

P161

Soybean DnaJ-like Protein Enhance Tolerance in *E. coli* during Heat Shock

Hyun-A So, Eunsook Chung, Chang-Woo Cho, Boh-Hyun Yun, Jee-Sook Kang,
Yao Ran, Kyoungmi Kim and Jai-Heon Lee*

Department of Genetic Engineering, Dong-A University, Busan, 604-714, Republic of Korea

We isolated wound-inducible genes using suppression subtractive hybridization (SSH) method and were able to obtain to clone *w123* gene encoding dnaJ like protein. The full-length cDNA of *w123* is 689 bp with an open reading frame (ORF) consisting of 163 amino acid (aa). Genomic southern blot confirmed that soybean genome has two copies of *w123* gene. Northern blot analysis was also carried out for the gene expression during heat, NaCl, drought, wounding stresses. The expression of *w123* gene specifically induced by heat, NaCl, wounding and drought stress. Using GFP fusion vector, *w123*-smGFP was targeted both to nucleus. For the functional analysis of *w123*, His-tagged *w123* recombinant protein was heterologously expressed in *E. coli*. The *w123* recombinant cells showed enhanced heat tolerance compared to that of vector control cells. We suggest that dnaJ-like *w123* protein function as molecular chaperone in the nucleus of the plant cell during various stresses.

Key words: Soybean, heat shock, DnaJ, His-tagged fusion protein

P162

Molecular Characterization of *SLTI98* Encoding Ribosomal Protein Genes S6 from Soybean (*Glycine max*)

Jee-Sook Gang, Eunsook Chung, Kyoungmi Kim, Chang-Woo Cho, Jee-Eun Heo,
Bo Hyun Yun, Hyun-A So, Yao Ran, Jung-In Kim¹, Young-Choon Lee²,
Young-Soo Chung and Jai-Heon Lee*

Department of Genetic Engineering, Dong-A University, Busan, 604-714, Republic of Korea

¹*School of Food and Life Science, Biohealth Products Research Center, Inje University, Kimhae 621-749, Republic of Korea*

²*Department of Biotechnology, Dong-A University, Busan, 604-714, Republic of Korea*

In an attempt to better understand translational control during abiotic stresses, we isolated and characterized a stress inducible gene designated as *SLTI98* encoding ribosomal protein S6 in soybean. The derived amino acid sequence of *SLTI98* showed the highest identity of 93% with ribosomal protein S6 from *Medicago truncatula* (ABD32373). The size of the full-length genomic clone of *SLTI98* is 2,701 bp containing 6 exons and 5 introns, of which structure is similar to that of *Arabidopsis* ribosomal protein S6. Genomic southern blot analysis confirmed that soybean genome has two copies of *SLTI98* gene. RNA expression of *SLTI98* was mildly induced by salt stress, ABA and wounding stress, but not by dehydration stress. According to localization study using GFP fusion expression system, we were able to confirm that *SLTI98*-smGFP was restricted mostly to nucleus and partly to cytoplasm. The present study implies that the nuclear *SLTI98*, ribosomal protein S6 play an important role in translational control during abiotic stresses.

Key words: Ribosomal protein, stress, soybean, northern blot, GFP