

# Arabidopsis Rho-Related GTPases: Differential Gene Expression in Pollen and Polar Localization in Fission Yeast<sup>1</sup>

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The Rho small GTP-binding proteins are versatile, conserved molecular switches in eukaryotic signal transduction. Plants contain a unique subfamily of Rho-GTPases called Rop (Rho-related GTPases from plants). Our previous studies involving injection of antibodies indicated that the pea Rop GTPase Rop1Ps is critical for pollen tube growth. In this study we show that overexpression of an apparent Arabidopsis ortholog of Rop1Ps, *Rop1At*, induces isotropic cell growth in fission yeast (*Schizosaccharomyces pombe*) and that green fluorescence protein-tagged Rop1At displays polar localization to the site of growth in yeast. We found that Rop1At and two other Arabidopsis Rops, Rop3At and Rop5At, are all expressed in mature pollen. All three pollen Rops fall into the same subgroup as Rop1Ps and diverge from those Rops that are not expressed in mature pollen, suggesting a coupling of the structural conservation of Rop GTPases to their gene expression in pollen. However, pollen-specific transcript accumulation for *Rop1At* is much higher than that for *Rop3At* and *Rop5At*. Furthermore, *Rop1At* is specifically expressed in anthers, whereas *Rop3At* and *Rop5At* are also expressed in vegetative tissues. In transgenic plants containing the *Rop1At* promoter:GUS fusion gene, GUS is specifically expressed in mature pollen and pollen tubes. We propose that Rop1At may play a predominant role in the regulation of polarized cell growth in pollen, whereas its close relatives Rop3At and Rop5At may be functionally redundant to Rop1At in pollen.

In angiosperms male gametophyte development can be divided into two major phases: microsporophyte development and pollen development. Microsporophyte development is the division of a diploid sporophytic cell, giving rise to the tapetal initial cell and the microspore mother cell. This diploid microspore mother cell undergoes meiosis to produce haploid microspores. Microspores then enter the phase of pollen development, which begins with an asymmetric mitotic division, resulting in the formation of a pollen grain containing a large, vegetative cell and a small, generative cell enclosed within it. In some species, such as Arabidopsis, the generative cell undergoes a second mitotic division in developing pollen before anthesis to produce a tricellular mature pollen grain. In other species, mature pollen grains are released as bicellular cells, and the second

mitotic division occurs during pollen tube growth within the style (Mascarenhas, 1993; McCormick, 1993).

Pollen development involves complex developmental control of gene expression by the haploid genome. It has been estimated that 10% of the 20,000 different genes expressed in pollen grains at anthesis are pollen specific (for review, see Mascarenhas, 1993; McCormick, 1993; Taylor and Helper, 1997). Pollen-specific genes can be divided into two groups: Genes expressed before the first pollen mitosis are referred to as “early” pollen genes and are believed to be involved in pollen development; genes activated after this mitosis are called “late” pollen genes and are presumably involved in pollen maturation and germination (Mascarenhas, 1993). At least 23 late pollen genes have been identified from different plant species (for review, see McCormick, 1993; Twell, 1994; Taylor and Helper, 1997). Several of these late pollen genes encode signaling proteins such as a Ca<sup>2+</sup>-dependent protein kinase involved in self-incompatibility in *Nicotiana glauca* (Kunz et al., 1996), a Ca<sup>2+</sup>-dependent calmodulin-independent protein kinase involved in pollen germination in maize (Estruch et al., 1994), a receptor-like kinase, PRK1, essential for normal pollen development in petunia (Lee et al., 1996), and a mitogen-activated protein kinase activated upon pollen hydration in *Nicotiana tabacum* (Wilson et al., 1997).

We previously reported a small GTP-binding protein, Rop1Ps, that preferentially accumulated in mature pollen of the garden pea (Lin et al., 1996). Rop1Ps belongs to the Rho family of small GTPases, which has become an important group of conserved signaling proteins in eukaryotes. Rho-dependent signaling controls a large variety of key cellular processes in animals and fungi, e.g. actin cytoskeletal reorganization, the establishment of cell polarity, polarized cell growth, membrane trafficking and organization (e.g. exocytosis and endocytosis), focal adhesion, and cell movement (Hall, 1994; Chant and Stowers, 1995; Lamaze et al., 1996; Laroche et al., 1996; Murphy et al., 1996; Nagata and Hall, 1996; Ridley, 1996).

Plants possess a family of genes encoding proteins closely related to Rop1Ps, including 10 reported genes from Arabidopsis (Yang and Watson, 1993; Delmer et al., 1995; Lin et al., 1996; Winge et al., 1997). Indirect immunofluorescence studies in pea suggest that Rop1Ps is localized to the tip of pollen tubes (Lin et al., 1996). We showed that

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Abbreviations: DAPI, 4',6-diamidino-phenylindole; GFP, green fluorescent protein; RT, reverse transcriptase.

injected anti-Rop1Ps antibodies inhibited pollen tube elongation in pea, and that this inhibition was independent of cytoplasmic streaming and potentiated by low extracellular  $\text{Ca}^{2+}$  and caffeine treatment (Lin and Yang, 1997). These results suggest that Rop1Ps plays a pivotal role in the control of pollen tube growth, probably by interacting with  $\text{Ca}^{2+}$  signaling (Lin and Yang, 1997). However, the precise function of these GTPases in pollen needs to be determined using a reverse-genetics approach. Such an approach is most feasible in Arabidopsis due to the recent development of homology-based gene replacement (Kempin et al., 1997) and PCR-mediated identification of T-DNA insertion into genes of known sequences (McKinney et al., 1995; Krysan et al., 1996).

In this paper we report the identification of a novel member of the Arabidopsis Rop gene family, *Rop1At*, the only Rop gene known to be specifically expressed in the anther. *Rop1At* appears to have a conserved function in regulating polarized cell growth in fission yeast (*Schizosaccharomyces pombe*). Analyses of promoter:GUS reporter fusion gene expression show that *Rop1At* is a late pollen gene. *Rop3At* and *Rop5At*, which are most closely related to *Rop1At*, are also expressed in mature pollen, although at a lower level, whereas other Rop genes divergent from *Rop1At* are not expressed in mature pollen. These results imply a functional constraint on the structural conservation of the Rop subfamily of GTPases, with the three most closely related members having a potential redundant function in the control of polarized cell growth in pollen.

## MATERIALS AND METHODS

### Plant Material

Arabidopsis ecotype Columbia plants were grown in growth chambers at 22°C under constant light. Rosette leaves from 4-week-old plants were harvested for genomic DNA isolation. For RNA extractions roots, stems, rosette leaves, open and closed flowers, siliques, and pollen grains were harvested from 4- to 6-week-old plants.

### cDNA and Genomic DNA Cloning and Sequencing

The Arabidopsis Columbia cDNA library,  $\lambda$ PRL-2 (Tom Newman, Michigan State University, obtained through the Arabidopsis Biological Resource Center, Ohio State University, Columbus), was screened with a  $^{32}\text{P}$ -labeled,

167-bp fragment of *Rop1Ps* cDNA, which corresponds to the most conserved region within the Rho gene family (Yang and Watson, 1993). Plasmids containing positive clones were excised in vivo from lambda phage, and inserts were subcloned into pBluescript II SK (Stratagene) and sequenced using the dideoxynucleotide chain-termination method (Sanger et al., 1977) and Sequenase version 2.0 (United States Biochemical). The cDNA library screen identified two distinct genes, *Rop1At* and *Rop2At*. To isolate additional *Rop1Ps*-related genes, an Arabidopsis genomic library (Voytas et al., 1990) obtained from the Arabidopsis Biological Resource Center was screened with a 369-bp *SacII* fragment of *Rop2At* cDNA under moderate hybridization stringency. Positive clones were subcloned and sequenced as described above.

### Computer Analysis of the Rop Subfamily

Predicted amino acid sequences for Arabidopsis *Rop1Ps*-related genes were compared with known members of the Rho family of GTP-binding proteins available from the GenBank database using computer software from DNASTAR, Inc. (Madison, WI). Alignments of these sequences were carried out using the MegAlign program (DNASTAR, Inc., Madison, WI). Phylogenetic analyses of the aligned sequences were conducted using PAUP (Phylogenetic Analysis Using Parsimony) software (version 3.1.1, D.L. Swofford [1993], Smithsonian Institution, Washington, DC).

### Reverse Transcription and PCRs

Total RNA was isolated from different Arabidopsis tissues as described previously (Logemann et al., 1987). The first-strand cDNAs were synthesized using murine leukemia virus RT (GIBCO-BRL) in a 50- $\mu\text{L}$  reaction containing 2.5  $\mu\text{M}$  oligo-dT primers (GIBCO-BRL), 5  $\mu\text{g}$  of total RNA, 10 mM DTT, 1 unit/ $\mu\text{L}$  RNase inhibitor (GIBCO-BRL), 0.20 mM dNTP mix, and 10 units/ $\mu\text{L}$  RT. Reverse-transcription reactions were carried out at 42°C for 60 min and were terminated by heating to 99°C for 5 min and chilling to 4°C for 5 min. Five microliters of the reaction mixture was used as a template for each of the PCRs described below. PCR reactions were carried out in 25  $\mu\text{L}$  of a mixture containing 2 mM  $\text{MgCl}_2$ , 0.25 unit of Taq polymerase (GIBCO-BRL), and 0.5  $\mu\text{M}$  gene-specific primers (see Table I). For *Rop1At*

**Table I.** *Rop* isogene-specific primers for RT-PCR

Genes	Sense Primers	Antisense Primers	Expected cDNA Length bp
Rop1At	5'-GAAATTAATAAACTTTGAGGGG-3' (-24 to -3)	5'-AGAGATTTCCAATCATCATAG-3' (+615 to +595)	639
Rop2At	5'-GCCGCAGAGATGGCGTCAAGG-3' (-9 to +11)	5'-CTTATCACAAGAACGCGCAACG-3' (+592 to 571)	601
Rop3At	5'-TACGTAGCTCCATTCTGGTGGAG-3' (-41 to -18)	5'-CCACAATCCAAGATTGACAGT-3' (+177 to 157)	218
Rop4At	5'-CATTATTATCTCTCATCGATTGG-3' (-184 to -161)	Same as above	361 <sup>a</sup>
Rop5At	5'-GTGACATATTTGGCTCGTCG-3' (-38 to -18)	Same as above	215 <sup>a</sup>
Rop6At	5'-CGTCCGTGAGGATGAGTAGT-3' (-172 to -153)	Same as above	349

<sup>a</sup> Expected genomic DNA lengths for Rop4At and Rop5At are approximately 800 and 335 bp, respectively.

and *Rop2At*, 25 cycles of PCR amplification were carried out at 94°C for 30 s (denaturation), at 60°C for 30 s (annealing), and at 72°C for 30 s (synthesis). Five microliters of each PCR product was loaded on a 1.5% agarose gel to visualize the amplified cDNAs.

The same procedures were used for *Rop3At* and *Rop4At*, except that the annealing temperature was 55°C. The same PCR reaction conditions were used for *Rop5At* and *Rop6At*, but the number of PCR cycles was increased to 45, and 25  $\mu$ L of the PCR reaction was loaded on a 2% low-melting agarose gel. As PCR amplification and loading controls, the same template cDNA was amplified using primers for the constitutive *Act2* gene (An et al., 1996b). *Act2* PCR amplification was conducted at 94°C for 30 s, at 55°C for 30 s, and at 72°C for 1 min for 25 cycles, and 5  $\mu$ L of the PCR reaction was loaded onto a 1% agarose gel. RT-PCR reactions were repeated twice using total RNAs extracted with a different isolation procedure (Thompson et al., 1983).

To ensure gene-specific PCR amplifications, at least one of the two PCR primers was designed according to sequences of divergent 5' untranslated regions (Table I). To confirm the specificity of each pair of primers, two sets of PCRs were performed separately, one containing 1 ng of a specific cDNA or a genomic DNA clone corresponding to the primers (positive control), and the other containing a mixture of an equal amount (1 ng of each) of cDNA or genomic DNAs for each of the other five Rop genes. PCR conditions were identical to those used for RT-PCR described above.

### Construction of *Rop1At* Promoter:GUS Fusion Gene and Plant Transformation

To direct the expression of *Rop1At* promoter:GUS fusion gene in Arabidopsis plants, a binary vector containing the fusion gene was constructed as follows. A 1.5-kb *XbaI/PstI* genomic fragment flanking the 5' end of the *Rop1At* coding sequence was subcloned into pUC19. This fragment was sequenced by the dideoxynucleotide chain-termination method (Sanger et al., 1977) to confirm that it contained 1.3 kb upstream of the *Rop1At* translation-initiation codon (see Fig. 4). The 1.5-kb genomic DNA fragment was then subcloned into the *EcoRI/PstI* sites of pBluescript II SK to allow the use of a *HindIII* site at the 5' end of the genomic sequence for further subcloning into a binary vector. To introduce a *SalI* site 20 bp downstream of the *Rop1At* ATG codon, the sense T7 sequencing primer and the antisense primer containing a *SalI* site were used for PCR amplification of the putative *Rop1At* promoter. The amplified fragment was digested with *HindIII* and *SalI*, and then translationally fused with the GUS gene in pBI101.2 vector (Clontech, Palo Alto, CA). This plasmid was designated pBR1P:GUS (Fig. 4).

pBR1P:GUS was mobilized into *Agrobacterium tumefaciens* strain LB4404 by the freeze-thaw method (An et al., 1988) and introduced into Arabidopsis (ecotype RLD) by the root-transformation method (Valvekens et al., 1988). Primary transgenic plants were selected on Murashige and Skoog medium (Sigma) containing 50  $\mu$ g/mL kanamycin.

Primary transgenic plants and their progenies were analyzed by histochemical GUS staining.

### Histochemical GUS Staining and DNA Staining

Histochemical assays for GUS activity in transgenic Arabidopsis plants were performed as described previously (Jefferson et al., 1987). To examine cell-specific GUS activity, tissues were photographed under a microscope (Zeiss) equipped with Nomarski optics. Nuclei of transgenic pollen grains were briefly costained with DAPI and visualized under an epifluorescence microscope (Coleman and Goff, 1985).

### Overexpression of the *Rop1At* Gene in Fission Yeast (*Schizosaccharomyces pombe*)

The *Rop1At* coding region was amplified by PCR using primers covering the translation start and stop codons, respectively. The PCR fragment was first cloned into the *EcoRV* site of pBluescript II SK and then subcloned into *SalI* and *SmaI* sites downstream of the *mtl1* promoter in the thiamine-repressible fission yeast expression vector pREP3X (Forsburg, 1993). The fission yeast strain FY254 (*h-*, *can1-1*, *leu1-32*, *ade6-M210*, and *ura4-D18*) was transformed by the electroporation method (Prentice, 1992), and transformants were plated on Edinburgh minimal medium/uracil plates containing 5 mM of thiamine, and were then incubated at 30°C. Overexpression of the *Rop1At* gene was induced by growing yeast cells in liquid medium lacking thiamine for 24 h. As a control, yeast cells containing the vector alone were treated in the same manner. Yeast morphology was examined under a microscope (Oxioskop, Zeiss) and recorded using a 35-mm camera.

### Expression of the Gene Encoding the Jellyfish GFP:*Rop1At* Fusion Protein in Fission Yeast

The mGFP4 coding region was amplified from pBIN-mGFP4 (Haseloff et al., 1997) by PCR using a sense primer containing an *XbaI* site upstream of the ATG codon and an antisense primer containing a *BglII* site in place of the GFP translation stop codon. The GFP fragment was cloned into *XbaI* and *SmaI* sites of pUC19. The *Rop1At* coding region was amplified by PCR using two primers containing a *BglII* site upstream of the ATG codon and a *SstI* site immediately following the UAG codon. This *Rop1At* fragment was then translationally fused with the mGFP4 gene at *BglII/SstI* sites in pUC19. The fusion gene was then cloned into pREP3X and introduced into the fission yeast strain FY254 as described above. To observe proper subcellular localization of the fusion protein, yeast cells containing the GFP:*Rop1At* fusion gene were grown in nonrepressive Edinburgh minimal medium/uracil for 24 h before transfer to a partially repressive medium containing 2 mM thiamine for 5 h. Green fluorescence was observed using an epifluorescent microscope (Oxioskop, Zeiss) and recorded using a 35-mm camera.

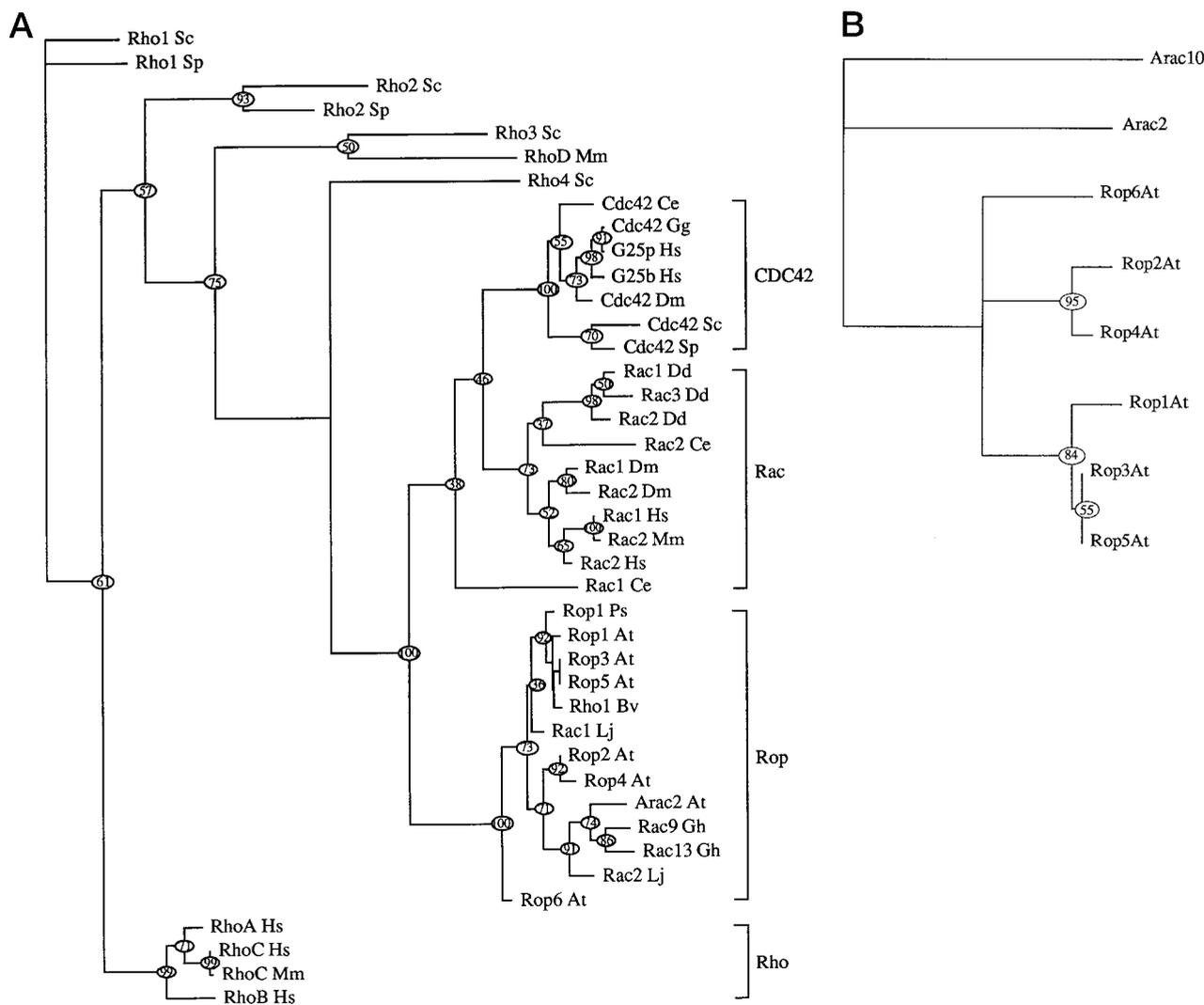
## RESULTS

## Identification of Rop1Ps Homologs in Arabidopsis

To identify *Rop1Ps* homologs, we screened an Arabidopsis cDNA library using a *Rop1Ps* probe (Yang and Watson, 1993). Two distinct *Rop1Ps*-related genes, *Rop1At* and *Rop2At*, were identified from this screen. Two additional genes, *Rop4At* and *Rop5At*, were isolated from an Arabidopsis genomic DNA library. Another *Rop1Ps*-related gene, *Rop6At*, was identified from an Arabidopsis expressed sequence tag database. A cDNA clone for *Rop3At* was obtained from Dr. Dring N. Crowell (Indiana University-Purdue University at Indianapolis). After this

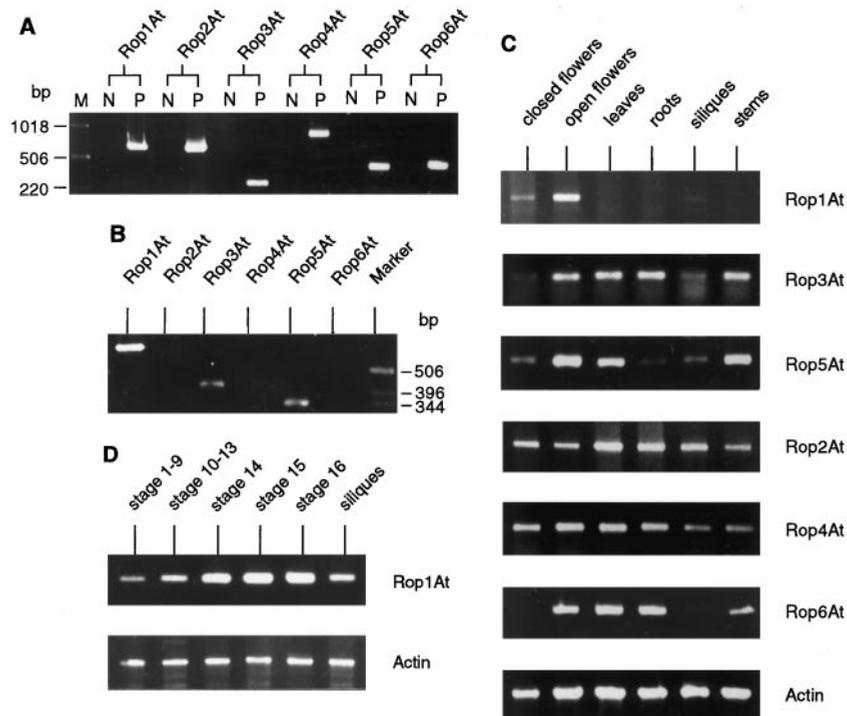
work was completed, a family of Arabidopsis genes related to *Rop1Ps*, designated *Arac*, was reported (Winge et al., 1997). Among the genes whose coding regions have been completely sequenced, four are identical to *Arac* genes: *Rop2At* (*Arac4*), *Rop3At* (*Arac1*), *Rop4At* (*Arac5*), and *Rop6At* (*Arac3*). However, *Rop1At*, which exhibits the highest homology to *Rop1Ps*, is a novel Rop member.

Phylogenetic analysis indicates that the predicted polypeptides encoded by these *Rop1Ps*-related genes belong to a subfamily of Rho-GTPases designated Rop, which is distinct from all major subfamilies of Rho-GTPases from animals and fungi (Fig. 1A). The Rop subfamily is more closely related to Rac (about 65% identity at the amino acid



**Figure 1.** Phylogenetic relationship between different Rho-GTPases. Unrooted trees were constructed using PAUP (Swoford, 1993). Amino acid sequences for various plant, animal, and yeast Rho-GTPases were obtained from GenBank using the Blast program. At, *Arabidopsis thaliana*; Bv, *Beta vulgaris*; Ce, *Caenorhabditis elegans*; Dd, *Dictyostelium discoideum*; Dm, *Drosophila melanogaster*; Gg, *Gallus gallus*; Gh, *Goosypium hirsutum*; Hs, *Homo sapiens*; Lj, *Lotus japonicus*; Mm, *Mus musculus*; Ps, *Pisum sativum*; Sc, *Saccharomyces cerevisiae*; Sp, *S. pombe*. A, Rho family tree showing phylogenetic relationships among major subfamilies from different eukaryotic kingdoms. This tree does not include all known members of Rho-GTPases, since several novel Rho-GTPases that do not fall within any of the major subfamilies are not included. B, Arabidopsis Rop family tree. Members shown include those described in this paper and those whose complete coding sequences are available in the Arabidopsis database.





**Figure 3.** RT-PCR analyses of *Rop* gene expression in various *Arabidopsis* tissues. A, Demonstration of *Rop* isogene-specific PCR amplification. Two sets of PCR reactions for each *Rop* isogene were performed. Lane M, DNA marker; lanes N, negative control, which involves the same primers and template DNAs containing a mixture of equal amounts of cDNA or genomic DNA for each of the other five *Rop* genes; lanes P, positive control, which involves a specific cDNA (for *Rop1At*, *Rop2At*, *Rop3At*, and *Rop6At*) or genomic DNA (for *Rop4At* and *Rop5At*) as templates and corresponding gene-specific primers. Expected cDNA lengths of amplified fragments are shown in Table I. The PCR reaction conditions are described in text. B, Accumulation of various *Arabidopsis Rop* transcripts in mature pollen. Total pollen RNA was isolated and the cDNA derived was amplified for 40 cycles using *Rop* gene-specific primers as described in the text. C, Organ distribution of various *Arabidopsis Rop* transcripts. RT-PCR was performed using *Rop* gene-specific primers described in A and total RNAs from different tissues as indicated. *Act2* RT-PCR was included as a constitutive control. The number of PCR cycles was: 25 for *Rop1At*, *Rop2At*, *Rop3At*, *Rop4At*, and *act2*, and 45 for *Rop5At* and *Rop6At*. D, Analyses of *Rop1At* mRNA accumulation during floral development. Total RNAs isolated from *Arabidopsis* floral buds and flowers at different stages were used for RT-PCR using the reaction conditions described in B. Flower stages were estimated as described previously (Smyth et al., 1990). Stages 1 to 9, Initiation and formation of floral primordia and organ differentiation; stages 10 to 13, organs fully developed, anthesis; stage 14, anthers extended above stigma, pollination; stage 15, stigma extends above anthers; stage 16, petals and sepals wither; siliques, developing siliques before seed maturation.

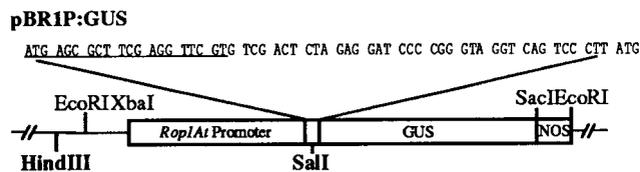
lationally fused with the GUS gene in pBI101.2 (Fig. 4) and was introduced into *Arabidopsis* plants.

Twelve primary transgenic plants were randomly chosen for histochemical staining of GUS activity (Jefferson et al., 1987) and all showed identical staining patterns, which were further confirmed in the next generation. As shown in Figure 5, GUS activity was specifically detected in anthers and in other parts of the plant. Very low levels of GUS activity were first detected in anthers undergoing microspore development. The GUS activity increases dramatically when all floral organs are differentiated and reaches a maximum at anthesis. Further analysis revealed that the strong GUS activity in the anther of open flowers was the result of GUS expression in mature pollen grains.

To define the stages of pollen development at which GUS activity was expressed, pollen was costained with 5-bromo-4-chloro-3-indolyl- $\beta$ -glucuronide cyclohexylamine salt and DAPI (Coleman and Goff, 1985). No GUS activity

was detected in microspores prior to the binuclear stage, and very weak GUS activity was detected in binuclear microspores (48 h of staining was required to detect any GUS activity). GUS expression started to increase in trinuclear pollen and reached a maximum in mature pollen (at anthesis), being detectable only after 6 h of staining. GUS was also found in transmitting tissue of carpels during pollination but not before or after pollination. Staining of in vitro-germinated transgenic pollen tubes suggests that the staining observed in the transmitting tissue resulted from GUS expression in pollen tubes (data not shown).

A weak GUS activity was also detected in the tapetum at the early stages of microspore development, which would be consistent with the accumulation of low levels of *Rop1At* transcripts during early flower development. This activity was only detectable following an extended staining (48 h). The tapetal GUS expression remained throughout pollen



**Figure 4.** Construction of *Rop1At* promoter:GUS fusion gene. *Rop1At* genomic clone containing the putative promoter region was cloned into the binary vector as described in the text. Shown is the region joining *Rop1At* and the GUS gene. The restriction sites used for translational fusion with the GUS gene are shown in bold; underlined sequences are the partial coding region for *Rop1At*. The last ATG codon shown is the translation-initiation codon for the GUS gene.

development until the degeneration of tapetal cells (see Fig. 5).

The GUS expression pattern described above is in accordance with the accumulation of *Rop1At* transcripts during flower development, indicating that the 1.3-kb genomic fragment truly represents the *Rop1At* promoter. The 1.3-kb fragment was sequenced to determine whether *Rop1At* promoter sequences contain the *cis*-acting elements required for the expression of pollen genes. It contained a region located 370 bp upstream of ATG (GTAATTGTGA) with a strong homology (9 of 10 bp matches to the 56/59 box) to a pollen-specific enhancer sequence shared by the *LAT56* and *LAT59* promoters (Twell et al., 1991). The GTGA motif within this box is essential for high levels of pollen-specific expression (Twell et al., 1991; Twell, 1994). At least two additional GTGA motifs, located 573 and 461 bp upstream of ATG, are present in the *Rop1At* promoter. Similar GTGA motifs are found in the promoters of several other pollen genes, e.g. *Zmg13* from maize, *chiA* from petunia, and GBAN215–6 and GBAN215–12 from Chinese cabbage (Hamilton et al., 1989; van Tunen et al., 1990; Kim et al., 1997). The *Rop1At* transcript accumulation and GUS fusion gene expression patterns, together with the presence of pollen-specific *cis*-elements, demonstrate that *Rop1At* is a late pollen gene.

### **Rop1At May Function in Fission Yeast to Regulate Polarized Cell Growth**

On the basis of its tip localization and involvement in pollen tube growth, we speculated that Rop1Ps and its homologs might be a molecular switch in the signal transduction pathway leading to polarized cell growth (Lin and Yang, 1997). The function of certain Rho-type GTPases is conserved across kingdoms, e.g. human CDC42 is able to complement the temperature-sensitive yeast *cdc42* mutants defective in polarity control. Therefore, we wanted to determine whether *Rop1At* is able to control polarized cell growth in fission yeast, which, like pollen tubes, elongates by tip growth. First, we overexpressed *Rop1At* in fission yeast under the control of the thiamine-repressible *nmf1* promoter. As shown in Figure 6, *Rop1At* overexpression induced dramatic morphological changes. The majority of cells become shorter and fatter and rounded or pear-shaped, in contrast to the elongated, rod-shaped wild-type

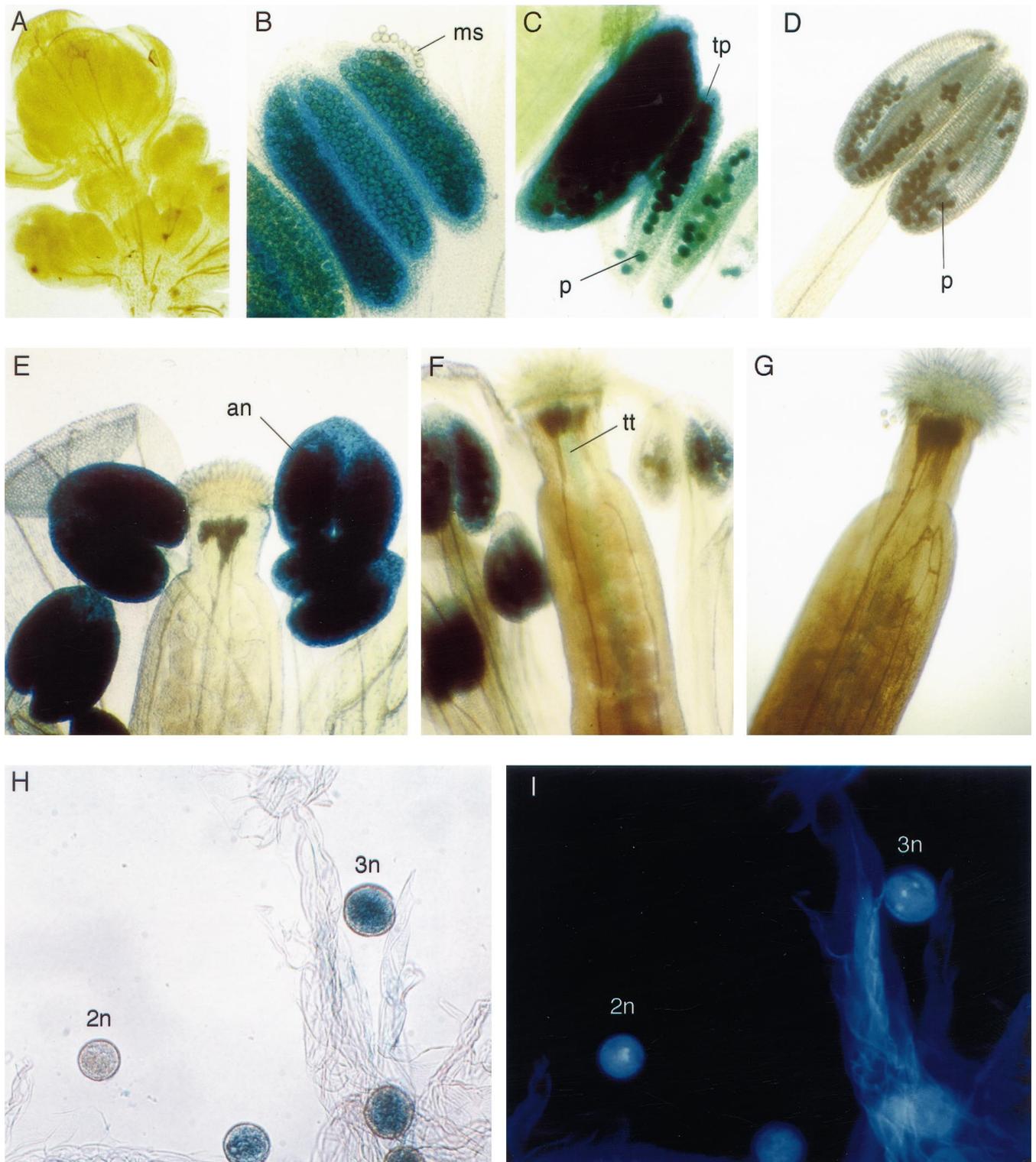
cells. When cells containing the *Rop1At* gene were grown in a repressive medium, or when cells containing the pREP3X plasmid alone were grown in a nonrepressive medium, the cell morphology was normal. The *Rop1At*-induced morphological changes were similar to those caused by the overexpression of constitutively active mutants of the fission yeast CDC42 gene, which is implicated in the control of polarized cell growth (Miller and Johnson, 1994). A *Rop1Ps*-related gene (designated here as *Rop6At*) was isolated during the screening of an Arabidopsis cDNA library for cDNA clones that induced nonpolarized growth in fission yeast (Xia et al., 1996).

We expected that *Rop1At* would be localized to polar sites if its function is to regulate polarized growth in fission yeast. We investigated the subcellular localization of *Rop1At* in living yeast cells using jellyfish GFP. The GFP-*Rop1At* fusion gene is expressed in fission yeast under the control of the thiamine-repressible *nmf1* promoter. Like the overexpression of the wild-type *Rop1At* gene, the fusion gene also induced nonpolarized phenotypes under nonrepressive conditions (data not shown). The severity of the phenotype was correlated with the level of fusion protein expression, as indicated by the intensity of the green fluorescence. Under partially repressive conditions (2 mM thiamine), most cells contained low fluorescence and were relatively normal in shape. Several fluorescence patterns were observed in these cells.

Overall, the fusion protein appeared to be localized to the plasma membrane, which is consistent with the presence of a polybasic domain at the C terminus of *Rop1At* (the polybasic domain has been shown to mediate targeting of isoprenylated proteins to the plasma membrane; Hancock et al., 1991; Adamson et al., 1992). Fluorescence was concentrated in the septum in dividing cells. Soon after cell division was completed, however, fluorescence shifted to old ends, where tip growth is re-initiated (unipolar growth). When cell elongation was shifted to a bipolar pattern, GFP was concentrated at both ends of the cell. Under partially repressive conditions, certain cells contained strong cytoplasmic fluorescence and were completely rounded and somewhat enlarged. In these cells GFP became uniformly distributed on the plasma membrane. Such cells most likely contained a high copy number of plasmids. In nonrepressive conditions most cells were rounded and did not show polarized localization of the fusion protein. The *Rop1At*-induced isotropic growth, together with its polar localization in fission yeast, suggest that *Rop1At* has a conserved function in regulating polarized cell growth.

### **DISCUSSION**

Although the Rho-GTPases are conserved in eukaryotic cells as key regulators of actin cytoskeletal organization, emerging evidence from fungi and mammals suggests that members of the Rho family have also diverged considerably in both structure and function as various eukaryotic phyla evolve. The current data suggest that the plant-specific Rop subfamily of Rho-GTPases has a conserved function in the regulation of polarized cell growth. How-



**Figure 5.** Histochemical localization of GUS expression in transgenic *Arabidopsis* plants carrying the *Rop1At* promoter:GUS fusion gene. Various parts of transgenic  $T_2$  plants were stained with 5-bromo-4-chloro-3-indolyl- $\beta$ -glucuronide cyclohexylamine salt, as described in the text. Typical staining patterns are shown. Anthers were costained with DAPI to determine the developmental stages of pollen. an, Anther; ms, microspore; p, pollen; tp, tapetum; tt, transmitting tissue; 2n, pollen at bicellular stage; 3n, pollen at tricellular stage. A, Early stages of floral buds; B, anthers at the stage of microspore development; C, anthers just prior to anthesis; D, anthers just after anthesis; E, stigma and anthers at anthesis; F, stigma and anthers after anthesis; G, stigma from stage-16 flowers; H, pollen at various developmental stages; I, pollen from H costained with DAPI.

ever, phylogenetically distinct subgroups of the Arabidopsis Rop subfamily exhibit different developmental expression patterns. One of these subgroups is of particular interest, in that all of its members are expressed in mature pollen implicating them in pollen tip growth.

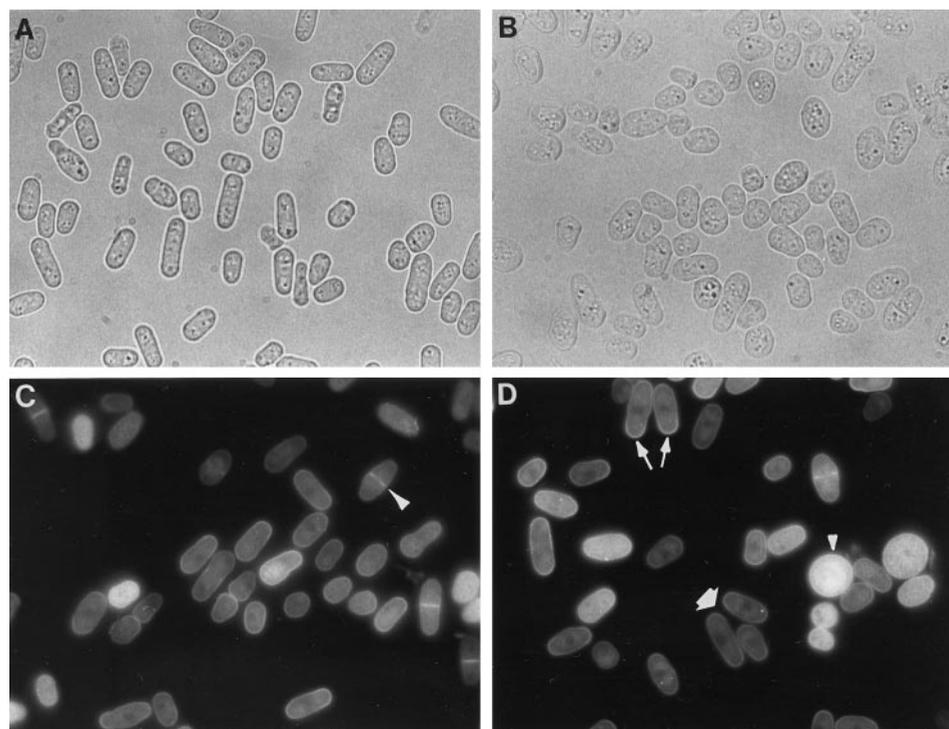
### Plants Have Evolved a Distinct Subfamily of Rho-GTPases

The Rho family of small GTP-binding proteins characterized in fungi and animals include three major subfamilies: CDC42, Rac, and Rho (Chardin, 1993; Hall, 1994; Nobes and Hall, 1995). In mammalian cells each of the three subfamilies controls a specific actin-dependent process (Ridley and Hall, 1992; Ridley et al., 1992; Luo et al., 1994; Chant and Stowers, 1995; Nobes and Hall, 1995). The distinct function for each subfamily is reflected in their amino acid sequence and in their ability to interact with specific downstream effector proteins (Chardin, 1993; Nagata and Hall, 1996; Ridley, 1996). For example, members within a subfamily share 80% or greater amino acid sequence identity, whereas sequence identity between subfamilies ranges from 45% to 70% (Chardin, 1993).

We propose that plants possess a distinct subfamily of Rho-GTPases called Rop. Although Rop is most similar to Rac, phylogenetic analysis suggest that Rop evolved prior to the divergence of Rac and CDC42. Rop members share

many unique motifs or residues that presumably determine functional specificity of these GTPases. Within the conserved effector domain (residues 29–49) exist several Rop-specific residues (T30, T33, F43, and V48). A Rho family-specific insert region consists of 12 amino acid residues (Thr-Arg-Arg-Glu-Leu-Ala-Lys-Met-Lys-Ala-Glu-Pro) in all fungal and animal Rho-GTPases (Chardin, 1993) and functions as an effector domain (Nisimoto et al., 1997). Although the Rop subfamily also contains a corresponding region of 10 residues (residues 128–137), its consensus amino acid sequence (Phe-Phe-Val-Asp-His-Pro-Gly-Ala-Val-Pro) is quite different. The fact that the Rop subfamily is absent from *S. cerevisiae* and has not yet been found in animals indicates that it is unique to plants.

Each eukaryotic kingdom has evolved a specific set of Rho-GTPases (Fig. 1A). The Rho subfamily is found both in fungi and animals and is most likely to exist in plants (Lin and Yang, 1997). The ancestor of the CDC42/Rac/Rop group may have diverged from the Rho subfamily and split into CDC42 and the plant-specific Rop. CDC42, which controls cell polarity and cortical actin formation in both fungi and animals, has not been found in plants. Rac, which appears to have split from CDC42 and become animal specific, regulates actin-dependent cell movement. Therefore, it is logical to speculate that Rop has retained certain conserved functions of the Rho family (e.g. cell



**Figure 6.** Overexpression of Rop1At and polar localization of GFP:Rop1At fusion in fission yeast. The *Rop1At* or *GFP:Rop1At* fusion genes were cloned into pREP3X under the control of a thiamine-repressible promoter and introduced into fission yeast, as described in the text. A, Yeast cells with pREP3X-Rop1At grown in a repressive medium containing 5 mM thiamine. Cells have a normal morphology. B, Yeast cells containing pREP3X-Rop1At grown in a nonrepressive medium lacking thiamine. Greater than 90% were abnormal in shape. C and D, Yeast cells containing pREP3X-GFP:Rop1At grown in a partially repressive medium containing 2 mM thiamine and examined under an epifluorescence microscope. Typical GFP localization patterns are indicated: long arrow, unipolar localization; thick arrow, bipolar localization; long arrowhead, localization to the septum; short arrowhead, nonpolar localization in GFP-Rop1At-overexpressing cells.

polarity control) and has also evolved to perform specific functions that are unique to plant cells. This hypothesis is clearly supported by our studies showing that Rop plays an essential role in pollen tube growth in pea (Lin et al., 1996; Lin and Yang, 1997), that Rop1At exhibits polarized localization and induces isotropic growth when overexpressed in fission yeast (Fig. 6), and that different subgroups of Rop GTPases exhibit distinct patterns of developmentally regulated gene expression in the male gametophyte and various vegetative tissues (Figs. 3 and 5).

### A Specific Rop Subgroup Is Conserved in Protein Structure and Developmental Gene Expression in the Male Gametophyte

Our studies show that the Rop subfamily can be further divided into several subgroups on the basis of primary structure and gene-expression patterns. Like Rop1Ps, all members of the Rop1 subgroup, Rop1At, Rop3At, and Rop5At, are expressed in mature pollen, whereas members of the second subgroup, Rop2At and Rop4At, are constitutively expressed in vegetative tissues but not in pollen. Furthermore, two other divergent Rop GTPases, Rop6At (this study) and Arac2 (Winge et al., 1997), exhibit differential expression patterns. Thus, the Rop subfamily may be divided into the reproductive class (the Rop1 subgroup expressed in pollen) and the vegetative class (Rop2At, Rop4At, Rop6At, Arac2), as do actins and the actin-binding proteins called profilins; Staiger et al., 1993; An et al., 1996a, 1996b; Christensen et al., 1996; Huang et al., 1996a, 1996b, 1997; McDowell et al., 1996). It is interesting that all three types of conserved proteins implicated in the organization of the actin cytoskeleton (actin, profilin, and Rho) exhibit a tight linkage between their structural conservation and developmental gene regulation in pollen. An important question is whether such a correlation reflects a possible functional conservation of these proteins in the regulation of certain pollen-specific arrays of the actin cytoskeleton. The ability to systematically knock out specific genes in Arabidopsis should allow this question to be addressed.

### Rop1At May Have a Distinct Role in the Control of Polarized Growth in Pollen

We have demonstrated that *Rop1At* displays a unique expression pattern associated with the development and the function of pollen. Although Rop3At and Rop5At are also expressed in pollen, their transcript levels are only a small fraction of *Rop1At* transcripts in pollen. This suggests that Rop1At has a predominant role in pollen development and function, whereas Rop3At and Rop5At may be functionally redundant to Rop1At.

Further insight into potential roles for Rop1At was gained by investigating its temporal and spatial expression patterns and by functional analyses in fission yeast. Analyses of GUS fusion gene expression in transgenic plants indicate that *Rop1At* is specifically expressed in pollen. *Rop1At* transcription is activated after the first postmeiotic cell division, reaches a maximum when pollen is mature,

and remains active during pollen tube growth. This expression, which is typical of a late pollen gene, is consistent with a role for *Rop1At* in pollen germination and tube growth. Nonetheless, the functional analyses in fission yeast suggest that Rop1At has a conserved function in the regulation of polarized cell growth. Polar localization of GFP-Rop1At fusion protein to the site of cell growth in yeast is analogous to that of Rop proteins in pea and Arabidopsis pollen tubes suggested by our immunofluorescence studies (Lin et al., 1996; Y. Lin and Z. Yang, unpublished results). These results led us to conclude that Rop1At and Rop1Ps may be orthologs in regard to their potential roles in the regulation of polarized cell growth in Arabidopsis and pea pollen tubes.

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The accession numbers for sequences reported in this study are: *Rop1At*, U49971; *Rop2At*, U49972, same as *Arac4* (U45236); *Rop3At*, same as *Arac1* (U41295); *Rop4At*, AF031428, same as *Arac5* (U52350); *Rop6At*, AF031427, same as *Arac3* (U43501) or clone AT43 (U62746); and *Rop1At* promoter (AF064082).

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