One of wild Solanum species, Solanum brevicaule, belongs to a taxonomically confusing group (called S. brevicaule Bitter complex) within Solanum sect. Petota. The species is originating from the south of Peru to the north of Argentina and its ploidy level is diverse from diploid to hexaploid (Miller and Spooner 1999; Hardigan et al. 2015). The plant material used in this study is wild tuber-bearing diploid originating from Argentina. Its EBN (Endosperm Balanced Number) value of two theoretically makes it directly crossable for breeding purposes with cultivated tetraploid potatoes (S. tuberosum) (Hawkes 1990; Ortiz and Ehlenfeldt 1992; Cho et al. 1997; Spooner et al. 2014). The species was identified as a source of resistance to a nematode, Globodera pallida and a soft rot, Pectobacterium carotovorum (Jackson et al. 2014). The species is taxonomically confusing and can be used for potato breeding, therefore, the information of plastid genome of the wild species obtained in this study will provide an opportunity to investigate more detailed evolution and breeding aspects.

The S. brevicaule (PI205394) was sampled from Highland Agriculture Research Institute, South Korea (37.7° N, 128.7° E). An Illumina paired-end (PE) genomic library was constructed with total genomic DNA according to the PE standard protocol (Illumina, San Diego, USA) and sequenced using an Illumina HiSeq2000 at Macrogen (http://www.macrogen.com/kor/). Low-quality bases with raw scores of 20 or less were removed and approximately 2.5 Gbp of high-quality of PE reads were assembled by a CLC genome assembler (CLC Inc, Rarhus, Denmark) (Kim et al. 2015). The reference chloroplast genome sequence of S. berthaultii (KY419708, Park 2017; Kim et al. 2018) was used to retrieve principal contigs representing the chloroplast genome from the total contigs using Nucmer (Kurtz et al. 2004). The representative chloroplast contigs were arranged in order based on BLASTZ analysis (Schwartz et al. 2003) with the reference sequence and connected to a single draft sequence by joining overlapping terminal sequences. DOGMA (Wyman et al. 2004) and BLAST searches were used to predict chloroplast genes.

The complete chloroplast genome of S. brevicaule (GenBank accession no. MK036507) was 155,531 bp in length including 25,599 bp inverted repeats (IRa and IRb) regions separated by small single copy (SSC) region of 18,352 bp and large single copy (LSC) region of 85,981 bp with the typical quadripartite structure of most plastids, and the structure and gene features were typically identical to those of higher plants. A total of 158 genes with an average size of 583.0 bp were annotated including 105 protein-coding genes with an average size of 764.6 bp, 45 tRNA genes and 8 rRNA genes with an average size of 223.4 bp. An overall GC content was 37.25%.

Phylogenetic analysis was performed using chloroplast coding sequences of S. brevicaule and 30 published species in Solanaceae family by a maximum likelihood method in MEGA 6.0 (Tamura et al. 2013). According to the phylogenetic tree, S. brevicaule belonged to the same clade in Solanum species as expected (Figure 1).
Figure 1. Maximum likelihood phylogenetic tree of *S. brevicaule* with 30 species belonging to the Solanaceae based on chloroplast protein coding sequences. Numbers in the nodes are the bootstrap values from 1000 replicates.

**Disclosure statement**

The author reports no conflicts of interest. The author alone is responsible for the content and writing of this article.

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