

Characterization of the complete chloroplast genome sequence of *Galinsoga parviflora* and its phylogenetic implications

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ABSTRACT

Galinsoga parviflora is an invasive weed in southwest of Chinese agricultural systems and commonly used as medicine and food. In this study, the complete chloroplast genome of the *G. parviflora* was assembled from the whole genome Illumina sequencing data. The circular genome is 151,811 bp in size, which composed of one large single-copy (LSC) and one small single-copy (SSC) regions of 83,594 bp and 18,141 bp, respectively, and separated by a pair of inverted repeat (IR) regions of 25,038 bp each. It encodes a total of 113 gene species (80 protein-coding, 29 tRNA, and four rRNA species), in which 19 of them with double copies. The overall GC content is 37.7% while the GC content of the LSC, SSC, and IR regions are 35.8%, 31.3%, and 43.1%, separately. Phylogenetic analysis indicated that *Galinsoga parviflora* was closely related to *Galinsoga quadriradiata*.

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Galinsoga parviflora is one of the important species of the genus *Galinsoga* within the family Asteraceae (Asterales) (Ferheen et al. 2009; Ali et al. 2017). It is commonly found in Southwest of China such as Yunnan, Guizhou, and Sichuan Provinces (Pan et al. 2007). Although it is considered an invasive weed, the plant usually can be utilized as medicinal herb for wound healing as well as for the treatment of blood coagulation problems, cold, flu, toothache, and dermatological and eye diseases due to the presence of diverse secondary metabolites (Pan et al. 2007; Ali et al. 2017). The plant is full of essential oil containing bioactive compounds (Pino et al. 2010). So far, 38 compounds classified into seven categories (flavonoids, aromatic esters, diterpenoids, caffeic acid derivatives, steroids, phenolic acid derivatives, and miscellaneous compounds) have been isolated from *G. parviflora* (Ali et al. 2017). It has been reported that the plant had antibacterial (Matu and Van Staden 2003; Damalas, 2008; Pino et al. 2010), antifungal (Ali et al. 2017), anti-inflammatory (Matu and Van Staden 2003; Damalas, 2008), antioxidant (Chipurura et al. 2009; Bazylko et al. 2012; Bazylko et al. 2015), hepatoprotective (Mostafa et al. 2013), and hypoglycemic activity (Mostafa et al. 2013; Ali et al. 2017).

To facilitate its genetic research and contribute to its utilization, in this study, the complete chloroplast genome of the *G. parviflora* was assembled from the whole genome Illumina sequencing data. Phylogenetic analysis was conducted, which will be useful for further studies on its chloroplast genetic engineering.

Total genomic DNA was isolated from fresh leaves of an individual of *G. parviflora* from Sichuan province in southwest of China, located at 101°53'44"E, 30°52'44"N, and was stored

in the Herbarium of Neijiang Normal University (accession number: 20190211GP03). The Illumina sequencing was conducted on Illumina HiSeq X Ten platform in Beijing Novogene Bioinformatics Technology Co., Ltd (Beijing, China). The complete chloroplast genome was assembled using the baiting and iterative mapping approach (Hahn et al. 2013), with that of its congener *Galinsoga quadriradiata* (GenBank accession number KX752097) (Wang et al. 2018) as the initial reference genome. The annotated genomic sequence has been submitted to GenBank with the accession number MK737938.

The circular chloroplast genome of *G. parviflora* was 151,811 bp in size, which comprised of one LSC and one SSC regions of 83,594 bp and 18,141 bp, respectively, and separated by a pair of IR regions of 25,038 bp each. It encodes a total of 113 genes (80 protein-coding, 29 tRNA, and four rRNA species), in which 19 of them with double copies. Intron-exon structure analysis indicated that 16 genes (10 protein-coding genes and six tRNA genes) contained intron, in which of them two protein-coding genes (*clpP* and *ycf3*) had two introns while the others had one intron. The total GC content is 37.7%, while the corresponding values of the LSC, SSC, and IR regions are 35.8%, 31.3%, and 43.1%, separately.

To identify the phylogenetic position of *G. parviflora*, phylogenetic analysis was conducted. The maximum-likelihood (ML) phylogenetic tree was generated using species within the family Asteraceae by MEGA 7.0 (Kumar et al. 2016), which showed the position of *G. parviflora* was situated as the sister of *Galinsoga quadriradiata* in Asteraceae (Figure 1). The results indicated that *G. parviflora* and the

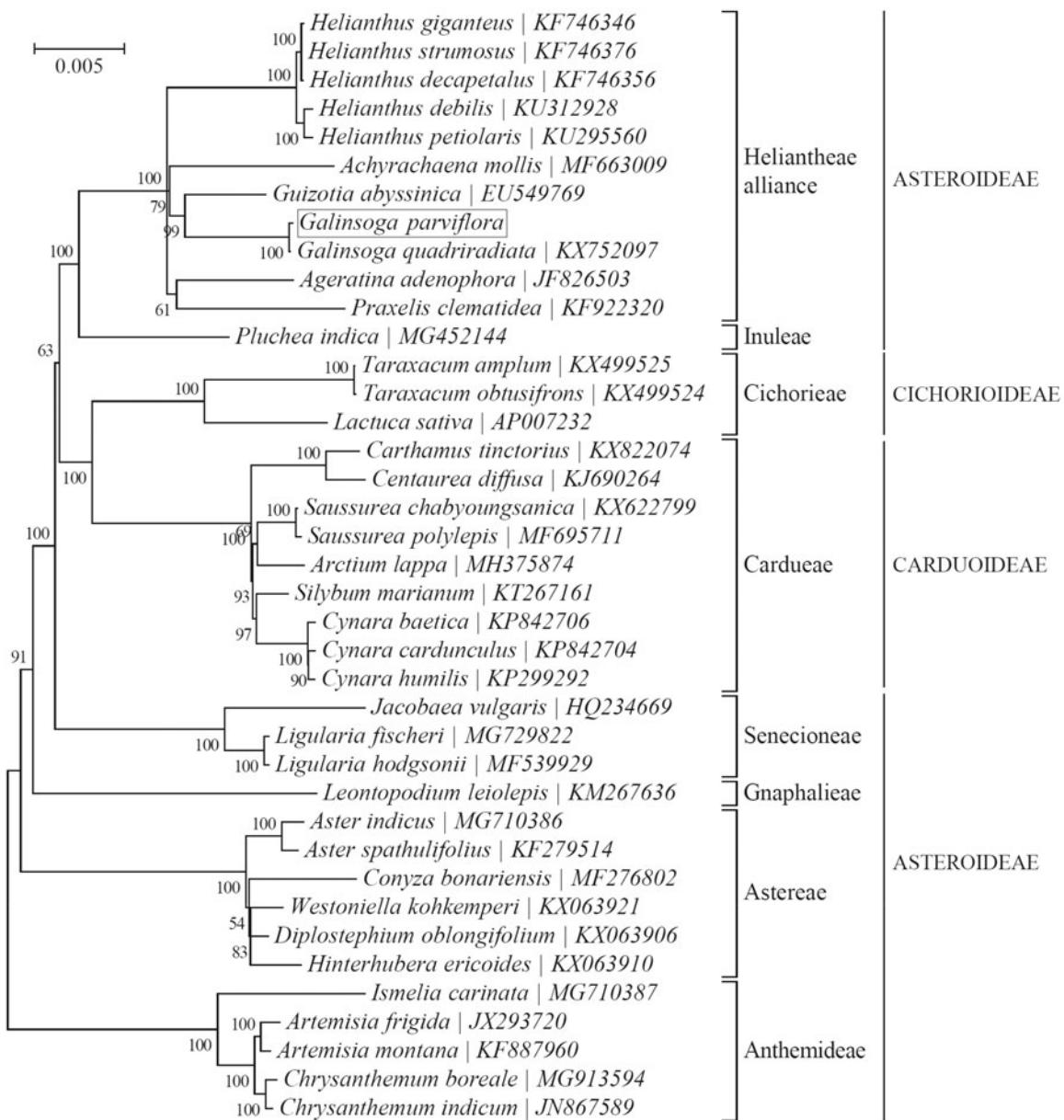


Figure 1. Phylogeny of 39 species within the family Asteraceae based on the maximum-likelihood (ML) of the concatenated chloroplast protein-coding sequences. The GTR + G + I model was employed as the best-fit nucleotide substitution model as suggested by MEGA7. The position of *Galinsoga parviflora* (GenBank accession number: MK737938) is shown in a box.

other 10 species were clustered into a clade. Our findings will provide a foundation for further investigation of chloroplast genome evolution in *Galinsoga*.

Disclosure statement

The authors declare no conflict of interest. The authors alone are responsible for the content and writing of the paper.

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