

The complete chloroplast genome sequence of the endangered Chinese endemic tree *Corylus fargesii*

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Abstract The complete chloroplast genome sequence of a Betulaceae species, *Corylus fargesii*, was mapped and determined based on Illumina sequencing data. The complete chloroplast genome was shown to be 159,856 bp, and comprises a pair of inverted repeat regions of 26,602 bp each, a large single-copy region of 88,313 bp, and a small single-copy region of 18,339 bp. It harbors 131 genes, including 94 protein-coding genes, 8 ribosomal RNA genes, and 29 transfer RNA genes. The A+T content of the entire genome is 63.49%. Phylogenetic analysis based on chloroplast genomes indicates that *C. fargesii* is closely related to *Ostrya rehderiana*.

Keywords *Corylus fargesii* · Chloroplast genome · Illumina sequencing · Phylogenetic analysis

Corylus fargesii, a Chinese endemic hazelnut species, grows in the mountain valleys from the elevation of 800–3000 m of the southwest China (Kuang et al. 1979). As a small arbor species in the genus *Corylus*, *C. fargesii* has been seriously threatened by human activities in the last few decades. Although several chloroplast (cp) DNA markers have previously been used for phylogenetic analysis (Erdogan and Mehlenbacher 2000; Palmé and Vendramin 2002; Boccacci and Botta 2009; Bassil et al. 2013;

Martins et al. 2013), except the *Corylus Chinensis* (GenBank accession number KX814336, Hu et al. 2016), little is known about the chloroplast genome in this genus. In the present study, we report the first complete chloroplast genome of *C. fargesii* based on Illumina pair-end sequencing data. The annotated chloroplast genome of *C. fargesii* has been deposited into GenBank under accession number KX822767.

Fresh leaves were collected from a single *C. fargesii* plant that grew at the resources nursery of the Institute of Forestry and Pomology (Beijing Academy of Agriculture and Forestry Sciences, Beijing, China). DNA extraction was performed using the Plant Genomic DNA Kit (TIANGEN Biotech Co., Beijing, China) according to the manufacturer's instructions, and high-throughput sequencing was carried out on the HiSeq2500 System (Illumina, San Diego, CA) by OriGene Technologies (Beijing, China). A total of 80.43 M of 126-bp raw reads were trimmed using SolexaQA (Cox et al. 2010), and approximately 9.577 G high-quality base pairs of sequence data were obtained and used for chloroplast genome assembly with SOAPdenovo software (Luo et al. 2012). Reference-guided assembly was then performed to reconstruct the chloroplast genome with the BLAST program (Altschul et al. 1990) using closely related species as references. After filling the gaps with GapCloser (<http://soap.genomics.org.cn/index.html>), a 159,856 bp chloroplast genome of *C. fargesii* was obtained. Annotation was performed using the CpGAVAS web server (Liu et al. 2012) to generate a physical map of the chloroplast genome.

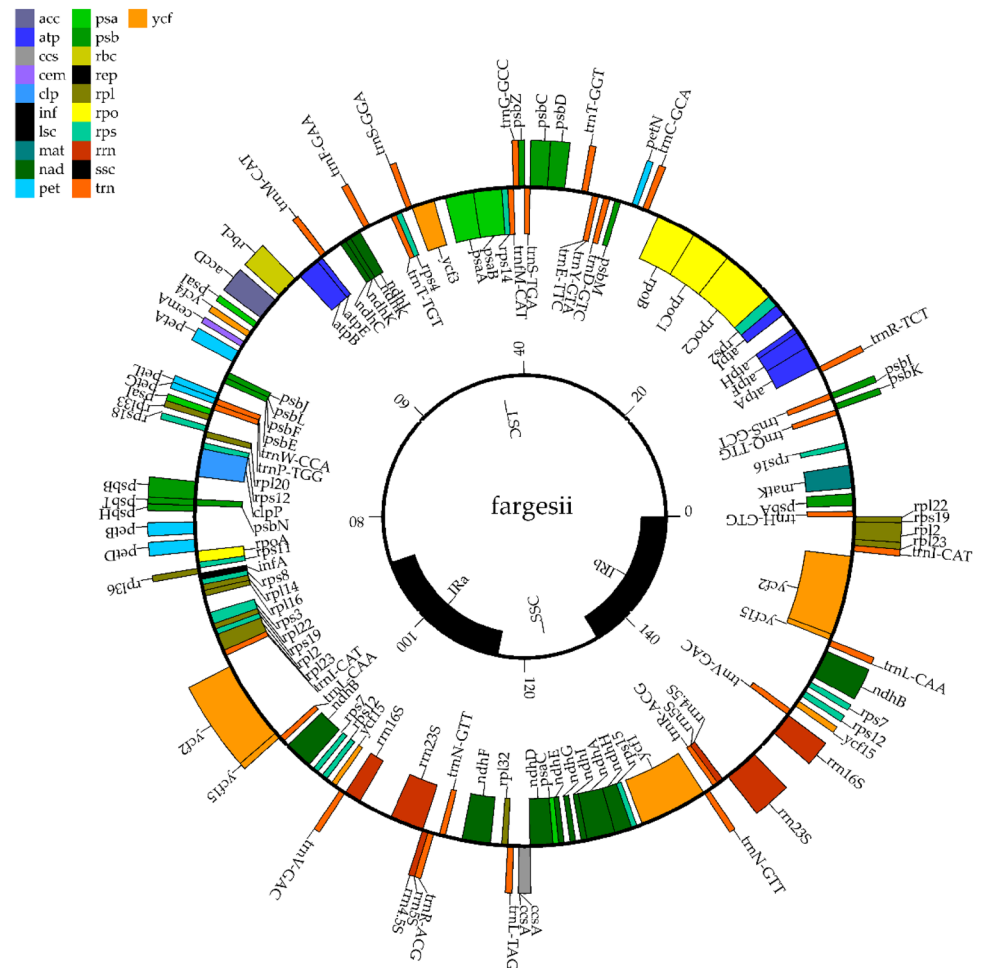
The circular chloroplast genome of *C. fargesii* contains a pair of inverted repeat (IRa and IRb) regions, each of 26,602 bp, a large single-copy (LSC) region and a small single-copy (SSC) region of 88,313 and 18,339 bp, respectively (Fig. 1). It comprises 131 genes, including 94

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Fig. 1 Gene map of the *Corylus fargesii* chloroplast genome



protein-coding genes, 8 ribosomal RNA genes, and 29 transfer RNA genes. Among the annotated genes, 11 protein-coding genes contain introns, including nine with a single intron each (*atpF*, *rpoC1*, *rpl2*, *ycf15*, *ndhB*, *ndhA*, *ndhB*, *ycf15*, and *rpl2*), and two with two introns each (*clpP* and *ycf3*). The overall AT content of the *C. fargesii* cp genome is 63.49%.

To identify the phylogenetic position of *C. fargesii*, maximum likelihood analysis was performed based on the 12 Fagales plant chloroplast genomes and seven other

outgroup plants using MEGA6 software (Tamura et al. 2013). The cp genome of *C. fargesii* was shown to be closely related to that of *Ostrya rehderiana* of the family Betulaceae (Fig. 2).

This complete chloroplast genome can be used for subsequent population, phylogenetic, and cp genetic engineering studies of *C. fargesii*. The new information will also be fundamental in formulating potential new conservation and management strategies for this important hazel species.

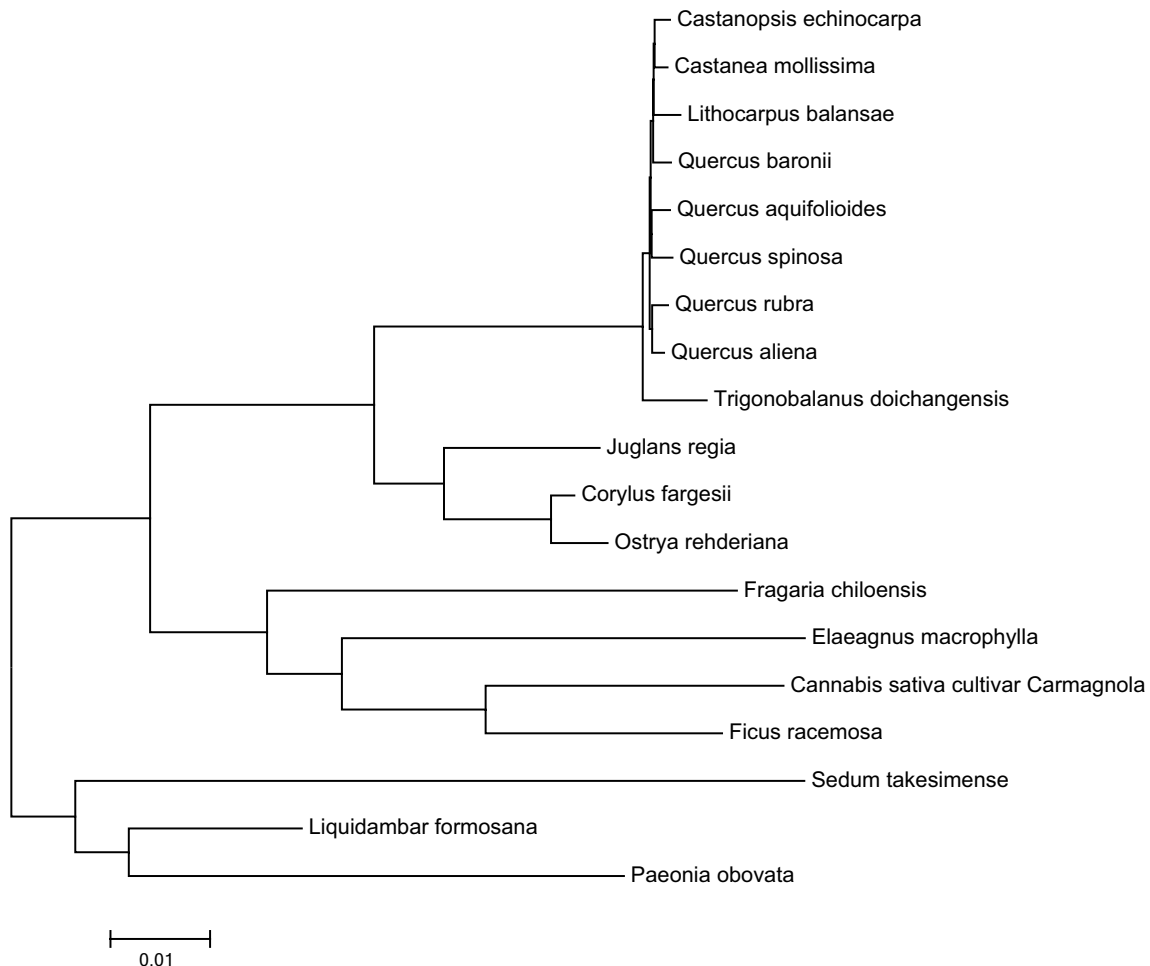


Fig. 2 Phylogenetic tree inferred using MEGA6 software from 19 complete cp genomes

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