

RESEARCH ARTICLE

Open Access

Analysis of promoter activity of members of the *PECTATE LYASE-LIKE (PLL)* gene family in cell separation in *Arabidopsis*

Lingxia Sun, Steven van Nocker*

Abstract

Background: Pectate lyases depolymerize pectins by catalyzing the eliminative cleavage of α -1,4-linked galacturonic acid. Pectate lyase-like (*PLL*) genes make up among the largest and most complex families in plants, but their cellular and organismal roles have not been well characterized, and the activity of these genes has been assessed only at the level of entire organs or plant parts, potentially obscuring important sub-organ or cell-type-specific activities. As a first step to understand the potential functional diversity of *PLL* genes in plants and specificity of individual genes, we utilized a reporter gene approach to document the spatial and temporal promoter activity for 23 of the 26 members of the *Arabidopsis thaliana* (*Arabidopsis*) *PLL* gene family throughout development, focusing on processes involving cell separation.

Results: Numerous *PLL* promoters directed activity in localized domains programmed for cell separation, such as the abscission zones of the sepal, petal, stamen, and seed, as well as the fruit dehiscence zone. Several drove activity in cell types expected to facilitate separation, including the style and root endodermal and cortical layers during lateral root emergence. However, *PLL* promoters were active in domains not obviously programmed for separation, including the stipule, hydathode and root axis. Nearly all *PLL* promoters showed extensive overlap of activity in most of the regions analyzed.

Conclusions: Our results document potential for involvement of *PLL* genes in numerous aspects of growth and development both dependent and independent of cell separation. Although the complexity of the *PLL* gene family allows for enormous potential for gene specialization through spatial or temporal regulation, the high degree of overlap of activity among the *PLL* promoters suggests extensive redundancy. Alternatively, functional specialization might be determined at the post-transcriptional or protein level.

Background

Pectins, a class of polysaccharide polymer typically characterized by a linear backbone of α -1,4-linked galacturonic acid (GalA) residues, are a major component of primary plant cell walls, and within the wall form a matrix in which a network of cellulose and hemicellulose is embedded [1-3]. Pectins contribute to the mechanical strength and physical properties of primary cell walls, but also function in intercellular adhesion [4-6], and can act as signaling molecules in morphogenesis and pathogen defense [2].

Plant growth and development is accompanied by dynamic remodeling of the cell wall, which in turn requires modifications of the various cell wall components including pectin. In accordance with the often complex structure of some pectins, an assortment of pectinase activities modify or degrade these polymers. Pectinesterases target methyl-esterified homogalacturonan, yielding substrates for polygalacturonases and pectate lyases, which cleave the GalA backbone [6]. Rhamnogalacturonases and rhamnogalacturonan lyases depolymerize branched regions of rhamnogalacturonan, whereas β -galactosidases and α -arabinosidases can degrade the galactan/arabinan/arabinogalactan side chains. Pectate lyases (EC 4.2.2.2) have been most extensively studied in *Erwinia chrysanthemi*, a major causal agent of soft-rot

* Correspondence: vannocke@msu.edu
Department of Horticulture, Michigan State University, East Lansing, MI, USA

diseases that affect a wide range of plant species [7-9]. Their action not only results in maceration of plant tissues but also can activate plant defense systems [10-12]. Plant *PECTATE LYASE-LIKE* (*PLL*) genes encode proteins with strong amino acid sequence homology with the PelC isoform of bacterial pectate lyases [13,14] and exist as large families in plants where studied [15]. The abundance of *PLL* genes in plants, including 26 in Arabidopsis, 12 in rice, and 22 in poplar, has arisen from multiple gene duplication events [16,17], a process that may enhance plasticity in adaptation to changing environments [18]. Models predict that gene redundancy is evolutionarily stable only when duplicated genes differentiate in some aspect of their function, suggesting that individual members of large families such as *PLL* may have some unique function [18].

Various analyses of *PLL* genes from plants revealed expression in a broad range of organs and plant parts including root, leaf, flower, pollen, filament, style, pistil, and ripening fruit [14,16,19-24]. *PLL* genes in several studies also showed elevated expression in response to auxin, wounding, and/or pathogen infection [16,25-28]. Transcriptional analysis in Arabidopsis revealed that a small subset of *PLL* genes were up-regulated during stamen abscission or in cortical cell separation during the emergence of the lateral root [16,29,30]. Reduced expression of strawberry *PL1* in transgenic strawberry plants suggested a natural role in tissue softening during fruit ripening [31]. All of these data implicate *PLL* genes in various plant growth and development events. However, the activity of these genes has been characterized only at the level of entire organs or plant parts, potentially obscuring important sub-organ or cell-type-specific activities.

During plant growth and development, there are many events in which adjacent cells separate in a coordinated manner. Cell separation resulting in organ abscission, or anther or fruit dehiscence, occurs in predetermined positions, called abscission zones (AZs) or dehiscence zones (DZs), respectively [32,33]. Intercellular space formed in leaves and stems can result from restricted separation of cells at the tricellular regions [34]. Fruit ripening also involves limited cell separation, in which only the middle lamella is degraded, with tricellular junctions and plasmodesmata often remaining intact [35,36]. Targeted cell separation is also involved in the processes of seed germination, lateral root emergence, pollen tube penetration through the transmitting tract, and shedding of columella root cap cells [37,38]. In this study, we examined spatial-temporal expression patterns of promoters of *PLL* family members, focusing on the various cell separation and wall loosening events that occur during Arabidopsis growth and development.

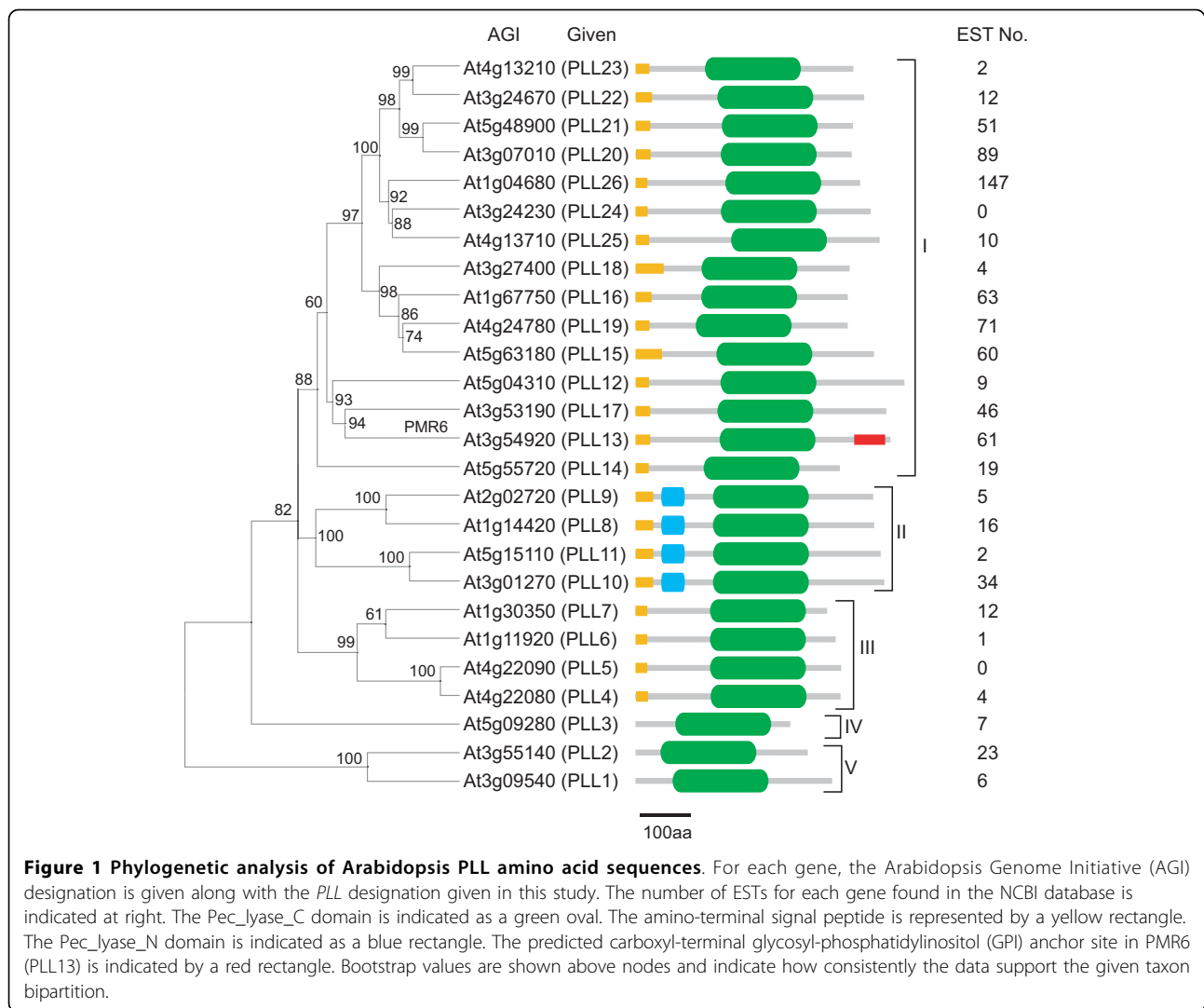
Results and Discussion

Phylogenetic analysis

Based on peptide sequence homology with the enzymatically characterized pectate lyase PL1 from banana and annotated protein domains, the Arabidopsis genome encodes for 26 pectate-lyase-like proteins. Neighbor-joining analysis partitioned these protein sequences into five subfamilies (**Figure 1**). This genomic content and phylogenetic organization is similar to that previously reported for *PLLs* from multiple plant species including Arabidopsis [16,21]. All Arabidopsis *PLLs* exhibited a region of sequence homology with pectate lyase C (pelC) from *Erwinia chrysanthemi*, annotated as the Pec_lyase_C (Pfam00544) domain (**Figure 1**) and 23 of the proteins contain a probable amino-terminal signal peptide. The four members included in Subfamily II (*PLL8-11*) also exhibit a Pec_lyase_N (Pfam04431) domain, a short region of high homology whose structure or function has not been described. The predicted carboxyl-terminal glycosyl-phosphatidylinositol (GPI) anchor site previously identified in PMR6 (POWDERY MILDEW RESISTANCE 6, also called *PLL13*) was not obviously present in other *PLLs*, suggesting a specialized function of PMR6 associated with powdery mildew susceptibility [28] (**Figure 1**).

Estimation of *PLL* gene expression through analysis of public transcriptome data

As a first step to assess expression pattern of *PLL* genes in Arabidopsis, we analyzed publicly available transcriptome data. Expressed sequence tags (ESTs) found in public databanks corresponding to individual *PLL* gene family members were found to be sourced from various tissues and stages across Arabidopsis growth and development, as well as various environmental conditions (not shown). Representational frequency of ESTs for individual *PLL* genes was highly variable both across the gene family and within subfamilies (**Figure 1**). Analysis of publicly available microarray data also suggested that there was at least some unique developmental pattern of expression for most *PLL* genes (see **Additional file 1**). A striking exception are the four members of Subfamily II (*PLL8-PLL11*), which showed a similar expression pattern localized predominately to pollen and stamens (see **Additional file 1**). Several genes (*PLL3, 4, 6, 7, 12, and 24*) were transcriptionally silenced across most sampled tissues, whereas *PLL2* was apparently expressed ubiquitously, suggesting a very general function (see **Additional file 1**). These data are generally consistent with the results of qualitative RT-PCR analysis of *PLL* genes [16] and support a collectively ubiquitous function for *PLLs* in growth and development and the potential for functional specialization by many of these genes.



Analysis of *PLL* promoter activity

Direct approaches for analysis of gene expression such as RNA-seq, microarray analysis and RT-PCR can be problematic when applied to large gene families, due to sequence homology among family members and/or inability to resolve expression at the cellular level. Accordingly, we utilized a histological reporter gene (*β-glucuronidase*, *GUS*) to determine the expression regulatory potential of individual *PLL* gene promoters. We engineered the *GUS* coding sequence adjacent to up to ~2 kb of 5' UTR/promoter sequence, preserving the authentic start codon of the *PLL* genes, and expressed the *PLL::GUS* fusions in transgenic Arabidopsis (see **Additional file 2**). For each *PLL* gene, we analyzed at least four independent transgenic lines. *GUS* activity patterns were generally consistent between independent lines, with the exception of three (*PLL6*, *PLL12*, and *PLL14*) that showed weak and variable activity and were

not analyzed further. No *GUS* activity was observed in transgenic plants transformed with a promoterless *GUS* construction, or in non-transgenic plants, in any plant part or under any condition used for analysis. Based on the known roles for pectins in cell adhesion and cell wall architecture, we focused our analysis on developmental events closely associated with cell separation.

PLL::GUS expression associated with cell separation

Floral organ AZs. Cell separation in Arabidopsis has been best characterized in the context of floral organ abscission [32]. Arabidopsis exhibits abscission of sepals, petals, and stamens following pollination. Abscission is conditioned by cell separation within the AZ, a tightly localized region at the base of the floral organs [39]. We analyzed developing flowers at various stages until Stage 18, when siliques began to yellow [40]. (**Figure 2** and see below). *GUS* activity within the AZs was observed

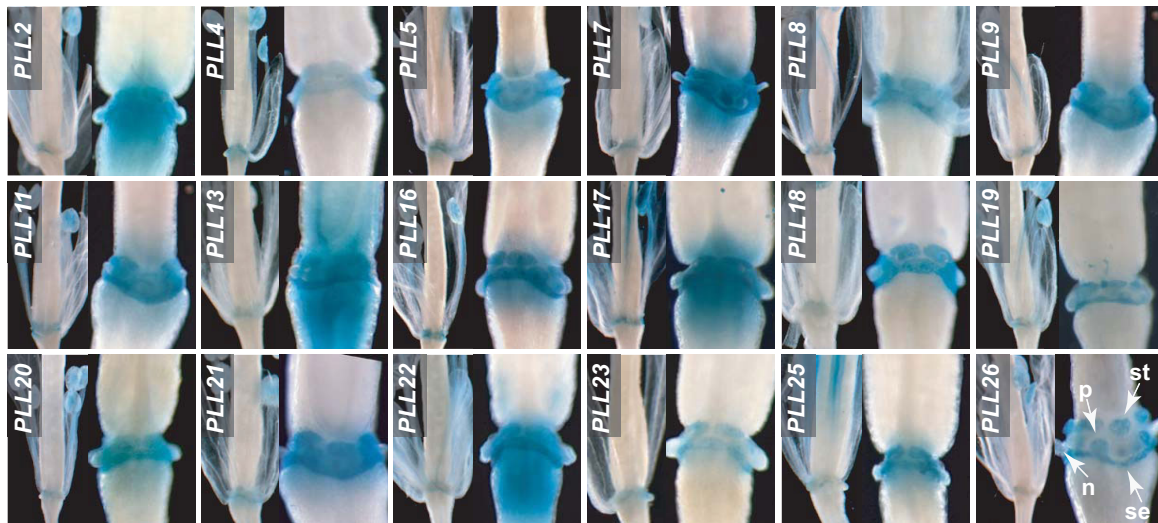


Figure 2 Spatial and temporal *PLL::GUS* expression in the AZs of sepals, petals, and stamens. Each frame is a composite of two independent photographs showing flowers at Stage 16 (left) and Stage 17 (right). Arrowheads in the *PLL26* panel exemplify the location of AZs of the sepal (se), petal (pe), and stamen (st). A nectary (n) is indicated.

for 18 *PLL* promoters (Figure 2 and see Additional file 3). For all of these, GUS activity was first detected in the AZs of sepals, petals, and stamens at Stage 16, which followed anthesis by about 2 days and was marked by the withering of sepals and petals (Figure 2) [40]. GUS activity had increased within the AZs of all three organ types by Stage 17, when perianth organs were abscising (Figure 2). For all *PLL* promoters, GUS activity in the AZs was detectable but very weak at Stage 18 (not shown). Activity for *PLL26* in floral organ AZs was apparent through Stage 19, marked by valve separation (not shown). These results are consistent with those of Cai and Lashbrook [29], who identified *PLL18* as substantially upregulated in the stamen AZ through transcriptional profiling of laser-capture-microdissected tissues. This upregulation was apparent in the latter part of Stage 15, following anthesis by approximately 36 h [29,40].

Fruit DZ and seed AZ

The Arabidopsis fruit consists of two valves separated by a replum. Mature fruits dehisce due to cell separation in the so-called DZ, which is distributed along the valve margins and includes a separation layer [41]. Mature seeds are released from the funiculus, a stalk-like structure connecting seeds to the replum, at a site referred to as the seed AZ. We analyzed GUS activity in these regions during development of flowers and fruit from Stage 16 through Stage 20, marked by seed abscission [40] (Figure 3A). We found that 17 of the *PLL* promoters

drove GUS activity within the apparent DZ of developing siliques. This subset of *PLLs* nearly completely overlapped with the subset driving expression in floral organ AZs (see Additional file 3). For all, GUS activity was evident at the onset of stage 18, which preceded separation of the valves by approximately 24 h (not shown). GUS activity was first seen at the basal and apical ends of siliques, where valve separation was initiated, and then became established along the entire length of the fruit as valve separation progressed (Figure 3A and not shown).

A mostly overlapping subset of 16 *PLL* promoters drove GUS activity within the apparent seed AZ (Figure 3B and see Additional file 3). For all of these, GUS activity was not observed at the beginning of stage 19, but was obvious in seeds that were still attached to the funiculus and septum before the beginning of Stage 20 (e.g., *PLL19* in Figure 3B).

Radicle emergence during seed germination

During seed germination in Arabidopsis, the radicle penetrates a single endosperm cell layer at the micropylar end of the seed [42]. To assess *PLL* promoter activity in this region, we stained seeds for GUS activity 24 h after imbibition, when approximately half of the seeds showed rupture of the testa but before radicle emergence was seen (not shown). Two *PLL* promoters (*PLL16* and *PLL22*) directed GUS activity in germinating seeds; in both cases GUS activity was observed in the micropylar end before obvious emergence of the radicle (Figure 4A and not shown).

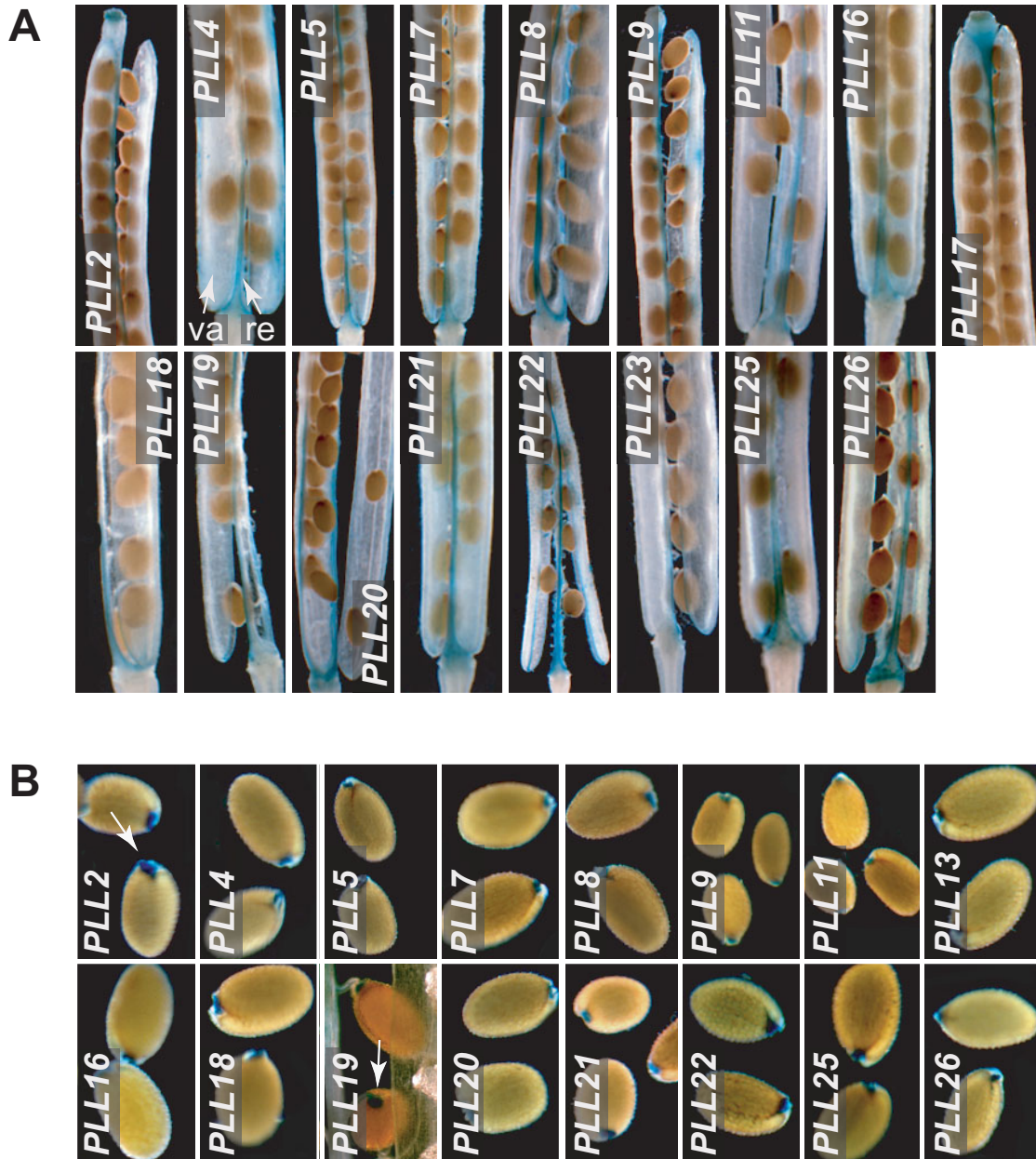
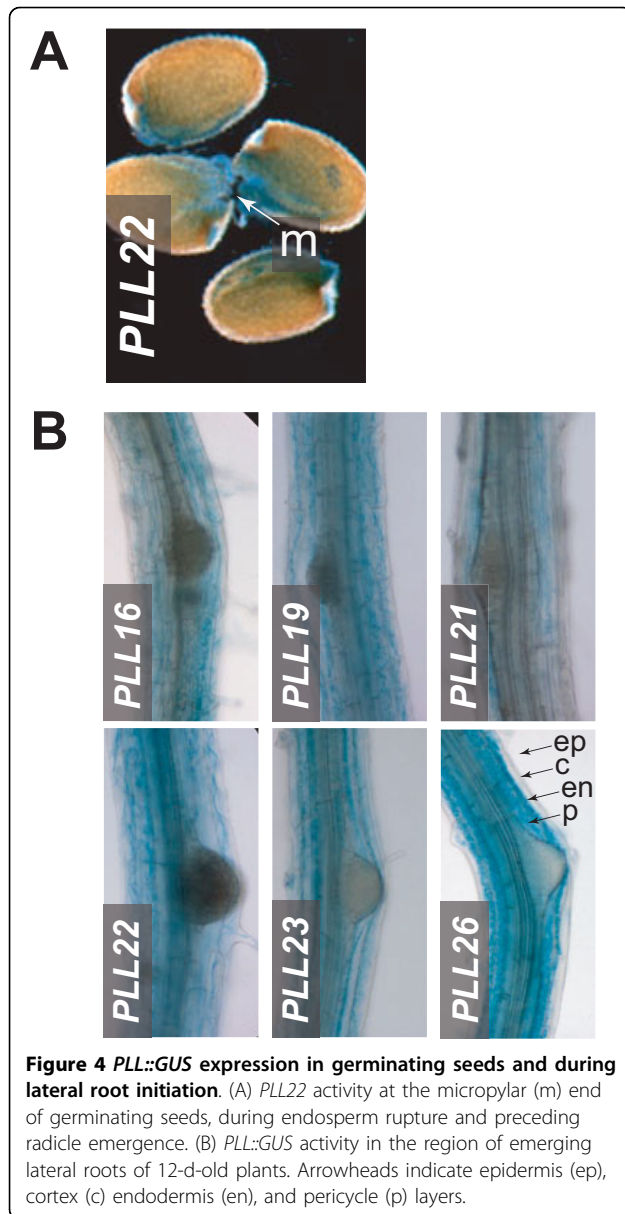


Figure 3 *PLL::GUS* expression in the silique DZ and seed AZ. (A) Siliques at Stage 19, beginning to dehisce and showing GUS activity in the DZ. The silique valve (va) and replum (re) are indicated with arrowheads in the *PLL4* panel. (B) Recently abscised, mature seeds showing GUS activity in the region of the seed AZ (arrowhead in *PLL2* panel). The arrowhead in the *PLL19* panel shows GUS activity in this region just before abscission.

Lateral root initiation

Arabidopsis lateral roots initiate from the pericycle cell layer and must penetrate the overlaying endodermal, cortical and epidermal layers during emergence [43]. Although several *PLL* promoters drove GUS activity at relatively weak levels along the axis of primary roots (see below), six (*PLL16*, *19*, *21-23*, *26*) showed strong activity in primary roots of 12-d-old plants that were

initiating lateral roots. This activity was confined to the endodermal and cortical layers in the region of initiation, but was not obviously restricted to cell layers directly overlaying the new lateral root primordia (**Figure 4B** and **not shown**). We did not observe strong GUS staining in the epidermis for any of the six promoters (**Figure 4B** and **not shown**). These findings are consistent with the increased transcriptional



accumulation previously reported for *PLL16* and *PLL26* during lateral root emergence induced by auxin [26,30].

PLL::GUS expression in other developmental contexts

To analyze *PLL* promoter activity in a broader developmental context, including processes potentially independent of cell separation, we undertook a comprehensive analysis of the activity of all of the studied *PLL* promoters during growth and development of Arabidopsis.

Seedlings

GUS activity was detected in 1-d-old seedlings for 22 *PLL* promoters. In seedlings, *GUS* staining was seen in the

cotyledons, hypocotyl, root/hypocotyl junction, root axes, and/or root tips (Figure 5A and see Additional file 3). A subset of 11 of these promoters also drove *GUS* activity along the root axes in 5-d-old seedlings (Figure 5B and see Additional file 3); this subset included the six that were strongly active during lateral root initiation (above). Four promoters, *PLL13*, *PLL17*, *PLL22*, and *PLL25*, were active in the root tips. *PLL25* was unique in that it drove *GUS* activity specifically in the root tip. Both *PLL22* and *PLL25* drove expression in a broad region that included the columella root cap (Figure 5B).

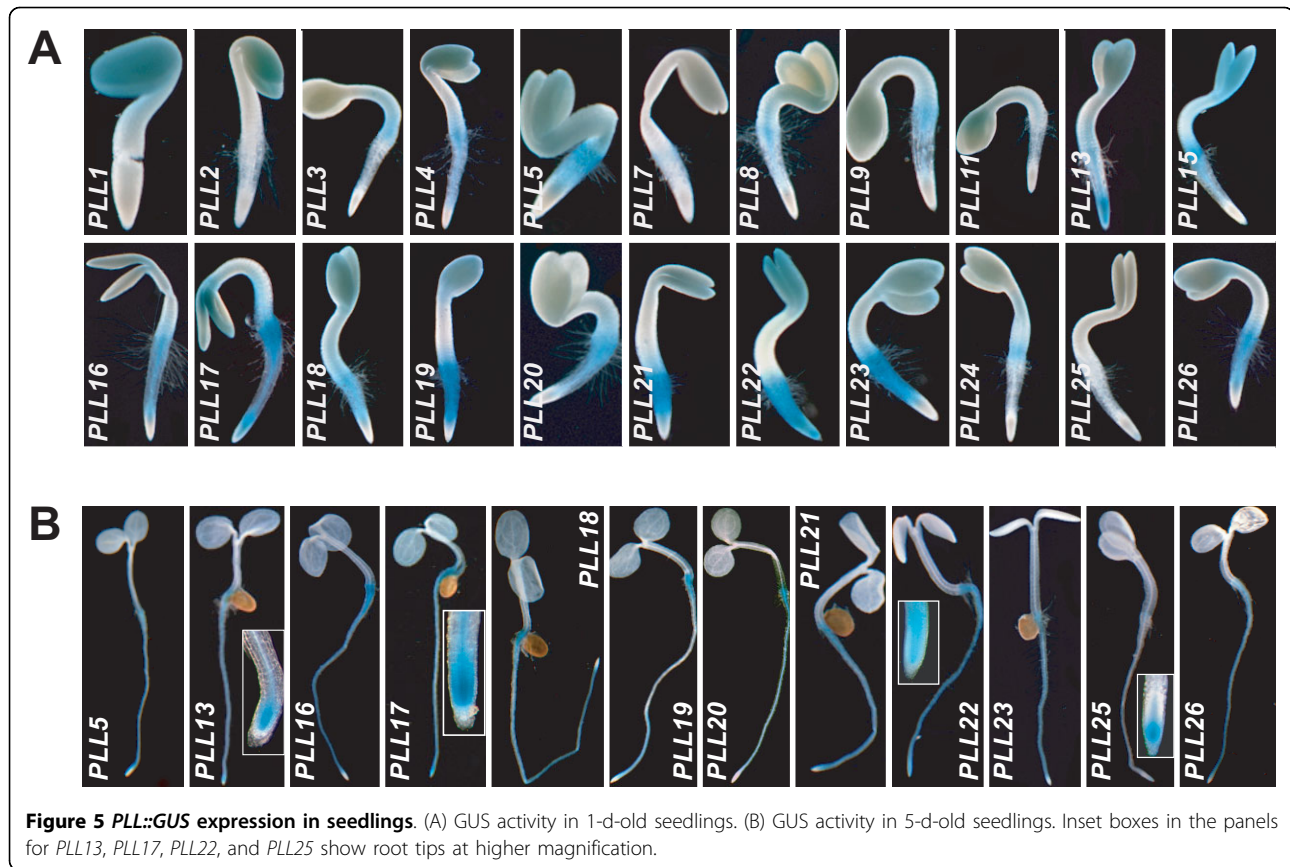
Hydathodes and Stipules

We observed *GUS* activity in fully expanded cotyledons and rosette leaves for seven *PLL* promoters (Figure 6A). For all seven, activity was restricted to the tip of the cotyledon/leaf. This corresponds to the position of the hydathodes, specialized pore-like structures that release water and solutes from the xylem [44].

GUS activity was apparent within the shoot apex of plants in the vegetative phase for 19 *PLL* promoters. Upon close examination, in all cases this activity appeared in tightly localized foci bordering the base of initiating leaves, including cauline leaves (Figure 6B and not shown), at the position of the stipules. In Arabidopsis, stipules are apparent on the newly formed leaves but become less obvious upon leaf expansion. For all *PLLs*, *GUS* activity was strongest in those leaves that had initiated and were undergoing expansion, and weakest or not apparent in fully expanded leaves (Figure 6C and not shown). The function of stipules in Arabidopsis development remains largely unknown. However, it has been demonstrated that stipules are primary sites for the accumulation of auxin associated with vascular differentiation and leaf morphogenesis [45,46]. No *GUS* activity was observed for any *PLL* promoter in the apex after the transition to flowering (not shown).

Base of flower/fruit pedicel

Two *PLL* promoters (*PLL15* and *PLL24*) drove *GUS* activity within a restricted region on the adaxial side of the base of pedicels (Figure 7A). For both, this was apparent after Stage 12 of floral development, just before anthesis, and through development of the green silique (not shown). The *PLL15* and *PLL24* promoters, in addition to *PLL1* and *PLL22*, also drove *GUS* activity at the base of trichomes on the leaf epidermis (Figure 7B and see Additional file 3). In all cases, *GUS* activity was limited to the stellate trichomes on the adaxial surface of rosette leaves, in the region roughly corresponding to basement cells, a group of specialized cells encircling the trichome and potentially serving as structural and/or biochemical support for the trichome (Figure 7B).



Flowers

GUS activity was detected in developing flowers for all 23 of the *PLL* promoters (Figure 8 and see Additional file 3). None of the promoters studied showed strong activity before Stage 11, which is marked by appearance of stigmatic papillae [40] (not shown). All except *PLL10* drove GUS activity in varying domains within the style and/or stigma (Figure 8A and 8B and see Additional file 3). Some of these showed localized expression in the style (e.g., *PLL1*), whereas others showed expression throughout the stigma (e.g., *PLL2*), or in the basal portion of the stigma extending into the style (e.g., *PLL20*). A potential function for *PLL* genes expressed in the stigma and style is to facilitate pollen tube growth, via separation of transmitting tract cells and/or remodeling of cell walls in this tissue. In the stamen, several promoters drove activity in the filament (e.g., *PLL2*), and/or throughout the anther and predominantly localized to pollen sacs and/or developing pollen (e.g., *PLL10*). The subset of *PLLs* driving expression in pollen is largely consistent with that determined based on transcriptome analysis (see Additional file 1) or RT-PCR [16]. *PLL3* and *PLL25* were distinguished by driving GUS activity in the stamen limited to the junction of the filament and anther (Figure 8B and see Additional file 3).

PLL13 drove expression in the style in immature flowers (Figure 8A), but by Stage 15 this activity was not apparent, and instead GUS staining was seen in developing seeds (Figure 8B). Activity of *PLL13* was not seen in ovules at Stage 12 or earlier (Figure 8A), or in developing seeds at Stage 16 or later (Figure 2). We hypothesize that this seed-associated pattern is associated with the cellularization of the endosperm [47].

Correlation or divergence of *PLL* promoter activity with patterns of RNA accumulation

To evaluate the potential for regulation of *PLL* genes outside of the context of promoter activity, we analyzed our findings in relation to previously reported transcriptional patterns based on microarray analysis of Arabidopsis organs or whole structures (see above and Additional file 1[48]).

Most notably, the strong promoter activity found in anthers for the four members of Subfamily II (*PLL8-PLL11*) (Figure 8) corresponded well with the stamen-associated pattern of mRNA localization (see Additional file 1). However, *PLL8*, *9*, and *11* also showed strong promoter activity variously outside of flowers in the hypocotyl (*PLL11*), root/hypocotyl junction (*PLL8* and *9*), and stipules (*PLL8* and *11*), regions where transcriptional

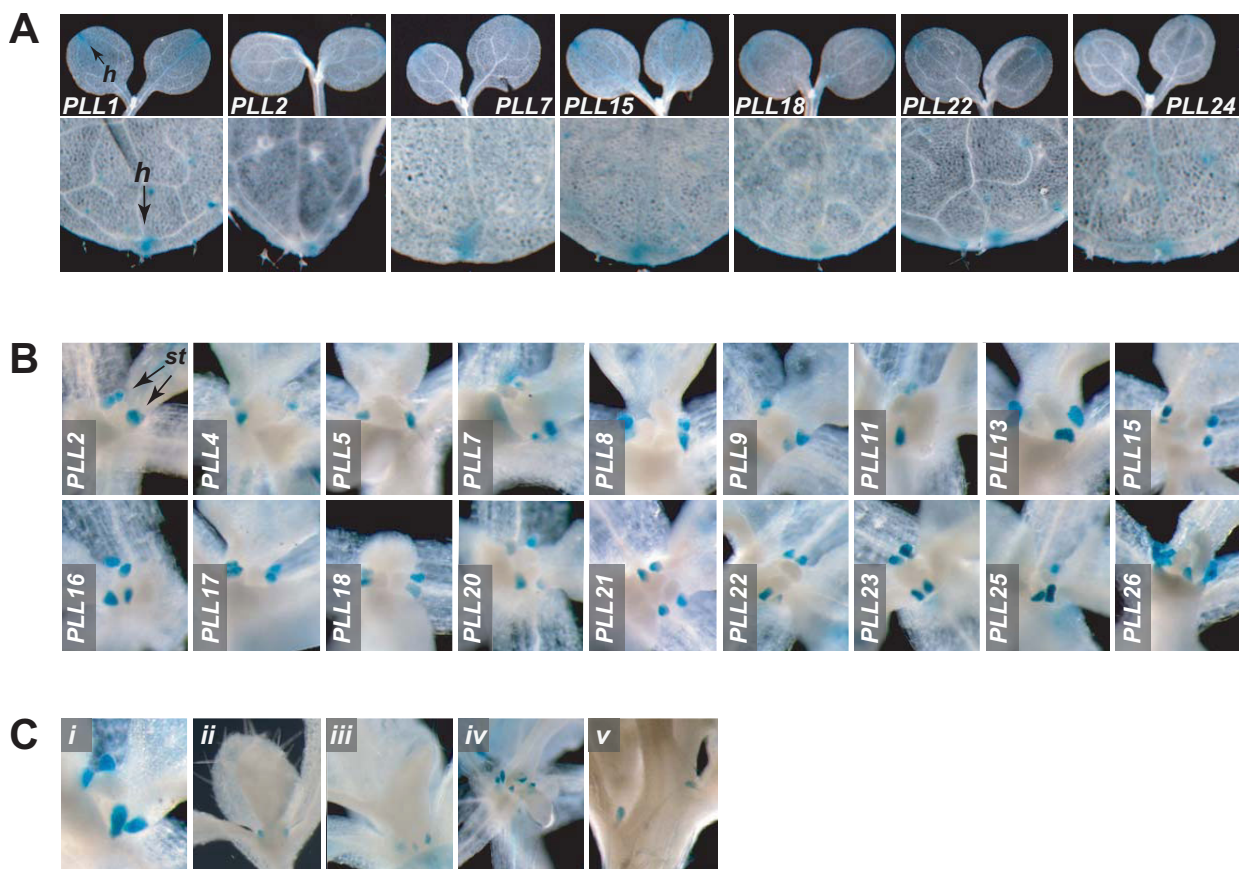


Figure 6 *PLL::GUS* expression in hydathodes and stipules. (A) *GUS* activity in hydathodes. In the upper row, 5-d-old seedlings with fully expanded cotyledons are shown. In the lower row, the distal tips of the first rosette leaf on 8-d-old seedlings are shown. The position of the hydathode (h) is indicated for the *PLL1* panels. (B) *GUS* activity in the stipules of an expanding leaf of 6-d-old seedlings. (C) *GUS* activity in stipules of *PLL19::GUS* plants at various stages of plant and leaf development: (i) expanding ~3 mm leaf of 6-d-old seedling; (ii) expanding ~6 mm leaf of 6-d-old seedling; (iii) nearly fully expanded leaf (~12 mm) of 8-d-old seedling; (iv) apex of 15-d-old plant showing staining in stipules of primordial cauline leaves; (v) flowering plant showing staining in the stipules of fully expanded cauline leaves

profiling did not obviously predict promoter activity. Genes expected to be transcriptionally quiescent based on microarray analysis, such as *PLL4* or *PLL24*, showed promoter activity that was relatively strong, and that was similar to genes that showed strong transcriptional signals such as *PLL8* or *PLL15*, respectively. *PLL2*, which might be expected to be constitutively expressed as a 'house-keeping' gene based on microarray analysis, showed promoter activity in only a subset of the regions analyzed (see **Additional file 3**). At least part of these apparent inconsistencies between promoter activity analysis and transcriptional profiling can be attributed to our qualitative representation of promoter activity, which is in reality a quantitative function. The lack of high values for mRNA accumulation in structures (e.g., the silique) that include tissues with strong promoter activity (e.g., the DZ) could be simply explained by the fact that such

tissues make up only a small portion of the bulk of the regions subject to analysis. Other cases where mRNA levels are low in spite of strong regional promoter activity might be attributed to rapid mRNA turnover. It is most difficult to explain the accumulation of mRNAs in tissues or structures that lack detectable promoter activity, for example the strong and ubiquitous mRNA expression of *PLL2*. In these cases regulatory elements required for appropriate expression might lay outside of the ~2 kb 5' region utilized to direct *GUS* expression.

Conclusions

In this study we determined the temporal and regional promoter activity for members of the *PLL* gene family during growth and development of *Arabidopsis*. Given the ubiquitous role of pectins in intercellular adhesion and cell wall architecture, and the various organismal

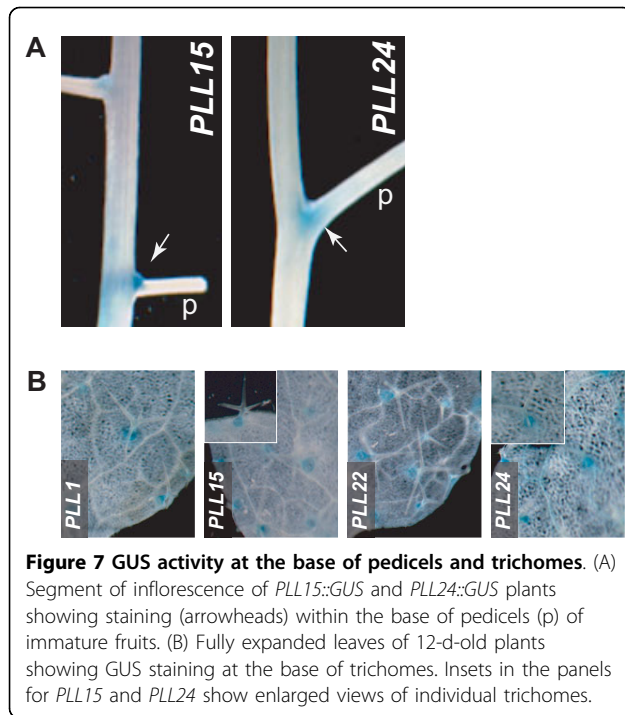


Figure 7 GUS activity at the base of pedicels and trichomes. (A) Segment of inflorescence of *PLL15::GUS* and *PLL24::GUS* plants showing staining (arrowheads) within the base of pedicels (p) of immature fruits. (B) Fully expanded leaves of 12-d-old plants showing GUS staining at the base of trichomes. Insets in the panels for *PLL15* and *PLL24* show enlarged views of individual trichomes.

processes that require targeted cell separation or cell wall remodeling, these genes are expected to be of fundamental importance.

We found that, as expected, promoter activity was common in cell types programmed for abscission, most obviously the perianth floral organ AZs, the DZ of the fruit, and the seed AZ. Activity was also seen in regions, such as cell layers in the region of initiating lateral roots, where cell separation is expected but has not been well characterized. Pectate lyase activity is expected to act in concert with other cell wall remodeling activities both to condition cell separation and to modify newly exposed cells for barrier functions. Interestingly, we saw more limited *PLL* promoter activity at the base of pedicels of flowers and fruits. Cells in these regions can be artificially programmed to undergo separation in *Arabidopsis*, in response to ectopic activity of the small abscission-associated peptide IDA [49], suggesting an abscission-related role. If these *PLL* genes do have a cryptic function in flower/fruit abscission, abscission may be normally limited by the degree of pectate lyase activity or deficiency of other, cooperative, cell wall-modifying activities. Interestingly, the two *PLL* promoters driving activity in this region also were active in cells at the base of trichomes, suggesting their pedicel- and trichome-associated roles may be analogous. Both regions are subject to mechanical stress imparted by movement of the pedicels/trichomes and an alternative function for

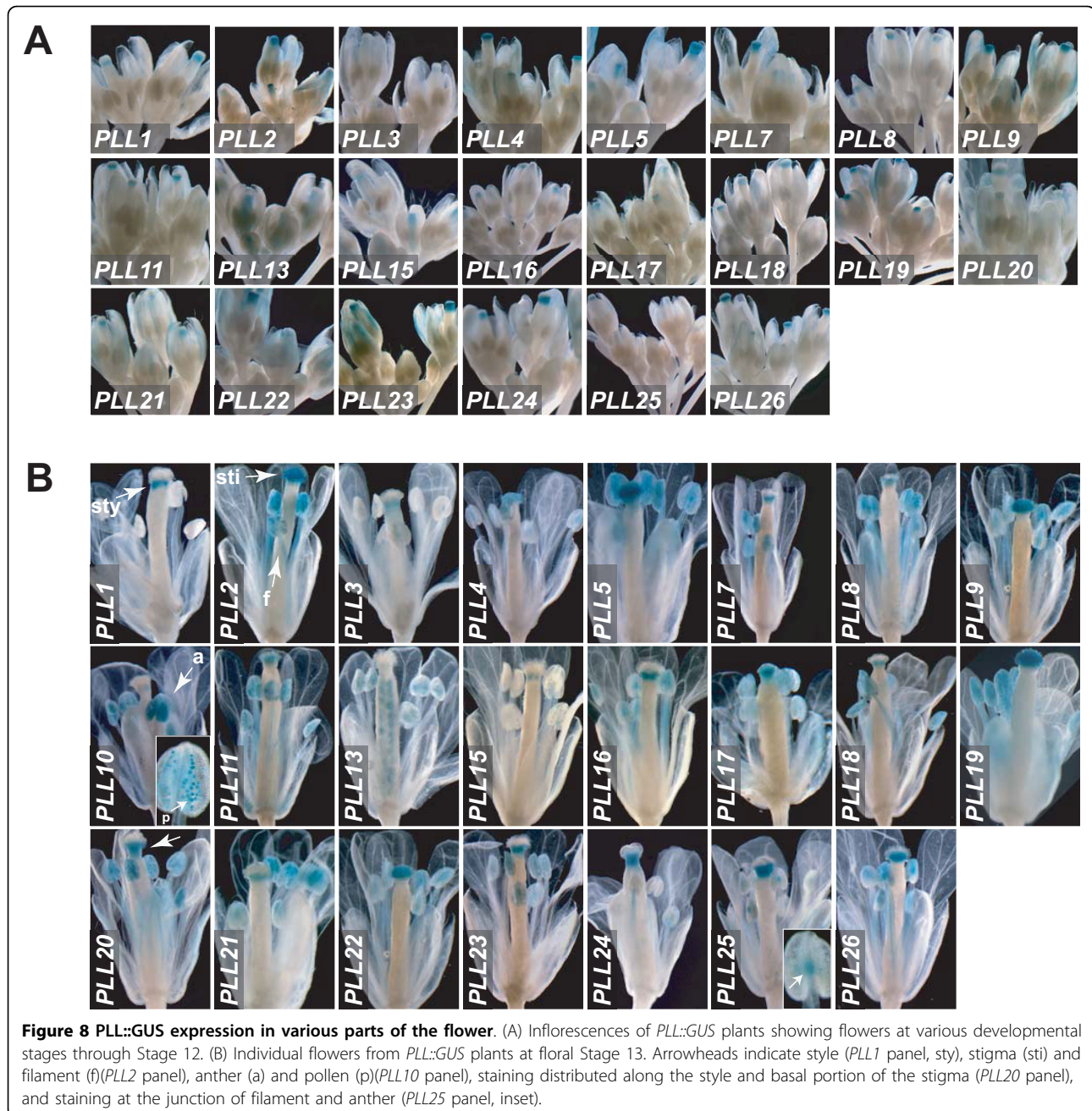
these *PLLs* may be to participate in a pathway of cell wall remodeling to accommodate such stress.

Other patterns of *PLL* activity were difficult to explain given a function limited to cell separation, and might instead be associated with general cell wall remodeling accompanying growth and differentiation. The extensive and various activities seen in the root is a good example of this. We can not provide a simple explanation for the strong and widespread promoter activity in stipules; however, these structures are absorbed into the leaf periphery during leaf expansion and this process could be accompanied by radical changes in cell wall morphology. Interestingly, both stipules and hydathodes were previously characterized as sites of high free auxin concentration during early leaf development [45]. A subset of *PLL* promoters in our study were found to be actively associated with lateral root emergence, or in the region of the root apical meristem, other sites associated with local auxin concentration maxima [30,50]. These observations suggest a close functional relationship between these *PLLs* and auxin.

We note that many promoter activities of the *PLL* genes are superficially similar to those exhibited collectively by several presumed polygalacturonase genes [51]. These include cases intuitively related to cell separation (perianth organ AZs, sites of lateral root emergence, fruit DZ, seed AZ, germinating seed) and cases in which activity may be limited to cell wall remodeling (developing pollen and seeds). This implies close functional association between *PLLs* and *PGs*, which is expected given that both should target pectins.

An obvious use of data from high-throughput analyses of promoter activity and/or transcriptional profiles is to guide reversed-genetics analyses, especially where phenotypes are difficult to anticipate and/or where potential redundancy of the targeted genes is a limitation. The original intent of this project was to identify individual or small subsets of *PLL* genes that might be targeted in this manner. However, the surprisingly high degree of overlap both in promoter activity and RNA accumulation observed among the *PLL* genes/promoters largely precluded this objective. We note that neither our promoter activity analysis nor previous microarray data would intuitively predict the documented non-redundant role for *PLL13* (*PMR6*) in facilitating infection by powdery mildew [28]. This role is accomplished where infection occurs in leaf cells, where neither transcriptional profiling nor our promoter analysis suggested strong or non-redundant expression.

Another obvious application of transcriptional data is the computational identification of DNA regulatory elements linked to regionally or temporally specific expression patterns. Here, it is noteworthy that the correlation



between the transcriptional profiles previously established for these genes [16,48] and their promoter activity (this study) was apparently very limited. This justifies the additional time and expense necessary to accomplish comprehensive promoter activity profiles, and/or cell-type specific transcriptional profiling to define transcriptional regulatory elements.

The *PLL* gene family is one of the largest and most complex in *Arabidopsis*, suggesting enormous potential for gene specialization through spatial or temporal regulation. However, we found a remarkably high degree of

overlap of activity among the *PLL* promoters. The activity of 19 of the 23 tested *PLL*s in the stipules is an excellent example of this, as is the activity of at least 18 in the apparent AZs of the sepals, petals and stamens at similar stages of development. This observation is surprising given that gene duplications should be evolutionarily stable only after having acquired a useful, non-redundant function. For *PLL*s, functional specificity may be bestowed by post-transcriptional and downstream mechanisms. Such specificity might include regulation of RNA stability. Specificity might also be directed at

the protein level, through potential differences in localization or mobility within the apoplast. Alternatively, given the expected dynamic pH environment of the apoplast, the proteins might function only within a narrow pH window. Finally, various PLL proteins might show preference for specific pectin substrates, an intriguing possibility given the structural complexity of pectins. Many of these possibilities could be addressed by the purification (or recombinant synthesis) and enzymatic characterization of individual PLLs.

Methods

Plant growth conditions

Wild type *Arabidopsis thaliana* (ecotype Col-0) was used for all experiments. Standard growth conditions were 16-h-cool white fluorescent light/8-h-dark photoperiods at 22°C. For growth in sterile culture, seeds were surface-sterilized and germinated on one-half-strength Murashige and Skoog (MS) medium supplemented with 10 mM MES [2-(N-morpholino) ethanesulfonic acid] (Sigma, St. Louis, MO), pH adjusted to 5.7, and solidified with 0.8% phytoagar (Invitrogen, Carlsbad, CA).

Phylogenetic analysis and protein domain identification

The peptide sequence of an enzymatically confirmed pectate lyase from banana, *Musa acuminata* *PL1* (GenBank accession AF206319) [22], was utilized as a query in BLASTP (Basic Local Alignment Search Tool Protein) analysis of *Arabidopsis* translated open reading frames [The *Arabidopsis* Information Resource (TAIR) Version 7] [52], and closely related proteins [Expect (E) value < 8E-36] were used for phylogenetic tree construction. Predicted amino acid sequences were aligned using ClustalW2 [53,54], and MEGA4 [55] was used to construct the phylogenetic tree. Annotated PLL protein domains were identified in the Pfam database [56,57] as follows: PF03211, Pectate_lyase; PF00544, Pec_lyase_C; PF09492, Pec_lyase; PF04431, Pec_lyase_N. Amino-terminal signal peptides were defined by SignalP 3.0 [58] using a maximum S-score larger than 0.95 and signal peptide probability greater than 0.80 [59]. The number of ESTs for each gene in the NCBI Unigene database was found at NCBI [60].

Transgenic plant construction

To engineer *GUS* as a reporter for *PLL* promoter activity, up to approximately 2 kb of genomic DNA, upstream from the start codon, was amplified by PCR using gene-specific primers (see **Additional file 4**) containing unique restriction sites. In general, where the upstream intergenic region was less than 2 kb, all of the region was included (see **Additional file 2**). All amplification products were sequenced to confirm lack

of PCR-induced mutation, and cloned into the pCAMBIA1305 vector (Cambia, Black Mountain, Australia) modified to contain an *NcoI* site at the start codon of the *GUS* gene and multiple cloning sites replacing the pCAMBIA Lac Z alpha and CaMV35S regions. This resulted in *PLL::GUS* fusions that preserve the authentic 5' UTR and start codon position. Plasmid DNAs were introduced into *Agrobacterium tumefaciens* strain GV3101 and transformed into wild-type plants using the floral dipping method [61]. Transgenic plants were subjected to selection with the herbicide glufosinate (Basta; Bayer CropScience, Barmen, Germany).

Histochemical GUS assays

Populations derived from at least four independently transformed transgenic lines were analyzed for each *PLL::GUS* construction. For histochemical GUS assays, plants or plant parts at various developmental stages were immersed in GUS staining solution [0.5 mM X-gluc, 0.5% (v/v) Triton X-100, 50 mM sodium phosphate buffer (pH adjusted to 7.2)] under vacuum infiltration for 5 min, and incubated at 37°C for various lengths of time. After staining, tissues were incubated in 70% ethanol for several hours to remove chlorophyll. Staining was visualized using a Nikon dissecting microscope equipped with a digital camera. Flower developmental stages annotated in this study were described by Smyth et al [40].

Analysis of AtGenExpress data

Microarray data were collected from the AtGenExpress Development data set [48]. Only results from wild-type plants were included in the analysis. The log₂-transformed absolute signal values were used in hierarchical clustering based on average linkage (Cluster version 3; [62]), and results were visualized in TreeView [62].

Additional material

Additional file 1: Hierarchical clustering of RNA accumulation for *Arabidopsis* *PLL* genes in various plant parts/developmental stages based on public microarray data. Plant parts and growth stages are labeled according to AtGenExpress [48]. Microarray signal value is indicated by color from yellow (lowest) to red (highest). *PLL5* was not represented on the microarray used in this analysis.

Additional file 2: Depiction of promoter region from individual *PLL* genes utilized to drive *GUS* expression. Genes are aligned with the predicted translational start (ATG) codon at right. The scale of upstream distance (kb) is given at top. The portion of the upstream region used is indicated with a heavy line. Upstream features (mapped 5' or 3' RNA ends and/or predicted translational start/stop codons) delineating the intergenic region are indicated with relative position given at left.

Additional file 3: Summary of *PLL::GUS* expression in various *Arabidopsis* parts.

Additional file 4: Primer sequences used for amplifying promoter region of *PLL* gene family members.

Acknowledgements

This work was funded by grants from the Michigan Agricultural Experiment Station (MAES) and Washington Tree Fruit Research Commission (WTRC) to SvN, and by funding from the USDA Multistate Research Project, NE1018, "Postharvest Biology of Fruit" to LS. We thank Dr. Art Cameron (Michigan State University) for assistance with microscopy and photography.

Authors' contributions

LS and SvN both participated in the planning and execution of the experiments, analysis of results, and writing of the manuscript.

Received: 2 March 2010 Accepted: 22 July 2010 Published: 22 July 2010

References

- Caffall KH, Mohnen D: **The structure, function, and biosynthesis of plant cell wall pectic polysaccharides.** *Carbohydr Res* 2009, **344**:1879-1900.
- Ridley BL, O'Neill MA, Mohnen D: **Pectins: structure, biosynthesis, and oligogalacturonide-related signaling.** *Phytochemistry* 2001, **57**:929-967.
- Willats WG, McCarney L, Mackie W, Knox PJ: **Pectin: cell biology and prospects for functional analysis.** *Plant Mol Biol* 2001, **47**:9-27.
- Iwai H, Masaoka N, Ishii T, Satoh S: **A pectin glucuronyltransferase gene is essential for intercellular attachment in the plant meristem.** *Proc Natl Acad Sci USA* 2002, **99**:16319-16324.
- Jarvis M, Briggs S, Knox J: **Intercellular adhesion and cell separation in plants.** *Plant Cell Environ* 2003, **26**:977-989.
- Knox JP: **Cell and developmental biology of pectins.** *Pectins and Their Manipulation* CRC press LLC Seymour GB, Knox JP 2002, 131-146.
- Barras F, van Gijsegem F, Chatterjee AK: **Extracellular enzymes and pathogenesis of soft-rot *Erwinia*.** *Annu Rev Phyto* 1994, **32**:201-234.
- Collmer A, Keen N: **The role of pectic enzymes in plant pathogenesis.** *Annu Rev Phytopathol* 1986, **24**:383-409.
- Kotoujansky A: **Molecular genetics of pathogenesis by soft rot *Erwinias*.** *Annu Rev Phytopathol* 1987, **25**:405-430.
- De Lorenzo G, Cervone F, Hahn MG, Darvill A, Albersheim P: **Bacterial endopectate lyase: evidence that plant cell wall pH prevents tissue maceration and increases the half-life of elicitor-active oligogalacturonides.** *Physiol Mol Plant Pathol* 1991, **39**:335-344.
- Fagard M, Dellagi A, Roux C, Périno C, Rigault M, Boucher V, Shevchik V, Expert D: ***Arabidopsis thaliana* expresses multiple lines of defense to counterattack *Erwinia chrysanthemi*.** *Mol Plant Microbe Interact* 2007, **20**:794-805.
- Norman-Setterblad C, Vidal S, Palva ET: **Interacting signal pathways control defense gene expression in *A. thaliana* in response to cell wall-degrading enzymes from *Erwinia carotovora*.** *Mol Plant Microbe Interact* 2000, **13**:430-438.
- Wing R, Yamaguchi J, Larabell S, Ursin V, McCormic S: **Molecular and genetic characterization of two pollen-expressed genes that have sequence similarity to pectate lyases of the plant pathogen *Erwinia*.** *Plant Mol Biol* 1990, **14**:17-28.
- Marín-Rodríguez MC, Orchard J, Seymour GB: **Pectate lyases, cell wall degradation and fruit softening.** *J Exp Bot* 2002, **53**:2115-2119.
- Henrissat B, Coutinho PM, Davies GJ: **A census of carbohydrate-active enzymes in the genome of *Arabidopsis thaliana*.** *Plant Mol Biol* 2001, **47**:55-72.
- Palusa SG, Golovkin M, Shin SB, Richardson DN, Reddy AS: **Organ-specific, developmental, hormonal and stress regulation of expression of putative pectate lyase genes in *Arabidopsis*.** *New Phytol* 2007, **174**:537-550.
- Zhang Z: **Evolution by gene duplication: an update.** *Trends Ecol Evol* 2003, **18**:292-298.
- Lynch M, Force A: **The probability of duplicate gene preservation by subfunctionalization.** *Genetics* 2000, **154**:459-473.
- Benítez-Burraco A, Blanco-Portales R, Redondo-Nevaldo J, Bellido ML, Moyano E, Caballero JL, Muñoz-Blanco J: **Cloning and characterization of two ripening-related strawberry (*Fragaria × ananassa* cv. Chandler) pectate lyase genes.** *J Exp Bot* 2003, **54**:633-645.
- Chourasia A, Sane VA, Nath P: **Differential expression of pectate lyase during ethylene-induced postharvest softening of mango (*Mangifera indica* var. Dashehari).** *Physiol Plant* 2006, **128**:546-555.
- Futamura N, Kouchi H, Shinohara K: **A gene for pectate lyase expressed in elongating and differentiating tissues of a Japanese willow (*Salix gilgiana*).** *J Plant Physiol* 2002, **159**:1123-1130.
- Marín-Rodríguez M, Smith D, Manning K, Orchard J, Seymour GB: **Pectate lyase gene expression and enzyme activity in ripening banana fruit.** *Plant Mol Biol* 2003, **51**:851-857.
- Mita S, Nagai Y, Asai T: **Isolation of cDNA clones corresponding to genes differentially expressed in pericarp of mume (*Prunus mume*) in response to ripening, ethylene and wounding signals.** *Physiol Plant* 2006, **128**:531-545.
- Pua E, Ong C, Liu P, Liu J: **Isolation and expression of two pectate lyase genes during fruit ripening of banana (*Musa acuminata*).** *Physiol Plant* 2001, **113**:92-99.
- Domingo C, Roberts K, Stacey NJ, Connerton I, Ruiz-Teran F, McCann M: **A pectate lyase from *Zinnia elegans* is auxin inducible.** *Plant J* 1998, **13**:17-28.
- Laskowski M, Biller S, Stanley K, Kajstura T, Prusty R: **Expression profiling of auxin-treated *Arabidopsis* roots: toward a molecular analysis of lateral root emergence.** *Plant Cell Physiol* 2006, **47**:788-792.
- Milioni D, Sado PE, Stacey NJ, Domingo C, Roberts K, McCann MC: **Differential expression of cell-wall-related genes during the formation of tracheary elements in the *Zinnia* mesophyll cell system.** *Plant Mol Biol* 2001, **47**:221-238.
- Vogel JP, Raab T, Somerville C: ***PMR6*, a pectate lyase-like gene required for powdery mildew susceptibility in *Arabidopsis*.** *Plant Cell* 2002, **14**:2095-2106.
- Cai S, Lashbrook CC: **Stamen abscission zone transcriptome profiling reveals new candidates for abscission control: enhanced retention of floral organs in transgenic plants overexpressing *Arabidopsis* ZINC FINGER PROTEIN2.** *Plant Physiol* 2008, **146**:1305-1321.
- Swarup K, Benková E, Swarup R, Casimiro I, Péret B, Yang Y, Parry G, Nielsen E, Smet ID, Vanneste S, Levesque MP, Carrier D, James N, Calvo V, Ljung K, Kramer E, Roberts R, Graham N, Marilionnet S, Patel K, Jones JDG, Taylor CG, Schachtman DP, Sandberg G, Benfey P, Friml J, Kerr I, Beeckman T, Laplace L, Bennett MJ: **The auxin influx carrier *LAX3* promotes lateral root emergence.** *Nat Cell Biol* 2008, **10**:946-954.
- Jiménez-Bermúdez S, Redondo-Nevaldo J, Muñoz-Blanco J, Caballero JL, López-Aranda JM, Valpuesta V, Pliego-Alfaro F, Quesada MA, Mercado JA: **Manipulation of strawberry fruit softening by antisense expression of a pectate lyase gene.** *Plant Physiol* 2002, **128**:751-759.
- Leslie ME, Lewis MW, Liljegren SJ: **Organ abscission.** *Annual Plant Reviews* Oxford: Blackwell Publishing Ltd Roberts JA, Gonzalez-Carranza Z 2007, **25**:91-106.
- Ostergaard L, Borkhardt B, Ulvskov P: **Dehiscence.** *Annual Plant Reviews* Oxford: Blackwell Publishing Ltd Roberts JA, Gonzalez-Carranza Z 2007, **25**:137-159.
- Jarvis M: **Intercellular separation forces generated by intracellular pressure.** *Plant Cell Environ* 1998, **21**:1307-1310.
- Roy S, Jauneau A, Vian B: **Analytical detection of calcium-ions and immunocytochemical visualization of homogalacturonic sequences in the ripe cherry tomato.** *Plant Physiol Biochem* 1994, **32**:633-640.
- Hallett IC, MacRae EA, Wegrzyn TF: **Changes in kiwifruit cell-wall ultrastructure and cell packing during postharvest ripening.** *Int J Plant Sci* 1992, **153**:49-60.
- Wen F, Laskowski M, Hawes M: **Cell separation in roots.** *Annual Plant Reviews* Oxford: Blackwell Publishing Ltd Roberts JA, Gonzalez-Carranza Z 2007, **25**:91-106.
- Mollet JC, Faugeron C, Morvan H: **Cell adhesion, separation and guidance in compatible plant reproduction.** *Annual Plant Reviews* Oxford: Blackwell Publishing Ltd Roberts JA, Gonzalez-Carranza Z 2007, **25**:69-90.
- Patterson SE: **Cutting loose. Abscission and dehiscence in *Arabidopsis*.** *Plant Physiol* 2001, **126**:494-500.
- Smyth DR, Bowman JL, Meyerowitz EM: **Early flower development in *Arabidopsis*.** *Plant Cell* 1990, **2**:755-767.
- Ferrández C: **Regulation of fruit dehiscence in *Arabidopsis*.** *J Exp Bot* 2002, **53**:2031-2038.
- Liu PP, Koizuka N, Homrichhausen TM, Hewitt JR, Martin RC, Nonogaki H: **Large-scale screening of *Arabidopsis* enhancer-trap lines for seed germination-associated genes.** *Plant J* 2005, **41**:936-944.

43. Malamy JE, Benfey PN: **Organization and cell differentiation in lateral roots of *Arabidopsis thaliana*.** *Development* 1997, **124**:33-44.
44. Esau K: **Hydathodes.** *Anatomy of Seed Plants* New York: John Wiley & Sons 1977, 205-206.
45. Aloni R, Schwalm K, Langhans M, Ullrich CI: **Gradual shifts in sites of free-auxin production during leaf-primordium development and their role in vascular differentiation and leaf morphogenesis in *Arabidopsis*.** *Planta* 2003, **216**:841-853.
46. Cheng Y, Dai X, Zhao Y: **Auxin synthesized by the YUCCA flavin monooxygenases is essential for embryogenesis and leaf formation in *Arabidopsis*.** *Plant Cell* 2007, **19**:2430-2439.
47. Berger F: **Endosperm: the crossroad of seed development.** *Curr Opin Plant Biol* 2003, **6**:42-50.
48. Schmid M, Davison TS, Henz SR, Pape UJ, Demar M, Vingron M, Schölkopf B, Weigel D, Lohmann JU: **A gene expression map of *Arabidopsis thaliana* development.** *Nat Genet* 2005, **37**:501-506.
49. Stenvik GE, Butenko MA, Urbanowicz BR, Rose JK, Aalen RB: **Overexpression of INFLORESCENCE DEFICIENT IN ABSCISSION activates cell separation in vestigial abscission zones in *Arabidopsis*.** *Plant Cell* 2006, **18**:1467-1476.
50. Sabatini S, Beis D, Wokenfelt H, Murfett J, Guilfoyle T, Malamy H, Benfey P, Leysner O, Bechtold N, Weisbeek P, Scheres B: **An auxin-dependent distal organizer of pattern and polarity in the *Arabidopsis* root.** *Cell* 1999, **99**:463-472.
51. González-Carranza ZH, Elliott KA, Roberts JA: **Expression of polygalacturonases and evidence to support their role during cell separation processes in *Arabidopsis thaliana*.** *J Exp Bot* 2007, **58**:3719-3730.
52. **The Arabidopsis Information Resource.** [<http://www.arabidopsis.org>].
53. **Clustal: Multiple Sequence Alignment.** [<http://www.clustal.org>].
54. Larkin MA, Blackshields G, Brown NP, Chenna R, McGettigan PA, McWilliam H, Valentin F, Wallace IM, Wilm A, Lopez R, Thompson JD, Gibson TJ, Higgins DG: **Clustal W and Clustal x version 2.0.** *Bioinformatics* 2007, **23**:2947-2948.
55. Tamura K, Dudley J, Nei M, Kumar S: **MEGA4: Molecular Evolutionary Genetics Analysis (MEGA) software version 4.0.** *Mol Biol Evol* 2007, **24**:1596-1599.
56. **Pfam Database.** [<http://pfam.sanger.ac.uk/>].
57. Finn RD, Mistry J, Tate J, Coggill P, Heger A, Pollington JE, Gavin OL, Gunasekaran P, Ceric G, Forslund K, Holm L, Sonnhammer EL, Eddy SR, Bateman A: **The Pfam protein families database.** *Nucleic Acids Res* 2010, **38**:D211-22.
58. **SignalP 3.0 Server.** [<http://www.cbs.dtu.dk/services/SignalP/>].
59. Emanuelsson O, Brunak S, von Heijne G, Nielsen H: **Locating proteins in the cell using TargetP, SignalP and related tools.** *Nat Protoc* 2007, **2**:953-971.
60. **National Center for Biotechnology Information.** [<http://www.ncbi.nlm.nih.gov/>].
61. Clough SJ, Bent AF: **Floral dip: a simplified method for *Agrobacterium*-mediated transformation of *Arabidopsis thaliana*.** *Plant J* 1998, **16**:735-743.
62. Eisen MB, Spellman PT, Brown PO, Botstein D: **Cluster analysis and display of genome-wide expression patterns.** *Proc Natl Acad Sci USA* 1998, **95**:14863-14868.

doi:10.1186/1471-2229-10-152

Cite this article as: Sun and van Nocker: Analysis of promoter activity of members of the *PECTATE LYASE-LIKE (PLL)* gene family in cell separation in *Arabidopsis*. *BMC Plant Biology* 2010 **10**:152.

**Submit your next manuscript to BioMed Central
and take full advantage of:**

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at
www.biomedcentral.com/submit

