The complete chloroplast genome of *Clivia miniata*

Wei Wang, Fan Zhang, Chenhe Li and Xudan Zhou

College of Horticulture, Jilin Agricultural University, Changchun, China

**ABSTRACT**

The complete chloroplast genome of *Clivia miniata* was assembled in this study. The genome comprised 158,114 bp in length. The GC content was 37.97%. A total of 133 genes are successfully annotated, including 87 protein-coding genes, 38 tRNA, and 8 rRNA genes. Seventeen protein-coding genes (atpF, ndhA, ndhB, petB, petD, rpl16, rpl2, rps16, trnA-UGC, trnG-UCC, trnI-GAU, trnK-UUU, trnL-UAA, trnV-UAC, clpP, rps12, ycf3) contained one or two introns. Phylogenetic tree analysis revealed that the *Clivia miniata* is the closest relative with *Lycoris radiata*, *Lycoris squamigera*, and *Narcissus poeticus*.

**ARTICLE HISTORY**

Received 4 January 2020
Accepted 19 January 2020

**KEYWORDS**

*Clivia miniata*; chloroplast genome; Amaryllidaceae

*Clivia miniata* belongs to *Clivia* of Amaryllidaceae. It originated in the deep forests of Natal Province, South Africa and was introduced to China in the early 20th century (Chen and Cheng 1990; Li 1988). *Clivia miniata* is widely used as a potted ornamental plant. It is also the city flower of Changchun City, Jilin province. At present, there is a great dispute about the phylogenetic position of the genus *Clivia miniata*, which intersects with Liliaceae and Asparagaceae. Amaryllidaceae and Liliaceae are closely related in system location. In taxonomy, different schools of thought hold different views on the boundary dividing the two families.

In this study, the complete chloroplast genome of *Clivia miniata* was sequenced and analyzed. Based on the complete chloroplast genome of *Clivia miniata* (GenBank accession number: MN857162), phylogenetic analysis was performed using the maximum likelihood method using MEGA7.0 to study the position of *Clivia miniata* in phylogenetic development (Kumar et al. 2016). The materials used in this experiment were taken from the horticulture laboratory of Jilin agricultural university (Changchun City, Jilin province, China, 43°48′37″N, 125°24′8″E). The specimens (NEFI20190802WW1) were kept in the horticulture laboratory of Jilin Agricultural University.

Pair-end Illumina raw reads were cleaned from adaptors and barcodes and then quality filtered using Trimmomatic (Bolger et al. 2014). All putative chloroplast reads mapped to the reference genome were then used for de novo assembly to reconstruct the chloroplast genomes using SPAdes 3.6.1 with iterative K-mer sizes of 55, 87, and 121 (Bankevich et al. 2012;Marcais and Kingsford 2011). De novo assembled chloroplast contigs were concatenated into larger contigs using Sequencher 5.3.2 Read coverage analysis was then conducted to determine the inverted repeat (IR) region boundaries and any misassembled contigs using Jellyfish v.2.2.3. Automatic annotation of the chloroplast genomes were generated by CpGAVAS and a circular representation of both sequences was drawn using the online tool OGDRAW. The draft annotations given by CpGAVAS were then manually corrected using the Artemis software and other plastid genomes for comparison (Liu et al. 2012).

The complete chloroplast genome of *Clivia miniata* is a typical four-segment structure with a total length of 158,114 bp, including a large single-copy region (LSC) with a length of 86,204 bp, a small single-copy region (SSC) with a length of 18,834 bp, and two reverse repeats of equal length but opposite direction (IRA/IRB) with a length of 26,788 bp. A total of 133 genes were encoded, including 87 protein-coding genes, 38 tRNA genes, and 8 rRNA genes with GC% of 37.97% and 36 duplicated genes in the IR region, which are atpF, ndhA, ndhB, petB, petD, rpl16, rpl2, rps16, trnA-UGC, trnG-UCC, trnI-GAU, trnK-UUU, trnL-UAA, trnV-UAC, clpP, rps12, and ycf3 contains two introns.

Phylogenetic analysis was performed using the complete cp genomes of *Clivia miniata* with those of 10 species in Figure 1. Phylogenetic tree based on chloroplast genome sequences from 16 plant species.
Liliaceae, 3 species of Amaryllidaceae, *Agave Americana*, and *Iris sanguinea* reported in Genbank of NCBI database by maximum likelihood method in MEGA version 7.0. The results showed that *Clivia miniata* and *Narcissus poeticus*, *Lycoris radiate*, *Lycoris squamigera* are in the same branch, belongs to Amaryllidaceae and is closely related to *Allium ampeloprasum* and *Allium obliquum* (Figure 1).

**Disclosure statement**

The authors declare no conflicts of interest. The authors alone are responsible for the content and writing of this article.

**Funding**

This work was supported by the Key Scientific and Technological Development Plan of Jilin Province [20190304008YY] and the Forestry Science and Technology Promotion Demonstration Project of Central Finance [ULT2019-17].

**References**

Bankevich A, Nurk S, Antipov D, Gurevich AA, Dvorkin M, Kulikov AS, Lesin VA, Nikolenko SI, Pham S, Prjibelski AD, et al. 2012. Spades: a new genome assembly algorithm and its applications to single-cell sequencing. J Comput Biol. 19(5):455–477.

Bolger AM, Lohse M, Usadel B. 2014. Trimmomatic: a flexible trimmer for Illumina sequence data. Bioinformatics. 30(15):2114–2120.

Chen JY, Cheng X.K. 1990. Chinese flower classics. Shanghai: Shanghai culture press.

Kumar S, Stecher G, Tamura K. 2016. MEGA7: molecular evolutionary genetics analysis version 7.0 for bigger datasets. Mol Biol Evol. 33(7):1870–1874.

Li ZQ. 1988. Chinese clivia. Chengdu: Sichuan Science and Technology Press. p. 20–47.

Liu C, Shi LC, Zhu YJ, Chen HM, Zhang JH, Lin XH, Guan XJ. 2012. Cpgavas, an integrated web server for the annotation, visualization, analysis, and Genbank submission of completely sequenced chloroplast genome sequences. BMC Genomics. 13(1):715.

Marcais G, Kingsford C. 2011. A fast, lock-free approach for efficient parallel counting of occurrences of k-mers. Bioinformatics. 27(6):764–770.