Genetic and genomic studies of shoot and flower growth in Arabidopsis

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in memory of my grandfather, Edwin Pratt Jordan

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## Abstract

This thesis is organized around the theme of modulation of transcriptional states in *Arabidopsis thaliana*. The two particular mechanisms on which this work focuses are (1) microRNA-mediated negative regulation of protein levels (either by mRNA cleavage or by repression of translation) and (2) transduction of extracellular signals into the cell to affect the transcription program.

Chapter 2 characterizes the role of the *EARLY EXTRA PETALS* (*EEP1*) microRNA in the regulation of organ formation in the flower and shoot. The *eep1* loss-of-function mutant has extra petals, and it enhances the shoot phenotype of the *pinoid* mutant, which has defects in auxin signaling and organ formation. *EEP1* is nearly identical to a pair of published miRNAs (*MIR164a* and *b*); all three are predicted to target the mRNAs of six genes in the NAC family of transcription factors. Two of these genes, *CUPSHAPED COTYLEDONS1* and 2 (*CUC1* and 2), are redundantly required in flower development. Phenotypic and molecular analysis of lines overexpressing *EEP1* are consistent with (1) negative regulation of *CUC1* and *CUC2* by *EEP1* and (2) cleavage of the *CUC2* mRNA promoted by *EEP1*.

Chapter 3 describes the investigation, by reverse genetics, of five proteins encoded by genes in the *CLV3/ERS* (*CLE*) family. Due to the similarity of these proteins to CLAVATA3 (CLV3), the likely secreted ligand for the CLAVATA1 receptor-like kinase, functional analyses were performed in order to determine whether these proteins might also function as ligands for CLV1 or other receptor-like kinases. The results presented here derive from experiments using overexpression, double-stranded RNA interference (dsRNAi), and promoter-glucuronidase (GUS) reporter expression.

## **Table of Contents**

Acknowledgmentsi	V
Abstractv	ii
List of Figures	x
Chapter 1 Transcription and modification of transcriptional states in	
Arabidopsis	1
1.1 Transcription in Arabidopsis development	1
1.2 Inputs from outside the cell can alter the transcription program	4
1.3 MicroRNAs modulate gene expression post-transcriptionally	8
1.4 New insight into microRNA function in plant development	.12
1.5 Advances in functional characterization of proteins in the CLE family	13
1.6 References	.14
Chapter 2 A role for the <i>EEP1</i> miRNA in <i>Arabidopsis</i> flower development	20
2.1 Introduction	20
2.2 Results	25
2.3 Discussion	46
2.4 Materials and methods5	54
2.5 Acknowledgments	58
2.6 References	58
Chapter 3 Functional analysis of five CLE proteins by reverse genetics	52
3.1 Introduction	52
3.2 Results	67
3.3 Discussion	76
3.4 Materials and methods	79

	ix
3.5 Acknowledgments	83
3.6. Patarancas	83
5.0 Rejerences	

## List of Figures

<b>Figure 2.1</b> <i>eep1</i> has extra petals in early flowers and slightly defective septa27
<b>Figure 2.2</b> Double mutants with <i>eep1</i> and other extra petal mutants30
Figure 2.3 <i>eep1</i> severely enhances the <i>pinoid</i> ( <i>pid</i> ) phenotype in both the shoot
and the flower
Figure 2.4 <i>eep1</i> partially suppresses the delayed flowering of <i>pinformed</i> ( <i>pin1</i> ) and
further reduces sepal number
<b>Figure 2.5</b> The mapping and identification of the <i>eep1</i> mutation
<b>Figure 2.6</b> <i>EEP1</i> is a putative microRNA closely related to <i>MIR164a</i> and <i>b</i> 39
Figure 2.7 The <i>EEP1</i> promoter drives <i>GUS</i> expression in young floral buds of
wild-type plants40
<b>Figure 2.8</b> A strong line of <i>35S::EEP1</i> is very similar to <i>cuc1; cuc2</i> double
mutants, and CUC1 and CUC2 mRNA levels are decreased in this line relative to
wild type45

Figure 3.1 CLAVATA3, a protein required for proper meristem regulation, is the
founding member of the CLV3/ESR (CLE) family66
Figure 3.2 Overexpression of several CLE genes results in specific
phenotypes69
Figure 3.3 The 35S::CLE12 phenotype requires wild-type CLV1 activity, but not
CLV370
<b>Figure 3.4</b> CLE12 can functionally replace CLV3 in the meristem72
Figure 3.5 The <i>CLE</i> genes have discrete but overlapping expression patterns in
the inflorescence