

## Complete plastid genome of *Gentiana trichotoma* (Gentianaceae) and phylogenetic analysis

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### ABSTRACT

The complete plastid genome of *Gentiana trichotoma* was determined and analyzed in this work. It had a circular-mapping molecule with the length of 144,759 bp, has similar gene composition with *G.* section *Cruciata* but contains 10 more genes than *G.* section *Kudoa*. Phylogenetic analysis showed that *G. trichotoma* clustered together with section *Kudoa* rather than section *Cruciata*. The plastome provided in this work would be useful for elucidation of *Gentiana* evolution.

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### KEYWORDS

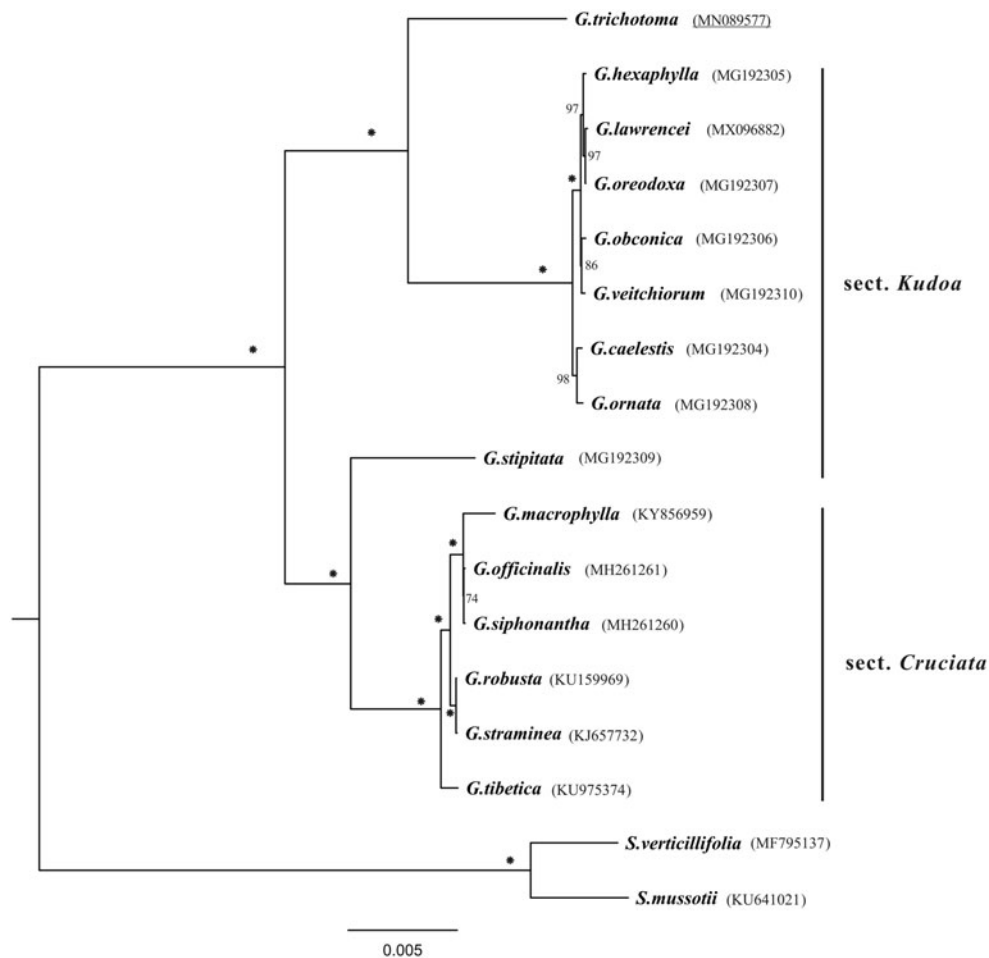
*Gentiana trichotoma*;  
phylogenetic ana-  
lysis; plastome

As a big genus containing 15 sections (Ho and Liu 2001), *Gentiana* plants are typically alpine and important parts of alpine shrub and meadow. *Gentiana trichotoma* Kusnezow, belonging to section *Frigidae* Kusnezow, is endemic to the Qinghai-Tibetan Plateau (Ho and Liu 2001). However, there has been no genomic studies in section *Frigidae*.

Herein, we reported and characterized the complete *G. trichotoma* plastome (MN089577). One *G. trichotoma* individual (specimen number: Fu2016163-6) was collected from Daocheng, Sichuan Province of China (29°27'09"N, 100°13'27"E) and its voucher specimens was deposited in the herbarium of School of Life Science, Luoyang Normal University. The fragmented genomic DNA was sequenced using Illumina HiSeq 2500 platform (Novogene, Tianjing, China), yielding approximately 5 Gb of 150-bp paired-end. The plastome was *de novo* assembled in NOVOPlasty 2.6.1 (Dierckxsens et al. 2016) and then annotated in GeSeq (Tillich et al. 2017) using the default parameters. Comparative analysis was conducted in mVISTA (Frazer et al. 2004) with *G. straminea* (Ni et al. 2016) and *G. lawrencei* var. *farreri* (Fu et al. 2016) which represents the only two plastome-available *Gentiana* sections, *Cruciata* Gaudin, and *Kudoa* (Masamune) Satake & Toyokuni ex Toyokuni, respectively. Shared protein-coding genes in plastomes of available *Gentiana* species were extracted and concatenated, then aligned using MAFFT (Katoh et al. 2002). The ML phylogeny was performed with

IQ-TREE (Nguyen et al. 2015) in PhyloSuite (Zhang et al. 2018) with 1000 replicates. *Swertia mussotii* (KC875852) and *S. verticillifolia* (MF795137) were served as the outgroups.

The complete *G. trichotoma* plastome is a circular-mapping molecule with the length of 144,759 bp. The LSC, IR, and SSC regions were 77,430, 25,162, and 17,005 bp, respectively. The overall GC content of the plastome was 37.8%. A total of 130 genes were annotated, containing 88 protein-coding genes, 34 tRNA genes, and 8 rRNA genes. Comparison analysis indicated that plastome of *G. trichotoma* has similar gene composition with section *Cruciata* (Ni et al. 2016; Zhou et al. 2018), with hotspots locating at intergenic regions such as *trnK*<sup>UUU</sup>-*rps16*, *atpH*-*atpI*, *petN*-*trnD*, and *trnL*<sup>UAG</sup>-*ccsA*. However, gene loss was not detected in *G. trichotoma*, which is different with section *Kudoa* which has almost lost 10 *ndh* genes (Sun et al. 2018). Phylogenetic analysis showed that *G. trichotoma* was clustered together with section *Kudoa* rather than section *Cruciata* (Figure 1), which is consistent with the previous study (Favre et al. 2016). *Gentiana stipitata* belonging to section *Kudoa* was clustered with section *Cruciata*, which is consistent with the previous study (Sun et al. 2018), indicating that more taxa should be involved for further study. The determination of the *G. trichotoma* plastome sequences provided new molecular data to illuminate the *Gentiana* evolution.



**Figure 1.** Phylogenetic tree (maximum likelihood) based on protein-coding genes of *Gentiana* plastomes. The asterisks along the branches mean 100% bootstrap supports based on 1000 replicates. The underline located in Genbank accession numbers indicates newly determined plastid genomes.

## Disclosure statement

There are no conflicts of interest for all the authors including the implementation of research experiments and writing this article.

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## References

- Dierckxsens N, Mardulyn P, Smits G. 2016. NOVOPlasty: *de novo* assembly of organelle genomes from whole genome data. *Nucleic Acids Res.* 45:e18.
- Favre A, Michalak I, Chen CH, Wang JC, Pringle JS, Matuszak S, Sun H, Yuan YM, Struwe L, Muellner-riehl A. 2016. Out-of-Tibet: the spatio-temporal evolution of *Gentiana* (Gentianaceae). *J Biogeogr.* 43: 1967–1978.
- Frazer KA, Pachter L, Poliakov A, Rubin EM, Dubchak I. 2004. VISTA: computational tools for comparative genomics. *Nucleic Acids Res.* 32: W273–W279.
- Fu PC, Zhang YZ, Geng HM, Chen SL. 2016. The complete chloroplast genome sequence of *Gentiana lawrencei* var. *farreri* (Gentianaceae) and comparative analysis with its congeneric species. *PeerJ.* 4:e2540.
- Ho TN, Liu SW. 2001. A worldwide monograph of *Gentiana*. Beijing: Science Press.
- Katoh K, Misawa K, Kuma K, Miyata T. 2002. MAFFT: a novel method for rapid multiple sequence alignment based on fast Fourier transform. *Nucleic Acids Res.* 30:3059–3066.
- Nguyen LT, Schmidt HA, von Haeseler A, Minh BQ. 2015. IQ-TREE: a fast and effective stochastic algorithm for estimating maximum-likelihood phylogenies. *Mol Biol Evol.* 32:268–274.
- Ni L, Zhao Z, Xu H, Chen S, Dorje G. 2016. Chloroplast genome structures in *Gentiana* (Gentianaceae), based on three medicinal alpine plants used in Tibetan herbal medicine. *Curr Genet.* 63:241–252.
- Sun SS, Fu PC, Zhou XJ, Cheng YW, Zhang FQ, Chen SL, Gao QB. 2018. The complete plastome sequences of seven species in *Gentiana* sect. *Kudoa* (Gentianaceae): Insights into plastid gene loss and molecular evolution. *Front Plant Sci.* 9:493.
- Tillich M, Lehwark P, Pellizzer T, Ulbricht-Jones ES, Fischer A, Bock R, Greiner S. 2017. GeSeq—versatile and accurate annotation of organelle genomes. *Nucleic Acids Res.* 45:W6–W11.
- Zhang D, Gao F, Li WX, Jakovlić I, Zou H, Zhang J, Wang GT. 2018. PhyloSuite: an integrated and scalable desktop platform for streamlined molecular sequence data management and evolutionary phylogenetics studies. *bioRxiv.* 20. DOI:10.1101/489088 [accessed 2018 Dec 7].
- Zhou T, Wang J, Jia Y, Li W, Xu F, Wang X. 2018. Comparative chloroplast genome analyses of species in *Gentiana* section *Cruciate* (Gentianaceae) and the development of authentication markers. *IJMS.* 19:1962.