
The Phragmoplast-Orienting Kinesin-12 Class Proteins Translate the Positional Information of the Preprophase Band to Establish the Cortical Division Zone in Arabidopsis thaliana

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The preprophase band (PPB) is a faithful but transient predictor of the division plane in somatic cell divisions. Throughout mitosis the PPBs positional information is preserved by factors that continuously mark the division plane at the cell cortex, the cortical division zone, by their distinct spatio-temporal localization patterns. However, the mechanism maintaining these identity factors at the plasma membrane after PPB disassembly remains obscure. The pair of kinesin-12 class proteins PHRAGMOPLAST ORIENTING KINESIN1 (POK1) and POK2 are key players in division plane maintenance. Here, we show that POK1 is continuously present at the cell cortex, providing a spatial reference for the site formerly occupied by the PPB. Fluorescence recovery after photobleaching analysis combined with microtubule destabilization revealed dynamic microtubule-dependent recruitment of POK1 to the PPB during prophase, while POK1 retention at the cortical division zone in the absence of cortical microtubules appeared static. POK function is strictly required to maintain the division plane identity factor TANGLED (TAN) after PPB disassembly, although POK1 and TAN recruitment to the PPB occur independently during prophase. Together, our data suggest that POKs represent fundamental early anchoring components of the cortical division zone, translating and preserving the positional information of the PPB by maintaining downstream identity markers.

INTRODUCTION

The plant microtubule (MT) cytoskeleton demonstrates remarkable plasticity in its ability to form complex arrays characteristic of distinct cell cycle phases and required for specific cellular functions. In expanding interphase cells, parallel aligned cortical MTs guide cellulose synthase complexes along transverse trajectories and permit cell expansion in a perpendicular direction (Paredez et al., 2006). Furthermore, vascular plants developed a specialized cytoskeletal array aiding the correct positional execution of cytokinesis. Entry into mitosis is characterized by the prophase band (PPB) (Pickett-Heaps and Northcote, 1986), an equatorial assembly of actin filaments and cortical MTs shaped by differential modulation of MT dynamic instability and selective MT stabilization in distinct regions of the cell cortex (Dhonukshe and Gadella, 2003; Vos et al., 2004). PPB formation requires the enzymatic action of a PP2A phosphatase holoenzyme complex containing FASS as a regulatory B subunit, complexed with the helper/assembly proteins TON1α and β, which is targeted to the MTs via the Tonneau Recruitment Motif proteins (Spinner et al., 2013). However, the PPB persists only transiently until pro-metaphase when its disassembly fuels the polymerization of spindle MT. After chromosome segregation and the subsequent condensation of daughter nuclei, the cytokinetic phragmoplast, a dual array of parallel-oriented MTs connecting at the cell division plane, evolves from the anaphase spindle remnants and assists in the synthesis of the cell plate, a de novo established membrane compartment generated by transport and fusion of endomembrane derived vesicles to the division plane, to bisect the daughter cells. The cell plate grows centrifugally and physically separates the daughter nuclei upon fusion with the plasma membrane at the end of plant cytokinesis. Intriguingly, the PPB anticipates the site of fusion between the cell plate and the parental cell wall despite the considerable time lag between the PPB’s disassembly at pro-metaphase and the end of cytokinesis when the position of the PPB is converted into the actual cell plate position (Gunning and Wick, 1985).

The correlation between PPB position and the site of cell plate fusion led to the hypothesis that the PPB determines the plane of cell division and recruits molecules that serve as positional information to establish and maintain the cortical division zone...
(CDZ) identity throughout mitosis (reviewed in Rasmussen et al., 2013). However, the first proteins described to serve as a reference beyond PPB disassembly were negative markers such as F-actin (Mineyuki and Palevitz, 1990; Cleary et al., 1992) and later the Arabidopsis thaliana kinesin-14 KCA1, both characterized by their low protein abundance or deficiency at the CDZ (Sano et al., 2005; Vanstraalen et al., 2006). In Arabidopsis, the microtubule binding protein TANGLER (TAN) and the Ran GTPase regulatory protein RanGAP1 both colocalize with the PPB and remain at the CDZ throughout mitosis (Walker et al., 2007; Xu et al., 2008). During the course of mitosis, the CDZ seems to alter from an initially wide zone (the CDZ) to a narrow site (cortical division site [CDS]) at the end of cytokinesis (Van Damme, 2009). The initial recruitment of both TAN and RanGAP1 to the cell cortex requires PPB formation (Walker et al., 2007; Xu et al., 2008), whereas TAN residency at the CDZ/CDS post PPB disassembly does not require MTs (Walker et al., 2007). In Arabidopsis, both TAN and RanGAP1 maintenance at the CDZ is abolished in a mutant background lacking a pair of kinesin-12 proteins, PHRAGMOPLAST ORIENTING KINESIN1 (POK1) and POK2 (Walker et al., 2007; Xu et al., 2008). Arabidopsis pok1 pok2 double mutants display severe cell wall positioning defects, reminiscent of the maize (Zea mays) tan1 phenotype (Smith et al., 1996). The majority of phragmoplasts in the pok1 pok2 double mutant do not correspond to a transverse orientation relative to the cell longitudinal axis, while PPBs were predominantly transverse with respect to the long axis of the cell supporting a role of POKs in the spatial control of cytokinesis (Müller et al., 2006). TAN and RanGAP1 have been shown to interact with the C-terminal domain of POK1 (Müller et al., 2006; Xu et al., 2008), suggesting that POK1, TAN, and RanGAP1 are part of a functional module required for CDZ identity maintenance. However, the exact nature of this module remains unclear due to the lack of positional information on the POK kinesins.

Here, we provide novel insight into POK function by characterizing a novel pok2 allele from a mutant screen for pok1-1 enhancers. In vivo observations of the novel allele combination pok1-1 pok2-3 revealed lengthy cytokinesis due to a reduced rate of phragmoplast expansion and frequent tilting of late phragmoplasts from their initial orientation. Nevertheless, fusion of the cell plate with the parental wall occurred at seemingly indiscriminate sites resulting in a highly disorganized framework of cell walls, underpinning POKs role in an efficient phragmoplast guidance mechanism and the timely progression through cytokinesis.

Localization studies using a functional yellow fluorescent protein (YFP)-POK1 fusion protein showed that POK1 is recruited to the PPB and remains at the cortex throughout mitosis. Similar to TAN, POK1 initially marks the CDZ and its localization domain narrows down to the CDS prior to cell plate fusion with the parental plasma membrane. POK1 is progressively and dynamically loaded onto the PPB from the cytosol in prophase, while its association with the plasma membrane at the CDZ following PPB degradation appears static. Throughout telophase, the ring-shaped localization narrows from an initial broad band to a sharp string-like pattern that disappears shortly after cell plate fusion. In pok1 pok2 mutants, the cortical division zone marker TAN disappeared from the CDZ upon entry into metaphase, emphasizing that the POK kinesins function as molecular anchors for other CDZ/CDS identity markers.

RESULTS

POK1 and POK2 Are Required for Guidance of the Phragmoplast toward the Cortical Division Site

Previously, we described two independent pok1 pok2 double mutant allele combinations, with similar phenotypic defects differing predominantly in their growth rate (Müller et al., 2006). In a sensitized ethyl methanesulfonate (EMS)-induced mutant screen to identify second site mutations in phenotypically wild-type looking pok1-1 single mutants, we discovered a severe mutant reminiscent of previously characterized pok1 pok2 double mutants. The novel mutant cosegregated with the pok1-1 genotype (Supplemental Figures 1A and 1B) and did not complement the pok1 pok2 double mutant phenotype in test crosses, suggesting that the mutation was located in POK2. Sequencing of POK2 revealed a C-to-T substitution in exon 16, 2197 bp of the coding sequence downstream of the ATG (Figure 1A). The mutation resulted in a premature STOP codon predicted to obliterate about three quarters of the POK2 protein. The allele was designated pok2-3. Due to the close linkage of POK1 and POK2 on chromosome III, pok2-3 single mutants were not analyzed.

Overall, the pok1-1 pok2-3 double mutant displayed an aggravated phenotype compared with the previously described allele combinations pok1-1 pok2-1 and pok1-2 pok2-2 (Supplemental Figure 1A; Müller et al., 2006). We reasoned that POK activity was further diminished in these plants and, thus, the novel mutant was used to expand on the in vivo characterization of POK function. The pok1-1 pok2-3 seedlings displayed shorter and wider roots and hypocotyls and succulent leaves (Figures 1B and 1D). Mature plants were dwarfed similar to weak ton2 alleles (Kirik et al., 2012; Spinner and Gadeyne et al., 2013), but in contrast to ton2, pok1-1 pok2-3 mutants were fertile (Supplemental Figure 1A). Double pok1-1 pok2-3 mutants grew significantly shorter roots (Figure 1C) and displayed smaller leaves than pok1-1 pok2-1 or pok1-1 (Figure 1D). Although organ formation apparently was not affected, at the cellular level, cell division patterns were dramatically disturbed in pok1-1 pok2-3 double mutants (Figure 1E, right panel), compared with wild-type and single mutant pok1-1 root meristems (Figure 1E, left panel).

Previously, immunolocalization of tubulin in root squashes established that phragmoplasts in pok1 pok2 did not expand in the direction of the former PPB (Müller et al., 2006). To investigate phragmoplast guidance in vivo, the MT reporter line green fluorescent protein (GFP)-MBD was introgressed into the pok1-1 pok2-3 double mutants. As in pok1-1 pok2-1 and pok1-2 pok2-2 (Müller et al., 2006), PPB, spindles, and phragmoplasts formed normally in the pok1-1 pok2-3 double mutant root meristem cells (Supplemental Figure 2). Time-lapse recordings of phragmoplast expansion, encompassing the stage of spindle-to-phragmoplast transition until completion of phragmoplast disassembly, revealed that phragmoplasts frequently tilted (Figures 2A, 2B, and 2D; Supplemental Movies 1 and 2 and Supplemental Figures 3A and 3B) and that the duration of cytokinesis was prolonged in pok1-1 pok2-3 double mutants (Figure 2C). While in the wild type cytokinesis lasted 6 min (n = 17) on average, in the double mutant, cytokinesis lasted on average 24 min (n = 24). The phenotype in test crosses, suggesting that the mutation was located in POK2. Sequencing of POK2 revealed a C-to-T substitution in exon 16, 2197 bp of the coding sequence downstream of the ATG (Figure 1A). The mutation resulted in a premature STOP codon predicted to obliterate about three quarters of the POK2 protein. The allele was designated pok2-3. Due to the close linkage of POK1 and POK2 on chromosome III, pok2-3 single mutants were not analyzed.

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antiparallel MTs connecting in the midzone (Figure 2A), which expands laterally throughout cytokinesis. In the wild type, periclinal root divisions, allowing a view of the phragmoplast torus/ring while it expands, occur with a very low frequency close to the root apical meristem as the Arabidopsis root contains a single layer of epidermal, cortical, and endodermal tissue. Remarkably, 23% of cytokinetic cells in pok1-1 pok2-3 double mutants displayed the phragmoplast ring view (Figure 2D), while only 2.4% of cytokinetic cells in the wild type exhibited the phragmoplast ring, in agreement with altered division plane orientations in pok1-1 pok2-3.

To determine whether the rate of phragmoplast expansion contributed to the prolonged cytokinesis in the pok1-1 pok2-3 double mutant, we compared the rate of phragmoplast expansion using kymograph analysis (Figures 2E and 2F). Regardless of the genotype, most phragmoplasts displayed an expansion asymmetry as reflected by varying inclines of the kymograph (Figure 2F, left panel). This demonstrated that lateral expansion of the phragmoplast did not occur at a steady velocity, also indicated by the high SD (Figure 2G). Nevertheless, in pok1-1 pok2-3 double mutants, the average expansion rate was moderately, but significantly reduced (0.21 ± 0.09 μm/min; *P < 0.02, n = 43) (Figure 2G), suggesting that slowing of the expansion rate in the double mutant contributes to the observed prolonged cytokinesis together with the frequent tilting of the phragmoplast.

POK1 Interacts with TAN through Its C-Terminal Domain and Is Required for TAN Maintenance at the CDZ but Not for Its Recruitment

As previously reported, TAN maintenance at the CDZ requires POK function, while recruitment to the PPB occurred, although inefficient, in pok1 pok2 mutants (Walker et al., 2007), in agreement with two distinct TAN protein domains mediating TAN recruitment to the CDZ before and after PPB disassembly (Rasmussen et al., 2011). To clarify the role of POKs in TAN recruitment, we analyzed TAN localization in the severe pok1-1 pok2-3 allele combination (Figure 3). In the wild type, the majority of cells displaying either the PPB (95%, n = 19 cells), pro-spindle (100%, n = 6 cells), spindle (92%, n = 12 cells), or phragmoplast (98%, n = 43 cells) also contained associated cortical TAN rings (Figures 3A to 3D). In contrast, the TAN recruitment in cells with PPs (55%, n = 11 cells) seemed less efficient or delayed, but not abolished in pok1-1 pok2-3 double mutants, consistent with results from previous studies. Interestingly, the cortical association of TAN in cells containing pro-spindles was high (89%, n = 9 cells; Figure 3E), suggesting that POK function was not critically required for TAN localization at prophase. However, the percentage of TAN rings in cells displaying spindles in metaphase dropped to 19% (n = 16 cells; Figures 3F, 3G, and 3I), indicating that binding of POK1 and TAN becomes essential during this cell cycle stage. Moreover, TAN rings were entirely absent from cells with phragmoplasts (0%, n = 18 cells; Figure 3H) in pok1 pok2 double mutants. This result further supports the partial POK dependence of TAN at the PPB and the essential POK1-dependent maintenance of TAN following prophase. In addition, we showed in planta interaction of nYFP-POK1 and cYFP-TAN in Arabidopsis protoplasts using bimolecular fluorescent complementation (BiFC) (Figures 3J and 3K), supporting that interaction between TAN and POK1 mediates TAN maintenance at the CDZ. In contrast to control experiments (Figures 3L to 3N), strong expressing protoplasts displayed a filamentous YFP pattern reminiscent of cortical MTs (Figure 3O), suggesting that stabilization of POK1 and TAN interaction increases their capacity for MT binding.

POK1 Localizes to the Cortical Division Zone

We analyzed the localization pattern of a functional YFP-POK1 fusion (Supplemental Figures 1C and 1D) in 4- to 6-d-old wild-type

**Figure 1.** Mutant Alleles and Mutant Phenotypes.

(A) Schematic representation of POK1 and POK2 gene structure and position of mutations. SALK-T-DNA insertions in alleles pok1-1 (exon 15), pok1-2 (exon 7), pok2-1 (exon 28), and pok2-2 (exon 23) are indicated by triangles. The ethyl methanesulfonate-induced nonsense mutation in allele pok2-3 is a C-to-T substitution (exon 16, at position 3845 bp), causing a premature stop at amino acid 733 (Gln to *).

(B) Seedling phenotypes of (right to left) the pok1-1 single mutant, pok1-1 pok2-1 double mutant, and pok1-1 pok2-3 double mutant. The ethyl methanesulfonate-induced nonsense mutation in allele pok2-3 is a C-to-T substitution (exon 16, at position 3845 bp), causing a premature stop at amino acid 733 (Gln to *).

(C) Comparison of average primary root length 10 d after germination (cm). Gray bar indicates the wild type (3.4 cm ± 0.8), dotted bar indicates pok1-1 (3.4 cm ± 0.8), black bar indicates pok1-1 pok2-2 (0.6 cm ± 0.2), and gray bar indicates pok1-1 pok2-3 (0.25 cm ± 0.09). The average root length of pok1-1 pok2-1 and pok1-1 pok2-3 is significantly (**P < 0.005, Student’s t test) shorter than in the wild type and pok1-1. Number of samples is indicated. Error bars indicate ±sd.

(D) Comparison of rosettes of 3-week-old pok1-1 single mutant, pok1-1 pok2-3 double mutant, and pok1-1 pok2-3 double mutant plants.

(E) Root meristems, stained with propidium iodide to visualize cell walls. Whereas the pok1-1 single mutant (left) shows a regular cell pattern of the root cell files, the pok1-1 pok2-3 double mutant (right) cell pattern is completely disorganized.

Bar = 0.5 cm in (B) and 20 μm in (E).
seedlings (Figure 4; Supplemental Figure 3). Only a subset of meristematic cells displayed a fluorescent YFP-POK1 signal, suggesting cell cycle–controlled expression or degradation of POK1. POK1-expressing cells displayed distinct fluorescent patterns of broad bands, sharp bands, or pairs of focused spots (Figure 4A) in single optical sections. Three-dimensional reconstruction of confocal image stacks revealed that YFP-POK1 formed continuous rings at the cell cortex.

To correlate the distinct YFP-POK1 rings with the corresponding cell cycle stages, we coexpressed YFP-POK1 and the MT reporter red fluorescent protein (RFP)-MBD. In root and leaf meristems, YFP-POK1 rings were associated with mitotic cells exhibiting PPBs, spindles, and phragmoplasts (Figure 4B; Supplemental Figures 4A and 4B). The observed YFP-POK1 patterns were reminiscent of the ring-shaped localization observed for the two POK1 interactors TAN and RanGAP1, which both colocalize with the PPB and continuously mark the CDZ throughout mitosis (Walker et al., 2007; Xu et al., 2008).

The short C-terminal fragment POK11683-2066 shown to be sufficient to bind TAN (Müller et al., 2006; this study) and Ran-GAP1 (Xu et al., 2008) fused to YFP (Pro35S:YFP-POK11683-2066; Figure 4C) accumulated mainly in aggregates in Arabidopsis root meristem cells (Supplemental Figure 4C), constraining detection of a potential cortex localization. Since POK11683-2066 showed substantial aggregation, we tested whether aggregation could be prevented by expressing longer fluorophore-tagged constructs of the POK1 C-term (Pro35S:GFP-POK11213-2066 and ProTP1:dTom-POK11265-2066; Supplemental Figure 4D). Both fusion proteins displayed a cell cycle–dependent subcellular localization pattern similar to the full-length YFP-POK1 and vastly diminished aggregation (Supplemental Figures 4E and 4F).

POK1 Cortical Recruitment Shows Cell Phase–Dependent Dynamicity

To further characterize POK1 cortical recruitment, fluorescence recovery after photobleaching (FRAP) analysis was performed.

(D) In the wild type (n = 49), 97.96% of the observed phragmoplasts displayed anticalinal orientation and only 2.04% were oriented with their torus parallel to the optical plane. In contrast, in pok1-1 pok2-3 mutants (n = 43), the proportion of phragmoplasts presenting the torus (ring view) rises to 23.26%.

(E) Projection of a time series throughout mitosis for wild type (45 min, 19 image stacks) and pok1-1 pok2-3 mutants (60 min, 25 images). At 2.5-min intervals, image stacks were taken from mitotic cells at 1-μm z-intervals throughout the duration of mitosis. Maximum z-projections of each time point were projected for the depicted images. The white dotted lines indicate the selection for the kymograph analysis.

(F) Space-time plots (kymographs) of respective phragmoplasts depicted in (E). The x-coordinate depicts the distance in micrometers. The y-coordinate indicates the time t. The kymograph plug-in in ImageJ was used to create kymographs and calculate the velocity of phragmoplast expansion. The slopes of the kymograph (contrast edge, selection is indicated by white dotted lines) are proportional to the velocity. In these examples, the two halves of the phragmoplasts expanded at different velocities. In addition, in one half the velocity decreased, indicated by the change in slope (two-phase expansion).

(G) The average velocity of phragmoplast expansion is significantly reduced in pok1-1 pok2-3 mutants (0.18 ± 0.09 μm/min; P < 0.02, n = 22, Student’s t test), compared with the expansion rate in the wild type (0.21 ± 0.09 μm/min; n = 43). Error bars indicate ±SD. Cell walls are stained with propidium iodide. Microtubules are visualized by GFP-MBD.
Figure 3. Loss of POK-Dependent TAN Localization at the Cortical Division Zone Correlates with PPB Disassembly.

(A) to (D) TAN-YFP is recruited to the preprophase band in the wild type (A) and pok1-1 pok2-3 double mutants (E). In the wild type, TAN-YFP rings coexist with the preprophase band (A) and (B), pro-spindle (B), spindle (B), and early late phragmoplasts (C) and (D).

(E) Cell with pro-spindle displaying the TAN-YFP ring.

(F) The spindle in the metaphase cell is not associated with cortical TAN-YFP nor does the signal reappear during cytokinesis (F) to (H) represent a time series. Images are z-projections of a varying number of images taken at 1-μm intervals. TAN-YFP rings are indicated by arrowheads. The nuclear localization of YFP-TAN is indicated by an asterisk. Bar = 10 μm.

(I) Presence of TAN-YFP rings during the cell cycle in wild type and pok1-1 pok2-3 mutants.

(J) to (K) Interaction of TAN and the POK1 C terminus, as determined by BiFC.
In the Arabidopsis root meristem, epidermis cells in pro- and metaphase were selected based on the presence or absence of condensed nuclei in bright field. In transverse sections, YFP-POK1 localized to both lateral cell sides (Figures 5A to 5D). YFP-POK1 was specifically bleached at one side of the confocal section and migration of fluorescent signal into the bleached region was subsequently monitored. In prophase cells, fluorescence clearly recovered during the observation period (Figures 5A, 5C, and 5E). By contrast, metaphase YFP-POK1 fluorescence hardly recovered (Figures 5B, 5D, and 5G).

Figure 4. POK1 Localizes at the Cortical Division Zone/Site.

(A) Maximum z-projection of a root meristem expressing YFP-POK1. In cross sections, YFP-POK1 localizes as broad bands or dots (arrowheads) and sharp bands (arrows). Occasionally, YFP-POK1 rings are visible (arrow with dotted shaft). Some cells accumulate cytoplasmic YFP-POK1 (asterisk).

(B) Coexpression of RFP-MBD and YFP-POK1 in different cell cycle stages in the root. YFP-POK1 colocalizes with the PPB in prophase and remains at the cell cortex throughout mitosis. Large yellow arrows indicate the longitudinal root axis.

(C) POK1 domain organization as indicated in the legend and overview of fluorescent protein fusions used for localization studies. The POK11683-2066 fragment corresponds to the TAN-interacting fragment used in the BiFC experiment (Figures 3J and 3K).

Bar = 20 μm in (A) and 10 μm in (B).

(Figures 5 and 6). In the Arabidopsis root meristem, epidermis cells in pro- and metaphase were selected based on the presence or absence of condensed nuclei in bright field. In transverse sections, YFP-POK1 localized to both lateral cell sides (Figures 5A to 5D). YFP-POK1 was specifically bleached at one side of the confocal section and migration of fluorescent signal into the bleached region was subsequently monitored. In prophase cells, fluorescence clearly recovered during the observation period (Figures 5A, 5C, and 5E). By contrast, metaphase YFP-POK1 fluorescence hardly recovered (Figures 5B, 5D, and 5G).
indicating a more dynamic YFP-POK1 recruitment or dispersal mechanism during prophase and a fairly static maintenance mechanism during subsequent cell cycle phases. The POK1 C-terminal fragment GFP-POK11213-2066 in BY-2 cells displayed differential dynamics in FRAP experiments similar to YFP-POK1 in Arabidopsis (Figures 6A to 6D). Also in prophase BY-2 cells, fluorescent signal recovered, although at a longer half-time (Figure 6D) than YFP-POK1 (Figure 5E), while BY-2 metaphase cells displayed sluggish GFP-POK11213-2066 signal recovery similar to YFP-POK1. Due to the superior imaging accessibility, we clarified whether the initial recruitment of POK1 to the CDZ requires MTs, using BY-2 cells. Prophase cells expressing ProS35:POK11213-2066-GFP were treated with oryzalin and subjected to FRAP analysis. The oryzalin treatment completely abolished fluorescence recovery to the CDZ following photobleaching (n = 6) (Figures 6E to 6L), suggesting the MT-dependent recruitment of POK11213-2066-GFP.

In Arabidopsis, a proportion of mitotic cells accumulated full-length YFP-POK1 signal in the cytoplasm, both in the absence and presence of YFP-POK1 cortical rings (Figure 4A, asterisk). We determined the proportion of cells containing cytoplasmic YFP-POK1 and/or YFP-POK1 at the CDZ for each cell cycle stage (Supplemental Figure 4A). In the presence of PPBs, YFP-POK1 accumulates in the cytoplasm and progressively associates with the PPB and the CDZ in pro-metaphase, suggesting that YFP-POK1 is dynamically recruited to the CDZ. From metaphase until the termination of cytokinesis, the majority of YFP-POK1 was found to be associated with the CDZ. Taken together, our data suggest a cell cycle-dependent relocalization from cytoplasmic to cortical, CDZ/CDS resident YFP-POK1.

Similar to TAN (Walker et al., 2007; Rasmussen et al., 2011; Supplemental Figures 4B and 4C) and RanGAP1 (Xu et al., 2008), the ring-shaped localization of POK1 narrowed during cytokinesis in BY-2 cells (Figures 6B and 6C; Supplemental Figures 5D and 5E) and Arabidopsis (Figures 5A and 5B) in a cell cycle progression-dependent manner. While the sharp punctate signal corresponds to the site of cell plate fusion, the CDS, the broad POK1 signal corresponded to the wider CDZ.

YFP-POK1 CDZ/CDS Localization Is Independent of Microtubules

To analyze the microtubule dependency of the subcellular YFP-POK1 localization, seedlings coexpressing YFP-POK1 and GFP-MBD were treated with the MT polymerization inhibitor oryzalin. The effect of oryzalin was visualized by the depolymerization of MT-dependent recruitment of POK11213-2066-GFP. ProS35:POK11213-2066-GFP were treated with oryzalin and subjected to FRAP analysis. The oryzalin treatment completely abolished fluorescence recovery to the CDZ following photobleaching (n = 6) (Figures 6E to 6L), suggesting the MT-dependent recruitment of POK11213-2066-GFP.

Protein interactions with other proteins are also important for cytokinesis. The proteins TAN and the POK kinesin-12 proteins were implicated in phragmoplast guidance due to the mispositioning of cell walls in the corresponding mutants with respect to the former PPB position (Smith et al., 1996; Müller et al., 2006), but how these proteins functionally interacted was unclear due to the lack of subcellular data regarding the POK kinesins. Our in vivo analysis confirmed the frequent mismatch of PPB and phragmoplast orientation in dividing pok1-1 pok2-3 cells, showing that the POK kinesins function downstream of PPB formation and division plane determination. In contrast, in mutant alleles of microtubule-associated proteins such as CLASP (Ambrose et al., 2007; Kirk et al., 2007), PP2A subunit mutants (Spinner and Gadeyne et al., 2013) and the putative membrane protein SABRE (Pietra et al., 2013), oblique cell walls and misguidance of the phragmoplast are a consequence of mispositioning of the PPB or lack thereof and, thus, these proteins act upstream of the POKs.

Consistent with phragmoplast expansion in BY-2 cells (Buschmann et al., 2010), we determined distinct expansion velocities for the left and right edge of wild-type and pok1-1 pok2-3 phragmoplasts. The rate of phragmoplast expansion was diminished in pok1-1 pok2-3 cytokinesis, but could not solely account for the difference with respect to wild-type cytokinesis duration. Tilting of the pok1-1 pok2-3 phragmoplast contributes to the extended duration of pok1-1 pok2-3 cytokinesis as well, since the tilted cell plate apparently does not follow the shortest distance rule (Rasmussen et al., 2013) in the
Figure 5. Differential POK1 Dynamicity during Mitosis.

(A) and (B) FRAP analysis of YFP-POK1. Representative time lapse of mitotic cells in the Arabidopsis root meristem. Transverse sections of (A) prophase and (B) metaphase cells recorded at different time points before (prebleach) and after photobleaching. Bleach regions are indicated by a dashed square (ROI) in first postbleach images (t = 0 s). Bar = 5 \( \mu m \).

(C) and (D) Kymographs of prebleach and postbleach time series corresponding to the dashed line selection as indicated in the prebleach images of (A) and (B). Prebleach kymograph (orange dashed line) of four prebleach images and postbleach kymograph of 30 postbleach images (magenta solid line) and corresponding profile plots are depicted. Profile plots show mean fluorescence intensities from the prebleach (orange dashed line) and postbleach (magenta solid line) kymographs, respectively. Note the recovery of YFP signal in prophase.

(E) An average of independent FRAP experiments is plotted to fit a single exponential curve with rise to a maximum \( y = a(1-\exp(-b\cdot x)) \). In prophase cells \( n = 10 \), the YFP-POK1 signal recovers after photobleaching with a half-time \( t_{1/2} \) of 154 s, while there is hardly any fluorescence recovery in metaphase cells \( n = 6 \). Error bars indicate ±SD.

(F) to (I) POK1 localization at the cortical division zone is independent of MTs. Maximum z-projections of an Arabidopsis root meristem prophase cell coexpressing the MT reporter RFP-MBD (F) and YFP-POK1 (G) before and after treatment with 10 \( \mu M \) oryzalin. The PPB is indicated by asterisks.

(F) and (G) Lower panels were recorded 15 min after oryzalin incubation and correspond to the cell in the upper panels. Due to MT depolymerization, the RFP-MBD reporter localization becomes cytosolic, while the YFP-POK1 signal remains present at the CDZ. Bar = 10 \( \mu m \).

(H) and (I) Fluorescence intensity profile plots from a rectangular selection in (F) and (G) depicting RFP-MBD (H) signal distribution before oryzalin treatment (continuous, magenta line) and after oryzalin treatment (dashed, black line) and YFP-POK1 (I) signal distribution before (continuous, green line) and after (dashed, black line) oryzalin treatment. Note that the RFP-MBD peaks disappear after oryzalin treatment, while the YFP-POK1 peaks remain.

[See online article for color version of this figure.]
Figure 6. Changes of POK1 Dynamicity during Mitosis.

(A) A tobacco BY-2 cell expressing Pro35S:GFP-POK11213-2066 and Pro35S:POK11213-2066-GFP and the corresponding bright-field image. Bar = 10 μm.

(B) to (D) FRAP analysis of GFP-POK1C.

(B) Representative time lapse of a prophase BY-2 cell after photobleaching of GFP-POK1C at the PPB (white box indicates the bleached area).
absence of POK-dependent phragmoplast guidance. The postulated direct correlation of cytokinesis duration and cell size (Gorst et al., 1986) is consistent with this assumption.

The cell cycle–dependent association of POK1 with the CDZ/CDS closely resembles the localization of CDZ/CDS identity markers TAN and RanGAP1 (Walker et al., 2007; Xu et al., 2008). Interaction of POK11683-2066 with TAN as determined by BIFC (Figure 3) and the phenotype similarity of the maize mutant tan (Smith et al., 1996; Cleary and Smith, 1998) and the pok1 pok2 double mutant in Arabidopsis further support a joint function of POK1 and TAN in the phragmoplast guidance mechanism.

POK1, TAN (Walker et al., 2007), and RanGAP1 (Xu et al., 2008) share the feature of narrowing from a broad toward a sharp ring during cytokinesis, suggesting that the proteins are coregulated and/or act in a complex. POK1 seems to be particularly important for CDZ identity maintenance throughout mitosis, since both TAN (this study; Walker et al., 2007) and RanGAP1 (Xu et al., 2008) fail to remain present in the narrowing of POK1 at the CDZ. With respect to TAN and RanGAP1 maintenance, POK1 provides a scaffolding function to retain CDZ resident proteins, thereby sustaining CDZ identity. Although the C-terminal POK11683-2066 is sufficient for TAN interaction, GFP-POK11683-2066 localization at the CDZ could not be unambiguously confirmed, in contrast to GFP-POK11213-2066, indicating that the necessary additional domain for efficient POK1 targeting to the PPB and CDZ lies directly upstream of POK11683-2066. With few exceptions, most kinesin class proteins act as homo- or heterodimers (Endow et al., 2010), mediated by multiple coiled coil domains, which are also predicted for POK1 and POK2 C termini (Figure 4C). Thus, the POK1C-terminal fusion proteins might associate with the PPB/CDZ through self-dimerization or with endogenous full-length POKs. Alternatively, the POK1C-terminal might be necessary and sufficient to recruit POK1 to the PPB and CDZ via interactions with yet unknown membrane-anchored binding partners.

Although the localization of POK2 was not directly investigated here, the rescue ability of the full-length YFP-POK1 fusion protein and the absence of aberrant single mutant phenotypes indicate functional redundancy of POK1 and POK2 and therefore also similarity in localization. In light of functional redundancy of POK1 and POK2, the aggravated phenotype of the novel pok1-1 pok2-3 allele combination can be explained by the fact that some POK2 activity might still be present in previously described double mutants.

Genetic evidence suggests that TAN arrives at the PPB independently of POK1; however, both POK1 and TAN1 (Walker et al., 2007) recruitment to the PPB require an intact microtubule cytoskeleton. To clarify the sequence of independent recruitment events at the PPB, time-resolved in vivo colocalization studies of POK1, TAN, and RanGAP1 proteins would prove useful, since genetic analysis in Arabidopsis is hampered by the lack of clear tan loss-of-function alleles as well as the pleotropic phenotypes of ran gap double knockdown mutants (Xu et al., 2008) and gametophytic lethality of ran gap double knockout mutants (Rodrigo-Peiris et al., 2011), respectively.

Based on homology of the motor domains, POK1 clusters with kinesin-12 class proteins, predicted to display MT plus end directed motility (Lee and Liu, 2004; Müller et al., 2006; Lipka and Müller, 2012). Although POK1 motor activity has not been investigated, given its localization at the plasma membrane, it potentially contributes to the phragmoplast guidance mechanism by directly binding to peripheral phragmoplast MTs and

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<th>Table 1. Association of YFP-POK1 with the Cortical Division Site Is Independent of Microtubules</th>
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While microtubules disassembled, YFP-POK1 signal was not affected upon oryzalin treatment.

Figure 6. (continued).

(C) Representative time lapse of a metaphase BY-2 cell after photobleaching the GFP-POK1 C terminus at the CDZ (indicated by white box).

(D) Graph representing FRAP data for prophase and metaphase BY-2 cells overexpressing GFP-POK11213-2066. In prophase cells, the GFP-POK1 signal recovers slowly after photobleaching with a half-time (t1/2) of 362 s, while there is hardly any fluorescence recovery in metaphase cells. An average of independent FRAP experiments is plotted to fit a single exponential curve with rise to a maximum (y = a*(1-exp(-b*x))) (prophase cells: n = 4; metaphase cells: n = 7). Error bars indicate ±SD.

(E) to (L) Recruitment of POK1C-GFP requires MTs, whereas PPB-localized POK1C-GFP statically associates with the CDZ independently of MTs in BY-2 cells.

(F) to (G) POK1C-GFP colocalizes with the preprophase band (E), POK1C-GFP; (F), RFP-MBD; (G), merged image.

(H) to (J) Upon oryzalin treatment, MTs depolymerize while POK1C-GFP remains associated with the CDZ (H), POK1C-GFP; (I), RFP-MBD; (J), merged image.

(K) Representative time lapse of POK1C-GFP FRAP performed on oryzalin-treated BY-2 cells in (E). In the absence of MTs, POK1C-GFP fluorescence did not recover at the bleached CDZ region. The bleached area is indicated by a white box.

(L) Graph representing the FRAP experiments performed on oryzalin-treated prophase BY-2 cells coexpressing the Pro35S:RFP-MBD microtubule marker and Pro35S:POK11213-2066-GFP (n = 6). GFP fluorescence did not recover at the CDZ upon MT depolymerization. Bar = 10 μm; error bars indicate ±SD.

[See online article for color version of this figure.]
exerting a motor activity–related force. Alternatively, POK, together with TAN, might simply function to stabilize outreaching MTs. Indeed, TAN is a highly basic protein that cosediments with MTs in vitro (Smith et al., 2001), showing that it has MT binding capacities. In this hypothesis, the motor activity required for orienting the phragmoplast might be delivered by phragmoplast-associated kinesins. Stabilization of the POK1-TAN protein complex by reconstitution of YFP in the BiFC experiment drew the complex to filamentous cellular structures, closely resembling MTs, thus likely unmasking a normally transient interaction of the POK1-TAN protein complex with MTs (Figure 3). This observation raises the exciting possibility that the CDZ resident POK1-TAN complex might indeed stabilize transient interactions with phragmoplast emanating MTs that approach the CDS to direct the phragmoplast/cell plate to its final destination.

In summary, we provide evidence for POK’s critical function for the maintenance of CDZ/CDS identity. In a MT-dependent
manner, dynamic POK1 is continuously recruited to the CDZ during prophase, most likely from a cytoplasmic pool that is depleted prior to metaphase. Upon its arrival at the CDZ, POK1 becomes immobilized and subsequently its localization narrows by a yet unknown mechanism as mitosis progresses (Figure 7). Independent from POK function, TAN and RanGAP1 arrive at the CDZ, where they become tethered by a POK-dependent mechanism possibly involving TON2-mediated dephosphorylation of TAN (Figure 7).

Together, these proteins provide positional information of the division plane and might even actively participate in the phragmoplast guidance mechanism. However, at this point it still remains to be elucidated whether POKs are actively engaged in the phragmoplast guidance mechanism or whether their function is primarily a scaffolding activity to provide a spatial reference for CDZ components following PPB disassembly. Future studies are geared toward resolving these open questions.

METHODS

Plant Material

Arabidopsis thaliana plants, accession Columbia, were used throughout the study unless otherwise indicated. SALK T-DNA insertion lines for pok1-1, pok1-2, pok2-1, and pok2-2, as well as the double mutants pok1-1 pok2-1 and pok1-2 pok2-2, have been described previously (Müller et al., 2006). Agrobacterium tumefaciens-mediated transformation to create diverse transgene lines was performed according to Clough and Bent (1998).

Growth Conditions

For examination of mutant phenotypes and localization studies at the seedling stage, seedlings were grown on plates containing Murashige and Skoog basal medium (Sigma-Aldrich) and 1% agarose or agar. For reproduction and crossing, plants were grown in soil. Plants and pots were incubated at 20 to 22°C on a 16-h-light/8-h-dark cycle.

Ethyl Methanesulfonate Mutagenesis and Identification of pok2-3

Plants homozygous for the pok1-1 mutation where imbibed in deionized water, mutagenized with 0.3% ethyl methanesulfonate overnight, and thoroughly washed subsequently. Five-hundred M1 progeny were scored for short roots on standard Murashige and Skoog medium plates, and seeds were collected individually for each M1 plant. Among eight pre-selected M2 families, one segregated the pok1-1 pok2-3 phenotype. We performed test crosses with pok1-1 pok2-1 and since the progeny (n = 120) showed the pok1 pok2 phenotype, we concluded that POK2 was mutated. Sequencing of the POK2 locus identified a C-to-T substitution in exon 16 leading to a premature stop codon.

Microtubule Marker Line

The MT binding domain vectors contained in a Gateway (Invitrogen) compatible destination vector pEG104-POK1C1213-2066 (Gutierrez et al., 2009) was recombined into pDONR221 via a Gateway BP reaction (Invitrogen). POK1C1213-2066 coding sequences with and without stop codon were amplified from entry clone pENTR-POK1 described by Xu et al. (2008).

The dTom-POK11265-2066 fusion construct was created in the pK7m34GW vector (Karimi et al., 2007) using the dTomato coding sequence and the POK cDNA fragment corresponding to amino acids 1265 to 2066 (Supplemental Table 1) encoding 50 nucleotides of POK1 homology arms as described (Tursun et al., 2009). In brief, recombineering steps were essentially performed using the recombination marker. The binary BAC 79I20 containing the genomic region of POK1 was transformed into Escherichia coli strain SW105, which allowed homologous recombination upon heat induction. Electro-competent SW105 containing BAC 79I20 were incubated at 42°C for 15 min to induce λ red recombinase and subsequently transformed with the PCR product amplified from the YFP-GalK cassette. Then cells were propagated for 3 h at 32°C and plated on galactose minimal medium. Colonies were streaked on McConkey indicator plates to screen for galactokinase activity, which lead to changes in pH of the medium turning colonies pink. Positive clones were further analyzed by PCR for the integration of the cassette into the BAC. Selected clones were cultured in liquid medium and split after propagation. One aliquot was supplied with 1/100 volume of 10% arabinoose solution for induction of FLP recombinase to remove the GalK selection marker. Both aliquots were propagated for two additional hours before DNA was extracted. Induced and noninduced samples were analyzed by PCR with primers (Supplemental Table 1) flanking the recombination site. Clones where GalK was efficiently excised were sequenced and verified for transformation into Agrobacterium strain GV3101. After Arabidopsis transformation, five independent T1 lines were recovered on nutrient agar plates containing 0.05% phosphotyrosin, and T2 plants were inspected for YFP-POK1 localization. Two independent T2 lines showed YFP fluorescence and were selected for further analysis. Both lines rescued the pok1 pok2 double mutant phenotype and showed an identical localization pattern of the YFP-POK1 fusion protein in Columbia and pok1 pok2 rescue plants.

TAN with and without stop codon was amplified from full-length cDNA clones obtained from the RIKEN institute (pda14314) (primers are listed in Table 3) under control of the constitutive promoter from the At3g16640 gene (Berkowitz et al., 2008).

Generation of YFP-POK1 Fusion Protein

A recombineering-based approach was used to genetically insert the YFP tag at the 5’ of POK1. All recombineering steps were essentially performed as described (Tursun et al., 2009). In brief, recombineering primers (Supplemental Table 1) encoding 50 nucleotides of POK1 homology arms immediately upstream and downstream of the ATG and ~50 nucleotides of a plasmid pBALU6 (Tursun et al., 2009), containing a YFP and GalK cassette, were used to amplify the fluorophore and the GalK selection marker. The binary BAC 79I20 carrying the genomic region of POK1 was transformed into Escherichia coli strain SW105, which allowed homologous recombination upon heat induction. Electro-competent SW105 containing BAC 79I20 were incubated at 42°C for 15 min to induce λ red recombinase and subsequently transformed with the PCR product amplified from the YFP-GalK cassette.

Confocal Imaging and Image Processing

Imaging was performed on a Leica SP2 equipped with a point scanner and a Leica SP8 confocal microscope equipped with a resonant scanner.
A 63× numerical aperture (NA) = 1.20, water immersion objective lens was used for image acquisition. YFP fluorescence was excited by the 514-nm laser line from an argon/krypton laser and detected with a standard PMT on the SP2 or a HyD detector on the SP8 set to a detection window between 520 and 550 nm. RFP was excited with a 561-nm He/Ne laser, and fluorescence was detected with a standard PMT or a HyD detector set at 570 to 650 nm. GFP was excited with a 488-nm laser line from an argon/krypton laser and detected with a PMT or a HyD detector at 500 to 550 nm.

BY-2 cells were imaged on an Olympus FV1000 inverted confocal microscope equipped with a water-corrected 60× objective (NA = 1.2) using a 488-nm laser excitation and a spectral detection bandwidth of 500 to 530 nm for enhanced GFP and a 559-nm laser excitation together with a spectral detection bandwidth of 570 to 670 nm for RFP detection and with a Zeiss 710 inverted confocal microscope with the ZEN 2009 software package and equipped with a 63× water-corrected objective (NA = 1.2). GFP was visualized using 488-nm laser excitation and 500- to 530-nm spectral detection; RFP was visualized using 458-nm laser excitation and 592- to 754-nm spectral detection.

Two-dimensional projections and three-dimensional reconstructions of Z stacks were generated with either ImageJ v.1.48a (http://rsb.info.nih.gov/ij/) or Leica LF Image processing. Color merges were performed with NIH ImageJ v.1.48s or Adobe Photoshop CS5 v12.0.4 (Adobe Systems).

To determine the duration of cytokinesis and the orientation of phragmoplasts (transverse versus torus view), a pok1-1 pok2-3 segregating population derived from a cross with the GFP-MBD marker line described by Marc et al. (1998) was used. Time-lapse imaging of cells displaying spindle stages was performed and continued throughout mitosis at 3- to 5-min intervals. Image intervals were shorter at the beginning of the time lapse to determine the transition from spindle to phragmoplast and at the end of cytokinesis, while longer interval times were chosen during phragmoplast expansion. The duration was timed from the spindle/phragmoplast transition until phragmoplast disassembly.

The rate of phragmoplast expansion was determined using kymograph analysis (Buschmann et al., 2010). We recorded time-lapse images of cells displaying spindles or earlier cell cycle stages at 2.5-min intervals until the phragmoplast made contact with the cell wall. In ImageJ, the segmented line tool was used to select the distance for the measurement, spanning the diameter of the phragmoplast. To determine relative velocities and to visualize the narrowing of the POK1-CAN1 or TAN localization at the CD2, the ImageJ kymograph plug-in (http://www.embl.de/earnet/html/body_kymograph.html) was used. Based on these measurements, the velocities of phragmoplast expansion were calculated in Excel.

Ratiometric BiFC

The ratiometric BiFC Gateway-compatible 2in1 system (Grefen and Blatt, 2012) was used to examine interaction between POK1-685-2066 and TAN. ROP2 was used as a negative control since, like POK1, it localized to the plasma membrane. The potential binding partners were expressed under the control of the Pro35S promoter on the same plasmid, which also carries RFP as an internal expression control. The ratio between YFP and RFP is used to estimate interaction strength. The primers used to amplify the cDNA of binding partners are listed in Supplemental Table 2. Amplicons were cloned via the BP reaction into pDONR221-P1P4 or pDONR221-P2P3. The resulting entry clones were used in subsequent LR reactions with destination vector pBIHt-2in1-N. All steps were performed according to Grefen and Blatt (2012). Protoplasts from Arabidopsis suspension culture were transformed with the pBIHt-2in1-N plasmids according to Schütze et al. (2009). YFP and RFP fluorescence were excited with 514- and 561-nm excitation wavelength by sequential scan using the Leica TCS SP8 resonant scanner and hybrid detector HyD2 set to 518 to 552 nm for YFP fluorescence and PMT4 set to 645 to 731 nm for RFP fluorescence. All images were taken at identical settings.

Quantification of fluorescence intensity was performed in ImageJ. The segmented line tool, set to a 7.5-pixel width, was used to trace the protoplasts and mean fluorescence intensity was measured along this line. Calculation of mean YFP/RFP ratio and the graph were performed in Excel. The mCherry-ROP2 control was generated using CreLox according to Geldner et al. (2009) via recombination of pUNI-ROP2 with pNIGL17. As localization control for TAN, the plasmid pEZRK:TAN-YFP described by Walker et al. (2007) was used.

BY-2 Transformation

Stable BY-2 transformation was performed as described before (Geelen and Inze, 2001). BY-2 cell lines expressing two fluorescent constructs were created by consecutive supertransformation of single transformed lines. Stably transformed calli were screened for fluorescence and localization patterns of tagged proteins were confirmed by analyzing several independent transformants.

FRAP Analysis

FRAP experiments for the POK1 C-terminal fragment GFP-POK1213-2066 were conducted on an Olympus Fluoview1000 inverted confocal microscope. One prebleach images was taken. A region of interest (ROI) was bleached for 10 s with 100% laser power. Subsequent images were taken every 60 s. BY-2 cells were mounted in a chambered cover glass system (Lab-Tek) in 1% low melting point agarose containing BY-2 medium.

The average fluorescence intensity of the bleached region (ROI1) was measured using ImageJ (I(t)). The position of ROI1 was adjusted manually during the time lapse to correct for cell drifting. The average intensity of a second ROI (ROI2) outside the cell was measured to compensate for background fluorescent signal (Ibase). Because Ibase approaches a constant value, we subtracted the average Ibase from I(t) to obtain the actual fluorescence intensity Ifrap(t) (I(t)-average(base) = Ifrap(t)). To estimate the amount of photobleaching due to image acquisition, we measured the average intensity of the whole cell (ROI3). From these values we also subtracted the background signal (average Ibase). These values were set out in a scatterplot and fitted to a linear trendline, which allowed us to determine the theoretical reduction in fluorescence intensity due to image acquisition for each time point (Ibleach(t)).

To obtain a normalized value for fluorescence recovery (Inorm(t)), we first determined the ratio of fluorescence loss due to image acquisition:

\[ \text{Ibleach}(t) = \text{I(t)}/\text{Ibleach}(t) \]  

Next, the remaining fluorescence intensity after photobleaching (Ifrap(post)) was set to zero and all intensity values were adjusted accordingly:

\[ \text{Ifrap}(t) = \text{Ifrap}(t) - \text{Ifrap}(post) \]  

\[ \text{Ibleach}(t) = \text{I(t)} - \text{Ibleach}(t) \]  

\[ \text{I(t)} = \text{I(t)} - \text{Ibleach}(t) \]

Finally, to obtain the normalized values for fluorescence recovery after photobleaching (Inorm(t)), the previous equations were combined as follows:

\[ \text{Inorm}(t) = (\text{Ifrap}(t)/\text{Ibleach}(t)) \]

The average Inorm(t) (FRAP, Figure 6D, prophase cells, n = 4; metaphase cells, n = 7; FRAP, Figure 6E, n = 6) was set out in a scatterplot using Sigmaplot 12. A single exponential regression with rise to a maximum (y = a*(1-exp(-b*t))) was fitted to the data points. The half-time (t1/2) for fluorescence recovery was calculated using the above formula with the fluorescence intensity value corresponding to half of the fitted maximum recovery at the latest time point as y.

FRAP experiments of full-length YFP-POK1 were performed using 4- to 7-d-old seedlings expressing ProPOK1:YFPgPOK1 in the pokf-1 pok2-3 background. Seedlings were mounted in water on solid half-strength nutrient (0.5× Murashige and Skoog salt, 1.5% agar). Using the
Leica SP8, four prebleach images were taken, and then the selected ROI (ROI1) was bleached with identical laser settings for each experiment. Fluorescent signal recovery was imaged at 60-s time intervals for 30 min. FRAP data analysis was performed as described for GFP-POK1C. The background was determined outside of the root (ROI2), and the average intensity of the whole cell (ROI3) was used to correct for photobleaching during image acquisition without trendline fitting.

**Oryzalin Treatment**

Four- to five-day-old seedlings coexpressing YFP-POK1 and RFP-MBD were imaged and subsequently treated by exchanging mounting medium with 10 μM oryzalin solution. Fluorescence of cells that had PPBs, spindles, or phragmoplasts and YFP-POK1 prior to oryzalin treatment were reanalyzed at 5-min intervals for 10 to 30 min. BY-2 cells were mounted in a chambered cover glass system (Lab-Tek). Cells were immobilized in a thin layer (1 mL) of BY-2 medium containing vitamins (Geelen and Inzé 2001), 0.8 to 1% low melting point agarose (Invitrogen) and 10 μM oryzalin. BY-2 cells were imaged to the point where MT depolymerization was evident followed by FRAP analysis.

**Accession Numbers**

Sequence data from this article can be found in the Arabidopsis Genome Initiative or GenBank/EMBL databases under the following accession numbers: At3g17360, POK1; At3g19050, POK2; and At3g05330, TAN1.

**Supplemental Data**

The following materials are available in the online version of this article.

Supplemental Figure 1. Adult Phenotype of pok1 pok2 and Rescue with YFP-POK1.

Supplemental Figure 2. Mitotic Microtubule Arrays in pok1-1 pok2-3 Mutants.

Supplemental Figure 3. Cell Cycle Progression.

Supplemental Figure 4. Subcellular Localization of Different Fluorescent POK1 Fusions.

Supplemental Figure 5. Subcellular Distribution of YFP-POK1 Signal in Arabidopsis Motic Cells.

Supplemental Figure 6. Examples of Transient Microtubules Reaching from the Phragmoplast toward the Putative Cortical Division Site.

Supplemental Table 1. Primers Used for Cloning Full-Length YFP-POK1 and Genotyping of pok1-1.

Supplemental Table 2. List of Primers Used for Cloning 2in1 Plasmids.

Supplemental Table 3. List of Primers Used for Cloning.

Supplemental Movie 1. Time Lapse of Wild-Type Cell Division.

Supplemental Movie 2. Time Lapse of pok1-1 pok2-3 Cell Division.

Supplemental Movie Legends.

Supplemental Movie 1. Time Lapse of Wild-Type Cell Division at 1.5 fps.

Supplemental Movie 2. Time Lapse of pok1-1 pok2-3 Cell Division at 1.5 fps.

**ACKNOWLEDGMENTS**

We acknowledge the ABRC and NASC for distribution of seed used in this study. The instructive advice and provision of E. coli SW105 and the recombineering plasmid collection by Baris Tursun is gratefully acknowledged. We thank Christopher Greifen for advice and the ratiometric BiFC plasmids. We thank Daniela Daumüller, Philipp Reichert, Hs Lee, Richard Gavidia, Angela Kirik, and Samantha Atkinson for help with data collection. We thank Katharina Brancato from the ZMBP Transformation service. We appreciate the support from the Department of Developmental Genetics lead by Gerd Jürgens. Funding for this work was provided by Deutsche Forschungsgemeinschaft grants to S.M. (DFG MU 3133/1-1 and SFB1101) and by a National Institutes of Health grant (R15GM102839)-01A1 (V.K.). A.G. was indebted to the Agency for Innovation by Science and Technology for a predoctoral fellowship.

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The Phragmoplast-Orienting Kinesin-12 Class Proteins Translate the Positional Information of the Preprophase Band to Establish the Cortical Division Zone in *Arabidopsis thaliana*

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