The complete chloroplast genome of the Euphorbia maculata L. (Euphorbiaceae): characterization and phylogeny

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\section*{ABSTRACT}

Euphorbia maculata is an important medicinal plants of the family Euphorbiaceae. The complete chloroplast genome reported here is 162,685 bp in length, including two inverted repeats (IRs) of 26,822 bp, which are separated by a large single-copy (LSC) and a small single-copy (SSC) of 90,514 bp and 18,527 bp, respectively. The whole chloroplast genome of E. maculata contains 111 genes, including 77 protein-coding genes, 4 transfer RNA, and 30 ribosome RNA. Phylogenetic analysis indicated that E. maculata is closely related to E. miliii and E. tirucalli.

\section*{ARTICLE HISTORY}

Received 8 September 2020
Accepted 2 October 2020

\section*{KEYWORDS}

Euphorbia maculata; chloroplast genome; phylogenetic analysis

\textbf{Euphorbia maculata} L. (Spotted spurge) is a fast-growing annual weed native to eastern North America and spread to Asia, Africa and Europe (Wu et al. 1994). It functions as a pioneer species with a prostrate growth habit in ecological succession. The milky sap of the plant is an irritant for many people. Like many members of the family Euphorbiaceae, it has been widely used as a folk medicine, which can produce anti-inflammatory and cancer chemopreventive agents of terpenoids (Yi et al. 2018). It is necessary to develop genomic resources for \textit{E. maculata} to provide intragenic information for its utilization and chloroplast genomes are valuable sources (Dong et al. 2020; Sun et al. 2020).

The fresh leaves of \textit{E. maculata} were collected from Luxi county, Hunan province, China (28°12'59"N, 110°13'11"E). Voucher specimens were deposited in Institute of Chinese Materia Medica (Specimen accession number: 430723LY0485), China Academy of Chinese Medical Sciences. Total genomic DNA was extracted with the modified cetyltrimethyl ammonium bromide (CTAB) method (Li et al. 2013). Paired-end libraries were prepared with the NEBNext Ultra DNA Library Prep Kit. The genome was sequenced using the HiSeq X Ten platform (Illumina, Santiago, CA, USA). All good quality paired reads were assembled using the Spades program to contigs (Bankevich et al. 2012). Chloroplast genome sequence contigs were selected by the program BLAST (Altschul et al. 1990) using \textit{E. miliii} (Genbank accession number: MNT713924) as a reference and the selected contigs were assembled using Sequencher 4.10 (Gene Codes Corporation, Ann Arbor, MI, USA, \url{http://www.genecodes.com}). Chloroplast genome annotation was performed with Plann (Huang and Cronk 2015) using the \textit{E. miliii} as reference sequence. The annotated sequence was submitted to the GenBank under the accession number MT830858.

The complete chloroplast genome reported here is 162,685 bp in length, including two inverted repeats (IRs) of 26,822 bp, which are separated by a large single-copy (LSC) and a small single-copy (SSC) of 90,514 bp and 18,527 bp, respectively. The overall GC-content of the chloroplast genome was 35.4%. The chloroplast DNA of \textit{E. maculata} comprised 111 distinct genes, including 77 protein-coding genes, 4 transfer RNA, and 30 ribosome RNA, but didn’t contain \textit{cemA} and \textit{rpl22} these two protein-coding genes. In these genes, 17 harbored a single intron, while two (\textit{ycf3} and \textit{clpP}) contained double introns.

In order to confirm the phylogenetic relationships of \textit{E. maculata} within the genus \textit{Euphorbia} and other related groups, total 26 complete cp genomes were obtained from Genbank and the genus \textit{Aristolochia} was taken as an out-group. All chloroplast genome sequences were aligned using MAFFT (Katoh et al. 2019) and ambiguous alignment regions were trimmed by Gblocks (Castresana 2000). Phylogenetic analysis was conducted based on maximum-likelihood (ML) analyses using RAxML (Stamatakis 2014), under the GTR + G model with 1000 rapid bootstrap replicates. The phylogenetic tree showed that all species of \textit{Euphorbia} form a monophyletic group with 100% support, and \textit{E. maculata} is closely related to \textit{E. miliii} and \textit{E. tirucalli} (Figure 1). The chloroplast genome of \textit{E. maculata} provided a lot of genetic information.
for species conservation and identification of genus *Euphorbia*.

**Disclosure statement**

No potential conflict of interest was reported by the author(s).

**Funding**

This work was supported by the National Key Research and Development Program of China [No. 2019YFC1711400 & No. 2019YFC1711402], and the National Natural Science Foundation of China [No. 81891014].

**Data availability statement**

The data that support the findings of this study are openly available in GenBank of NCBI [https://www.ncbi.nlm.nih.gov/], reference number MT830858, raw data BioProject ID: PRJNA662166.

**References**

Altschul SF, Gish W, Miller W, Myers EW, Lipman DJ. 1990. Basic local alignment search tool. J Mol Biol. 215(3):403–410.

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

Castresana J. 2000. GBLOCKS: selection of conserved blocks from multiple alignments for their use in phylogenetic analysis. Version 0.91b. Mol Biol Evol. 17:540–552.

Dong W, Xu C, Wen J, Zhou S. 2020. Evolutionary directions of single nucleotide substitutions and structural mutations in the chloroplast genomes of the family Calycanthaceae. BMC Evol Biol. 20(1):96.

Katoh K, Rozewicki J, Yamada KD. 2019. MAFFT online service: multiple sequence alignment, interactive sequence choice and visualization. Brief Bioinform. 20(4):1160–1166.

Li JL, Wang S, Jing Y, Wang L, Zhou SL. 2013. A modified CTAB protocol for plant DNA extraction. Chin Bull Bot. 48:75–78.

Stamatakis A. 2014. RAxML version 8: a tool for phylogenetic analysis and post-analysis of large phylogenies. Bioinformatics. 30(9):1312–1313.

Sun J, Wang Y, Liu Y, Xu C, Yuan Q, Guo L, Huang L. 2020. Evolutionary and phylogenetic aspects of the chloroplast genome of *Chaenomeles* species. Sci Rep. 10(1):11466.

Wu ZY, Raven PH, Hong DY. 1994. Flora of China. 44.

Yi S, Liang-Liang G, Meng-Yue T, Bao-Min F, Yue-Hu P, Ken Y. 2018. Triterpenoids from *Euphorbia maculata* and their anti-inflammatory effects. Molecules. 23:2112.

Figure 1. Phylogenetic tree reconstruction of 27 taxa using maximum likelihood (ML) methods in the chloroplast genome sequences. ML bootstrap support value presented at each node.