The complete plastid genome of *Selaginella erythropus* (Selaginellaceae), a species with distinctive giant chloroplasts

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**ABSTRACT**

The plastid genome of the deep-shade plant *Selaginella erythropus*, which has highly unusual chloroplasts, was characterized using Illumina pair-end sequencing. This plastome is 140,151 bp in length with a large single-copy region (LSC) of 56,133 bp, a small single-copy region (SSC) of 61,268 bp, and two direct repeats (DRs) of 11,375 bp. The overall GC content is 50.68%, while those of LSC, SSC, and DR are 48.96%, 50.3%, and 55.96%, respectively. The plastome contains 102 genes, including 76 protein-coding, 15 tRNA (12 tRNA species), and 8 rRNA genes (4 rRNA species). The phylogenetic analysis shows that *S. erythropus* is closely related to *S. moellendorfii* and *S. doederleini*. This result is consistent with the previous phylogenetic relationship inferred from multiple plastid and nuclear loci. However, only *S. erythropus* has the two-zoned giant chloroplast, the bizonoplast. The plastome provides an excellent reference for understanding the unique chloroplast differentiation in Selaginellaceae.

In most land plants, chloroplasts are located in the mesophyll cells, with 50–250 chloroplasts per cell (Pyke 2009). *Selaginella* is unusual in having several variations on this typical chloroplast structure (Liu et al. 2020) including giant chloroplasts of several different forms. The bizonoplast, found in every leaf dorsal epidermal cell of some *Selaginella* species, is a cup-shaped giant chloroplast with dimorphic ultrastructure (Sheue et al. 2007). The upper zone is characterized by groups of 2–4 thylakoid membranes parallel to each other, while the lower zone consists of grana and stroma thylakoid membranes, similar to normal chloroplasts.

The bizonoplast, first reported from *S. erythropus* (Mart.) Spring (Sheue et al. 2007), and only reported in *Selaginella*, originates from a proplastid, developing its zoned structure after exposure to low light (Sheue et al. 2015). *Selaginella*, with about 750 species occurring globally in various habitats (Jermy 1990), is an ideal model genus to understand the diversity of chloroplasts and their adaptive significance. Given the unique structure of the bizonoplast and its environmental correlates, the chloroplast genome of a bizonoplast-containing species is of special interest. Here we assembled and annotated the plastid genome of *S. erythropus* from a specimen growing in the nursery at National Chung Hsing University, Taiwan (24° 07’N, 120° 40’E) to contribute to the bioinformatics and genome structure of the bizonoplast.

The total genomic DNA was extracted from fresh shoots of *S. erythropus* (voucher # Liu JW-05, in the herbarium of National Chung Hsing University, TCB, Taiwan; Chiu-Rong Sheue, crsheue@nchu.edu.tw) using the CTAB method (Doyle and Doyle 1990). The genomic DNA was fragmented and libraries were constructed with the insertion sizes 180 bp, 350 bp, 500 bp, and 700 bp. These libraries were paired-end sequenced on an Illumina HiSeq platform (Illumina Inc., San Diego, CA). The de novo assembly was performed using the GENIOUS Prime Velvet plugin (Zerbino and Birney 2008), and subsequently, the plastid contigs were arranged based on the plastome of *S. moellendorfii* (HM173080) (Banks et al. 2011). PCR and Sanger sequencing were conducted to confirm the sequences from the SC-DR junctions and highly variable regions. The *S. erythropus* plastome was annotated using the software PGA (Qu et al. 2019) and GENEOUS Prime (Kearse et al. 2012) by comparing it with the plastomes of *S. moellendorfii* (HM173080) (Banks et al. 2011) and *S. doederleini* (MH598532) (Zhang et al. 2019).

Typically, plastomes contain two inverted repeats (IRs) separated by a large single-copy region (LSC) and a small single-copy region (SSC) (Mower and Vickrey 2018). However, the *S. erythropus* plastome features a set of direct repeats (DRs), similar to those from some other *Selaginella* species (Mower et al. 2019; Zhang et al. 2019) (Figure 1). The plastome is 140,151 bp in length and has two DRs of 11,375 bp.
(8.12%), which are separated by an LSC of 56,133 bp (40.05%) and an SSC of 61,268 bp (43.72%). The DR in Selaginella species is presumably caused by a large inversion containing a former IR copy (i.e. IRb). This inversion also relocated a partial LSC region into the former SSC region, thereby resulting in the relatively short LSC as compared to the SSC in other Selaginella (Shim et al. 2021).

The S. erythropus plastome GC content is 50.68%, with the LSC, SSC, and DR regions GC contents being 48.96%, 50.3%, and 55.96%, respectively. The plastome GC content of S. erythropus is on the low end of the range (50.75%–56.49%) from 16 other Selaginella species (Shim et al. 2021) but is much higher than the average for 3,507 plastome sequences from algae to seed plants (37.38 ± 2.26%, Kwon et al. 2020). The S. erythropus plastome comprises 102 genes, including 76 protein-coding (76 PCG species), 8 ribosomal RNA (4 rRNA species) and 15 transfer RNA genes (12 tRNA species). Nine PCG genes (atpF, clpP, ndhA, ndhB, petB, petD, rpl2, rpl16 and rpoC1) harbor a single intron, and one (ycf3) contains two introns. In addition, accD, infA and rpl20 are likely pseudogenes because of incomplete open reading frames.

To construct a phylogenetic tree, 51 shared protein-coding genes of 20 plastomes were extracted and aligned individually in MAFFT (Katoh and Standley 2013) implemented in GENEIOUS Prime (Kearse et al. 2012). The maximum likelihood (ML) tree was determined using GARLI v.2.0 (Zwickl 2006), with 1000 bootstrap replicates and the best model GTR + G + I model was selected based on Akaike Information Criterion (AIC) in jModeltest (Posada 2008). In the ML tree, S. erythropus is most closely related to S. moellendorffii (Figure 1). This result is consistent with the previous phylogenetic relationship inferred from three gene regions (rbcL, pgIc, SQD1) and morphological features (Weststrand and Korall 2016). The plastome of S. erythropus provides an excellent reference for elucidating the evolution and functional divergence of the giant chloroplasts and bizonoplasts in Selaginellaceae.

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No potential conflict of interest was reported by the authors.

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Data availability statement
The genome sequence data that support the findings of this study are openly available in GenBank of NCBI at (https://www.ncbi.nlm.nih.gov/) under the accession no. MZ392854. The associated BioProject, SRA, and
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