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SLTI 114 gene has matrix mrtallo proteinase activity from Soybean (Glycine max)

<u>Jee-Eun Heo</u>, Jung-Haw Kang, Kyung-Mee Kim, Suk-Hee Ji, Dae-Kun Jin Young-Seob Yang, Jai-Heon Lee¹

Dept. of Plant Biotechnology, Dong-A University, Busan 604-714, Korea

Objectives

Functional analysis of SLTI114 gene.

Materials and Methods

1. Material

Plant - Glycine max. cv Sinpaldal 2, E.coil BL21, vector pET

2 Methods:

Construction of expression plasmid, Expresstion of the SLTI114 gene in E.coil

Results and Discussion

An electrophoresis in polyacrylimide gel (SDS-PAGE) was performed with SLTI 114 protein to get basic information of the activity of the metalloproteinase. The recombiant SLTI 114 protein purified by affinity chromatography on HisTrapTM Chelating HP, 1 ml column (Amersham) bound with Ni2⁺ions in 20mM phosphate buffer containing 0.2M NaCl and 10mM imidazole with a flow rate of 1 ml/min.

We found that the suspension-cultured cell infected with SLTI114 exhibited a higher proteolytic activity in degesting Azocoll, substrate for metalloproteinase activity assay (Graham et al.,1991)

The present study suggests that SLTI114 may have metalloproteinase activity.

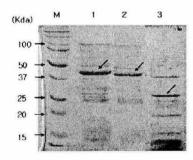


Fig. 1. SDS-PAGE analysis of the recombiant proteins produced in E. *coil* cells.

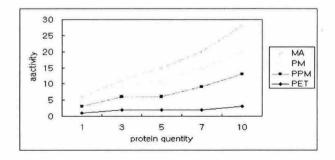


Fig. 2. The effect of metalloproteinase activity on differ protein quantity.

^{*}Corresponding author: Jai-Heon Lee, Tel: 051-200-7592, E-mail: jhnlee@dau.ac.kr