Cortical domain correction repositions the polarity boundary to match the cytokinesis furrow in C. elegans embryos

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SUMMARY
In asymmetrically dividing cells, a failure to coordinate cell polarity with the site of cell division can lead to cell fate transformations and tumorigenesis. Cell polarity in C. elegans embryos is defined by PAR proteins, which occupy reciprocal halves of the cell cortex. During asymmetric division, the boundary between the anterior and posterior PAR domains precisely matches the site of cell division, ensuring exclusive segregation of cell fate. The PAR domains determine the site of cell division by positioning the mitotic spindle, suggesting one means by which cell polarity and cell division might be coordinated. Here, we report that cell polarity and cell division are coordinated through an additional mechanism: the site of cell division repositions the PAR-2 boundary. Gc-mediated microtubule-cortex interactions appear to direct cortical flows of PAR-2 and myosin toward the site of cell division, which acts as a PAR-2 and myosin sink. Embryos with defects in PAR-2 boundary correction undergo mis-segregation of cortical polarity and cytoplasmic determinants, suggesting that PAR domain correction might help prevent cell fate transformation.

KEY WORDS: PAR polarity, Asymmetric cell division, Cytokinesis, C. elegans

INTRODUCTION
Asymmetric cell divisions play essential roles throughout development in all organisms. They provide the basis of cellular differentiation and thus underlie numerous cell fate determination events. Three basic processes ensure that the resulting daughter cells have different fates (Gonczy and Hyman, 1996; Nelson, 2003). First, a polarity axis is established in response to a cue; the formation of a cortical domain marks the polarity axis. Second, fate determinants segregate according to the cortical polarity axis. Third, the mitotic spindle aligns parallel to the polarity axis and positions the cytokinesis furrow. The cortical polarity domain and fate determinants are restricted to one daughter cell.

The importance of coordinating cortical polarity domains with the site of cell division is emphasized by the finding that spindle misorientation relative to the polarity axis can prove tumorigenic in Drosophila (for a review, see Gonzalez, 2007; Knoblich, 2008; Zhong and Chia, 2008). Mutations in the tumor suppressors lethal giant larvae (lgl), scribble (scrib) and discs large (dlg) and regulators of centrosome duplication and spindle orientation [overexpressed Polo kinase and partner of inscuteable (pins) mutants] exhibit defects in asymmetric cell division and undergo overproliferation of stem cell lineages in the brain, leading to massive tumors that can continue to divide indefinitely (Basto et al., 2008; Caussinus and Gonzalez, 2005; Gonzalez, 2007; Knoblich, 2008). In wild-type neuroblasts, a crescent of the fate determinant Numb and its adaptors Pon and Miranda is generated at the basal cortex upon the onset of mitosis through activation of the conserved Bazooka/Par3–Par6–aPKC complex. The cortical polarity defined by the PAR complex is also responsible for spindle orientation, ensuring parallel alignment of the spindle and polarity axes. In the mutant neuroblasts, the spindle axis is randomized relative to the cell polarity axis and cell polarity is distributed randomly during division. In some cases, both daughter cells inherit polarity that would normally be restricted to one self-renewing daughter cell. The downstream effect of this mis-segregation of polarity is continued cell division in both of the daughters and their progeny.

Spindle misorientation does not always lead to symmetric proliferative divisions and tumors. Drosophila PAR polarity mutants such as bazooka (baz or Par3), aPKC and inscuteable (insc) use a correction process known as telophase rescue to position cortical polarity according to the spindle (Cai et al., 2003; Siegrist and Doe, 2007). At the onset of mitosis in these mutant neuroblasts, spindle orientation is random and the Numb crescent is absent. In late mitosis, Numb is recruited to the cortex overlying one of the spindle poles through interactions between the astral microtubules and the cortex. The kinesin motor protein Khc73 and the microtubule-binding protein Mud facilitate formation of a cortical domain of Gz3 and its regulator Pins, thereby establishing cell polarity (for a review, see Knoblich, 2008; Siegrist and Doe, 2005). After telophase rescue, the spindle and polarity axes lie in parallel and the mutant neuroblasts can divide asymmetrically. It remains unclear why some polarity defects are corrected by telophase rescue but others are not.

We sought to address, first, if polarity correction mechanisms exist in other asymmetrically dividing cells; second, where the positional information for correction comes from and how it is transduced; and finally, what circumstances eliminate the ability to correct cortical polarity during cell division. Previous work proposed that C. elegans spn-4 mutant embryos coordinate the cell polarity axis with spindle orientation even when spindle orientation is defective (Gomes et al., 2001) but the mechanism remained unexamined. C. elegans embryos provide several advantages for investigating how polarity domains are corrected during

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asymmetric cell division, offering high spatial and temporal resolution as well as the possibility of both genetic and physical manipulations of the cell division machinery.

_C. elegans_ embryos undergo a series of asymmetric cell divisions to establish the founding cells of five developmental lineages (for a review, see Gonczy and Rose, 2005). One-cell embryos establish a polarity axis shortly after fertilization (for a review, see Cowan and Hyman, 2007), thereby defining the anterior and posterior of the embryo. One-cell embryos divide: the anterior blastomere of the two-cell embryo is already restricted in its developmental potential, whereas the posterior blastomere is not. The posterior blastomere divides three more times in a series of stem cell-like divisions (reviewed by Strome, 2005).

These asymmetric divisions depend on cell polarity defined by PAR proteins (reviewed by Cowan and Hyman, 2004b; Gonczy and Rose, 2005; Schneider and Bowerman, 2003). PAR-3, PAR-6 and aPKC (PKC-3) localize to the anterior half of the one-cell embryo and PAR-2 and PAR-1 localize to the posterior half. The anterior and posterior PAR domains are mutually exclusive. The PAR proteins control the segregation of fate determinants and position the spindle. During division, the anterior PAR proteins are inherited by the anterior daughter cell and the posterior PAR proteins are inherited by the posterior daughter cell. This exclusive segregation of polarity requires that the position of the boundary between the two PAR domains is coordinated with the site of cell division. It has been assumed that because the PAR domains position the spindle, they ensure their own exclusive segregation, but this has not been demonstrated.

We addressed the question of how polarity domains are segregated during asymmetric division by examining the behavior of the PAR domains in response to different genetic or mechanical perturbations. Using mutants with either large or small posterior domains, we have found that a domain correction process requiring Gcr and its regulators GPR-1/2 and LIN-5 acts during division to reposition the boundary between the PAR domains to match the cytokinesis furrow. The Gcr pathway regulates microtubule-cortex interactions and thereby large-scale cortical reorganization that moves PAR-2 toward the furrow. In cases of extreme spindle positioning defects, cortical polarity is mis-segregated and cells divide symmetrically, suggesting the corrective function can be distorted. Correction of cortical polarity domains is likely to be a general mechanism in asymmetrically dividing cells and might help differentiate between symmetric and asymmetric cell divisions in more complex systems.

**MATERIALS AND METHODS**

### C. elegans strains

All _C. elegans_ strains were maintained at 16°C on NGM (nematode growth media) with _E. coli_ OP50 as a food source. Strain genotypes and details of their construction can be found in Table 1.

### RNAi and mutant analysis

RNAi was performed either by injection or feeding as detailed in Table S1 in the supplementary material. Our previous work showed that approximately half of cye-1(RNAi) and cdk-2(RNAi) embryos exhibit no cortical PAR-2 domain throughout the entire cell cycle in one-cell embryos (Cowan and Hyman, 2006). Only cye-1(RNAi)/cdk-2(RNAi) embryos that exhibited cortical PAR-2 prior to anaphase onset were included in the analysis. For analysis of temperature-sensitive zyg-8(b235ts) and spd-1(a5ts) alleles, worms were shifted to 25°C at the fourth larval (L4) stage and incubated at the non-permissive temperature for at least 24 hours prior to recording. For generating partial centrosome duplication phenotypes, mixed-stage zyg-1(b1ts) males and hermaphrodites were shifted to 25°C for 16-24 hours, at which point, L4 and young adult males were selected and mated to GFP::PAR-2; GFP::SPD-2 hermaphrodites for at least 20 hours at 25°C. A homozygous deletion mutant of _F26H9.2_ is available, but the mutant does not reproducibly exhibit the large posterior domain phenotype (C.R.C., unpublished observations). We have not characterized the mutant in any further detail.

### Mechanical manipulations

Displacement of the spindle and cytokinesis furrow using a needle was performed as described (Bringmann et al., 2007). Briefly, using a motorized micromanipulator, a microinjection capillary needle was laid on the free surface of the egg, roughly perpendicular to the long axis of the embryo. The needle was gradually pressed onto the embryo until the eggshell surface was deformed and normal spindle movement was prevented. To generate posterior spindles and furrows, the needle was applied at the time of pronuclear meeting. To generate anterior spindles and furrows, the needle was applied at the maximum anterior position of the centrosome-pronuclei complex, during centration.

The laser ablation experiments were performed on a custom-built setup (developed by Katrin Heinz, IMP) based on an inverted epifluorescence microscope (Zeiss Axiovert 200M) and a UV-VIS-IR 63×I-C-Apochromat 1.25 NA lens (Zeiss), and equipped with a cooled CCD camera (CoolSnap HQ, Photometrics) and LED illumination (465 nm and 565 nm, CoolLED, Precise Excite). The stage was controlled by an XYZ driver (ASI Inc.). The ablation laser (365 nm, 50 mW, Teem Photonics) was focused through the rear slider port. An external pulse generator (1000 Hz, _Pc Lab2000SE, Velleman Components_) was used to control the laser, and ablation was regulated by an external shutter (Thor Labs) placed immediately before the laser path entered the microscope and controlled via the computer. Continuous pulses were used to cut the anterior kinetochore microtubules during early anaphase (determined by posterior spindle pole movement toward the cortex) and then destroy the anterior centrosome to generate maximum anterior displacement of the astral microtubule furrow (Bringmann and Hyman, 2005).

### Time-lapse microscopy, drug treatments and image display

Worms were dissected in 0.1 M NaCl 4% sucrose solution on coverslips to release early embryos and mounted directly on 2% agarose (0.1 M NaCl 4% sucrose) pads on standard microscope slides. For nocodazole treatments, worms were cut open at the vulva in a solution of 7.5 μg/ml nocodazole. 0.1 M NaCl 4% sucrose on a coverslip. The cut worms were incubated in a humid chamber for 10-20 minutes and then further dissected to release embryos and mounted as described. All images were acquired with a Zeiss Axioscope equipped with a Hamamatsu C9100-13 camera attached to a CoolLED light source, and images were captured and analyzed using ZEN (Zeiss) software.

**Table 1. Worm strains used in this study**

<table>
<thead>
<tr>
<th>Strain</th>
<th>Genotype</th>
<th>Reference</th>
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</thead>
<tbody>
<tr>
<td>GFP::PAR-2</td>
<td>JH1380</td>
<td>axEx1094 [ppie-1::gfp::par-2::3::pie-1::pRF4;N2genomic]</td>
</tr>
<tr>
<td>GFP::PAR-2; mCHERRY::PAR-6</td>
<td>TH119</td>
<td>axEx1094; dds26 [ppie-1::mcherry::par-6::3::pie-1::unc-119(+)]</td>
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<tr>
<td>GFP::PAR-2; GFP::γ-tubulin; GFP::histone</td>
<td>UE20</td>
<td>axEx1094; dds6 [ppie-1::gfp::bg-1::3::pie-1::unc-119(+)]; ruls32 [ppie-1::gfp::histoneH2B::3::pie-1::unc-119(+)]</td>
</tr>
<tr>
<td>GFP::PAR-2; GFP::SPD-2</td>
<td>TH49</td>
<td>axEx1094; dds10 [ppie-1::gfp::spd-2::3::pie-1::unc-119(+)]</td>
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<tr>
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<td>DH1</td>
<td>zyg-1(b1ts)</td>
</tr>
<tr>
<td>GFP::PAR-2; zyg-8(b235ts)</td>
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<td>axEx1094; zyg-8(b235ts)</td>
</tr>
<tr>
<td>PIE-1::GFP; zyg-8(b235ts)</td>
<td>UE23</td>
<td>axEx1327 [ppie-1::pie-1::gfp::3::pie-1::pRF4;N2genomic]; zyg-8(b235ts)</td>
</tr>
</tbody>
</table>
coordination of cortical polarity and cytokinesis

RESULTS

The site of cell division repositions the PAR domain boundary

How are the PAR domains segregated in C. elegans embryos so that one cell inherits the anterior PAR domain and the other cell the posterior PAR domain? In one-cell embryos, the boundary between the anterior and posterior PAR domains is located midway along the embryo axis but cell division is asymmetric, displaced toward the posterior (Fig. 1A,B). To investigate how cell polarity and cell division are coordinated, we looked through RNAi-based screen data (Gunalsal et al., 2004; Piano et al., 2000; Sonnichsen et al., 2005; Zipperlen et al., 2001) to find phenotypes in which the PAR boundary is displaced from the middle of the embryo. The position of the PAR boundary is coincident with the position of the pseudocleavage furrow, a deep but transient cortical ingression that reaches its maximum in the middle of the embryo, roughly ten minutes after polarity is initiated (for a review, see Cowan and Hyman, 2004b). We identified several gene products that, when depleted, lead to either anterior or posterior displacement of the pseudocleavage furrow. Further analysis of these RNAi phenotypes using time-lapse microscopy of GFP::PAR-2, a component of the posterior PAR domain, and mChERRY::PAR-6, a member of the anterior PAR complex, indicated that the PAR domain boundary was displaced away from the embryo middle (Fig. 1A; see Movie 1 in the supplementary material). After cell division in these domain-size mutants, however, PAR-2 occupied only the cortex of the posterior cell and PAR-6 was located only in the cortex of the anterior cell, as in wild type. Two possibilities could explain the exclusive inheritance of cell polarity in PAR domain-size mutants: the site of cell division moved to match the altered position of the PAR domain boundary or the PAR domain boundary moved to match the site of cell division.

To determine how the PAR domain boundary and the site of cell division were coordinated, we first examined whether the change in the position of the PAR boundary resulting from mis-sized PAR domains affected the site of cell division. We measured anaphase spindle displacement and the position of the cytokinesis furrow from time-lapse movies of wild type and PAR domain-size mutant embryos. We selected one large domain mutant, F25H9.2(RNAi), and two indistinguishable small domain mutants, cye-1(RNAi) and cdk-2(RNAi) (Cowan and Hyman, 2006), for further analysis. Embryos depleted of the predicted actin and phosphatase regulator F26H9.2 (www.wormbase.org) exhibited large posterior domains extending to approximately 70% embryo length (Fig. 1A,B; see Movie 1 in the supplementary material). However, spindle displacement and the site of cytokinesis were not significantly affected in F26H9.2(RNAi) embryos compared with wild type (Fig. 1B,C). Similarly, in cyclin E- or CDK-2-depleted [cye-1(RNAi) or cdk-2(RNAi)] embryos that established polarity late in the cell cycle and thus had small posterior domains at the time of mitosis (Fig. 1A,B; see Movie 1 in the supplementary material), spindle displacement and the position of cytokinesis were similar to wild-type embryos (Fig. 1B,C). The position of the PAR domain boundary, as determined by domain size, had little influence on the site of cell division.

It seemed probable that the PAR domain boundary was repositioned to match the site of cell division. We analyzed the position of the PAR domain boundary relative to the site of cell division by time-lapse microscopy of GFP::PAR-2. In F26H9.2(RNAi) embryos, the PAR boundary was located in the anterior embryo half prior to cell division (Fig. 1). As the cell began to divide, the PAR boundary shifted posteriorly to match the site of cell division (Fig. 1A,D; see Movie 1 in the supplementary material). In cye-1(RNAi) or cdk-2(RNAi) embryos exhibiting small posterior domains, the PAR boundary was in the posterior embryo half prior to cell division (Fig. 1). The PAR-2 domain expanded during mitosis until the domain boundary matched the site of cell division.
division (Fig. 1A,D; see Movie 1 in the supplementary material). Although the PAR-2 domain expanded throughout the cell cycle in cye-1(RNAi) and cdk-2(RNAi) embryos owing to delayed polarity establishment, the rate at which PAR-2 moved changed significantly during division (Fig. 1D). Similar PAR domain size correction can be observed in other mutants with enlarged posterior domains, such as rga-3/4(RNAi) embryos (Schmutz et al., 2007; Schonegg et al., 2007), in two-cell embryos with spindle orientation defects (Gomes et al., 2001) and in subsequent divisions of the germline lineage in wild-type C. elegans embryos (see Figs S1 and S2 in the supplementary material; see Movie 2 in the supplementary material). C. elegans embryos appear to match the PAR domain boundary to the site of cell division. 

The PAR domain boundary responds to the position of the cytokinesis furrow
What is the positional information that corrects the PAR domain boundary during cell division? Two general mechanisms could provide a spatial signal: a cell division-independent landmark or a cell division-dependent cue, such as the spindle or cytokinesis

Fig. 1. Mis-sized PAR domains are corrected during cell division. Wild-type (left), F26H9.2(RNAi) big PAR-2 domain (middle) and cye-1(RNAi)/cdk-2(RNAi) small PAR-2 domain (right) embryos. (A) Time-lapse images of GFP::PAR-2 and mCHERRY::PAR-6, marking the posterior and anterior PAR domains, respectively. F26H9.2(RNAi) embryos exhibit increased cortical activity in the anterior domain as the domain size decreases. Higher temporal resolution time-lapse sequences (30 second intervals) of GFP::PAR-2 are shown to the right. (B) Posterior PAR-2 domain size in domain-size mutants. Striped green, maximum PAR-2 domain; light green, PAR-2 domain at nuclear envelope breakdown; dark green, PAR-2 domain after cell division; black triangles, cytokinesis furrow position; vertical dotted line, wild-type PAR boundary position at nuclear envelope breakdown; vertical solid line, wild-type PAR boundary position at nuclear envelope breakdown. (C) Spindle pole position in domain-size mutants. Kymographs were generated by tracking centrosome position in embryos expressing GFP::PAR-2;GFP::γ-tubulin;GFP::histone or GFP::PAR-2;GFP::SPD-2. Gray x’s, individual positions; gray lines, average position (wild type, n=7; F26H9.2(RNAi), n=5; cye-1(RNAi), n=6). (D) PAR-2 boundary position relative to the cell cycle. Green lines, PAR-2 boundary (the domain extends to the posterior pole); red dots, centrosomes; blue x’s, pronuclei/chromosomes; black triangles, cytokinesis furrow position. In cye-1(RNAi) embryos, the PAR-2 domain continues to expand throughout the cell cycle but the rate of PAR-2 boundary movement increases significantly at anaphase onset. In A, C and D, times are indicated relative to anaphase onset. A, anterior; P, posterior; posterior pole, 0.0. Scale bars: 10 μm.
We tested the existence of a static cellular landmark by asking whether a misplaced site of cell division could reposition a correctly positioned PAR domain boundary. Because the posterior PAR-2 boundary is sharper than the PAR-6 boundary and thus easier to identify, we focused on the PAR-2 domain in subsequent experiments. First, we displaced the site of cell division toward either the anterior or posterior by pushing the spindle with a glass needle (see Materials and methods). The cleavage furrow formed in the anterior- or posterior-third of the embryo. After anaphase onset, regardless of the direction or extent of displacement, the PAR-2 domain boundary shifted to match the site of cell division (anterior spindle, n=4; posterior spindle, n=7; Fig. 2A; see Movie 3 in the supplementary material). Second, we generated spindles with varying spindle pole sizes by partially inhibiting centriole duplication through partial inactivation of ZYG-1, an essential regulator of this process (O’Connell et al., 2001), using a conditional mutant allele. The resulting asymmetric spindles led to a range of cleavage sites. Following more complete zyg-1 inactivation, centriole duplication did not occur, leaving only one centrosome, and the furrow was displaced to the anterior. In all cases, the PAR-2 domain boundary matched the site of cytokinesis furrow initiation (n=14; Fig. 2B; see Movie 4 in the supplementary material). The PAR-2 domain boundary can be repositioned to match the site of cell division, regardless of location within the cell.

The PAR-2 domain boundary matched the furrow in embryos with a single centrosome [zyg-1(b1)], thus lacking a mitotic spindle. Similarly, embryos depleted of microtubules do not form a spindle but the posteriorly located centrosomes can still induce furrow ingression in the anterior (Werner et al., 2007). The PAR-2 boundary matched the furrow position in embryos depleted of microtubules (Fig. 3A), indicating that the cytokinesis furrow, rather than the spindle, is important for PAR domain correction. When cytokinesis furrow ingression was prevented by depletion of the actin nucleation factor CYK-1, the PAR-2 domain boundary fluctuated after anaphase onset and eventually shrank toward the posterior pole (Fig. 3A). Together, these experiments indicate that the cytokinesis furrow positions the PAR domain boundary during cell division.

In C. elegans embryos, the cytokinesis furrow is positioned by two signals: one from the spindle pole asters and one from the spindle midzone (Bringmann and Hyman, 2005; Dechant and Glotzer, 2003; Verbrugghe and White, 2004; Werner et al., 2007). Normally these two signals converge, generating one furrow. The positions of the two signals can be separated by asymmetrically cutting the spindle with a laser, leading to two spatially and temporally distinct furrows (Bringmann and Hyman, 2005). The first furrow forms in response to the asters and the second in response to the midzone. To see whether the PAR domain boundary distinguished between the aster and midzone furrows, we followed the PAR-2 boundary position relative to the two furrows following asymmetric spindle cutting (n=3; Fig. 3B; see Movie 6 in the supplementary material). The PAR-2 boundary shifted anteriorly to match the first furrow specified by the asters. When the second midzone-specified furrow ingressed, the PAR-2 boundary retracted slightly from the astral furrow but did not precisely match the midzone furrow. The astral cytokinesis furrow appears to dominate PAR-2 boundary positioning.

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**Fig. 2.** The site of cell division repositions the PAR-2 domain boundary. (A) Time-lapse images and kymographs of the GFP::PAR-2 domain following manual spindle displacement using a glass needle (pink in the first images). The spindle was pushed toward the posterior (left panel) or the anterior (right panel). Green solid lines, PAR-2 boundary; green dotted lines, cortical PAR-2; black triangle, cytokinesis furrow position; pink triangle, needle position. Times are indicated relative to anaphase onset. (B) Linear position of the PAR-2 domain (green bar), cytokinesis furrow (black triangle) and centrosomes (red dots) in zyg-1(b1ts) embryos with asymmetric (partial loss-of-function) or monopolar (complete loss-of-function) spindles. Bars represent individual embryos. The size of centrosome dots indicates the relative size of GFP::SPD-2 centrosomes (normal versus small centrosome) but not absolute size. The vertical gray bar indicates the range of furrow positions in wild-type embryos. A, anterior; P, posterior; posterior pole, 0.0. Scale bars: 10 μm.
PAR domain correction depends on Gα, GPR-1/2 and LIN-5

What are the molecular requirements for PAR domain correction? The PAR-2 domain boundary appeared to respond to the astral cytokinesis furrow and so we asked whether molecules involved in positioning the furrow are required to position the PAR domain boundary. At the onset of anaphase in embryos depleted of two redundant Gα subunits \([\text{goa-1} + \text{gpa-16}(\text{RNAi})]\) required for the astral signal (Bringmann et al., 2007; Dechant and Glotzer, 2003), the PAR-2 domain boundary shrank toward the posterior pole of the embryo, and after division, the PAR-2 domain occupied only a subset of the posterior cell circumference (Fig. 4A,B; see Movie 7 in the supplementary material). A similar failure in domain correction was observed in embryos lacking the conserved Gα regulator GPR-1/2 (AGS3/Pins) (Gotta et al., 2003; Colombo et al., 2003; Srinivasan et al., 2003) and the Gα effector LIN-5 (NuMA/Mud) (Srinivasan et al., 2003) (see Movie 7 in the supplementary material). A similar failure in domain correction was observed in embryos lacking the conserved Gα signaling pathway, GPR-1/2 and LIN-5.

Gα-dependent cytokinesis furrow specification involves the reorganization of the cell cortex, including a domain of the Gα regulator GPR-1/2 (Bringmann et al., 2007). By tracking the cortical intensity of PAR proteins during domain correction, we found that the PAR-2 domain also underwent reorganization: the maximum PAR-2 concentration shifted from the posterior pole to the nascent furrow position during anaphase (Fig. 5A; see Fig. S3 in the supplementary material). The morphology of the PAR-6 domain did not appear to change (see Fig. S3 in the supplementary material). At the same time, cortical myosin moved toward the nascent furrow, predominantly from the posterior (see Fig. S4 in the supplementary material). The morphology of the PAR-6 domain did not appear to change (see Fig. S3 in the supplementary material). At the same time, cortical myosin moved toward the nascent furrow, predominantly from the posterior (see Fig. S4 in the supplementary material). The morphology of the PAR-6 domain did not appear to change (see Fig. S3 in the supplementary material). At the same time, cortical myosin moved toward the nascent furrow, predominantly from the posterior (see Fig. S4 in the supplementary material).
cortical domain correction, Gα is required to shift the distribution of PAR-2 toward the cytokinesis furrow. This cortical reorganization is accompanied by a coincident Gα-dependent myosin flow.

The coincidence of PAR-2 redistribution and furrow-directed myosin flows, both of which required Gα, suggested a role for cortical flow in domain correction. We therefore analyzed PAR-2 domain reorganization in embryos with reduced acto-myosin contractility, the mechanical force thought to drive cortical flow. The myosin regulatory light chain MLC-4 is required for cortical contractility, cortical flow and furrow ingression. Embryos strongly depleted of mlc-4 fail to establish polarity (Shelton et al., 1999). Embryos partially depleted of mlc-4 [mlc-4(partialRNAi)] could establish a normal PAR-2 cortical domain (Fig. 5B; see Fig. S2 in the supplementary material), although the polarity axis was sometimes tilted with respect to the long axis of the cell in contrast to wild-type embryos. During anaphase in mlc-4(partialRNAi) embryos, the PAR-2 domain did not exhibit an anterior shift in PAR-2 intensity (Fig. 5A). If the PAR-2 domain boundary was within 0.25 embryo length of the cytokinesis furrow, the boundary shifted to match the furrow position. If the PAR-2 domain boundary was further from the cytokinesis furrow, the boundary...
did not correct (Fig. 5B,C). The limited contractility remaining in mlc-4(partialRNAi) embryos might be sufficient for small shifts in the position of the PAR-2 domain boundary. Large-scale reorganization of the PAR-2 domain and bigger shifts in boundary position both appear to depend on myosin-driven cortical contractility, consistent with the finding that Gα-directed cortical flows are required for domain correction.

During cell division, Gα has an established role controlling spindle position through regulation of microtubule-cortex interactions. To see whether Gα-dependent cortical reorganization is controlled through Gα-dependent microtubule regulation as opposed to direct regulation of contractility, we asked whether we could rescue the lack of domain correction in Gα-depleted embryos by depolymerizing microtubules. We treated goa-1+gpa-16(RNAi) embryos with the microtubule poison nocodazole and assessed the position of the PAR-2 domain boundary and cortical myosin during cell division. Whereas Gα-depleted embryos do not undergo cortical reorganization during anaphase, Gα-depleted embryos treated with nocodazole underwent PAR-2 domain correction and furrow-directed myosin flow. Thus, Gα controls microtubule-cortex interactions that facilitate large-scale cortical reorganizations, the probable mechanical basis of PAR-2 domain repositioning during cell division.

**PAR domain size and spindle orientation might limit domain correction**

Not all embryos with altered domain sizes repositioned the PAR domain boundary to match the site of cell division. We found that cye-1(RNAi) and cdk-2(RNAi) embryos did not correct PAR domains if the PAR-2 domain occupied less than ~20% cell circumference (n=7 out of 20; Fig. 6A,B; see Movie 9 in the supplementary material). A similar size limit was seen in mex-5(RNAi) embryos, which often exhibited small PAR-2 domains (S. Reiter and C.R.C., unpublished observations). In embryos in which the PAR-2 domain extended into both nascent daughter cells owing to spindle misorientation [zyg-8(RNAi) or zyg-8(b235ts)], PAR-2 expanded around the cortex of both cells (Fig. 6C), but only when the PAR-2 domain occupied 30% or more of the cortex bounded by the cytokinesis furrow (n=25 out of 28 cortical segments). The mis-segregation of PAR-2 in zyg-8(–) embryos often correlated with mis-segregation of cell fate determinants, such that both daughter cells expressed the germline marker PIE-1 ([zyg-8(RNAi) or zyg-8(b235ts)], PAR-2 expanded around the cortex of both cells (Fig. 6C), but only when the PAR-2 domain occupied 30% or more of the cortex bounded by the cytokinesis furrow (n=25 out of 28 cortical segments). The mis-segregation of PAR-2 in zyg-8(–) embryos often correlated with mis-segregation of cell fate determinants, such that both daughter cells expressed the germline marker PIE-1 (n=4 out of 8 zyg-8(b235ts) embryos; Fig. 6C). Equal segregation of PAR-2 also occurred if the cytokinesis furrow was induced to bisect the domain by physically manipulating spindle orientation in genetically wild-type embryos (n=2; see Fig. S5 and Movie 10 in the supplementary material). Thus, either a small PAR-2 domain or spindle misorientation relative to the PAR-2 domain reduced the ability of the embryo to undergo domain correction and exclusively segregate polarity. Although we cannot rule out a size-independent mechanism, these results are consistent with a model in which PAR-2 domain size influences whether or not the domain boundary will be corrected: if the domain size is above a threshold, it expands to match the furrow, and if it is below a threshold, it does not.

Together, our experiments show that C. elegans embryos posses a cortical domain correction mechanism that ensures the PAR boundary matches the site of cell division. The PAR boundary responds to the position of the cytokinesis furrow by following cortical myosin flows into the furrow. Gα-mediated microtubule-cortex interactions facilitate cortical flow and thus domain correction. Embryos defective for Gα exhibit cell fate...
transformations in subsequent embryonic divisions ($n=5$ out of 8; Fig. 4C), suggesting that domain correction might be important to maintain cell fates. Domain correction fails either when the PAR-2 domain is small or when the spindle axis is misoriented relative to the polarity axis, indicating that the corrective capacity is limited.

**DISCUSSION**

Our data demonstrate that the boundary between PAR polarity domains is repositioned during anaphase to match the site of cell division. This correction of cortical polarity domains ensures that the two resulting cells inherit the appropriate complement of PAR proteins and thus the capacity for differentiation or for continued asymmetric division. Initially, the PAR domains determine the position of the spindle (Grill et al., 2001; Kemphues et al., 1988). Later, the spindle indirectly determines the position of the cytokinesis furrow. Finally, the cytokinesis furrow controls the position of the PAR domain boundary. This circular regulatory pathway ensures the proper segregation of PAR domains during cell division.

PAR domain correction entails large-scale reorganization of the cortex, such that cortical myosin and PAR-2 exhibit bulk movement toward the cytokinesis furrow. These flows of cortical myosin and PAR-2 never moved through the furrow but rather the furrow appeared to act as a myosin and PAR-2 sink. Cortical flows depended on $G_{\alpha}$, GPR-1/2 and LIN-5, members of a conserved $G_{\alpha}$ signaling complex. This molecular requirement for cortical flow is largely shared by cytokinesis furrow specification in response to signals from the astral microtubules (Fig. 4) (Bringmann et al., 2007; Dechant and Glotzer, 2003; Werner et al., 2007), suggesting that PAR domain correction and aster-dependent cytokinesis might result from the same cortical reorganization. By contrast, midzone-specified furrowing is thought to involve local control of the cortex and thus might have a limited capacity to influence the PAR-2 domain.

$G_{\alpha}$ appeared to control cortical flows indirectly through its role in regulating microtubule-cortex interactions as depolymerization of microtubules could restore cortical flows in embryos lacking $G_{\alpha}$. Stable microtubules generally suppress contractility (for a review, see Mandato et al., 2000). In the posterior, $G_{\alpha}$ might help promote microtubule instability, which could facilitate spindle movement and increase cortical contractility. The spatial asymmetry in contractility arising from the displacement of the spindle toward the posterior might be sufficient to induce cortical flow along the contractility gradient, away from the posterior pole (Bray and White, 1988) (reviewed by Cowan and Hyman, 2004b).
It remains to be determined how PAR-2 localizes to the cell cortex and the mechanical basis by which PAR-2 is coupled to cortical myosin flows.

In *C. elegans* embryos and *Drosophila* neuroblasts, spindle orientation and polarity correction mechanisms both depend on heterotrimeric Gα, its regulator GPR-1/2/Pins and LIN-5/Mud (Siegrist and Doe, 2005). In vertebrates, spindle orientation is coordinated with cell polarity (Chang et al., 2007; Fleming et al., 2007; Konno et al., 2008; Llamprecht et al., 1986; Lechler and Fuchs, 2005; Yingling et al., 2008), as in *C. elegans* and *Drosophila*. Likewise, regulators of heterotrimeric G proteins have been shown to mediate spindle orientation during asymmetric divisions in birds and mammals (Morin et al., 2007; Sanada and Tsai, 2005). In both the epidermal and neural progenitor divisions in the mouse brain, the orientation of the mitotic spindle relative to polarity cues correlates with the fate of the daughter cells. Inhibition of G protein regulators interferes with correct spindle orientation and alters cell fates (Sanada and Tsai, 2005). The emphasis in these studies has been on the alignment of the spindle relative to the existing cell polarity. The contribution of the Gα pathway to correcting cell polarity remains unexplored in vertebrate systems.

Given that cell polarity can be corrected to match the site of cell division, why does mis-segregation of cell polarity occur? We have demonstrated that in *C. elegans* embryos, limitations in the correction process might contribute to the mis-segregation of cortical polarity in two ways: when correction cannot act or when correction is active in the inappropriate cell. In cases where the PAR-2 domain size was small or in embryos depleted of Gα, the PAR-2 domain boundary did not match the furrow. In cases where the spindle was misaligned with respect to the polarity axis, PAR-2 domains exceeded the apparent size threshold in both daughter cells and PAR-2 expanded around both to match the furrow. Either type of mis-segregation could alter cell fate segregation (Figs 4, 6). By analogy, *Drosophila* mutant neuroblasts that exhibit symmetric proliferative divisions might result when domain thresholds were surpassed in both nascent cells, either owing to spindle misorientation, changes in domain size, or both, leading to correction acting in both daughters.

Two distinct outcomes were observed in embryos that accumulated PAR-2 in the anterior cortex. In large PAR-2 domain *F26H2.2(RNAi)* embryos, PAR-2 disappeared from the anterior and was inherited by only one cell. In *zyg-8(–)* embryos, PAR-2 expanded around the anterior and was inherited by both daughter cells. In both cases, the PAR-2 domain boundary appeared to move toward the cytokinesis furrow, suggesting that correction was active. One difference between the two conditions was the presence of PAR-2 on the cortex overlapping the spindle pole, raising the possibility that astral microtubule interactions with the PAR-2 cortex determine domain expansion. Gα-regulated microtubule-cortex interactions indeed appear to be essential for domain correction (Fig. 4) and PAR polarity is thought to direct Gα activity (Colombo et al., 2003; Gotta et al., 2003), suggesting a mechanistic connection between PAR-2 domain size and Gα-induced cortical flow. Alternatively, another domain property that we could not detect, such as protein amount, might be monitored. Further experiments will be necessary to understand what cortical properties determine domain correction and how those properties are sensed.

Cortical domain size thresholds provide a useful basis for considering how stem cells maintain the capacity for both proliferation through symmetric division and differentiation through asymmetric division (for a review, see Morrison and Kimble, 2006). The domain size perceived during division appears to be determined by the orientation of cell division relative to the polarity axis. Cells can ensure either symmetric or asymmetric division by controlling the site of cell division relative to polarity, for instance, by regulating spindle orientation. Such a domain size threshold-based mechanism might be relevant in the embryonic mouse brain, for example, where a precise distinction between spindle orientation during symmetric and asymmetric divisions has been difficult to determine (for a review, see Zhong and Chia, 2008).

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Competing interests statement
The authors declare no competing financial interests.

Supplementary material
Supplementary material for this article is available at http://dev.biologists.orglookup/suppl doi:10.1242/dev.040346-DC1

References


Cortical PAR intensity (fraction of maximum)

Position along anterior-posterior cortex (fraction of cortex length)

wild type

goa-1+gpa-16(RNAi) + nocodazole
### Wild Type

**NMY-2::GFP along anterior-posterior cortex (fraction of cortex length)**

- Time relative to anaphase onset (min:sec):
  - $0:00$
  - $1:40$
  - $3:20$

### goa-1+ gpa-16(RNAi)

**NMY-2::GFP along anterior-posterior cortex (fraction of cortex length)**

- Time relative to anaphase onset (min:sec):
  - $0:00$
  - $2:40$

### goa-1+ gpa-16(RNAi) nocodazole

**NMY-2::GFP along anterior-posterior cortex (fraction of cortex length)**

- Time relative to nuclear envelope breakdown (min:sec):
  - $0:00$

### gpr-1/2(RNAi)

**NMY-2::GFP along anterior-posterior cortex (fraction of cortex length)**

- Time relative to anaphase onset (min:sec):
  - $0:00$
  - $2:40$

### goa-1+ lin-5(RNAi)

**NMY-2::GFP along anterior-posterior cortex (fraction of cortex length)**

- Time relative to anaphase onset (min:sec):
  - $0:00$
  - $2:40$

### gpr-1/2(RNAi)

**NMY-2::GFP along anterior-posterior cortex (fraction of cortex length)**

- Time relative to anaphase onset (min:sec):
  - $0:00$
  - $2:40$
PAR-2 occupied entire cell cortex during cleavage
PAR-2 disappeared from cell cortex during cleavage

Size of PAR-2 domain at metaphase (percent of cortical segment defined by nascent furrow bisection)
### Table S1. dsRNA primer sequences, clone information, RNAi method and conditions

<table>
<thead>
<tr>
<th>Gene</th>
<th>Primer 1 sequence</th>
<th>Primer 2 sequence</th>
<th>Feeding clone reference</th>
<th>Method</th>
</tr>
</thead>
<tbody>
<tr>
<td>cdk-2</td>
<td>CTTCAACTGCAAATCCAGCA</td>
<td>TGAATTCCTGGTTCCTGGG</td>
<td>GS I-1D09</td>
<td>I 22 hours</td>
</tr>
<tr>
<td>cye-1</td>
<td>TATCCGGGATTTTCCAAGC</td>
<td>GAAGGATCGAATGTAGC</td>
<td>I 22 hours / F 30 hours</td>
<td></td>
</tr>
<tr>
<td>lin-5</td>
<td>GCGTTCTCTCTGCTGAAGG</td>
<td>CGAGCAAAGAAGTCTGGAGG</td>
<td>GS I-1D09</td>
<td>I 22 hours / F 30 hours</td>
</tr>
<tr>
<td>goa-1+ gpa-16</td>
<td>Y9588A.5</td>
<td></td>
<td>Colombo et al., 2003</td>
<td>F 48 hours</td>
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<tr>
<td>cye-1</td>
<td>K03E5.3</td>
<td></td>
<td></td>
<td>F 24 hours</td>
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<tr>
<td>cdk-2</td>
<td>C37A2.4</td>
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<tr>
<td>lin-5</td>
<td>T09A5.10</td>
<td></td>
<td></td>
<td>I 22 hours / F 30 hours</td>
</tr>
<tr>
<td>goa-1+ gpa-16</td>
<td>C26C6.2+</td>
<td></td>
<td></td>
<td>F 48 hours</td>
</tr>
<tr>
<td>cye-1</td>
<td>GS III-3P11</td>
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<td></td>
<td>F 24 hours</td>
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<td>cdk-2</td>
<td>F22B7.3</td>
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<tr>
<td>tbb-1</td>
<td>F44F4.11</td>
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<td>F 48 hours</td>
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<td>tbb-2</td>
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<td>zyg-8</td>
<td>Y79H2A.11</td>
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<tr>
<td>cyk-1</td>
<td>F11H8.4</td>
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<td>F 48 hours</td>
</tr>
<tr>
<td>mlc-4 partial</td>
<td>C56G7.1</td>
<td></td>
<td></td>
<td>I 12 hours</td>
</tr>
</tbody>
</table>

Production of dsRNA in vitro was performed as described (Gönczy et al., 2000). The gene-specific primer sequences used are indicated in the table; either a T3 or T7 polymerase recognition sequence was added to the 5' end of the sequence listed. To generate feeding clones, the primers indicated were used to amplify a portion of the gene, which was cloned into the L4440 vector and transformed into HT115 E. coli (Timmons et al., 2001). Genomic N2 DNA was used as the template for both in vitro RNA production and feeding clone construction. The sources of other feeding clones are indicated. The tbb-1 and cdk-2 clones were obtained from Geneservice Ltd (Cambridge, UK, ‘GS’); the identification numbers are provided. RNAi was performed either by injection of dsRNA transcribed in vitro (injection, I) or by feeding worms with bacteria producing dsRNA (feeding, F) as described (Kamath et al., 2001). The method indicates the time elapsed from injection or placing the worms on feeding bacteria until recording. Times should be considered as averages (approximately ± 1 hour). All worms were kept at 25°C during RNAi.

