 2015. RASTtk: a modular and extensible implementation of the RAST algorithm for building custom annotation pipelines and annotating batches of genomes. *Sci Rep* 5:8365. <https://doi.org/10.1038/srep08365>.
6. Aziz RK, Bartels D, Best AA, DeJongh M, Disz T, Edwards RA, Formsma K, 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. <https://doi.org/10.1186/1471-2164-9-75>.
7. 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>.
8. Chaumeil P-A, Mussig AJ, Hugenholtz P, Parks DH. 2020. GTDB-Tk: a tool-kit to classify genomes with the Genome Taxonomy Database. *Bioinformatics* 36:1925–1927. <https://doi.org/10.1093/bioinformatics/btz848>.
9. Jain C, Rodriguez-R LM, Phillippy AM, Konstantinidis KT, Aluru S. 2018. High throughput ANI analysis of 90K prokaryotic genomes reveals clear species boundaries. *Nat Commun* 9:5114. <https://doi.org/10.1038/s41467-018-07641-9>.
10. Lee MD. 2019. GToTree: a user-friendly workflow for phylogenomics. *Bioinformatics* 35:4162–4164. <https://doi.org/10.1093/bioinformatics/btz188>.
11. Eddy SR. 2011. Accelerated profile HMM searches. *PLoS Comput Biol* 7:e1002195. <https://doi.org/10.1371/journal.pcbi.1002195>.
12. Edgar RC. 2004. MUSCLE: a multiple sequence alignment method with reduced time and space complexity. *BMC Bioinformatics* 5:113. <https://doi.org/10.1186/1471-2105-5-113>.
13. Capella-Gutiérrez S, Silla-Martínez JM, Gabaldón T. 2009. trimAl: a tool for automated alignment trimming in large-scale phylogenetic analyses. *Bioinformatics* 25:1972–1973. <https://doi.org/10.1093/bioinformatics/btp348>.
14. Nguyen L-T, Schmidt HA, von Haeseler A, Minh BQ. 2015. IQ-TREE: a fast and effective stochastic algorithm for estimating maximum-likelihood phylogenies. *Mol Biol Evol* 32:268–274. <https://doi.org/10.1093/molbev/msu300>.
15. Tange O. 2020. GNU Parallel 20201122 ('Biden'). <https://doi.org/10.5281/zenodo.4284075>.
16. Place RB, Hiestand D, Burri S, Teuber M. 2002. *Staphylococcus succinus* subsp. *casei* subsp. nov., a dominant isolate from a surface ripened cheese. *Syst Appl Microbiol* 25:353–359. <https://doi.org/10.1078/0723-2020-00130>.
17. Lambert LH, Cox T, Mitchell K, Rossello-Mora RA, Del Cueto C, Dodge DE, Orkand P, Cano RJ. 1998. *Staphylococcus succinus* sp. nov., isolated from Dominican amber. *Int J Syst Bacteriol* 48:511–518. <https://doi.org/10.1099/00207713-48-2-511>.
18. Foster T. 1996. *Staphylococcus*. In Baron S (ed), *Medical microbiology*, 4th ed. University of Texas Medical Branch at Galveston, Galveston, TX.