Identification of Panax species and Panax ginseng cultivars using SNP markers

Yurry Um*, Eunjung Cho, Chan-Moon Chung, Yi Lee[†]

Department of Industrial Plant Science & Technology, Chungbuk National University Cheongju, Korea

Objectives

The single nucleotide polymorphism (SNP) has been widely used for species or variety identification from many crops. Direct sequencing of DNA fragments amplified by PCR from several individuals is one of direct way to identify single nucleotide polymorphisms. In this experiment, we carried out SNP discovery by direct sequencing of PCR fragments for identification of *Panax* species and *Panax ginseng* cultivars.

Materials and Methods

We used 2,549 genome survey sequence (GSS) of *P. ginseng* for design of 96 random PCR primer sets to amplify 500~700 bp fragments. DNA was extracted from leaf tissue of five *Panax* cultivars, Chunpoong, Yunpoong, Jakyungjong, *Panax quinquefolius* L., and *Panax notoginseng* Wall, and used for PCR amplification. The PCR products were directly sequenced with one of each primer set. SNPs were discovered from the sequence data with CodonCode Aligner software. We tested the reproducibility of the SNP markers using *Panax* cultivars.

Results

Sequence data of five *Panax* cultivars were obtained from the fragments amplified using 96 random PCR primer sets. We identified 251 SNPs or INDELs by aligning of the sequences using CodonCode Aligner program. Most of the SNPs showed polymorphism from different *Panax* species, but, only 6 SNPs showed polymorphism from *Panax ginseng* cultivars. We tested the reproducibility of the 6 SNPs using multiple samples of each *Panax ginseng* cultivars. Among the 6 SNPs, 944 showed reproducibility and differentiate Yunpoong from other *Panax ginseng* cultivars, Chunpoong and Jakyungjong. The other SNP positions were not reproducible.

[†] 주저자 연락처(Corresponding author): 이 이 E-mail: <u>leeyi22@cbnu.ac.kr</u> Tel: 043-261-3373

Table 1. Primer sets for SNP marker discovery in Panax ginseng cultivars

Marker #	GenBank #	Primer sequence($5' \rightarrow 3'$)	Tm(℃)	
903	BZ958992	Forward: TAATTCCTTGGGCACGTAGTAT	56	
		Reverse: TGATCCGTAAAGATCTCCACTT		
910	BZ958926	Forward: CGCCCCTAAAATAGGTATACAA	56	
		Reverse: AGGGATAAAATATTGATGATGGA		
937	BZ958782	Forward: GGAAAGTTATTTTCAACAAATGC		
		Reverse: AATTTCAGTATCTTCCATCTCTGAA	56	
944	BZ958754	Forward : GAAGTAAGAGAACGTCGAATGC	56	
		Reverse : CGTTTTGAAGAAAAAGAAGTGC		
951	BZ958717	BZ958717 Forward : GCATCGAGAACATCTCTGTAAAA Reverse : TTATGACATGTTACTGTTGCGA		56

Table 2. Haplotypes of Panax cultivars

Marker _	Panax ginseng C. A. Meyer			Panax	Panax	D 1 11. 11/4
	Chunpoong	Yunpoong	Jakyungjong	quinquefolius L.	notoginseng Wall	Reproducibility
903	G	A or G	G	A or G	A	NR
910	T	T	С	Т	С	NR
910	A	G	G	A	A	NR
937	G	T	G	Т	G	NR
944	A	G	A	G	G	R
951	A	_	A	-	A	NR
955	G	A	G	A	G	NR

R: Reproducible, NR: Not Reproducible