The complete chloroplast genome of *Leptochilus hemionitideus*, a traditional Chinese medical fern

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**ABSTRACT**

The complete chloroplast genome of *Leptochilus hemionitideus* was sequenced. Its length is 156,083 bp with 44.2% GC content. The genome exhibits typical quadripartite with two inverted repeat regions (24,594 bp, each) separated by a large single-copy (LSC, 81,403 bp) region and a small single-copy (SSC, 25,492 bp) region. It has 131 genes, including 87 protein-coding genes, 34 tRNA genes, eight rRNA genes and two pseudogenes. Maximum-likelihood phylogenetic tree indicated that *L. hemionitideus* was closely related to *Lepisorus clathratus*. The complete chloroplast genome of *L. hemionitideus* would provide very valuable molecular information for further inferring the relationships of the microsoroid ferns.

*Leptochilus hemionitideus* is terrestrial fern belonging to the subfamily Microsoroideae in Polypodiaceae (Zhang et al. 2013). Different from congeneric other species, it has orbicular to elongate sori on tertiary veins parallel to secondary veins (Zhang et al. 2013). Its fronds are not or only slightly dimorphic. The plant has rock habit, usually growing on stones in streams at an altitude of 700–2000 m (Zhang et al. 2013). Its main distribution areas are concentrated in China, Bhutan, India, Japan, Nepal and Thailand (Zhang et al. 2013). In China, *L. hemionitideus* is a traditional Chinese medical fern with clearing heat and detoxification (State Administration of Traditional Chinese Medicine ‘Chinese Materia Medica Committee’ 1999). In addition, whether *Leptochilus* were merged with *Colysis* occurs controversy due to lack of the obvious generic delimitation (Nooteboom 1997; Shi and Zhang 1999). Phylogenetic relationships of *Leptochilus* to some genera in Microsoroideae such as *Microsorum* are also needed to further explore (Christenhusz and Chase 2014; PPG I 2016). Therefore, sequencing complete chloroplast genome of *L. hemionitideus* will contribute to deal with these issues and lay solid foundations for further phylogenomic investigation.

We sampled fresh and young leaves of *L. hemionitideus* from South China Botanical Garden, Chinese Academy of Sciences (CAS; 23°11′3.56″N, 113°21′43.28″E). The voucher specimen was conserved in Herbarium of Sun Yat-sen University (SYS; voucher: SS Liu 20161014). After total genomic DNA extraction, we built up ~300 bp genomic library and sequenced in Illumina Hiseq 2500 platform (Illumina Inc., San Diego, CA). Total 8,119,002 raw reads were retrieved and trimmed by Trimmomatic (Bolger et al. 2014). A subset of 6,801,257 trimmed reads was used for reconstructing the chloroplast genome by Velvet v1.2.07 (Zerbino and Birney 2008) with 33× coverage. The chloroplast genome was annotated using DOGMA (Wyman et al. 2004) and tRNAscan-SE (Schattner et al. 2005) programs with default settings and finally validated with BLAST searches and manually corrected for intron/exon boundaries. The complete chloroplast genome sequence of *L. hemionitideus* was aligned with 11 representative ferns including *Marsilea crenata* as outgroup using MAFFT v7.311 (Katoh and Standley 2013). A maximum likelihood (ML) phylogenetic tree was constructed using RAxML v8.0 with 1000 bootstrap replicates (Stamatakis 2014).

We determined complete chloroplast genome of *L. hemionitideus*, which possesses a total length of 156,083 bp with 44.2% GC content (GenBank accession number: MH319943). The circular cp genome exhibits typical quadripartite with two inverted repeat regions (IRa and IRb) of 24,594 bp separated by a large single-copy (LSC) region of 81,403 bp and a small single-copy (SSC) region of 25,492 bp. It was predicted to contain 131 genes, including 87 protein-coding genes, 34 tRNA genes, eight rRNA genes and two pseudogenes (*ndhB* and *rpoC1*). Among them, 115 genes occur as a single copy, whereas 14 genes are duplicated in the IR regions. Fourteen genes contain one intron, especially, the gene *ycf3*, *clpP*, and *rps12* have two introns. ML tree indicated that *L.*
hemionitideus is closely related to *Lepisorus clathratus* (Figure 1). The complete chloroplast genome of *L. hemionitideus* will provide very valuable molecular information for further inferring the relationships of the microsoroid ferns.

**Disclosure statement**

No potential conflict of interest was reported by the authors.

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**References**

Bolger AM, Lohse M, Usadel B. 2014. Trimmomatic: a flexible trimmer for Illumina sequence data. Bioinformatics. 30:2114–2120.

Christenhusz MJ, Chase MW. 2014. Trends and concepts in fern classification. Ann Bot. 113:571–594.

Katoh K, Standley DM. 2013. MAFFFT multiple sequence alignment software version 7: improvements in performance and usability. Mol Biol Evol. 30:772–780.

NP I 2016. A community-derived classification for extant lycophytes and ferns. J Syst Evol. 54:563–603.

Shi L, Zhang XC. 1999. Taxonomic studies of the genus *Colysis* C. Presl (Polypodiaceae) from China and neighboring regions. Acta Phytotaxon Sin. 37:54–80.

State Administration of Traditional Chinese Medicine ‘Chinese Materia Medica Committee’. 1999. Zhong Hua Ben Cao [Chinese Materia Medica], Vol. 4. Shanghai: Shanghai Science and Technology Publisher, p. 223–224. [Chinese]

Schattner P, Brooks AN, Lowe TM. 2005. The trRNAscan-SE, snoscan and snoGPS web servers for the detection of tRNAs and snoRNAs. Nucleic Acids Res. 33:W686–W689.

Nooteboom H. 1997. The microsoroid ferns. Blumea. 42:261–395.

Stamatakis A. 2014. RAxML version 8: a tool for phylogenetic analysis and post-analysis of large phylogenies. Bioinformatics. 30:1312–1313.

Wyman SK, Jansen RK, Boore J. 2004. Automatic annotation of organelar genomes with DOGMA. Bioinformatics. 20:3252–3255.

Zerbino DR, Birney E. 2008. Velvet: algorithms for de novo short read assembly using de Bruijn graphs. Genome Res. 18:821–829.

Zhang XC, Lu SG, Lin YX, Qi XP, Moore S, Xing FW, Wang FG, Hovenkamp PH, Gilbert MG, Nooteboom HP, et al. 2013. Polypodiaceae. In: Wu ZY, Raven PH, Hong DY, eds., Flora of China. Vol. 2–3(Pteridophytes). Beijing: Science Press; p. 758–850.