The complete mitochondrial genome of medicinal fungus *Taiwanofungus camphoratus* reveals gene rearrangements and intron dynamics of *Polyporales*

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*Taiwanofungus camphoratus* is a highly valued medicinal mushroom that is endemic to Taiwan, China. In the present study, the mitogenome of *T. camphoratus* was assembled and compared with other published *Polyporales* mitogenomes. The *T. camphoratus* mitogenome was composed of circular DNA molecules, with a total size of 114,922 bp. Genome collinearity analysis revealed large-scale gene rearrangements between the mitogenomes of *Polyporales*, and *T. camphoratus* contained a unique gene order. The number and classes of introns were highly variable in 12 *Polyporales* species we examined, which proved that numerous intron loss or gain events occurred in the evolution of *Polyporales*. The Ka/Ks values for most core protein coding genes in *Polyporales* species were less than 1, indicating that these genes were subject to purifying selection. However, the *rps3* gene was found under positive or relaxed selection between some *Polyporales* species. Phylogenetic analysis based on the combined mitochondrial gene set obtained a well-supported topology, and *T. camphoratus* was identified as a sister species to *Laetiporus sulphureus*. This study served as the first report on the mitogenome in the *Taiwanofungus* genus, which will provide a basis for understanding the phylogeny and evolution of this important fungus.

*Taiwanofungus camphoratus*, belonging to *Polyporales*, Basidiomycota, is an endemic mushroom of Taiwan, China, which is restricted to the endemic aromatic tree *Cinnamomum kanehirai*. In the last few decades, *T. camphoratus* was included in folk medicine in Asia and has shown remarkable effectiveness in the treatment of inflammatory disorders, cancer, hypertension and hepatitis. The total market value of *T. camphoratus* products has grown rapidly, with market value reported to exceed $100 million annually in 2014. Many active secondary metabolites were isolated from the mycelium or fruiting bodies of *T. camphoratus*, such as triterpenoids, lipids, and benzenoids, showing excellent anti-inflammatory, hepatoprotective, antioxidant, radioprotective, and chemoprevention activities. *T. camphoratus* shares many common characters with *Antrodia* and *Antrodiella*, which caused its taxonomic confusion in the past. Phylogenetic analysis based on LSU rDNA sequence indicated that *T. camphoratus* should be proposed as a new genus of lignicolous polypore. The genome of *T. camphoratus* has been published to reveal the sexual development and metabolic synthesis of *T. camphoratus*. However, the characterization of the mitogenome of *Taiwanofungus camphoratus* has not been conducted to date.

Mitochondria generate most of the cell’s supply of adenosine triphosphate (ATP) in eukaryotes. Most eukaryotes have their own mitochondrial genomes (mitogenomes), which are believed to be derived through endosymbiosis from the ancestral alpha-proteobacterium. As the “second genome” of eukaryotes, mitogenomes have many characteristics different from nuclear genomes. For example, each eukaryotic cell contains a large number of mitochondrial organelles, and each mitochondrial organelle has multiple mitochondrial genomes.

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In addition, its rapid evolutionary rate, and multiple available molecular markers have made the mitochondrial genome a powerful tool for studying phylogeny and population genetics. With the rapid development of next-generation sequencing technology, more and more mitochondrial genomes have been obtained. So far, more than 10,000 complete mitochondrial genomes have been released in the NCBI reference database, including animal, plant and fungal mitogenomes. These available mitogenomes promote the phylogeny and taxonomy of eukaryotes. However, compared with its animal counterparts, the mitochondrial genome of fungi has been less studied. As an important group of eukaryotes, fungus is widely distributed in the world. It is estimated that more than 2.2 million of fungal species exist in nature, which play an important role in maintaining the carbon and nitrogen cycle in nature, maintaining the healthy development of forest ecosystems, and providing food sources for human beings. However, fewer than 800 fungal mitogenomes are available, with available mitogenomes of Basidiomycetes are even less (< 130). The few available mitogenomes limit our overall understanding of evolution, phylogeny, and taxonomy of the fungal lineages, because phylogenies based on mitochondrial genes and nuclear genes are not always consistent.

Up to now, only 11 complete mitogenomes of Polyporales species have been published, including seven species in Ganoderma genus, one in Trametes, one in Phlebia, one in Laetiporus, and one in Fomitopsis. The sizes of these mitogenomes varied greatly, ranging from 60,635 to 156,348 bp. This is thought to be due to variations of intron numbers. Introns in the mitogenome of fungi can be divided into two groups, namely the group I and the group II. Group II intron excision occurs in the absence of GTP and involves the formation of a lariat, which were different from group I intron. These introns were found distributed in the cox, cob, nad and tRNA genes of the fungal mitogenomes, which could modify the organization and size of mitogenome. In addition, a variety of homing endonuclease genes have been found in fungal introns, such as the LAGLIDAG endonuclease and GIY endonucleases, the products of which have homing endonuclease activities. However, the origin, evolution, and functions of the mitochondrial introns in the order Polyporales are still unknown.

The mitogenomes of fungi are significantly different from that of other eukaryotes. The genome size, gene arrangement, repetitive sequences, introns, intergenic regions, and open reading frames (ORFs) of fungal mitogenomes varied greatly between species and even between intraspecific species. Despite the large variations, most Basidiomycete mitogenomes were found containing 14 core protein coding genes (atp6, atp8, atp9, cox1, cox2, cox3, cob, nad1, nad2, nad3, nad4, nad4L, nad5, and nad6) for energy metabolism, one rps3 gene for transcriptional regulation, 22–36 tRNA genes, and two rRNA genes. Repeated sequences in the mitogenome, gene arrangement, and structures of tRNAs can provide important information to reveal the evolutionary and phylogenetic relationships of fungi.

In the present study, the mitogenome of T. camphoratus were assembled and annotated. The aims of this study are (1) to reveal variations and conservation of gene content, genome organization, and gene order between T. camphoratus mitogenome and other Polyporales species; (2) to provide insights into the evolution and dynamic changes of introns in Polyporales mitogenomes; (3) to reveal the phylogenetic status of T. camphoratus among various Agaricomycetes species based on the combined mitochondrial gene set. The mitogenome of T. camphoratus will allow further study of the taxonomy, phylogenetics, and evolutionary studies of this important medical fungus and other closely related species.

Results

Genome features and protein coding genes. The complete mitogenome of T. camphoratus was composed of circular DNA molecules, with the total size of 114,922 bp (Fig. 1). GC content of the T. camphoratus mitogenome was 26.01%. The AT skew was negative in the mitogenome of T. camphoratus, where the GC skew was positive (Table S1). Fifteen core protein coding genes were detected in the T. camphoratus mitogenome, including atp6, atp8, atp9, cob, cox1, cox2, cox3, nad1, nad2, nad3, nad4, nad4L, nad5, and nad6 for energy metabolism, and one rps3 gene for transcriptional regulation. All of these introns belong to the group I. Twenty-three intronic ORFs were found located in these introns, which encoded LAGLIDAG and GIY-YIG homing endonucleases.

RNA genes in the T. camphoratus mitogenome. Two tRNA genes were detected in the T. camphoratus mitogenome, namely the large subunit ribosomal RNA (rnl), and the small subunit ribosomal RNA (rns) (Table S2). The mitogenome of T. camphoratus contained 27 tRNA genes, which were folded into classical cloverleaf structures (Fig. 2). The mitogenome of T. camphoratus contained 2 tRNAs with different anticodons coding for serine, arginine, leucine and 3 tRNAs with different anticodons coding for glutamate. In addition, the T. camphoratus mitogenome contained 3 tRNAs with the same anticodon that coded for methionine. The length of individual tRNAs ranged from 71 to 86 bp, which was mainly due to the variation of extra arm. A ribonuclease P RNA (rnpB) gene was found in the T. camphoratus mitogenome, with the length of 329 bp.

Overlapping nucleotides and mitogenome composition. Only one overlapping nucleotide was detected in the mitogenome of T. camphoratus, which was located across the neighboring genes nad4L and nad5 (1 bp) (Table S2). We detected 41,887 bp intergenic sequences in the mitogenomes of T. camphoratus. The length of these intergenic sequences ranged from 0 to 2903 bp, and the longest intergenic sequence was located between the trnW and orf292 gene. Intergenic regions accounted for the largest proportion of the T. camphoratus mitogenome, reaching 36.45% (Fig. 3), which showed that the mitogenome of T. camphoratus had a relatively relaxed structure. The intronic region occupied the second largest proportion of the T. camphoratus
mitogenome, accounting for 29.94%. The protein coding region accounted for 27.56% of the *T. camphoratus* mitogenome. The RNA region (rRNA + tRNA + rnpB) accounted for 6.05% of the entire mitogenome.

**Codon usage analysis.** ATG was the most commonly used start codon for the 15 core PCGs in the 12 *Polyporales* species we examined (Table S3). Eleven of the 15 core PCGs used ATG as start codons in all the 12 *Polyporales* mitogenomes we detected. GTG was the most commonly used start codon in *cob* gene of the 12 *Polyporales* mitogenomes. In addition, ATA and TTG were also used as start codons in *cox2*, *nad1* and *nad6* gene of some *Polyporales* species. TAA was most commonly used as stop codons of core PCGs in 12 *Polyporales* species, followed by TTG and AGT. We found that all core PCGs in *T. camphoratus* used ATG as start codons and TAA as stop codons.

Codon usage analysis indicated that the most frequently used codons in the *T. camphoratus* mitogenome were TTT (for phenylalanine; Phe), AAA (for lycine; Lys), TTA (for leucine; Leu), AAT (for asparagine; Asn),
ATT (for isoleucine; Ile), and TAT (for tyrosine; Tyr) (Fig. 4 and Table S4). The frequent use of A and T in codon contributed to the high AT content in the *T. camphoratus* mitogenome (average: 73.99%).

**Repetitive sequences in the mitogenome.** Comparing the whole mitogenome of *T. camphoratus* with itself via BLASTn search, we identified 45 repeat sequences in the mitogenome of *T. camphoratus* (Table S5). The length of these repeat sequences ranged from 28 to 902 bp, with pair-wise nucleotide similarities ranging from 74.72 to 100%. The largest repeats were observed in the third introns and fourth introns of the *rnl* gene. The second largest repeats were found located in the intergenic region between orf145 and *rnl*, as well as in intergenic region between orf388 and orf245, with each repeating sequence 840 bp long. Repeat sequence accounted for 9.44% of the *T. camphoratus* mitogenome.

Using the Tandem Repeats Finder, we identified 55 tandem repeats in the *T. camphoratus* mitogenome (Table S6). The longest tandem repeat sequence was observed in the intergenic region between *nad4* and *trnA*, comprising 79 bp. Most of the tandem repeat sequences were duplicated once or twice in the *T. camphoratus* mitogenome, with the highest replication number 15. Tandem repeat sequences accounted for 2.04% of the *T. camphoratus* mitogenome. REPuter identified 27 forward, 7 palindromic, and 16 reverse repeats in the mitogenome of *T. camphoratus*, accounting for 6.33% of the entire mitogenome (Table S7).

**Comparative genome analysis and gene rearrangement.** The mitogenome of *T. camphoratus* was the fourth largest among the published mitogenomes in Polyporales, which was smaller than *Phlebia radiata*26, *G. calidophilum*, and *G. applanatum*, and larger than *Trametes cingulata*35, *Fomitopsis palustris*38, *Laetiporus sulphureus*27 and other *Ganoderma* spp. species (Table S1). The GC content of the *T. camphoratus* mitogenome was

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**Figure 2.** Putative secondary structures of the 27 tRNA genes identified in the mitochondrial genome of *Taiwanofungus camphoratus*. All genes are shown in order of occurrence in the mitochondrial genome of *Taiwanofungus camphoratus*, starting from *trnV*. tRNA structures were determined using MITOS.
**Figure 3.** The protein-coding, intronic, intergenic, and RNA gene region proportions of the entire mitochondrial genome of *Taiwanofungus camphoratus*.

**Figure 4.** Codon usage in the mitochondrial genome of *Taiwanofungus camphoratus*. Count of codon usage is plotted on the y-axis.
in the middle among all Polyporales mitogenomes. Of the 12 Polyporales we tested, only T. camphoratus and L. sulphureus had negative AT skews, while the others were positive. The GC skew of the T. camphoratus mitogenome was positive, just like most of the Polyporales. Thirty-four non-intronic ORFs and 24 introns were found in the T. camphoratus mitogenome, which were smaller than the 79 non-intronic ORFs and 36 introns in the P. radiata mitogenome that made P. radiata the largest mitochondrial genome in the Polyporales. All 12 Polyporales mitogenomes detected contain 2 rRNA genes. The number of tRNA genes in the mitogenomes of Polyporales ranged from 25 to 29.

Genomic collinearity analysis showed that the mitogenome of T. camphoratus had 28 homologous regions with other Polyporales mitogenomes (Fig. 5). The relative positions of these homologous regions were highly variable between Polyporales species, suggesting that frequent gene rearrangements had occurred during the mitogenome evolution of Polyporales. In addition, we found that the gene arrangement was highly conserved in the genus Ganoderma and highly variable at family or order levels. Gene arrangement analysis involving all core PCGs, rRNA genes and tRNAs indicated that T. camphoratus had a unique gene order, which showed its unique evolutionary characteristics in Polyporales species.

Dynamics of introns in cox1 and cob genes of Polyporales. The number and size of introns were highly variable in mitogenomes of Polyporales, which promoted the size and organization dynamic changes of Polyporales. A total of 271 introns were detected in 12 published mitogenomes of Polyporales, with individual species containing 6–36 introns. These introns distributed in several host genes, including cox1, cob, rnl, cox2, and nad1. The cox1 gene contained 130 introns, accounting for 47.97% of the total number of introns, followed by cob gene, which contained 48 introns, accounting for 17.71% of the total number of introns. According to previous studies, each intron was considered as a mobile genetic element. Group I introns in fungi could be classified into different position classes (Pcls) according to their insertion site in the coding region of cox1 or cob gene. The same Pcls were considered as orthologous, and had high nucleotide similarities. Different Pcls had lower sequence similarities, and with non-homologous homing endonucleases. In the present study, we classified group I introns in cox1 genes of the 12 Polyporales species into 26 Pcls according to methods described by Férandon et al. As shown in Fig. 6, the same letters denoted the same Pcls. T. cingulata contained the most Pcls (15) in the cox1 gene, followed by G. calidophilum and G. applanatum. G. lucidum contained the least Pcls, suggesting that intron loss had occurred in the evolution of G. lucidum. Pcls P was the most common intronic classes in the cox1 genes of Polyporales, which presented in 10 of the 12 Polyporales species. Pcls T and D were also widely distributed in cox1 genes of Polyporales species, which existed in 9 and 8 Polyporales species, respectively. In addition, some rare introns were found in the cox1 gene of Polyporales, e.g. Pcl Z, which was observed only in the cox1 gene of L. sulphureus. However, this Pcls has been observed in the mitogenome of Agaricus bisporus from Agaricales, suggesting potential horizontal gene transfer events between the two species. Pcl R was observed only in P. radiata from Polyporales, but was observed in cox1 gene of Rhizophydium sp. 136. In addition, Pcl Y was also considered to be a rare intron class in Polyporales, which only existed in T. camphoro-
**Figure 6.** Pcl information of *cox1* gene (a) and *cob* gene (b) of the 12 *Polyporales* species. The same Pcl (orthologous intron) is represented by the same letter or the same roman numeral. The phylogenetic positions of 12 *Polyporales* species were established using the Bayesian inference (BI) method and Maximum Likelihood (ML) method based on 15 concatenated mitochondrial core proteins and 2 rRNA genes. UN indicates that the intron is different from the known introns<sup>44</sup> in insertion site and sequence similarity. The II in the figure above shows that the intron belongs to the group II intron. Species ID are shown in Supplementary Table S8.

tus and *G. calidophilum*. The Pcl Y was also observed in distant species *Chlorokybus atmophyticus*. In addition, three unknown Pcls were found in the mitogenome of *Polyporales* with no significant nucleotide similarities to reported Pcls.

The introns of *cob* genes in *Polyporales* were classified into 18 Pcls according to the insertion sites in the coding region of *cob* gene (Fig. 6). *T. camphoratus*, *G. applanatum* and *G. calidophilum* contained the most Pcls in *cob* gene. The *cob* gene of *T. cingulata* and *G. lucidum* was intronless. Pcls VI and VII were considered to be the common Pcls in *Polyporales*, which existed in at least 6 of the 12 *Polyporales* species detected. While Pcls IV, VIII, IX, XII, XV, and XIX only existed in one of 12 species, which were considered to be rare introns in the *cob* gene. However, these rare Pcls showed highly nucleotide similarities with distant species, suggesting frequent gain or loss of *cob* introns in *Polyporales*.

**Evolution rate of core genes and phylogenetic analyses.** Of the 15 core PCGs detected, the *rps3* gene had the highest mean K2P genetic distance among the 12 *Polyporales* species, followed by *nad3* (Fig. 7). The pairwise K2P genetic distance of *rps3* and *nad3* gene between *Polyporales* species varied largely. The mean genetic distance of *nad4L* gene between the 12 *Polyporales* species was the smallest among the 15 core PCGs detected, indicating that this gene was highly conserved across the mitogenomes. The *nad3* gene had the highest nonsynonymous substitution rate (Ka) of the 15 core PCGs detected, while *atp9* had the lowest rate. The highest synonymous substitutions rate (Ks) was observed in the *nad3* gene, while *atp9* exhibited the lowest Ks value of the 15 PCGs detected. The Ka/Ks values for most core PCGs were less than 1, indicating that these genes were subject to purifying selection. However, the Ka/Ks value of *rps3* gene was observed more than 1 between some species, such as between *T. camphoratus* and *L. sulphureus*, between *T. camphoratus* and *F. palustris*, as well as between *G. sinense* and *G. applanatum*, indicating that *rps3* gene was under positive selection pressure in some *Polyporales* species.

Phylogenetic analyses based on the combined mitochondrial gene set (15 core PCGs + 2 rRNAs), using bayesian inference (BI) and maximum likelihood (ML) methods with the GTR+I+G model of nucleotide substitution, yielded identical and well-supported tree topologies (Fig. 8). All major clades of the trees were found with good
support (BPP = 1.00 and BS ≥ 75). Based on the phylogenetic analyses, the 25 Agaricomycetes species could be divided into four major clades, corresponding to the orders Polyporales, Agaricales, Russulales, and Cantharellales (Table S8). The 12 Polyporales species could be divided into three groups; one group was composed of one species in the Phlebia genus; the second group was composed of one species in the Trametes genus and seven species in the Ganoderma genus; the third group was recovered as (F. palustris + (T. camphoratus + L. sulphureus)). T. camphoratus was identified as a sister species to L. sulphureus. Phylogenetic studies showed the evolutionary status of the T. camphoratus in the Agaricomycetes class. The results indicated that the combined mitochondrial gene set was suitable as reliable molecular marker for analysis of the phylogenetic relationships among Agaricomycetes species.

Discussion

As the ‘second genome’ of eukaryotes, mitochondrial genome plays an important role in aging, death, disease occurrence and stress resistance of eukaryotes. Mitochondrial genome organization, core protein coding genes, repetitive sequences, gene arrangement, the number and structure of tRNAs, and open reading frames provided rich genetic information for understanding the genitics and evolution of eukaryotes. Mitochondrial genomes have been extensively studied in animals. However, as an important group of eukaryotes, the mitochondrial genomes of fungi have been less studied, especially in Agaricomycetes. As the largest mushroom-forming fungal group, less available Agaricomycetes mitochondrial genomes have limited our understanding of the origin, evolution, and phylogenetic relationships of mushroom-forming fungi. The mitochondrial genome of fungi was thought to have a moderate mutation rate, intermediating to that of plants (highest mutation rate) and animals (lowest mutation rate). It was reported that the mitogenome size, gene arrangement, and repetitive sequences of fungi varied greatly between and within species. The largest known Agaricomycetes mitochondrial genome was Rhizoctonia solani, which was 235.85 kb long and contained 127 PCGs, 2 rRNA genes and 26 tRNA genes. In Polyporales order, the mitogenome size varied greatly, ranging from 60,635 to 156,348 bp. Accordingly, the non-intronic ORFs ranged from 15 to 79; intron numbers ranged from 6 to 36, and the number of tRNA genes ranged from 25 to 29. Variations in the number and size of these genes contributed to huge variations in the size of the mitochondrial genomes in Polyporales. Previous studies showed that the size variation of mitogenome was

Figure 7. Genetic analysis of 15 protein coding genes conserved in 12 Polyporales mitogenomes. K2P the Kimura-2-parameter distance, Ka the mean number of nonsynonymous substitutions per nonsynonymous site, Ks the mean number of synonymous substitutions per synonymous site.
closely associated with intron number variation\(^{33,44}\), which was consistent with this study. In the present study, the *T. camphoratus* genome size, open reading frames, GC content, GC skew and AT skew were not consistent with other *Polyporales* species, indicating the unique evolutionary characteristics of *T. camphoratus* mitogenome.

In the present study, we found that most introns in *Polyporales* belonged to the group I, which was different from plants\(^{61}\). A total of 271 introns were found in the 12 *Polyporales* species, most of which were distributed in the protein coding genes *cox1* and *cob*. The dynamic changes of these introns contributed to variations of mitochondrial genome size and organization. Introns were considered to be a mobile genetic element\(^{65}\), while homologous introns had the same insertion site in the coding region of PCGs\(^{43}\). The same Pcls had high sequence similarities, while different Pcls showed low similarities in nucleotide sequences. In the present study, the classes and number of Pcls were inconsistent in *Polyporales*. Some Pcls were widely distributed in *Polyporales* species, suggesting that these Pcls may be obtained from the common ancestor. However, some Pcls, e.g. Pcls Z and R, were observed only in one or two *Polyporales* species, which were also found in distant species such as *A. bisporus* and *C. atmophyticus*\(^{43}\), suggesting the possibility of horizontal gene transfer events. In general, the number and classes of introns were highly dynamic changes in the *Polyporales* species, which proved that numerous intron loss
or gain events occurred in the evolution of Polyporales. The result was consistent with previous studies that the fungal group I introns appears to result from numerous losses and gains of the mobile genetic elements.48, 63, 66, 67.

The arrangement of mitochondrial genes can also serve as an important reference for revealing phylogenetic relationships among species.69–71 In this study, we found that the arrangement of mitochondrial genes was highly variable in Polyporales. Gene orders between Ganoderma species were found highly conserved, while large-scale gene rearrangements were observed at family levels. Gene order analysis revealed that large-scale gene rearrangements occurred in T. camphoratus mitogenome compared with other Polyporales, involving protein-coding genes, tRNA genes, and rRNA genes, resulting in unique gene arrangement in T. camphoratus mitogenome. Previous studies have shown that the accumulation of repetitive sequences in the mitochondrial genome of fungi contributed to gene recombination of the fungal mitogenome, which in turn led to the rearrangement of mitochondrial genes.83 In this study, high proportion of repeats (9.44%) were observed in the mitogenome of T. camphoratus. It is reasonably speculated that the accumulation of repetitive sequences in the mitogenome of T. camphoratus led to gene rearrangement in T. camphoratus. The effect of the repeat sequences accumulation in T. camphoratus mitogenome on the evolution and variation of T. camphoratus mitogenome needs to be further revealed.

Mitogenomes are thought to be derived from the common ancestral alpha-proteobacterium through endosymbiosis.72, 73. During evolution, most mitochondrial genes have been transferred into the nuclear genome, and this phenomenon was considered with multiple advantages.74 However, fungal mitogenomes also retained a number of genes for energy metabolism and transcriptional regulation, including atp6, atp8, atp9, cob, cox1, cox2, cox3, nad1, nad2, nad3, nad4, nad4L, nad5, nad6, and rps3, which we call the core PCGs. We found that the core PCGs of Polyporales varied greatly in base composition and gene length. The effect of these variations on fungal adaptation to environment and energy metabolism remains unknown. In addition, a series of ORFs were found in the T. camphoratus mitogenome, which show low sequence similarities to known proteins in public database, suggesting that there are many unknown protein-coding genes in the T. camphoratus mitogenome that need to be revealed, which will facilitate understanding of the origin, evolution and function of fungal mitogenomes. Genetic distance analysis showed that nad4L gene was highly conserved among Polyporales species, while rps3 gene was highly variable in Polyporales. We found that most of the core PCGs in Polyporales were subjected to purifying selection. However, rps3 gene was under strong positive selection pressure between some species, which may be to better adapt to their lifestyles or surrounding environment.49.

Because of the limited and confusing macroscopic and microscopic morphological characteristics of macrofungi, it is difficult to precisely classify fungal species and subspecies, which limits the development and utilization of fungi. The introduction of molecular markers promotes the taxonomy of fungi. Mitochondrial genomes have been widely used in phylogenetics, evolutionary and population genetics because of rapid evolution rates and many available molecular markers.5, 75, 76. Mitochondrial cox1 gene and rRNA genes have been widely used in phylogenetic studies of animals.75, 76. However, there are few reports on the application of mitochondrial genome in fungal taxonomy or phylogenetics. In the present study, we obtained a high-support phylogenetic tree based on the combined mitochondrial gene set, which divided the 25 Agaricomycetes species into four major clades that is consistent with previous studies.37, 50. T. camphoratus was found to be a sister species of L. sulphureus and closely related to F. palustris. The results showed that mitochondrial genes were suitable as molecular markers for phylogenetic analysis of Agaricomycetes. More fungal mitochondrial genomes need to be revealed in the future to facilitate the study of fungal taxonomy, phylogenetics and population genetics.

Materials and methods
Assembly and annotation of T. camphoratus mitogenome. The whole genome sequencing reads of T. camphoratus used for mitogenome assembly were downloaded from NCBI SRA database under the accession number (SRX124102). De novo assembly of the T. camphoratus mitogenome was performed as implemented by SPAdes 3.9.99 with a kmer size of 17, using the downloaded data. We used MITOhm V1.9.90 to fill gaps among contigs. The obtained complete mitogenome of T. camphoratus was initially annotated combining the results of mfannot tool (https://megasun.bch.umontreal.ca/cgi-bin/dev_mfa/mfannotInterface.pl)41 and MITOS82, both based on the genetic code 4. At this step, protein coding genes (PCGs), tRNA genes and rRNA genes are preliminarily annotated. PCGs were also predicted or modified using the NCBI Open Reading Frame Finder (https://www.ncbi.nlm.nih.gov/orffinder/) and annotated via BLASTP searches against the NCBI non-redundant protein sequence database.83 PCGs which have no significant similarity to previously characterized proteins were annotated as hypothetical proteins. Intron–exon borders of PCGs were verified using exonerate v2.84. tRNAGenes were also predicted using the tRNAscan-SE 1.3.1 program.85

Sequence analysis. DNASTAR Lasergene v7.1 (https://www.dnastar.com/) was used to analyze the base composition of the T. camphoratus mitogenome. Strand asymmetry of the mitogenome was assessed according to the following formulas: AT skew = |A – T|/[A + T], and GC skew = |G – C|/[G + C].86 We used the DnaSP v6.10.01 software to calculate the synonymous substitution rate (Ks) and the nonsynonymous substitution rate (Ka) for all core PCGs in the T. camphoratus mitogenome, as well as in the previously published Polyporales mitogenomes.

The genetic distances between each pair of the 14 core PCGs (atp6, atp8, atp9, cox1, cox2, cox3, nad1, nad2, nad3, nad4, nad4L, nad5, nad6, and cob) and the ribosomal protein S3 (rps3) gene were calculated with MEGA v6.06,87 using the Kimura-2-parameter (K2P) model. Codon usage were analyzed using the Sequence Manipulation Suite88, based on genetic code 4. Genome synteny of the T. camphoratus mitogenome and its closely related species were analyzed with the Mauve v2.4.089.
Identification of repetitive elements. BLASTn searches of the whole mitogenome against itself were performed to determine whether there was intra-genomic duplication of large fragments and interspersed repeats in the *T. camphoratus* mitogenome at an E value of $<10^{-10}$. In addition, tandem repeats (>10 bp in length) were detected using the online program Tandem Repeats Finder with default parameters. Repeated sequences were also searched by REPuter to identify forward (direct), reverse, complemented, and palindromic (revere complemented) repeats.

Comparative mitogenomic analysis and intron analysis. The genome sizes, GC content, base composition, and gene numbers were compared among different *Polyporales* species to assess variations and conservation among mitogenomes. Group 1 introns in *cox1* and *cob* genes of the 12 *Polyporales* species we detected were classified into different position classes (Pcls) according to the method described by Férandon et al. Each Pcl was constituted by introns inserted at the same position in the coding region of the *cox1* or the *cob* gene. The Pcls of *cox1* gene were named in letter according to the similarity with the described Pcls. Pcls of *cob* gene were named in Latin number according to the insert position in the coding region of the *cob* gene.

Phylogenetic analysis. In order to investigate the phylogenetic status of *T. camphoratus* among *Agaricomycetes* class, we constructed a phylogenetic tree of 25 *Agaricomycetes* species based on the combined mitochondrial gene set (15 core PCGs + 2 rRNA genes). Single mitochondrial gene was first aligned using MAFFT v7.037. Then we and concatenated these alignments to gene set using SequenceMatrix v1.7.8. In order to detect potential phylogenetic conflicts between different genes, we carried out a preliminary partition homogeneity test. Best-fit models of evolution and partitioning schemes for the gene set were determined according to PartitionFinder 2.1.1. MrBayes v3.2.6 was used to construct the phylogenetic tree using Bayesian inference (BI) method based on the combined gene set. Two independent runs with four chains (three heated and one cold) each were conducted simultaneously for $2 \times 10^6$ generations. Each run was sampled every 100 generations. We assumed that stationarity had been reached when estimated sample size (ESS) was greater than 100, and the potential scale reduction factor (PSRF) approached 1.0. The first 25% samples were discarded as burn-in, and the remaining trees were used to calculate Bayesian posterior probabilities (BPP) in a 50% majority-rule consensus tree. We also constructed the phylogenetic tree using maximum likelihood (ML) method based on the combined gene set using RAxML v8.0.111.

Ethical approval. This article does not contain any studies with human participants performed by any of the authors.

Data availability

The *T. camphoratus* mitochondrial genome sequences were submitted to GenBank under accession number MH745717.

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**Author contributions**

Conceived and designed experiments: Q.L., X.W., and N.L. Performed the experiments: L.J., M.W., and H.Y. Analyzed the data: Q.L., M.C., X.L., and H.L. Wrote and revised the paper: Q.L., and X.W.

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Competing interests
The authors declare no competing interests.

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