



Draft Genome Sequence of the Decomposition Fungus *Aquanectria penicillioides* Strain NNIBRFG19

Jaeduk Goh,^a Hye Yeon Mun,^a Yoosun Oh,^a Namil Chung^a

^aFungal Resources Research Division, Nakdonggang National Institute of Biological Resources, Sangju, Republic of Korea

ABSTRACT *Aquanectria penicillioides* is a common aquatic fungal species. Here, we report the 53.7-Mb draft genome sequence of *A. penicillioides* strain NNIBRFG19, which has an overall G+C content of 47.93%, comprising 13 scaffolds with an N_{50} value of 4.932 Mb.

Aquanectria penicillioides (basionym: *Flagellospora penicillioides*, teleomorph: *Nectria penicillioides*) is a common aquatic fungus and is one of the decomposition fungi of leaf litter in freshwater ecosystems (1, 2). Strain NNIBRFG19 was first reported as *A. penicillioides*, isolated from plant litter in Samcheok, Republic of Korea (3). Here, we report the first draft genome sequence of this fungus as one of the representative aquatic fungi in a freshwater environment.

A mycelial plug grown on potato dextrose agar (PDA; BD, Franklin Lakes, NJ) at 25°C, was used as the initial inoculum for the liquid culture. Genomic DNA and total RNA were extracted from mycelia grown in potato dextrose broth by shaking at 180 rpm for 1 week at 25°C under dark conditions using the DNeasy minikit (Qiagen, Valencia, CA) and the Easy-spin total RNA extraction kit (iNtRON, Sungnam, Republic of Korea), respectively. The genome sequence of NNIBRFG19 was obtained through a combination of four PacBio RS II single-molecule real-time (SMRT) cells (total 731,038 reads and 6,066,032,880 bp) and one paired-end (total 42,974,130 reads and 4,340,387,130 bp) and two mate pair libraries (total 93,212,966 reads and 9.41 Gb) on the HiSeq 2500 Illumina platform (Theragen Etex Bio, Suwon, Republic of Korea). The PacBio library and a paired-end Illumina library with an insert size of 400 bp were constructed after DNA fragmentation, A-tailing, phosphorylation, and adapter ligation. Two mate pair Illumina libraries with an insert size of 10 kb were constructed with the Nextera mate pair library preparation kit. An RNA-seq library was constructed by paired-end sequencing after fragmentation by random hexamer priming. Reads of RNA-seq were mapped to the assembled genome using TopHat v2 (4). The quality of the reads was checked using FastQC (<http://www.bioinformatics.babraham.ac.uk/projects/fastqc>). The reads were preprocessed by mHAP in Canu v1.6 (default parameters) for PacBio raw data (5), Trimmomatic v0.30 for paired-end sequencing (default parameters) (6), and NextClip v1.3 for mate pair sequencing (7) (min_length, 46; trim_ends, 0; removal of PCR duplication). The reads were assembled by Canu v1.6 for PacBio reads (default parameters) and SOAPdenovo v2 for Illumina reads (default parameters, k-mer frequency = 65) (8), and the two assemblies were merged by HaploMerger2 v20151124 with default parameters (9).

The draft genome of NNIBRFG19 consisted of 13 contigs with an N_{50} value of 4.932 Mb covering 341.72-fold of the genome (the average depth of the genome calculated using all reads). The total length of the assembled genome was 53,763,490 bp with a G+C content of 47.93%. The maximum and minimum scaffold lengths were 8,280,041 bp and 17,759 bp, respectively. A total of 13,658 genes were predicted by AUGUSTUS v3.2.1 using transcript alignment and protein-based alignment among similar species (10). The average length of

Citation Goh J, Mun HY, Oh Y, Chung N. 2019. Draft genome sequence of the decomposition fungus *Aquanectria penicillioides* strain NNIBRFG19. *Microbiol Resour Announc* 8:e01349-18. <https://doi.org/10.1128/MRA.01349-18>.

Editor Christina Cuomo, Broad Institute of MIT and Harvard University

Copyright © 2019 Goh 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 Jaeduk Goh, jdgo@nnibr.re.kr.

Received 28 September 2018

Accepted 27 November 2018

Published 10 January 2019

the predicted genes was 1,604 bp, and the average number of exons was 2.86. This draft genome sequence will support functional genetic research on the mechanism of leaf litter decomposition by aquatic fungi.

Data availability. The draft genome sequence of *A. penicillioides* strain NNIBRFG19 has been deposited in GenBank under accession number [PYYU00000000](https://doi.org/10.1093/bioinformatics/btt702). The SRA accession numbers are [SRR8067720](https://doi.org/10.1093/bioinformatics/btt702), [SRR8067732](https://doi.org/10.1093/bioinformatics/btt702), [SRR7800907](https://doi.org/10.1093/bioinformatics/btt702) and [SRR8204250](https://doi.org/10.1093/bioinformatics/btt702). This paper describes the first version of the genome for this strain.

ACKNOWLEDGMENT

This study was supported by The Survey and Discovery of Freshwater Bioresources (NNIBR2017, NNIBR2018) research program of the Nakdonggang National Institute of Biological Resources, funded by the Ministry of Environment of the Republic of Korea.

REFERENCES

1. Ingold CT. 1942. Aquatic hyphomycetes of decaying alder leaves. *Trans Br Mycol Soc* 25:339–417. [https://doi.org/10.1016/S0007-1536\(42\)80001-7](https://doi.org/10.1016/S0007-1536(42)80001-7).
2. Nikolcheva LG, Cockshutt AM, Bärlocher F. 2003. Determining diversity of freshwater fungi on decaying leaves: comparison of traditional and molecular approaches. *Appl Environ Microbiol* 69:2548–2554. <https://doi.org/10.1128/AEM.69.5.2548-2554.2003>.
3. Mun HY, Goh J, Oh Y, Chung N. 2016. New records of three aquatic fungi isolated from freshwater in Samcheok and Yeongju, Korea. *Kor J Mycol* 44:247–251.
4. Trapnell C, Pachter L, Salzberg SL. 2009. TopHat: discovering splice junctions with RNA-Seq. *Bioinformatics* 25:1105–1111. <https://doi.org/10.1093/bioinformatics/btp120>.
5. Koren S, Walenz BP, Berlin K, Miller JR, Bergman NH, Phillippy AM. 2017. Canu: scalable and accurate long-read assembly via adaptive k-mer weighting and repeat separation. *Genome Res* 27:722–736. <https://doi.org/10.1101/gr.215087.116>.
6. 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>.
7. Leggett RM, Clavijo BJ, Clissold L, Clark MD, Caccamo M. 2014. NextClip: an analysis and read preparation tool for Nextera Long Mate Pair libraries. *Bioinformatics* 30:566–568. <https://doi.org/10.1093/bioinformatics/btt702>.
8. Luo R, Liu B, Xie Y, Li Z, Huang W, Yuan J, He G, Chen Y, Pan Q, Liu Y, Tang J, Wu G, Zhang H, Shi Y, Liu Y, Yu C, Wang B, Lu Y, Han C, Cheung DW, Yiu SM, Peng S, Xiaoqian Z, Liu G, Liao X, Li Y, Yang H, Wang J, Lam TW, Wang J. 2015. Erratum: SOAPdenovo2: an empirically improved memory-efficient short-read de novo assembler. *Gigascience* 4:30. <https://doi.org/10.1186/s13742-015-0069-2>.
9. Huang S, Chen Z, Huang G, Yu T, Yang P, Li J, Fu Y, Yuan S, Chen S, Xu A. 2012. HaploMerger: reconstructing allelic relationships for polyploid diploid genome assemblies. *Genome Res* 22:1581–1588. <https://doi.org/10.1101/gr.133652.111>.
10. Stanke M, Steinkamp R, Waack S, Morgenstern B. 2004. AUGUSTUS: a Web server for gene finding in eukaryotes. *Nucleic Acids Res* 32:W309–W312. <https://doi.org/10.1093/nar/gkh379>.