RESEARCH ARTICLE



Characterization of the complete chloroplast genome of *Nitraria* tangutorum, a desert shrub

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Abstract. The chloroplast genome sequence of *Nitraria tangutorum*, a desert shrub, was sequenced using high-throughput sequencing technology and analysed phylogenetically in the present study. The chloroplast genome is 159,414 bp in length, including a large single copy region of 87,924 bp and small single copy region of 18,318 bp, and a pair of inverted repeat regions of 26,586 bp. The chloroplast genome contains 110 unique genes, including 77 protein-coding genes, four ribosomal RNA genes, and 29 tRNA genes. Most of these genes are present as a single copy and in two or more copies 19 genes occurred. Seventeen genes have one intron, and *clpP* and *ycf3* genes contain two introns. A total of 81 simple sequence repeats (SSRs) were identified, most of them were found to be mononucleotide repeats composed of A/T. In addition to SSRs, 66 repeats were identified, including 41 tandem repeats, 10 palindromic repeats, and 15 forward repeats. The phylogenetic analysis based on 54 protein-coding genes demonstrated a close relationship between *N. tangutorum* and other plant species in Sapindales. The complete chloroplast genome sequence of *N. tangutorum* will provide important data for further study of taxonomy and systematics of the genus *Nitraria*.

Keywords. chloroplast genome; simple sequence repeats analysis; phylogenetic analysis; Nitraria tangutorum.

Introduction

Nitraria tangutorum, a member of genus Nitraria, is a wild shrub distributed in the desert and semi-desert areas of northwest China. It exhibits high tolerance to high salinity and drought stresses, and plays a key role in maintaining the fragile ecosystems in the desert areas of central Asia (Yang et al. 2010). In addition, N. tangutorum is of great economic value for the local people (Liu et al. 2014), e.g., fruits and seeds are used to make medicines and drinks (Zhao et al. 2017), and the dry branches are often used as firewood by locals.

As a shrub with ecological and economic importance in harsh environment, *N. tangutorum* has attracted the attention of many researchers in recent years. Studies have been conducted to investigate the ecological adaption and stress tolerance mechanism by using molecular biological technology and biochemistry methods (Yang *et al.* 2013; Zheng *et al.* 2014; Yan *et al.* 2018). However, the phylogenetic relationship of the genus *Nitraria* remains an open question. *N. tangutorum* was classified into different family by Liu and Zhou (2003), and Xu and Huang (1998).

(http://frps.iplant.cn/frps/Nitraria). *Nitraria* is one of the six genus in Zygophyllaceae of Geraniales (Xu and Huang 1998), but by Liu and Zhou (2003, http://foc.iplant.cn/), genus *Nitraria* was classified into Nitraria of Sapindales. Thus, more molecular evidences are needed to clarify the evolutionary position of genus *Nitraria*.

Recent studies have shown that the chloroplast genome sequences are essential data for plant phylogenetic and genetic population analyses (Parks *et al.* 2009). Thus, the phylogenetic analysis using the complete chloroplast genome of *N. tangutorum* should be an appropriate way to get a better understanding of the evolution of this plant species. Here, we present a complete chloroplast genome of *N. tangutorum* based on the next-generation sequencing data.

Materials and methods

Plant materials

Leaf sample of a wild individual of *N. tangutorum* were collected by Fei Gao from Mengxi Town, Erdos City, Inner

91 Page 2 of 6 Merhaba Abla et al.

Table 1. Chloroplast genome sequences used for phylogenetic tree construction.

Species name	GenBank ID	Family	Order
Acer buergerianum	NC_034744.1	Sapindaceae	Sapindales
Atuna racemose	NC_030546.1	Chrysobalanaceae	Malpighiales
Averrhoa carambola	NC_033350.1	Oxalidaceae	Oxalidales
Citrus aurantiifolia	NC_024929.1	Rutaceae	Sapindales
Dipteronia dyeriana	NC_031899.1	Sapindaceae	Sapindales
Erodium absinthoides	NC_026847.1	Geraniaceae	Geraniales
Erodium carvifolium	NC_015083.1	Geraniaceae	Geraniales
Erodium chrysanthum	NC_027065.1	Geraniaceae	Geraniales
Erythroxylum novogranatense	NC_030601.1	Erythroxylaceae	Malpighiales
Geranium incanum	NC_030045.1	Geraniaceae	Geraniales
Geranium maderense	NC 029999.1	Geraniaceae	Geraniales
Larrea tridentata	NC_028023.1	Zygophyllaceae	Zygophyllales
Linum usitatissimum	NC 036356.1	Linaceae	Malpighiales
Litchi chinensis	NC_035238.1	Sapindaceae	Sapindales
Pistacia vera	NC 034998.1	Anacardiaceae	Sapindales
Rhus chinensis	NC 033535.1	Anacardiaceae	Sapindales
Sapindus mukorossi	NC_025554.1	Sapindaceae	Sapindales
Spondias bahiensis	NC_030526.1	Anacardiaceae	Sapindales

Mongolia Autonomous Region (106° 79′E, 39°83′N). The sample (PM20181001-Nta-1) was deposited in College of Life and Environmental Sciences, Minzu University of China, Beijing.

Genome sequencing and annotation

The genomic DNA was extracted from the leaves using the modified CTAB method (Doyle 1987). DNA sequencing was performed using an Illumina Hiseq2500 (Illumina, San Diego, USA) at Shenzhen Huitong biotechnology (Shenzhen, China). After adapter trimming and filtering of the low quality reads (read has >5\% unidentified nucleotides and >50% of its bases with a quality value of <20.), the resulting clean reads were assembled into contigs using the assembler SPAdes v3.9.0 (Bankevich et al. 2012) using the default parameters. The contigs were aligned to chloroplast genome sequences of Arabidopsis thaliana and Nicotiana tabacum using BLAST program (E value $< 1e^{-10}$) to find the fragments of chloroplast genome of N. tangutorum and the contigs supported by higher sequencing depth were used for assembling chloroplast genome of N. tangutorum.

The genes in chloroplast genome of *N. tangutorum* were annotated using the DOGMA tool with default parameters (Wyman *et al.* 2004). Online program OGDRAW (https://chlorobox.mpimp-golm.mpg.de/OGDraw.html; Lohse *et al.* 2013) was used to draw the gene map of the *N. tangutorum* chloroplast genome. The finally annotated chloroplast genome of *N. tangutorum* was deposited in GenBank with the accession number MK341053.

Repeat and SSR analysis

Simple sequence repeats (SSR) are some DNA repeats formed by one or several tandemly arranged nucleotides, which spread widely in eukaryotic genomes. Perl script MISA (http://pgrc.ipk-gatersleben.de/misa/misa.html) was used to detect microsatellites with minimal repeat numbers of 10, 5 and 4 for mononucleotide, dinucleotide and trinucleotide repeats, respectively.

Phylogenetic analysis

To get more knowledge about phylogenetic analysis of *N. tangutorum*, we chose 18 plant species (table 1) from five orders (Sapindales, Geraniales, Malpighiales, Zygophyllales and Oxalidales) which belong to the same evolutionary branch (rosids) as *N. tangutorum*, and the phylogenetic relationships of these species and *N. tangutorum* were analysed. To analyse the phylogenetic tree of *N. tangutorum*, we downloaded their whole chloroplast genome sequences from the NCBI Organelle Genome and Nucleotide Resources. The software MAFFT 7.380 (Katoh and Standley 2013) was used to align the genome sequence, and RAxML 8.2.4 (Stamatakis 2014) was used to analyse the evolution of these species. The significance level for the phylogenetic tree was assessed by bootstrap testing with 1000 replications.

Results and discussion

Organization and features of the N. tangutorum chloroplast genome

The chloroplast genome of *N. tangutorum* was assembled using \sim 3.98 G sequencing reads. The length of

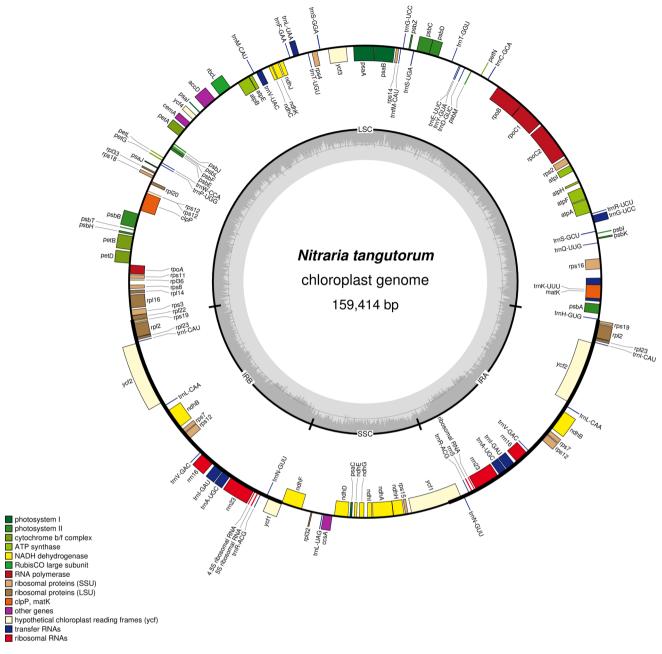


Figure 1. Gene map of the N. tangutorum chloroplast genome. The genes shown inside of the circle indicates transcriptional direction is clockwise, while those shown outside are counterclockwise. Genes belonging to different functional groups are labelled with different colours. The GC content of the genome shown with grey histogram in the inner circle, and the grey line depicts the 50% threshold line.

chloroplast genome of N. tangutorum was 159,414 bp, and the average sequencing depth was 2494.8X. The large single copy (LSC) region, small single copy (SSC) region and the two inverted repeat regions (IRs), IRa and IRb, were 87,924, 18,318, and 26,586 bp in length, respectively (figure 1). A total of 110 unique genes were annotated from the chloroplast genome of N. tangutorum, including 77 protein-coding genes, four ribosomal RNA genes, and 29 tRNA genes. Most of these genes were present as single copy and in two or more copies 19 genes occurred. Of the 110 unique genes, 58 were involved in self-replication of chloroplast genome, 12 genes encode ribosomal small subunit proteins, nine genes encode ribosomal large subunit proteins, and four genes encode RNA polymerase subunits. Forty-three genes in N. tangutorum chloroplast genome encode proteins associated with photosynthesis, including six ATP synthase subunits, 11 subunits of NADH dehydrogenase complex, six components of cytochrome b/f complex, five subunits of photosystem I, 14 subunits of photosystem II, and one large chain of rubisco Page 4 of 6 Merhaba Abla et al.

Table 2. List of the annotated genes in *N. tangutorum* chloroplast genome.

Category of genes	Subcategory of genes	Gene names
Self-replication	rRNA genes	rrn4.5*, rrn5*, rrn16*, rrn23*
•	tRNA genes	29 tRNA (trnI-CAU*, trnL-CAA*, trnV-GAC*, trnI-GAU*, trnA-UGC*, trnR-ACG*, trnN-GUU*, 7 tRNA genes in IR regions)
	Small subunit of ribosome	rps2, rps3, rps4, rps7*, rps8, rps11, rps12*, rps14, rps15, rps16, rps18, rps19*
	Large subunit of ribosome	rpl2*, rps14, rpl16, rpl20, rpl22, rpl23*, rpl32, rpl33, rpl36
	RNA polymerase subunits	rpoA, rpoB, rpoC1, rpoC2
Photosynthesis	ATP synthase gene	atpA, $atpB$, $atpE$, $atpF$, $atpH$, $atpI$
·	NADH dehydrogenase	ndhA, ndhB*, ndhC, ndhD, ndhE, ndhF, ndhG, ndhH, ndhI, ndhJ, ndhK
	Cytochrome b/f complex	petA, $petB$, $petD$, $petG$, $petL$, $petN$
	Photosystem I	psaA, psaB, psaC, psaI, psaJ
	Photosystem II	psbA, psbB, psbC, psbD, spbE, psbF, psbH, psbI, psbJ, psbK, psbL, psbM, psbT, psbZ
	Large chain of rubisco	rbcL
Other genes	ATP-dependent protease	clpP
_	Cytochrome c biogenesis	ccsA
	Acetyl-CoA carboxylase	accD
	Membrane protein	cemA
	Maturase	matK
Unknown function	Hypothetical chloroplast reading frame	ycf1*, ycf2*, ycf3, ycf4

^{*}Duplicated gene.

(table 2). A total of 17 genes with introns were found in the chloroplast genome of *N. tangutorum*. Among these genes, *atpF*, *ndhA*, *ndhB*, *petB*, *petD*, *rpl16*, *rpl2*, *rpoC1*, *rps16*, *trnK-UUU*, *trnG-UCC*, *trnL-UAA*, *trnV-UAC*, *trnI-GAU*, and *trnA-UGC* had one intron, and *clpP* and *ycf3* gene contained two introns. The overall GC content of the *N. tangutorum* chloroplast genome was 37.3%.

Repeat and SSR analysis

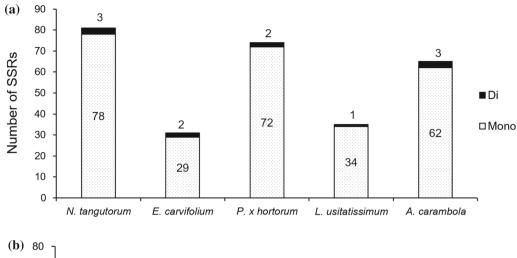
SSR markers in the N. tangutorum chloroplast genome were predicted using MISA, and compared with the chloroplast genomes of E. carvifolium, Pelagronium x hortorum, L. usitatissimum and A. carambola. In total, 81 SSRs were identified in the N. tangutorum chloroplast genome, including 78 mononucleotide repeats and three dinucleotide repeats, and no other type of SSR markers was found. The total numbers of the SSR repeats were 31, 74, 35 and 65 in E. carvifolium, P. x hortorum, L. usitatissimum and A. carambola, respectively (figure 2a). Similar to N. tangutorum, only mononucleotide repeats and dinucleotide repeats were found in the chloroplast genomes of E. carvifolium, P. x hortorum, L. usitatissimum and A. carambola. The total number of SSRs predicted from the N. tangutorum chloroplast genome were comparable to the chloroplast genomes of P. x hortorum and A. carambola, and were higher than those of E. carvifolium and L. usitatissimum.

A total of 66 repeats were identified from *N. tanguto-rum* chloroplast genome, including 41 tandem repeats,

10 palindromic repeats, and 15 forward repeats. The distribution of the repetitive sequences of different species in the chloroplast genomes of *N. tangutorum*, *E. carvifolium*, *L. usitatissimum* and *A. carambola* were similar: the tandem repeats is also the most abundant repeat category, followed by forward repeats and palindromic repeats. However, in the chloroplast genome of *P. x hortorum*, the numbers of the three categories of repeats (tandem repeat, forward repeat and palindromic repeat) are very similar. In addition, no reverse repeat was found in all these chloroplast genomes (figure 2b).

Phylogenetic analysis of N. tangutorum based on conserved protein sequences

A phylogenetic analysis was performed based on 19 complete chloroplast genomes of plant species in Sapindales, Geraniales, Malpighiales, Zygophyllales, and Oxalidales. The phylogenetic tree was constructed from the 54 proteincoding genes presented in all the 13 species using maximum likelihood (ML). RAxML was used to construct the ML tree with 1000-bootstrap replicates (Stamatakis 2014). The results indicated that *N. tangutorum* was clustered into a monophyletic group with the other eight plant species in Sapindales, and the five species in Geraniales were clustered in another clade (figure 3). Our results supported the taxonomic status of *N. tangutorum* defined in Liu and Zhou (2003, http://foc.iplant.cn/) and Angiosperm phylogeny website (Stevens 2001). In brief, the present study



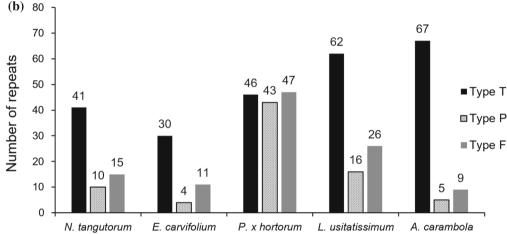


Figure 2. Repeat and SSRs analysis. (a) Numbers of SSRs in the cp genome of *N. tangutorum* compared with other four species. (b) Numbers of repeats in the cp genome of N. tangutorum compared with other four species. Type F, forward repeat; Type P, palindromic repeat; Type T, tandem repeat.

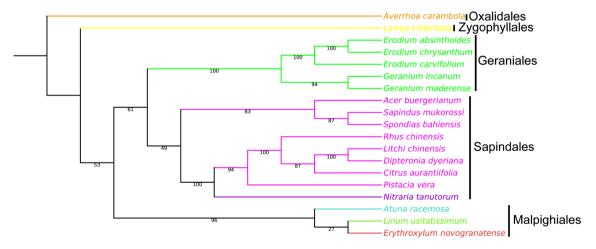


Figure 3. ML phylogenetic tree inferred from 19 chloroplast genome sequences. Numbers at nodes indicate bootstrap values.

characterized the complete chloroplast genome structure of N. tangutorum, and clarified phylogenetic relationships of N. tangutorum and relative taxa in rosids, which may be useful for further study of taxonomy and systematics of genus Nitraria.

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