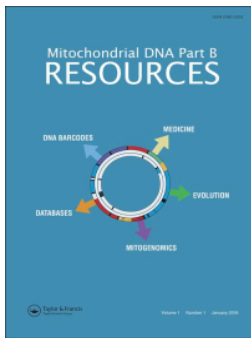


The complete mitochondrial genome of the white-rot fungus *Phanerochaete sordida* YK-624

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MITOGENOME ANNOUNCEMENT



The complete mitochondrial genome of the white-rot fungus *Phanerochaete sordida* YK-624

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ABSTRACT

The white-rot fungus *Phanerochaete sordida* (Karsten) Eriksson and Ryvarden 1978 is known for its excellent ligninolytic activity and capability to degrade various recalcitrant organic pollutants. In this study, we determined the complete mitochondrial genome sequence of *P. sordida* YK-624. The mitochondrial genome is 129,567 bp in length with a GC content of 28.9%, and contains two ribosomal RNA genes, 26 transfer RNA genes, and 50 open reading frames, including 14 conserved proteins. Phylogenetic analysis based on the mitochondrial genome confirmed that *P. sordida* belongs to the family Phanerochaetaceae in the order Polyporales, and showed the general phylogenetic relationships.

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KEYWORDS

Ligninolytic fungi;
mitochondrial genome;
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Introduction

White-rot fungi are a type of wood-decay fungi that can degrade lignin in wood. *Phanerochaete sordida* YK-624 (ATCC 90872) collected in Yakushima, Kagoshima, Japan (30°18'17"N, 130°34'30"E) is a typical saprotrophic white-rot basidiomycetous fungus belonging to the family Phanerochaetaceae that shows excellent lignin-degrading activity (Hirai et al. 1994). The fungus has been reported to have biodegradative activity for various organic pollutants, such as neonicotinoid insecticides (Mori et al. 2017; Wang et al. 2019). Among Phanerochaetaceae fungi, the mitochondrial DNA (mtDNA) sequence of only *Phanerochaete carnososa* has been available on GenBank to date. Although fungi belonging to the genus *Phanerochaete* are among the most intensively studied white-rot fungi, little information is available on their genome. Therefore, we newly determined the mtDNA sequence of *P. sordida*, and analyzed its phylogenetic relationships with other basidiomycetes. The mitogenome data provided in the present study will be useful for the investigation of detailed phylogenetic relationships of Phanerochaetaceae fungi.

Materials and methods

P. sordida YK-624 which was isolated from decaying wood collected from Yakushima, Japan, was deposited at American Type Culture Collection (ATCC; <https://www.atcc.org/>, tech@atcc.org) as the voucher number ATCC 90872. The genomic DNA of this fungus was extracted using Isoplant II (Nippon Gene Co., Ltd., Tokyo, Japan). Purified genomic DNA by ethanol precipitation was fragmented by using a Focused-ultrasonicator M220 (Covaris Inc., Woburn, MA), then the

genomic library was prepared using a TruSeq DNA PCR-free Library Preparation Kit (Illumina, San Diego, CA), and sequenced using MiSeq (Illumina, San Diego, CA) (Mori et al. 2021). Raw sequence reads (DRR311115 and DRR311116) were cleaned using Trimmomatic ver. 0.38 (Bolger et al. 2014) as described previously (Mori et al. 2021), and the resulting 8,146,783 read pairs totaling approximately 4,157 Mb were assembled using NOVOPlasty ver. 4.3.1 (Dierckxsens et al. 2017) with the nucleotide sequence of the cytochrome c oxidase subunit 1 gene from *P. carnososa*

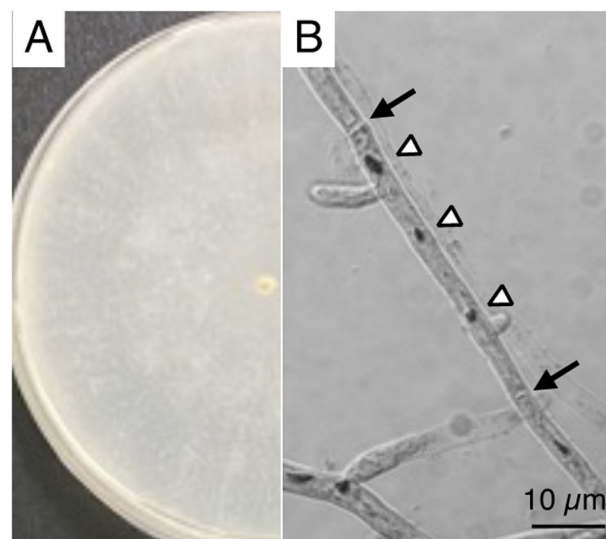


Figure 1. *P. sordida* YK-624 mycelium and hyphae observation. (A) Mycelia were grown on potato dextrose agar medium (i.d. 9 cm). (B) Hyphae were stained by HCl-Giemsa method following method described in previous study (Mori et al. 2020).

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(GenBank accession no. MT090080.1, nt. 1–29,351) as the seed sequence. The obtained circular mtDNA was 129,567 bp in length with a GC content of 28.9% (GenBank accession no. LC707859). DNA annotation was conducted using the MFannot tool (<http://megasun.bch.umontreal.ca/cgi-bin/mfannot/mfannotInterface.pl>), then manually curated. The ribosomal RNA genes were predicted by aligning the mtDNA sequences to those of *P. carnosa* (NC_057606). Subsequently, a phylogenetic analysis based on the 14 conserved

mitochondrial proteins in 14 Agaricomycetes and an Ustilaginomycetes was performed. The concatenated amino acid sequences were aligned using MAFFT ver. 7.427 (Katoh and Standley 2013), and the phylogenetic tree was constructed by the neighbor-joining method (Saitou and Nei 1987).

Results

A white-rot basidiomycetes fungi *P. sordida* YK-624 forms white-colored filamentous fungi, and exhibits multi-nuclei hyphae without clamp connections (Figure 1) (Mori et al. 2020). Overall, 78 genes, including 50 putative protein-coding genes, two ribosomal RNAs (rnl and rns), and 26 tRNAs (for all 20 standard amino acids) were predicted in the mtDNA of *P. sordida* YK-624 (Figure 2). The 50 protein-coding genes encoded 14 conserved mitochondrial proteins (*cox1–3*, *cob*, *nad1–6*, *nad4L*, *atp6*, *atp8*, and *atp9*), DNA polymerase type B (*dpo*), ribosomal protein S3 (*rps3*), and 34 hypothetical proteins, most of which showed similarities to LAGLIDADG homing endonuclease. The phylogenetic tree showed that *P. sordida* is closely related to *P. carnosa* belonging to the family Phanerochaetaceae in the order Polyporales (Figure 3). The phylogenetic relationships between the mtDNA of *P. sordida* and other Agaricomycetes fungi are consistent with those based on nuclear genes.

Discussion

In the present study, complete mitochondrial genome sequence of *P. sordida* YK-624 was newly determined. Although Phanerochaetaceae family included well studied groups like *Phanerochaete* and *Bjerkandera* fungi, classification study is not sufficiently progressed. In particular, the genus *Phanerochaete* is often very similar in morphology, and ITS (internal transcribed sequences) in the closely related species, so classification between each species is still uncertain

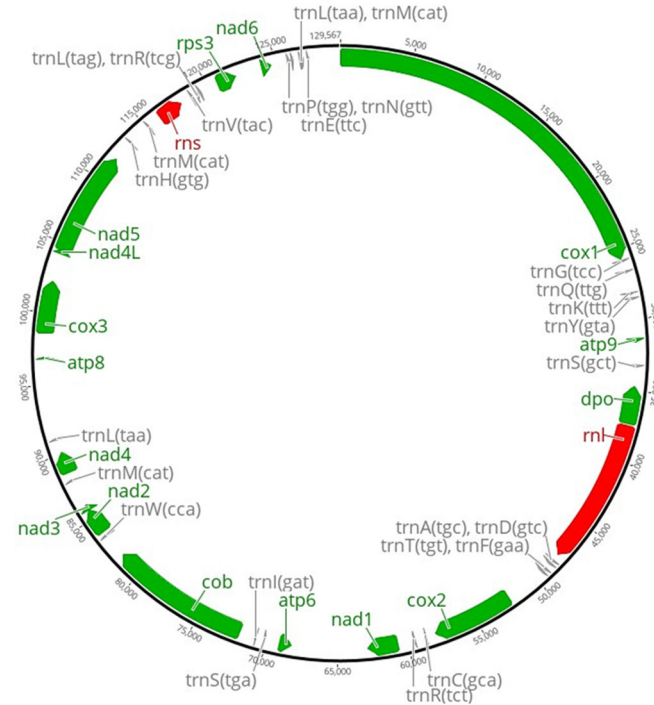


Figure 2. Mitochondrial DNA of *P. sordida* YK-624. The map of the 129,567 bp shows 14 conserved mitochondrial proteins (*cox1–3*, *cob*, *nad1–6*, *nad4L*, *atp6*, *atp8*, and *atp9*), DNA polymerase type B (*dpo*), ribosomal protein S3 (*rps3*), and tRNAs. Other 34 hypothetical proteins are not shown.

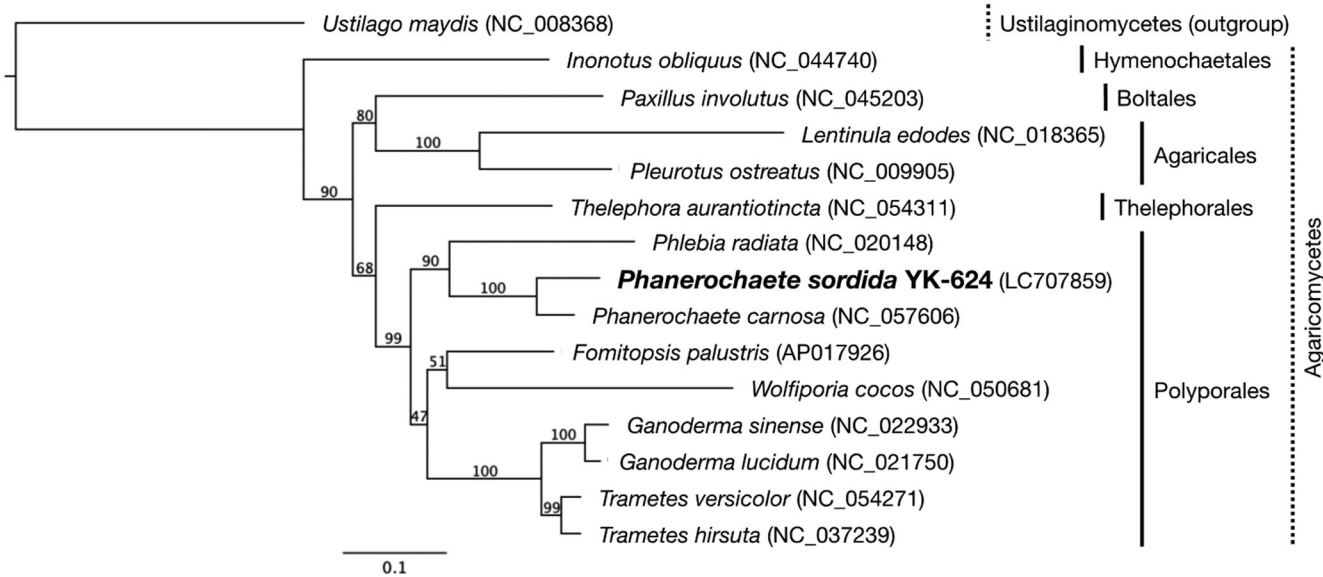


Figure 3. Phylogenetic relationships among 14 Agaricomycetes and an Ustilaginomycetes. The phylogenetic analysis was conducted by the neighbor-joining method based on the concatenated amino acid sequences of 14 conserved protein-coding genes (*atp6*, *atp8*, *atp9*, *cob*, *cox1–3*, *nad1–6*, and *nad4L*). *Ustilago maydis* (NC_008368) served as an outgroup. The bootstrap support values are shown at each node. The scale bar indicates the number of substitutions per site.

(Floudas and Hibbett 2015). Although mitochondrial genome information on the genus *Phanerochaete* is currently very limited, the present study will assist in future taxonomic studies using mitochondrial genome information.

Ethics statement

Ethical approval does not apply to this paper because the research material is mushrooms (fungus), and not human, animals, plants, or threatened/endangered species that require ethical approval.

Author contributions

Conceived and designed the research: TM and HH. Performed the experiments and data analysis: TM and HD. Wrote the manuscript: TM and HH. Contributed to data interpretation, discussions, and critically reviewed the manuscript: HK. All authors reviewed the manuscript and approved the final version to be published.

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Disclosure statement

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Data availability statement

The genome sequence data that support the findings of this study are openly available in GenBank of NCBI (<https://www.ncbi.nlm.nih.gov>) under the accession number LC707859. The associated BioProject, SRA, and BioSample accession numbers are PRJDB11801, DRR311115, DRR311116, and SAMD00328051, respectively.

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