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The promoter of the gene encoding extensin like protein drives root-specific expression in transgenic plants

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Objectives

Isolation and functional characterization of root-specific promoter in transgenic plants

Materials and Methods

1. Material

Plant - *Arabidopsis thaliana* Columbia

Agrobacterium strain - GV3101

2. Methods:

The promoter region(2kb) of gene encoding extensin-like protein was amplified by PCR and cloned into binary vector pBGWFS7 using gateway cloning system and transformed into *Arabidopsis* by floral spray. The T1 transformants were obtained after basta treatment and T2 seedlings were subject for GFP/GUS expression analysis. Total RNA was isolated from roots, leaves, stem, flower, and silique and cDNA was made from total RNA using the SuperScript first-strand synthesis system(Gibco BRL).

Results and Discussion

The promoter region of the gene encoding extensin like protein was isolated by PCR using whole genome databases of *Arabidopsis thaliana*. The promoter region was fused to a reporter gene encoding green fluorescent protein(GFP) and beta-glucuronidase(GUS) in a binary vector pBGWFS7 using gateway cloning system and was introduced into *Arabidopsis* plants. GUS staining was detected in root tissue of *Arabidopsis* T2 seedlings. The transient expression assays in carrot and sweet potato indicated that GUS activity of this promoter is comparable to its activity of CaMV 35S promoter in root. RT-PCR revealed that the extensin-like protein gene was specifically expressed in roots of *Arabidopsis* plant. A series of 5'-deletions of this promoter were cloned to GFP/GUS reporter vector and transformed into *Arabidopsis*. We are now going to find potential cis elements that account for the tissue-specificity of this promoter.

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