A Green Fluorescent Protein Fusion to Actin-Binding Domain 2 of Arabidopsis Fimbrin Highlights New Features of a Dynamic Actin Cytoskeleton in Live Plant Cells1[w]

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The actin cytoskeleton coordinates numerous cellular processes required for plant development. The functions of this network are intricately linked to its dynamic arrangement, and thus progress in understanding how actin orchestrates cellular processes relies on critical evaluation of actin organization and turnover. To investigate the dynamic nature of the actin cytoskeleton, we used a fusion protein between green fluorescent protein (GFP) and the second actin-binding domain (fABD2) of Arabidopsis (Arabidopsis thaliana) fimbrin, AtFIM1. The GFP-fABD2 fusion protein labeled highly dynamic and dense actin networks in diverse species and cell types, revealing structural detail not seen with alternative labeling methods, such as the commonly used mouse talin GFP fusion (GFP-mTalin). Further, we show that expression of the GFP-fABD2 fusion protein in Arabidopsis, unlike GFP-mTalin, has no detectable adverse effects on plant morphology or development. Time-lapse confocal microscopy and fluorescence recovery after photobleaching analyses of the actin cytoskeleton labeled with GFP-fABD2 revealed that lateral-filament migration and sliding of individual actin filaments or bundles are processes that contribute to the dynamic and continually reorganizing nature of the actin scaffold. These new observations of the dynamic actin cytoskeleton in plant cells using GFP-fABD2 reveal the value of this probe for future investigations of how actin filaments coordinate cellular processes required for plant development.

The actin cytoskeleton coordinates numerous cellular processes required for plant development, including targeted-vesicle transport, to establish hormone gradients (Muday, 2000), polarity determination to direct localized-wall biosynthesis (Baluska et al., 2003), and organelle positioning in response to environmental cues (Takagi, 2003) or to achieve unbiased organelle inheritance (Sheahan et al., 2004). Actin has no known enzymatic activity, and thus its participation in such diverse cellular processes resides in its ability to provide a dynamic filamentous scaffold upon which many regulatory and motor proteins interact (Staiger et al., 2000, and refs. therein). Actin cytoskeleton function, therefore, is intricately linked to its dynamic structure, and thus progress in understanding how actin orchestrates cellular processes will depend in part on critical evaluation of actin organization and turnover in plant cells.

The recent discovery of the Arp2/3 complex as a regulator of plant cell morphogenesis (Le et al., 2003; Li et al., 2003; Mathur et al., 2003a, 2003b; El-Assal et al., 2004) provides a pertinent example of structure function relationships relevant to the plant actin cytoskeleton. The Arp2/3 complex in nonplant systems nucleates the formation of dendritic actin networks by promoting nucleation and growth of new actin filaments (AFs) to the sides of existing AFs (Svitkina and Borisy, 1999; Blanchoin et al., 2000). Filament nucleation by this mechanism facilitates plasma membrane protrusion at the leading edge of migrating cells and other motility related phenomena (for review, see Welch and Mullins, 2002). The ability to image detailed AF organization in these systems has contributed substantially to understanding how the Arp2/3 complex functions (e.g. Svitkina and Borisy, 1999). In contrast, while genetic evidence clearly establishes a role for the plant Arp2/3 complex in cellular morphogenesis, some disagreement exists as to the precise effects different mutations in Arp2/3 components have on actin organization (Le et al., 2003; Li et al., 2003) and thus how Arp2/3 may regulate such processes.

Unlike microtubules, the plant actin cytoskeleton can be difficult to preserve by conventional fixation...
and embedding techniques (Kost et al., 1999; Vitha et al., 2000). Microinjection of fluorescent-phalloidin conjugates has provided novel insights into the dynamic nature of AFs (Schmit and Lambert, 1990) but is not applicable to most cell types and may stabilize actin dynamics. These difficulties have led to the use of various green fluorescent protein (GFP) fusions to image the actin cytoskeleton in live plant cells (Kost et al., 1999; Klahre et al., 2000; Timmers et al., 2002). The most commonly used is GFP tagged to the carboxy-terminal actin-binding domain (ABD) of mouse talin (GFP-mTalin; Kost et al., 1998), or recently human talin (GFP-hTalin; Takemoto et al., 2003; Kwock and Hanson, 2004). The GFP-mTalin fusion, however, does not label all identifiable actin arrays in plant cells (Le et al., 2003; El-Assal et al., 2004), and some images of actin networks labeled with this probe suggest that the GFP-mTalin fusion may cause artificial aggregation of actin networks (e.g. see Jedd and Chua, 2002; Oikawa et al., 2003). These observations suggest GFP talin fusions may not accurately depict actin organization and turnover in plant cells.

To explore alternatives for GFP tagging of the plant actin cytoskeleton, we have used a fusion between GFP and ABD2 of the Arabidopsis (Arabidopsis thaliana) fimbrin AtFIM1 (fABD2). AtFIM1 binds plant AFs both in vitro (Kovar et al., 2000) and in vivo (Kovar et al., 2001), and both protein sequence analysis (McCurdy and Kim, 1998) and in vitro binding data (Kovar et al., 2000) suggest that AtFIM1 contains two independent ABDs. Recently, we used this GFP-fABD2 fusion to demonstrate detailed features of the actin cytoskeleton in protoplasts derived from mesophyll tissue of stably transformed Nicotiana tabacum (Sheahan et al., 2004). This fusion protein labeled a highly dynamic and dense actin cytoskeleton in protoplasts, suggesting that it reveals more structural detail than alternative GFP-fusion constructs. GFP-fABD2 has also been used to analyze actin cytoskeletal organization in Arabidopsis lines with modified levels of actin-interacting protein 1 (Ketelaar et al., 2004), and recently a series of GFP fusion constructs exploiting the ABDs of AtFIM1 has been used to image the actin cytoskeleton in Arabidopsis roots (Wang et al., 2004).

We report here quantitative, dynamic analysis of the plant actin cytoskeleton by stable expression of GFP-fABD2 in different species and cell types. In comparison to GFP-mTalin (Kost et al., 1998), GFP-fABD2 consistently labels finer and denser networks of cortical and subcortical actin, and does so without adverse effects on plant morphology or development. Fluorescence recovery after photobleaching (FRAP) in cells from GFP-fABD2-expressing lines revealed that AF bundles are highly dynamic structures, composed of AFs capable of sliding in opposite directions. Our study provides new insights into the complexity of the actin cytoskeleton and its dynamic properties in plant cells, and we conclude that GFP-fABD2 currently provides the best available GFP-fusion protein for investigating dynamic actin networks in live cells.

RESULTS

Expression of GFP-fABD2 Highlights Elaborate Actin Networks without Adverse Effect on Plant Morphology

Expression of GFP-fABD2 in stably transformed Arabidopsis plants revealed dense networks of free and bundled AFs in the cell types examined (Fig. 1). Such arrays were present as dense filamentous networks in the cortex (Fig. 1, B-D) and perinuclear region (Fig. 1, B and D) of hypocotyl, pavement epidermal, and trichome cells, or as masses of bundled filaments in transvacuolar strands of pavement epidermal cells (Fig. 1C) and cortex of mature root epidermal cells (Fig. 1A). Guard cells had notably higher fluorescence intensities compared with surrounding epidermal cells, and AFs were more difficult to visualize. However, intricate cortical and perinuclear meshworks (Fig. 1E) or radially aligned AF bundles (Fig. 1E inset) were commonly observed. AF bundles were frequently arrayed longitudinally along the axis of expansion in elongating cells while in diffuse-growing pavement epidermal cells, bundles were arrayed more randomly. In contrast to previous findings (Li et al., 2003), diffuse fluorescence, interpreted to represent fine AF arrays, in lobed regions of expanding pavement epidermal cells from both Arabidopsis (Fig. 1C, inset) and N. tabacum (data not shown) was rare and appeared to be present equally in regions of lobe expansion as it was in constrictions. Similar diffuse labeling was frequently observed at the ends of growing root hairs (Fig. 1A, inset) and trichome tips (Fig. 1D, inset). We interpret this diffuse fluorescence to represent fine AF arrays; however, our attempts to further resolve this fluorescence by conventional confocal microscopy were unsuccessful.

In addition to GFP-fABD2, we tested GFP fusions to the amino terminus of full-length AtFIM1 (GFP-AtFIM1) and to either the carboxy or amino terminus of the first ABD (fABD1) of AtFIM1 (fABD1-GFP, GFP-fABD1; Fig. 2A). Transient expression of GFP-AtFIM1 in N. tabacum leaf epidermal cells showed predominantly filamentous labeling, but labeling became cytoplasmic in regions where expression levels were locally higher (Fig. 2B). Expression of either fABD1-GFP or GFP-fABD1 consistently resulted in cytoplasmic labeling (Fig. 2, C and D), in contrast to GFP-fABD2, which labeled distinctly filamentous structures in all but the most highly expressing cells (Fig. 2E). Similar results were obtained for GFP-AtFIM1 and GFP-fABD2 with stably expressing N. tabacum mesophyll protoplasts, confirming these findings (data not shown). We therefore conclude that, of the constructs tested, GFP-fABD2 provides the best GFP fusion protein for consistent labeling of the actin cytoskeleton.

Our analysis of Arabidopsis plants stably transformed with the GFP-mTalin fusion protein (Kost et al., 1998) revealed broadly similar structures to those in GFP-fABD2 plants (Fig. 1, F-J); however, close examination revealed that AFs in GFP-mTalin plants


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displayed increased bundling and were more randomly arranged (compare Fig. 1, B, G, and insets). AF networks in these plants generally displayed increased branching, being composed of shorter, more convoluted filaments, while in root hairs and trichomes, longitudinal AF bundles tended to form into helical arrays (Fig. 1, F and I). Plants expressing GFP-mTalin also showed fluorescence localized to the nucleoplasm (arrows in Fig. 1, F and I), whereas cells expressing GFP-fABD2 exhibited neither organellar nor nucleoplasmic staining. Notably, due to small cell size, tissue organization, and low levels of expression, we were unable to satisfactorily visualize AFs in the root meristem of plants stably transformed with either GFP-fABD2 or GFP-mTalin (data not shown). This observation is consistent with the generally poor visualization of actin networks seen in meristematic cells by Wang et al. (2004) using fimbrin-based fusion proteins.

We compared plant development in transgenic Arabidopsis seedlings expressing either GFP-mTalin or GFP-fABD2. Although reported to have no effects on plant morphology or development (Kost et al., 1998), plants expressing GFP-mTalin in our experiments showed retarded growth compared to wild-type or GFP-fABD2 plants. In 4-d-old etiolated seedlings, plants expressing GFP-mTalin showed retarded hypocotyl elongation but only slightly reduced root growth (Fig. 3A). However, 3 d after transfer to light, differences in the growth of GFP-fABD2- and GFP-mTalin-expressing plants were clear, with reduced growth in all organs in GFP-mTalin plants (Fig. 3B). Mature GFP-mTalin-expressing plants were stunted and had short inflorescences and stout siliques (Fig. 3C) that carried only 45% ± 7% (mean ± se; n = 4) of the seed of GFP-fABD2-expressing siliques. Further analysis of epidermal cells from hypocotyls, inflorescences, siliques, and leaves indicated that cells from plants expressing GFP-mTalin had smaller cell areas compared with GFP-fABD2 plants (Table I). Cell perimeter was also reduced in GFP-mTalin plants at 69% ± 4% (mean ± se; n = 385), the length of cell peripheries in GFP-fABD2-expressing plants (Table I). Most distinct, however, was the difference in the length of cells, with GFP-mTalin-expressing plants having cells on average 66% ± 4% (mean ± se; n = 385) the length of GFP-fABD2 cells (Table I). Average cell width in the two lines, however, was similar. Stunted growth in GFP-mTalin plants is therefore primarily due to reduced cell elongation.

GFP-mTalin expression affected diffuse-growth processes in leaves, hypocotyls, inflorescences, and siliques. However, no significant differences were found between GFP-mTalin- and GFP-fABD2-expressing root hairs, suggesting that tip-growth processes remain unaffected, consistent with the finding that GFP-mTalin expression does not perturb pollen tube growth in N. tabacum (Kost et al., 1998). In contrast to our general finding that GFP-mTalin expression causes reduced cell elongation, expression of GFP-fABD2 in Arabidopsis had no measurable effect on plant growth, suggesting...
that the GFP-fABD2 fusion protein interferes minimally with native AF function.

Estimation of GFP levels in GFP-fABD2- and GFP-mTalin-expressing plants, based on quantification of total GFP fluorescence in individual cells (Table II), indicated that GFP-mTalin expression levels were roughly twice that of GFP-fABD2 in the plants used in this study (Table II). In agreement with this analysis, semiquantitative immunoblotting using an anti-GFP antibody and whole 7-d-old seedlings revealed that the abundance, relative to α-tubulin, of GFP-mTalin was 1.6 ± 0.9-fold (mean ± se; n = 3) greater than GFP-fABD2 in the respective transgenic lines (see Supplemental Fig. 1, available at www.plantphysiol.org). We were unable to generate stable lines of Arabidopsis expressing GFP-fABD2 to levels comparable to that seen in GFP-mTalin plants. However, when GFP-fABD2 was transiently expressed in N. tabacum pavement epidermal cells to levels comparable to GFP-mTalin plants, as determined by measurement of total cellular GFP fluorescence, GFP-fABD2 accumulated in the cytoplasm (data not shown). These observations suggest fundamental differences in the mechanisms of GFP-mTalin and GFP-fABD2 binding in plant cells, and the consequent effects this may have on plant development. Our conclusion that expression of GFP-fABD2 does not perturb actin functionality is supported by the observation that chloroplast repositioning during N. tabacum protoplast culture proceeds Figure 3. Expression of GFP-mTalin affects plant growth and morphology. A, Root and hypocotyl elongation of 4-d-old, dark-grown seedlings; elongation is retarded in GFP-mTalin, but not GFP-fABD2 plants. B, Retarded growth in 7-d-old GFP-mTalin but not GFP-fABD2 plants. C, Altered silique morphology in GFP-mTalin plants. Error bars shown in A, se (n = 20); scale bar = 5 mm (B and C).
at similar rates in both wild-type and GFP-fABD2-expressing cells (Sheahan et al., 2004).

**GFP-fABD2 Labels Actin Networks in Diverse Cell Types from Different Species**

We investigated the usefulness of the GFP-fABD2 fusion protein to label actin networks in diverse cell types from different species. In both stably transformed *N. tabacum* BY-2 and *Medicago truncatula* suspension-cultured cells, we observed intricate networks of nucleus-associated AF arrays and dense, at times transversely oriented, cortical networks (Fig. 4, A and C). In BY-2 cells, transverse bundles of cortical actin were more prominent in smaller, expanding cells compared with older, expanded cells (Fig. 4A). In cells predicted to be entering division, actin was arranged prominently as complex transvacuolar strands linking the subcortical-perinuclear region with the cortex (Fig. 4, A and B). This observation indicates that reorganization of the actin cytoskeleton occurs during the transition from cell expansion to division. In mesophyll tissue from *N. tabacum* and *Lycopersicon esculentum*, dense cortical arrays and chloroplast actin baskets were evident (Fig. 4D and insets). These results demonstrate that expression of the GFP-fABD2 fusion protein provides detailed visualization of the actin network in live cells, across different species and cell types.

**GFP-fABD2 Highlights Dynamic Networks of AFs**

We analyzed the dynamic nature of the actin network with time-lapse imaging (see supplemental videos), Latrunculin B (LatB) washout experiments, and FRAP. Time-lapse imaging of epidermal cells in leaves, hypocotyls, and roots of 7-d-old plants revealed that the AF network labeled with GFP-fABD2 underwent continual reorganization. Closer analysis established that the highly dynamic nature of this network arose predominantly from the small movements of a large number of filaments or bundles. Transvacuolar bundles abutting the cell periphery underwent lateral migrations, while regions where bundles of AFs intersected, termed focal junctions, moved rapidly around the cell (Supplemental Video 1). Measurement of focal junction movement in pavement epidermal cells using centroid tracking revealed irregular, yet directional movement, with speeds ranging from 80 to 800 nm s⁻¹ (mean, 248 ± 38; n = 20) and velocities from 25 to 600 nm s⁻¹ (mean, 140 ± 32; n = 20). Both lateral AF movement and dynamic movements of focal junctions were difficult to observe in plants expressing GFP-mTalin (Supplemental Video 2), suggesting that this fusion protein stabilizes an otherwise highly dynamic network.

To estimate the contribution of filament turnover to dynamic processes, we used LatB washout experiments. Treatment of GFP-fABD2-expressing *N. tabacum* protoplasts for 24 h with 1 μM LatB caused complete depolymerization of AFs in most cells, with GFP-fABD2 visualized as localized accumulations of fluorescence in the cell cortex (Fig. 5A and insets). Monitoring recovery of the actin network following LatB washout indicated that most cells had at least a partially repolymerized network within 1 h of washing (Fig. 5A). Treatment of GFP-fABD2-expressing Arabidopsis seedlings with 2 μM LatB for 8 h also caused an almost complete depolymerization of the actin network in near-fully expanded pavement epidermal cells (Fig. 5B, section a). Recovery following thorough washing was swift, and within 1.5 h, most cells had actin networks (Fig. 5B, section c) similar to dimethyl sulfoxide (DMSO)-treated controls (Fig. 5B, section b). These networks were dynamic, as indicated by time-lapse imaging (Supplemental Video 3); however, bundles of AFs appeared somewhat thicker and shorter than those in DMSO controls. Parallel analysis of GFP-mTalin plants similarly indicated an almost complete depolymerization of the actin network in the presence of LatB. In contrast, however, reassembly of the actin network following LatB washout was substantially retarded, with few cells exhibiting polymerized filaments but many exhibiting short, rod-like structures (Fig. 5B, section d). Time-lapse analysis indicated that these rods moved slightly, but overall, dynamic movement was substantially reduced compared to GFP-fABD2 plants (Supplemental Video 3). These results suggest that filament assembly/disassembly processes contribute significantly to observed

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**Table I. Epidermal cell characteristics in GFP-fABD2- and GFP-mTalin-expressing Arabidopsis plants**

Epidermal cells from hypocotyls, inflorescences, siliques, and leaves of cleared GFP-fABD2 and GFP-mTalin-expressing plants were imaged, and measurements of individual cells obtained. Values are means ± se, n = 385.

<table>
<thead>
<tr>
<th>Measurement</th>
<th>GFP-fABD2</th>
<th>GFP-mTalin</th>
</tr>
</thead>
<tbody>
<tr>
<td>Plan Area (μm²)</td>
<td>2,639 ± 95</td>
<td>2,127 ± 98</td>
</tr>
<tr>
<td>Cell Perimeter (μm)</td>
<td>363 ± 9</td>
<td>249 ± 8</td>
</tr>
<tr>
<td>Cell Length (μm)</td>
<td>139 ± 3</td>
<td>92 ± 3</td>
</tr>
<tr>
<td>Cell Width (μm)</td>
<td>26 ± 1</td>
<td>28 ± 1</td>
</tr>
<tr>
<td>Circularity</td>
<td>0.285 ± 0.006</td>
<td>0.454 ± 0.008</td>
</tr>
</tbody>
</table>

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**Table II. GFP fluorescence levels in GFP-fABD2- and GFP-mTalin-expressing Arabidopsis plants**

<table>
<thead>
<tr>
<th>Actin Label</th>
<th>Root hairs</th>
<th>Hypocotyl Epidermis</th>
<th>Leaf Epidermis</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Fluorescence Intensity μm⁻³</td>
<td></td>
<td></td>
</tr>
<tr>
<td>GFP-fABD2</td>
<td>9.7 ± 0.4</td>
<td>5.7 ± 0.1</td>
<td>10.1 ± 0.5</td>
</tr>
<tr>
<td>GFP-mTalin</td>
<td>22.7 ± 0.6</td>
<td>11.5 ± 0.7</td>
<td>16.6 ± 0.5</td>
</tr>
</tbody>
</table>

Quantification of GFP fluorescence in individual root hairs and epidermal cells from hypocotyls and leaves of GFP-fABD2- and GFP-mTalin-expressing plants. Values are means ± se, n ≥ 20.
sections. Scale bar 71 (B), 15 (C and D top inset), or 4 (D and D bottom inset) optical mesophyll (bottom inset). Cells predicted to be entering division are tum... suspension cultures. D, Cortical AF arrays in N. tabacum... GFP-fABD2 fluorescence recovery, we first analyzed FRAP in expanding epidermal cells. By comparing the ratio of fluorescence recovery at the ends of the bleached bundle (left and right boxes in Fig. 5C) with the central region of the bundle (center box in Fig. 5C), we found that fluorescence recovered along the whole bundle, but recovery initially occurred faster and to a greater extent at either end of the bundle (Fig. 5D). We also compared the mean fluorescence intensity plotted along the horizontal axis of the bleach window over time for both GFP-fABD2-expressing cells and FDA-stained cells (Fig. 5, E and F). This analysis confirmed that fluorescence recovery in bleached AF bundles was greater and faster at each edge of the bleached AF bundle (Fig. 5E). In addition, recovery appeared to be directional in several cases, in that recovery was slightly faster at one end of the bleached bundle compared to the other (Fig. 5E). In contrast, recovery of mean fluorescence intensity was comparatively uniform across the bleach window in FDA-stained cells (Fig. 5F). By comparing recovery in GFP-fABD2 bleached AF bundles and FDA-stained cytoplasmic strands, we found that the ratio of change in fluorescence intensity between the edge and center of the bleach window over time was on average 5.9-fold greater in GFP-fABD2-expressing cells. This analysis confirmed that fluorescence recovery is limited by the exchange of bleached units on AFs with those in the cytoplasm. Fluorescence recoveries were similar in different cell types of GFP-fABD2-expressing plants (Table III), suggesting that the dynamic nature of actin networks are comparable in different cell types.

In FRAP experiments using GFP-fABD2, we did not observe translocation of the bleach window, however the size of the bleached region progressively decreased (Fig. 5C). Similar observations of YFP-expressing and FDA-stained cells indicated that the bleach window sometimes translocated in the direction of net cytoplasmic streaming, but that fluorescence recovery was reasonably uniform across the bleach window (data not shown). To understand the contribution of diffusion to the observed recovery patterns, we analyzed how fluorescence recovered along bleached AF bundles in hypocotyl epidermal cells. By comparing the ratio of fluorescence recovery at the ends of the bleached bundle (left and right boxes in Fig. 5C) with the central region of the bundle (center box in Fig. 5C), we found that fluorescence recovered along the whole bundle, but recovery initially occurred faster and to a greater extent at either end of the bundle (Fig. 5D). We also compared the mean fluorescence intensity plotted along the horizontal axis of the bleach window over time for both GFP-fABD2-expressing cells and FDA-stained cells (Fig. 5, E and F). This analysis confirmed that fluorescence recovery in bleached AF bundles was greater and faster at each edge of the bleached AF bundle (Fig. 5E). In addition, recovery appeared to be directional in several cases, in that recovery was slightly faster at one end of the bleached bundle compared to the other (Fig. 5E). In contrast, recovery of mean fluorescence intensity was comparatively uniform across the bleach window in FDA-stained cells (Fig. 5F). By comparing recovery in GFP-fABD2 bleached AF bundles and FDA-stained cytoplasmic strands, we found that the ratio of change in fluorescence intensity between the edge and center of the bleach window over time was on average 5.9-fold greater in GFP-fABD2-expressing cells. Thus, diffusion alone cannot account for the more rapid recovery of fluorescence at the edges of bleached AF bundles, suggesting that rapid-edge recovery may be the result of lateral sliding in opposite directions of GFP-fABD2-labeled filaments or bundles into the bleach window. Consistent with potential filament sliding, bleached GFP-fABD2-labeled AFs moving out of the bleach window caused fluorescence intensities immediately outside the bleach window to decrease an additional 0.7 ± 0.1-fold (mean ± se, n = 7) more than could be accounted for by bleaching caused by laser scanning during image collection.

Comparison of fluorescence recovery in bleached GFP-fABD2- and GFP-mTalin-labeled AF bundles indicated that the ratio of change in fluorescence intensity between the edge and the center of the bleach window over time was, on average, 1.3-fold greater in
Figure 5. Expression of GFP-fABD2 reveals a dynamic framework of AFs. A, Recovery from LatB treatment in washed N. tabacum protoplast populations; rapid recovery of filamentous actin within 1.5 h of washing. Insets show half-cell (approximately 15 μm), confocal projections of protoplasts treated with 1 μM LatB or 0.1% DMSO for 24 h. B, Recovery from LatB treatment in washed Arabidopsis leaf epidermis; 8 h of LatB treatment (a) caused near complete depolymerization of the network, contrasting normal arrays in 0.1% DMSO-treated controls (b). Substantial recovery of the actin network 1.5 h after LatB washout in GFP-fABD2-expressing plants (c) but not in plants expressing GFP-mTalin (d). C, Fluorescence recovery along bleached AF bundles occurs rapidly and in a pattern resembling the prebleach image; images show typical fluorescence recovery in bleached bundle over 45 s, with left, center, and right boxes used in recovery analysis shown. Fluorescence recovers to 84% of prebleach intensity by 45 s in this example. D, Detailed analysis of fluorescence recovery following bleaching of bundled AFs; graph shows recovery ratio for one end compared with the center; arrows on graph correspond to time point for images displayed on right (n = 7). E, Plotted mean fluorescence intensity over time along bleach window in a GFP-fABD2-tagged transvacuolar actin bundle. F, Plotted mean fluorescence intensity over time along bleach window in an FDA-stained transvacuolar strand; fluorescence recovery is uniform over the bleach window with FDA, but is irregular and occurs fastest at the edges of the bleach window in GFP-fABD2-expressing cells. Insets in A are confocal projections (half-cell) and images in B are single-plane images with a z-depth of 2.5 μm. Scale bar = 10 μm (A–C).
GFP-fABD2-expressing cells, suggesting that filament sliding occurs to a greater extent in these cells. Similar fluorescence-recovery (FRAP) rates and half-times for FRAP taken in the GFP-fABD2 and GFP-AtFIM1 fusion proteins (see Fig. 2), may be important for stabilizing the actin structure and plant growth. The suppressed recovery of AFs following LatB washout and reduced dynamic behavior of actin networks in GFP-mTalin-expressing plants indicate that GFP-mTalin may affect plant growth by interfering with AF polymerization. Evidence from profilin mutants suggests that hypocotyl elongation requires a dynamic actin network, since underexpression of profilin decreases hypocotyl cell elongation (Ramachandran et al., 2000). In plants, anisotropic growth but not polarity determination requires dynamic AFs (Smith, 2003) and, typically, fine AF arrays are associated with rapid, polar cell growth (Waller et al., 2002). Disruption of the actin network with LatB (Baluska et al., 2001) or by overexpression of certain ABDs can alter function of the actin network. Similarly, Wang et al. (2004) found that overexpression of the BD1/2-GFP construct altered actin structure and plant growth. The suppressed recovery of AFs following LatB washout and reduced dynamic behavior of actin networks in GFP-mTalin-expressing plants indicate that GFP-mTalin may affect plant growth by interfering with AF polymerization. Evidence from profilin mutants suggests that hypocotyl elongation requires a dynamic actin network, since underexpression of profilin decreases hypocotyl cell elongation (Ramachandran et al., 2000). In plants, anisotropic growth but not polarity determination requires dynamic AFs (Smith, 2003) and, typically, fine AF arrays are associated with rapid, polar cell growth (Waller et al., 2002). Disruption of the actin network with LatB (Baluska et al., 2001) or by overexpression of a reductive actin (ACT1; Kandasamy et al., 2002) creates plants with a dwarf phenotype but no significant changes in morphology. AFs probably direct vesicular transport, but may also influence cell shape by determining microtubule alignment (Kost et al., 1999). In contrast to GFP-mTalin, which is expressed at high levels, several undetermined factors presumably might contribute to the different outcomes observed for these fusion constructs. In vitro binding assays predict two different actin-binding activities for AtFIM1 (Kovar et al., 2001), and recent biochemical analysis of L-plastin, a human fimbrin homolog, shows that ABD2 binds with lower affinity to actin than does ABD1 (Lebart et al., 2004; but see Nakano et al., 2001). Thus, the potential lower affinity ABD2 of AtFIM1 may provide superior characteristics for non-invasive labeling of AFs. Furthermore, recent analysis of the crystal structure of AtFIM1 suggests that the carboxy-terminal extension of AtFIM1, which is retained in the GFP-fABD2 and GFP-AtFIM1 fusion proteins (see Fig. 2), may be important for stabilizing AF interactions (Klein et al., 2004). Our observations agree with recent work by Wang et al. (2004), who show that different GFP fusions, including ABD2 of AtFIM1, produce filamentous labeling when expressed in Arabidopsis, whereas fusion to ABD1 of AtFIM1 yields cytoplasmic labeling. Expression of GFP-fABD2 in Arabidopsis revealed elaborate and intricate actin networks in the cell types examined. Cortical AFs were organized into linear arrays of predominantly longitudinally aligned AFs in elongating tissues such as hypocotyl and root epidermis. In pavement epidermal cells, cortical actin tended to have a more random alignment. Subcortical actin bundles radiated from the perinuclear AF basket through the vacuole were common to all cell types examined. Similar structures were observed in N. tabacum and L. esculentum cells, and in BY-2 and M. truncatula suspension cultures expressing GFP-fABD2.

Expression of GFP-fABD2 in Arabidopsis did not disrupt plant growth or morphology when compared with nontransgenic controls. In contrast, however, plants expressing GFP-mTalin showed reduced cell elongation, resulting in plants with stunted growth and reduced seed set. Actin networks visualized by the GFP-mTalin construct differed to those visualized by GFP-fABD2, being composed of comparatively shorter, more branched and convoluted filament networks with increased bundling and greatly reduced dynamism.

Our observations indicate that the altered morphological and growth phenotypes of GFP-mTalin-expressing plants are likely to be caused by changes in actin network structure, suggesting that overexpression of certain ABDs can alter function of the actin network. Similarly, Wang et al. (2004) found that overexpression of the BD1/2-GFP construct altered actin structure and plant growth. The suppressed recovery of AFs following LatB washout and reduced dynamic behavior of actin networks in GFP-mTalin-expressing plants indicate that GFP-mTalin may affect plant growth by interfering with AF polymerization. Evidence from profilin mutants suggests that hypocotyl elongation requires a dynamic actin network, since underexpression of profilin decreases hypocotyl cell elongation (Ramachandran et al., 2000). In plants, anisotropic growth but not polarity determination requires dynamic AFs (Smith, 2003) and, typically, fine AF arrays are associated with rapid, polar cell growth (Waller et al., 2002). Disruption of the actin network with LatB (Baluska et al., 2001) or by overexpression of a reductive actin (ACT1; Kandasamy et al., 2002) creates plants with a dwarf phenotype but no significant changes in morphology. AFs probably direct vesicular transport, but may also influence cell shape by determining microtubule alignment (Kost et al., 1999).

In contrast to GFP-mTalin, which is expressed at high relative levels but continues to label highly bundled actin structures, cells showing similar levels of GFP fluorescence in GFP-fABD2-expressing cells caused
cytoplasmic accumulation. This finding may reflect differences in the affinity of the carboxy-terminal ABD of talin and ABD2 of AtFIM1 for plant AFs, or reflect a reduced number of available binding sites for GFP-fABD2 given the presence of endogenous fimbrins. The tendency of GFP-mTalin to cause increased bundling may be due to a high density of binding along an AF causing a localized high concentration of GFP. GFP has a weak tendency to dimerize \( (k_d = 100 \mu M; \text{Phillips, 1997}) \) and thus under such conditions may cause cross-linking of AFs in vivo similar to the way McCann and Craig (1997) observed in vitro cross-linking of AFs when the single carboxy-terminal ABD of mTalin was fused to glutathione S-transferase. Interestingly, Wang et al. (2004) observed that plants expressing their independently created GFP-mTalin construct also displayed increased bundling and reduced dynamic behavior of AFs compared to AtFIM1-based GFP fusions.

Potential filament stabilization by GFP-mTalin expression in Arabidopsis plants may also occur, and indeed a carboxy-terminal 63-kD fragment of talin A has been used as a stabilizer of actin networks in Dictyostelium cells (Weber et al., 2002). The apparent inability of plants to regulate the binding of the talin ABD to AFs may also result in the failure of endogenous actin-binding proteins to interact correctly with AFs. Our findings highlight the delicate nature of the actin network in plant cells and emphasize the potential for GFP-fusion approaches to compromise the dynamic nature of actin networks. Based on the data presented, we conclude that the GFP-fABD2 fusion represents the best available probe for visualization of dynamic AFs in live plant cells, and emphasize that low levels of expression are vital for accurate imaging of the actin network in plants. Ultimately, GFP-actin fusions may provide the best approach to investigating dynamic turnover of the plant actin cytoskeleton. However, given the sensitive nature of plant actin networks demonstrated in this study and the large pool of unpolymerized actin in plant cells implicated in studies of pollen tubes (Gibbon et al., 1999) and soybean suspension cultures (Lee et al., 2003), it is likely that GFP-actin fusions may also encounter difficulties.

Several Processes Contribute to Actin Dynamics Observed in Plant Cells

Time-lapse imaging of GFP-fABD2-expressing cells revealed that several, simultaneous dynamic processes contribute to continual reorganization of the actin cytoskeleton. Large-scale, lateral movements of AF bundles occurred where transvacuolar AF bundles abutted cortical bundles and at the intersections of transvacuolar AF bundles (focal junctions) in the population of subcortical actin. Dynamic turnover of the actin network is also likely to contribute appreciably to overall cytoskeletal reorganization as implicated by LatB washout experiments. Detailed FRAP-based analysis suggested that AF bundles represent an actively moving collection of AFs, where either individual AFs or small bundles of AFs slide past one another, probably in opposite directions. Taken together, these results portray the plant actin network as a highly fluidic system of interacting filament-based structures. The ability of the actin cytoskeleton to change its form at several different levels of organization would presumably allow the cell to simultaneously perform the multitude of AF-dependent functions known in plants. Clearly, further analysis into the role of individual actin isovariants and the influence of specific actin-binding proteins on this dynamic behavior is required to understand these processes and their regulation at the molecular level.

MATERIALS AND METHODS

GFP-fABD2 Fusion Construct and Agrobacterium Transformation

Fusions between modified GFP and full-length AtFIM1 (McCurdy and Kim, 1998; GFP-AtFIM1, ABD1 (amino acids 101–570 of AtFIM1; ABD1-GFP and GFP-ABD1), and ABD2 of AtFIM1 (amino acids 325–687 of AtFIM1; GFP-ABD2) were tested (see Fig. 2A). The GFP-AtFIM1 and GFP-ABD2 constructs were supplied by Boris Voigt and Diedrik Menzel (Universität Bonn) in pCAT (see Ketelar et al., 2004) and the ABD1-GFP and GFP-ABD1 constructs by Steven Pirlo (The University of Newcastle) in pART7 expression vector (Gleave, 1992). In the GFP-AtFIM1 and GFP-ABD2 constructs, the appropriate AtFIM1-derived DNA sequence was fused directly with GFP-encoding sequences without an intervening-linker sequence, while the ABD1-GFP and GFP-ABD1 constructs were created with a 5 \( \times \) Gly linker. The 3′SS-GFP-fABD2.35SpA or 35SS-GFP-AtFIM1.35SpA region of pCAT was moved into pGEM-T by restriction digestion with Sall and then into pART27 (Gleave, 1992) using Noll digestion. The 35SS-GFP-fABD1ocs3′ or 35SS/fABD1-GFPocs3′ of pART7 was moved into pART27 using Noll digestion. Each recombinant pART27 vector was transfected into Agrobacterium tumefaciens strain LBA4404 by electroporation and bacteria selected on Luria-Bertani agar with 50 \( \mu \)g ml\(^{-1}\) kanamycin (Sigma, St. Louis).

Plant Transformation

The GFP-fABD2 fusion construct in pART27 was transformed into Arabidopsis (Arabidopsis thaliana; Col) by floral dipping (Clough and Bent, 1998) and transformants selected on 50 \( \mu \)g ml\(^{-1}\) kanamycin. Nicotiana tabacum was stably transformed using the leaf disc procedure of Horsch et al. (1985) as described by Sheahan et al. (2004). \( N. \) tabacum BY-2 cells were stably transformed by the method of Sullivan and Green (1996) with selection on 100 \( \mu \)g ml\(^{-1}\) kanamycin and 500 \( \mu \)g ml\(^{-1}\) carbenicillin (Sigma). The highly re-generable Medicago truncatula line 2HA (Rose et al., 1999) was stably transformed as described by Thomas et al. (1992). \( N. \) tabacum and Lycopersicon esculentum leaves were transiently transformed using the agroinfiltration method (Yang et al., 2000). Arabidopsis seed derived from the GFP-mTalin-expressing line reported by Kost et al. (1998) was used in this study. Madeline Rashbrooke (Australian National University) supplied transgenic Arabidopsis seed expressing cytoplasmic YFP.

Plant Growth

Arabidopsis plants were grown on vertical or horizontal plates containing 0.5 \( \times \) Murashige and Skoog salts, 0.8% (w/v) agar, and 1% (w/v) Suc. Surface-sterilized seeds were positioned on plates and vernalized for 2 d at 4°C before being moved to a growth room (25°C, 16/8 h photoperiod at 50 \( \mu \)mol photons m\(^{-2}\) s\(^{-1}\)). Mesophyll protoplasts were isolated and cultured from axenic shoot cultures of transgenic GFP-fABD2 \( N. \) tabacum as described by Sheahan et al. (2004). \( N. \) tabacum and \( L. \) esculentum plants used for agroinfiltration were grown under glasshouse conditions as described for...
N. tabacum in Stehan et al. (2004). Stably transformed BY-2 cultures were subcultured weekly and maintained in liquid Murashige and Skoog medium (plus 3% [w/v] Suc, 0.2 mg mL\(^{-1}\) 2,4-dichlorophenoxyacetic acid, 255 mg L\(^{-1}\) KH\(_2\)PO\(_4\), 100 \(\mu\)g mL\(^{-1}\) kanamycin, and 250 \(\mu\)g mL\(^{-1}\) Timentin [Glaxo Smith-Kline, Melbourne, Australia], pH 5.7) on a rotary shaker (130 rpm) at 27°C in darkness. M. truncatula suspension cultures were freshly established for each experiment using callus grown on plates containing P4 medium (Thomas et al., 1992). Callus was used to inoculate liquid Murashige and Skoog medium (plus 3% [w/v] Suc, 0.2 mg mL\(^{-1}\) 2,4-dichlorophenoxyacetic acid, 100 \(\mu\)g mL\(^{-1}\) kanamycin, and 250 \(\mu\)g mL\(^{-1}\) Timentin, pH 5.7) and cultures grown as described for BY-2 cells.

Drug Treatments
Protoplasts were treated in culture for 24 h with 1 \(\mu\)M LatB (Merck, Rahway, NJ) or 0.1% (v/v) DMSO before washing three times in modified Nagata and Takebe medium (Thomas and Rose, 1983). Whole 14-d-old Arabidopsis plants were incubated in petri dishes with wetted filter paper containing 2 \(\mu\)M LatB. After 8 h, plants were washed thoroughly with distilled water and transferred to petri dishes containing filter paper wetted with distilled water. Arabidopsis seedlings were stained for 30 min with 0.5% (w/v) FDA (Calbiochem-Novabiochem, San Diego) in petri dishes containing filter paper wetted by distilled water.

Microscopy and Cell Measurements
The morphology of epidermal cells from wild type, GFP-FABD2- and GFP-mTalin-expressing inflorescences, siliques, and rootste leaves was analyzed using cleared intact tissues as described previously (Fu et al., 2002). Cleared tissues and hypocotyls of 7-d-old plants were stained with toluidine blue, washed, and observed with a light microscope (Axioskop, Zeiss, Sydney) equipped with a CCD camera (Axiscam MR, Zeiss). Images were captured using Axiovision 4.0 (Zeiss) and cell plan-area, length, width, perimeter, and circularity measurements performed in the public domain NIH ImageJ. 1.31v program (developed at the United States National Institutes of Health and available on the Internet at http://rsb.info.nih.gov/ij).

Confocal Imaging and Analysis
Arabidopsis seedlings, N. tabacum, and L. esculentum leaf sections were mounted in water under a coverslip while N. tabacum protoplasts, BY-2, and M. truncatula cells were embedded in agar by mixing 50 \(\mu\)L of cells with 100 \(\mu\)L of 0.5% (w/v) low-melt agarose (type VII; Sigma-Aldrich) in wellled slides before mounting under a coverslip. Images of GFP-FABD2- and GFP-mTalin-expressing cells or FDA-stained cells were acquired as z-series with a 1- 

Immunoblotting
Protein was extracted from whole seedlings into 2X SDS sample buffer and equal total protein loaded and run on a 10% SDS-polyacrylamide gel before transfer to nitrocellulose overnight. Membranes were blocked for 1 h in 5% (w/v) skim milk powder in phosphate-buffered saline before incubation with a 1:500 dilution of anti-GFP (raised against native GFP; Peter Lewis, personal communication) or 1:500 anti-a-tubulin (Sigma; clone B-5-1-2). Washed membranes were then incubated with secondary antibody conjugated to alkaline phosphatase. Bands developed with Western Blue substrate (Promega, Madison, WI; no. S3848) were scanned and the integrated intensities from all bands quantified in ImageJ.

Upon request, all novel materials described in this publication will be made available in a timely manner for noncommercial research purposes, subject to the requisite permission from any third-party owners of all or parts of the material. Obtaining any permission will be the responsibility of the requestor.

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