

Asymmetric cortical extension shifts cleavage furrow position in *Drosophila* neuroblasts

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ABSTRACT The cytokinetic cleavage furrow is typically positioned symmetrically relative to the cortical cell boundaries, but it can also be asymmetric. The mechanisms that control furrow site specification have been intensively studied, but how polar cortex movements influence ultimate furrow position remains poorly understood. We measured the position of the apical and the basal cortex in asymmetrically dividing *Drosophila* neuroblasts and observed preferential displacement of the apical cortex that becomes the larger daughter cell during anaphase, effectively shifting the cleavage furrow toward the smaller daughter cell. Asymmetric cortical extension is correlated with the presence of cortical myosin II, which is polarized in neuroblasts. Loss of myosin II asymmetry by perturbing heterotrimeric G-protein signaling results in symmetric extension and equal-sized daughter cells. We propose a model in which contraction-driven asymmetric polar extension of the neuroblast cortex during anaphase contributes to asymmetric furrow position and daughter cell size.

Monitoring Editor
Fred Chang
Columbia University

Received: Mar 1, 2011
Revised: Sep 1, 2011
Accepted: Sep 12, 2011

INTRODUCTION

During development, asymmetric cell division is used repeatedly to generate daughter cells that differ in size and fate (Knoblich, 2008). Daughter cell size asymmetry, which may be important for maintaining progenitor growth potential (Jorgensen and Tyers, 2004), can result from asymmetric positioning of the cleavage furrow (Glotzer, 2004). The site on the cortex where the cleavage furrow forms can be specified by the mitotic spindle (Oliferenko *et al.*, 2009; von Dassow, 2009). For example, in the *Caenorhabditis elegans* zygote the spindle is displaced posteriorly at the end of metaphase and the furrow forms accordingly, leading to large anterior and small posterior daughter cells (Albertson, 1984; Keating and White, 1998; Glotzer, 2004). However, the position of the furrow depends not only on the site of furrow selection, but also on the relationship between the site

of spindle specification and its position relative to the poles of the cell. Thus it is possible that an asymmetric furrow could result from specification of a furrow site at the center of the cell followed by asymmetric movement of the cortex at the cell poles. The morphology changes during mitosis can be dramatic, prompting us to explore the role of polar cortical movements in furrow position.

The study of furrow positioning has focused on how the site on the cortex that will become the cleavage furrow is selected by the mitotic spindle (Glotzer, 2004; von Dassow, 2009). The spindle pathway for furrow site selection is initiated at the central spindle by the centralspindlin complex consisting of the kinesin Pavarotti (ZEN-4 in *C. elegans*), the RACGAP50 Tumbleweed (CYK-4 in *C. elegans*), and the RhoGEF Pebble (ECT-2 in *C. elegans*). At the cell cortex, centralspindlin activates a narrow band of GTP-loaded Rho GTPase (Bement *et al.*, 2006), ultimately leading to recruitment and activation of actomyosin to initiate cleavage furrow constriction. Astral microtubules can sharpen the site of furrow selection by inhibiting activation of Rho at the poles.

In addition to the spindle-directed equatorial constriction that occurs during cleavage furrowing, other morphological changes can happen late in mitosis. Symmetrically dividing cells, such as cultured S2 cells, round up at the beginning of mitosis but elongate at the poles late in anaphase (Hickson *et al.*, 2006; Kunda *et al.*, 2008; Rosenblatt, 2008; Figure 1A). The elongation that results from polar extension (i.e., outward displacement of the cortex) allows the spindle to expand into the polar regions as

This article was published online ahead of print in MBoC in Press (<http://www.molbiolcell.org/cgi/doi/10.1091/mbc.E11-02-0173>) on September 21, 2011.
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Abbreviations used: *Dlg*, discs large; GFP, green fluorescent protein; GMC, ganglion mother cell; moe, moesin; mRFP, monomeric red fluorescent protein; NB, neuroblast; *pins*, partner of inscuteable; *rod*, rough deal; *sqh*, spaghetti squash (*Drosophila* myosin II).

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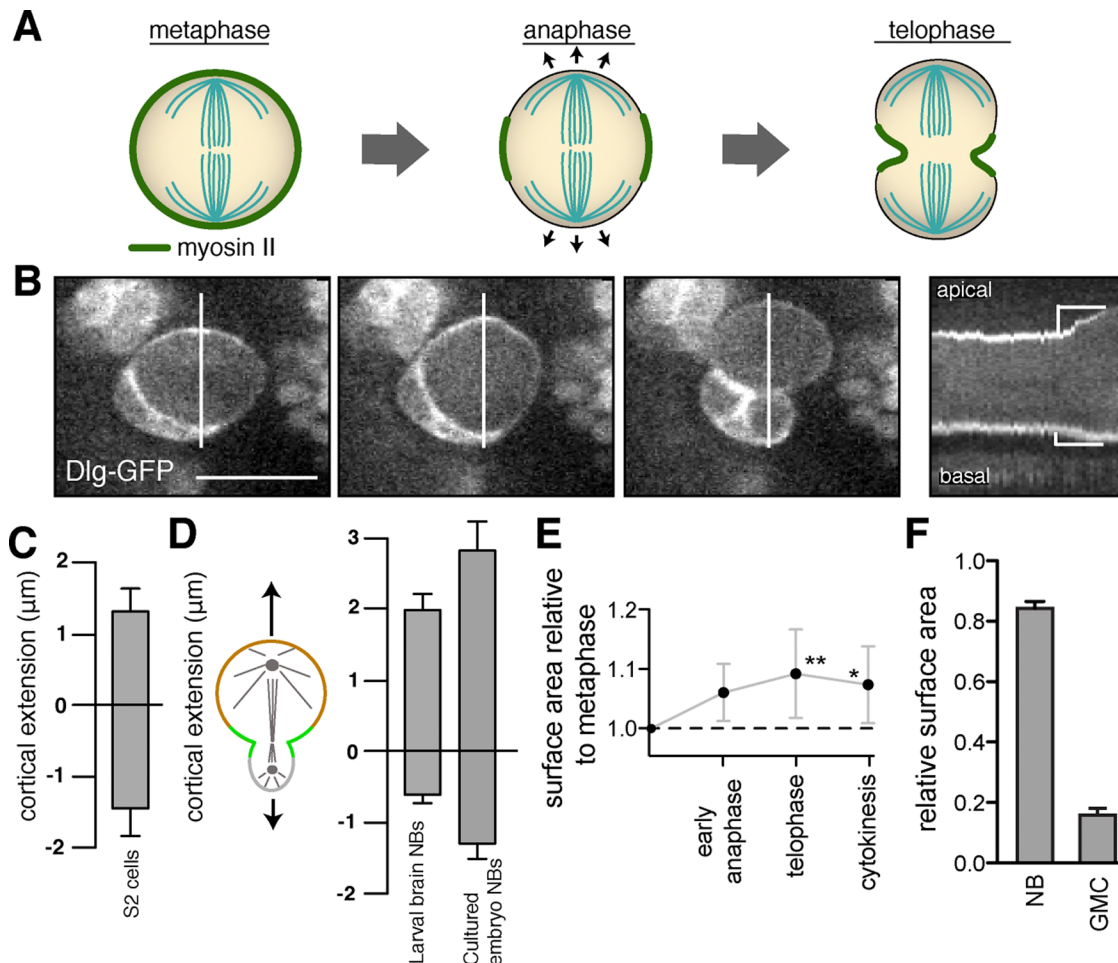


FIGURE 1: *Drosophila* neuroblasts undergo asymmetric polar elongation during anaphase. (A) Schematic of a symmetric division in which cortical extension is equal at both poles during anaphase. Myosin II is localized uniformly early in mitosis but becomes restricted to the equatorial region during anaphase. (B) Neuroblast cortical dynamics during mitosis using Dlg-GFP as a cortical marker. Selected frames from the movie are shown along with a kymograph of the entire division at 6-s intervals. The lines in the movie frames denote the section of the frame used for the kymograph. Cortical extension during anaphase is denoted by yellow brackets in the kymograph. The signal is enriched at the basal cortex because of contact with GMCs from previous divisions that also express GFP-Dlg (Supplemental Figure S1). (C) Mean anaphase polar extension in cultured *Drosophila* S2 cells transiently expressing Cherry-Zeus. The edge of the cell was marked at the point at which cytoplasmic fluorescence was no longer observed. Error bars, 1 SD. (D) Quantification of anaphase cortical extension in wild-type neuroblasts. The mean cortical extension from metaphase to the end of anaphase is shown for the apical (top) and basal (bottom) cortexes (NB, neuroblast). (E) The surface area of dividing neuroblasts measured using three-dimensional reconstruction normalized to that at the end of metaphase. The time points for measurements were early anaphase (completion of cortical extension), telophase (initiation of furrowing), and cytokinesis (completion of furrowing). (F) Mean relative surface areas of the daughter neuroblast (NB) and GMC that results from a neuroblast asymmetric cell division measured as in E at the completion of furrowing.

anaphase progresses (Rosenblatt *et al.*, 2004; Hickson *et al.*, 2006). The degree to which the polar cortex extends in cells that divide asymmetrically has been less characterized. Here, we use neuroblast asymmetric cell division as a model system for investigating the role of polar extension in cleavage furrow position and daughter cell size.

Drosophila neuroblasts are progenitors of the CNS, dividing to generate a larger apical cell that retains the neuroblast fate and a smaller basal ganglion mother cell (GMC) that assumes a differentiated fate (Doe, 2008; Knoblich, 2008). Neuroblasts divide rapidly, and daughter cell size asymmetry may be a mechanism for retaining sufficient resources to allow neuroblast self-renewal. The difference in fate of the two cells results from the polarization of fate determi-

nants into separate apical and basal cortical domains that are precisely separated by the cleavage furrow (Knoblich, 2008). For example, the protein Miranda is localized to the basal cortex in metaphase and becomes segregated into the basal daughter cell as part of the machinery that confers GMC fate (Rolls *et al.*, 2003; Lee *et al.*, 2006; Atwood *et al.*, 2007).

In addition to the spindle-directed pathway that controls targeting of furrow components such as centralspindlin to the equatorial cortex, neuroblasts possess a spindle-independent pathway that targets furrow components to the basal cortex (Cabernard *et al.*, 2010). Shortly before the spindle directs recruitment to the equator, the spindle-independent pathway initiates contraction of the basal cortex (which becomes the GMC following abscission). A similar

polar domain containing myosin II has recently been identified in *C. elegans* Q neuroblasts (Ou *et al.*, 2010), which also divide to generate unequal-sized progeny, suggesting that the domain could be part of a common mechanism for daughter cell size asymmetry. Although the “asymmetric contraction” pathway is active in both *Drosophila* and *C. elegans*, and thus may be widely used in other systems, little is known about its mechanism. In particular, we know little about the cortical properties of the myosin-enriched (basal) and myosin-depleted (apical) neuroblast cortex.

RESULTS

Neuroblasts elongate asymmetrically during anaphase

We imaged larval brain neuroblasts expressing a green fluorescent protein (GFP) fusion to the cortical marker Discs Large (Dlg-GFP) or myosin II regulatory light chain (Spaghetti squash in *Drosophila*; Sqh-GFP) as a cortical marker to measure polar extension during mitosis (where “polar extension” refers to displacement of the cortex at the poles, without regard to the underlying mechanism). In contrast to the equal polar extension observed in symmetrically dividing cells (Rosenblatt *et al.*, 2004; Hickson *et al.*, 2006), neuroblast cortical extension is highly asymmetric (Figure 1, B and C). The neuroblast apical cortex (associated with the larger daughter cell that retains the neuroblast fate) extends significantly during anaphase, whereas the basal cortex (associated with the smaller differentiated cell) undergoes very little extension (Figure 1, B and D, and Supplemental Movie S1). On average, the apical pole extends over three-fold more than the basal pole (2.0 vs. 0.6 μm). Thus polar extension in neuroblasts in intact tissue is highly asymmetric.

To ensure that asymmetric cortical extension was not a consequence of physical constraints imposed by the surrounding tissue, we measured cortical extension in cultured embryonic neuroblasts. These cells are dissociated from their surrounding tissue but continue to undergo asymmetric divisions (Siegrist and Doe, 2006). We found that cultured embryonic neuroblasts underwent asymmetric polar extension similar to their counterparts in the larval brain (Figure 1D and Supplemental Movie S2). We conclude that neuroblast asymmetric extension is an intrinsic property of the asymmetric cell division. Thus anaphase cortical extension differs significantly between symmetrically dividing cells and asymmetrically dividing neuroblasts. Whereas symmetrically dividing cells expand equally at both poles, neuroblast asymmetric divisions preferentially expand at the pole that becomes the larger daughter cell.

Asymmetric cortical extension is not caused solely by membrane synthesis

Neuroblast asymmetric cortical extension could occur by the preferential creation of new membrane at the apical surface. In this model the total surface area of the two daughter cells should be significantly larger than the surface area of the neuroblast before asymmetric cortical extension. To test this model, we measured the total surface area of the cell using three-dimensional reconstruction as a function of the cell cycle. We observed that the total surface area of the two daughter cells at the completion of cytokinesis is only ~10% larger than the surface area of the metaphase neuroblast (Figure 1E). In contrast, the difference in surface area of the two daughter cells is much more extreme (Figure 1F). For example, the surface area of a representative neuroblast at metaphase was 923 μm^2 , and upon completion of cytokinesis the resulting neuroblast and GMC had surface areas of 828 and 163 μm^2 , respectively. Thus preferential membrane synthesis at the apical cortex is not sufficient to explain the asymmetric cortical extension that we observe, although it could contribute to the effect. Note

that our measurements do not rule out a role for biased membrane flux.

Asymmetric cortical extension is independent of astral microtubules

To identify the cellular components responsible for cell-intrinsic asymmetric polar extension, we first focused on a possible role for the mitotic spindle, as it controls many of the morphological changes that occur during mitosis (von Dassow, 2009). At the poles, astral microtubules contact the cortex and could be responsible for controlling the asymmetric polar extension observed in neuroblasts. For example, asymmetric growth of the apical spindle could “push” the apical cortex outward. We examined whether astral microtubules are required for the difference in polar extension seen at the apical and basal neuroblast cortex by examining *sas4* mutants, which lack astral microtubules (Basto *et al.*, 2006). As shown in Figure 2A, the cortical extension in *sas4* occurs predominantly at the apical pole, similar to wild-type neuroblasts. We conclude that astral microtubules are not required for asymmetric cortical extension.

Cortical extension occurs at the onset of apical myosin II depletion

We examined the localization of the cortical factor myosin II to determine whether it could be important for polar extension in neuroblasts. In symmetrically dividing cells, myosin II is uniformly cortical in prophase but becomes confined to the equatorial region by late anaphase (Rosenblatt *et al.*, 2004). Neuroblasts exhibit a similar pattern of myosin II localization, except that myosin II is retained at the basal cortex during anaphase along with Anillin and Pav, which are normally restricted to the furrowing region (Cabernard *et al.*, 2010). In examining the localization of myosin II, we noticed a striking correlation with the loss of Sqh-GFP signal at the apical cortex and the onset of cortical extension, whereas myosin II remained at the basal cortex where extension was limited (Figure 2, B–D). Preferential apical clearing is not observed for the control proteins Dlg-GFP and Moe-RFP (Supplemental Figure S2). Furthermore, asymmetric cortical extension is not dependent on Sqh overexpression, as it also occurs in *sqh* mutants expressing Sqh-GFP (Figure 2C). The retention of myosin II on the basal cortex, along with the limited cortical extension at this location, prompted us to hypothesize that the basal domain containing furrow components inhibits cortical extension, limiting cortical extension to the apical cortex.

Asymmetric cortical extension requires a G-protein/Partner of Inscuteable–regulated basal furrow domain

As the onset of cortical extension is highly correlated with the loss of myosin II, we hypothesized that the “basal furrow domain” containing myosin II, Anillin, and Pavarotti, might be responsible for the preferential extension of the apical cortex. This domain is not regulated by the spindle but is instead controlled by cortical polarity factors such as Partner of Inscuteable (Pins; Cabernard *et al.*, 2010). We examined cortical extension in symmetrically dividing *pins* mutants to determine how loss of the basal myosin domain influences extension. We confirmed that neuroblasts lacking Pins fail to form the basal myosin domain (Cabernard *et al.*, 2010; Figure 2E). We find that these neuroblasts extend equally at both the apical and basal poles during anaphase (Figure 2, F and G). We conclude that Pins, which is required for the basal furrow domain, is also required for asymmetric polar extension. The known role of Pins in regulating asymmetric daughter cell size (Yu *et al.*, 2000) suggests that these three processes—the basal furrow domain, asymmetric polar elongation, and daughter cell size asymmetry—are tightly coupled.

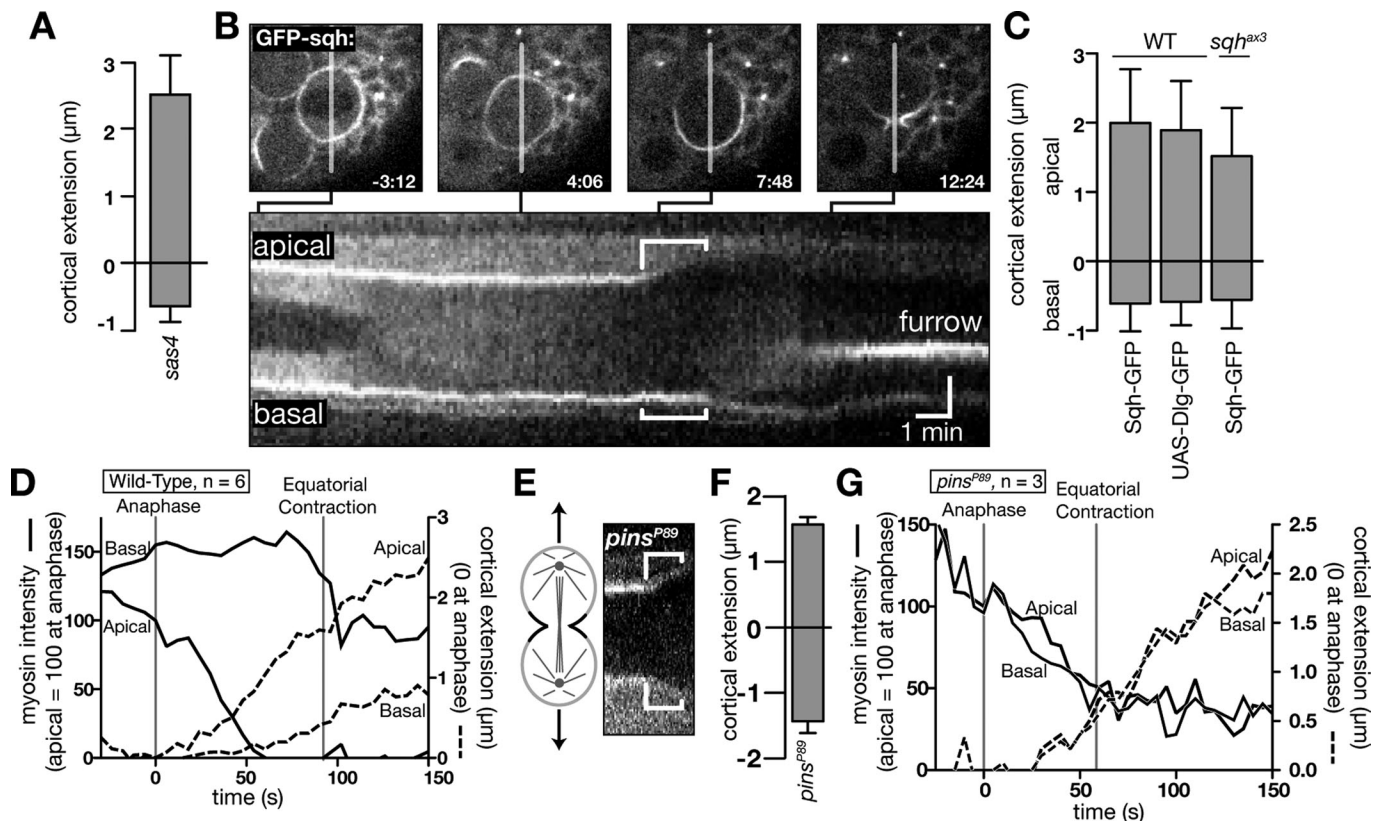


FIGURE 2: Basal myosin II is required for asymmetric polar elongation. (A) Mean anaphase polar extension in *sas4* mutant neuroblasts. Error bars, 1 SD. (B) Kymograph of myosin II (Sqh-GFP) in a wild-type larval neuroblast imaged at 6-s intervals. Top, selected frames with time relative to nuclear envelope breakdown. A line marks the section of the frame used for the kymograph. Anaphase cortical extension is denoted by brackets in the kymograph. (C) Quantification of apical and basal cortical extension in *sqh^{ax3}*; *sqh*-GFP (larval neuroblasts), *sqh*-GFP, and *worniu*-Gal4; UAS-Dlg-GFP neuroblasts. Error bars, 1 SD. Anaphase onset was determined using spindle (Jupiter-cherry) or chromosome (His2A-mRFP) markers. (D) Time dependence of cortical myosin signal and cortical position for wild-type neuroblasts. Dashed lines indicate the cortical position at each pole relative to the position at anaphase start. Solid lines indicate the intensity at each pole relative to the apical cortical intensity at anaphase start (determined as in C). Equatorial contraction indicates the time point at which the initiation of furrow ingression was observed. (E) Kymograph of Sqh-GFP in *pins^{P89}* mutant neuroblasts. Brackets denote polar extension during anaphase. (F) Quantification of anaphase cortical extension in *pins* mutant neuroblasts. Error bars, 1 SD. (G) Time dependence of cortical myosin signal and cortical position for *pins^{P89}* mutant neuroblasts. Annotations as in D.

To further test the relationship between the basal furrow domain, asymmetric cortical elongation, and daughter cell size, we examined another genetic background in which neuroblasts divide symmetrically. Overexpression of the Pins-binding G α i protein induces a high rate of symmetric divisions (Yu *et al.*, 2000; Schaefer *et al.*, 2001; Nipper *et al.*, 2007). We imaged larval brain neuroblasts expressing G α i under the control of the neuroblast-specific *worniu*-GAL4 driver to determine whether elevated G α i levels concurrently alter the basal furrow domain and cortical extension prior to symmetric cell division. We found that the basal furrow domain failed to form in approximately half the neuroblasts, and neuroblasts lacking the domain expanded equally at both poles during anaphase (Figure 3, A–C and E, and Supplemental Movie S3). To determine whether the effect on the basal furrow domain and cortical extension is due to a greater pool of activated G α i, we examined cortical extension in neuroblasts expressing the “activated” Q205L G α i mutant (UAS-G α iQ205L driven with *worniu*-GAL4), which does not bind Pins or the G β \gamma subunits (Schaefer *et al.*, 2001). These neuroblasts divided normally, suggesting that the ability to bind Pins is required for G α i-induced symmetric cell division (Figure 3, C and D, and Supplemental Movie S4). Thus heterotrimeric G-proteins and

Pins are required for the basal furrow domain and asymmetric polar extension and daughter cell size.

Asymmetric cortical extension requires alignment of the spindle with the basal furrow domain

We examined cortical extension in *mud* mutants to examine the consequence of decoupling the spindle-directed and basal furrows on polar extension. Mud orients the spindle with the polarity axis, and loss of *mud* function leads to randomization of spindle position (Guan *et al.*, 2000; Bowman *et al.*, 2006; Izumi *et al.*, 2006; Siller *et al.*, 2006). In *mud* mutant neuroblasts with misaligned spindles, structures resembling polar bodies form at the basal furrow domain prior to equatorial contraction from the spindle-directed furrow (Cabernard *et al.*, 2010). We examined cortical extension in this context to determine whether the basal furrow domain must be aligned with the spindle for asymmetric cortical extension. We found that although the cortex opposite the basal furrow domain begins to expand, subsequent spindle-induced equatorial furrowing overcomes this extension and ultimately leads to symmetric extension at the spindle poles and daughter cell size (Figure 3F). Thus coupling of both spindle-independent and

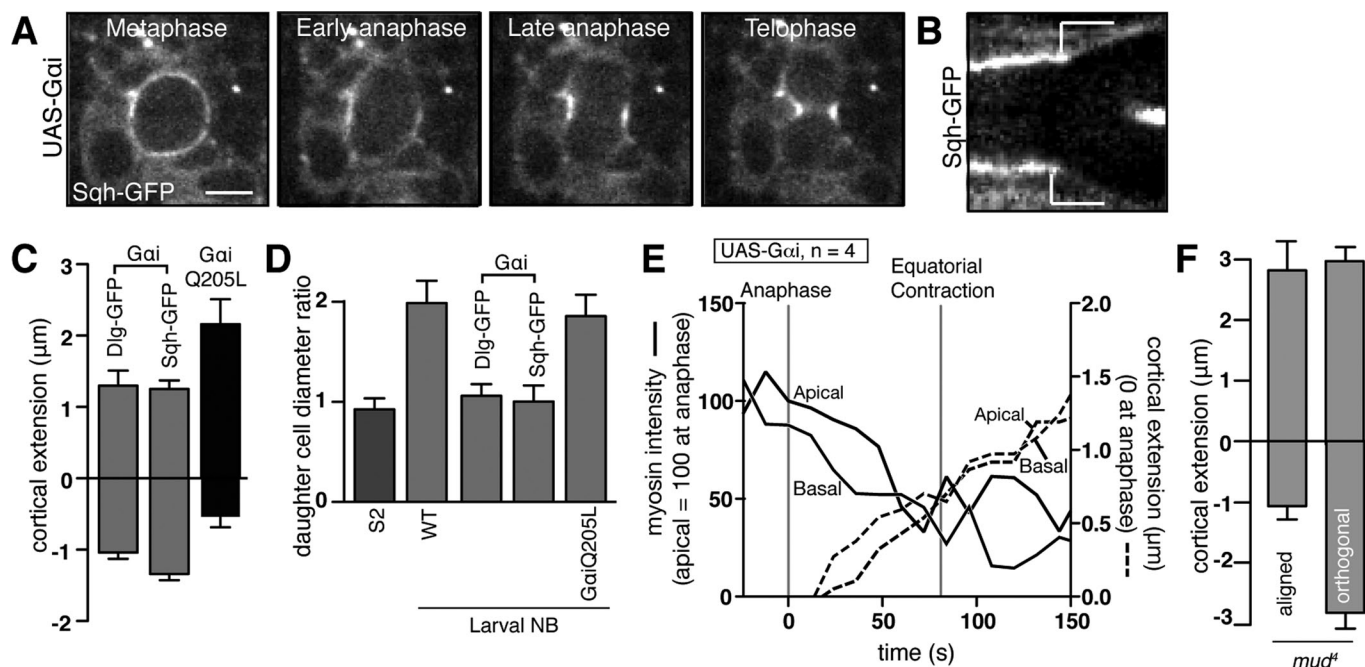


FIGURE 3: G-protein signaling regulates the basal furrow domain. (A) Myosin II (Sqh-GFP) localization in larval brain neuroblasts expressing *Gαi* using *worniu-Gal4*; *UAS-Gαi*. Images shown were taken 12 s apart. Scale bar, 5 μm. (B) Kymograph of Sqh-GFP signal across the poles from movie in A. Cortical extension is marked by the white lines. (C) Mean polar elongation for neuroblasts expressing *Gαi* or *Gαi* Q205L, a constitutively active variant that does not bind Gβγ or Pins. Cortical extension for *Gαi* is shown with two different cortical markers (Sqh-GFP or Dlg-GFP). Error bars, 1 SD. (D) Comparison of daughter cell size ratio for various cell types examined here. For asymmetrically dividing cells, this ratio was determined by dividing the diameter of the apical cell by the diameter of the basal cell. (E) Time dependence of cortical myosin signal and cortical position for neuroblasts expressing *Gαi*. Dashed lines indicate the cortical extension at each pole, whereas solid lines indicate the normalized intensity at each pole (as in Figure 2D). (F) Anaphase cortical extension for *mud^Δ* mutants where the spindle was aligned with basal myosin domain or orthogonal to it.

spindle-dependent contractile pathways is required for asymmetric polar extension.

Asymmetric cortical extension does not require spindle-directed equatorial contraction

Our results suggest that the basal furrow domain is required for asymmetric cortical extension. The basal domain could bias cortical extension toward the apical pole in at least two possible ways. The basal domain could increase the rigidity of the basal cortex, thereby focusing cortical extension induced by spindle-induced equatorial contraction to the apical cortex. Alternatively, the basal domain itself could be responsible for extension at the apical cortex. To distinguish between these models, we treated neuroblasts with Colcemid to depolymerize microtubules, thereby inhibiting formation of the spindle and the spindle-directed furrow (Brinkley *et al.*, 1967; Cabernard *et al.*, 2010). To allow these neuroblasts treated with Colcemid to pass through the spindle checkpoint without a spindle, the Colcemid treatments were done in a *rod* background (Basto *et al.*, 2000; Chan *et al.*, 2000; Savoian *et al.*, 2000; Cabernard *et al.*, 2010). The presence of the basal furrow domain alone is sufficient for asymmetric cortical extension, as anaphase cortical extension is indistinguishable from wild-type neuroblasts (Figure 4, A and B). Thus we conclude that spindle-directed equatorial contraction is not required to produce asymmetric cortical extension.

DISCUSSION

Morphological changes that occur during mitosis involve both positive and negative signals that emanate from the mitotic spindle. For

example, in *C. elegans* the first division yields unequal-sized daughter cells (Albertson, 1984; Keating and White, 1998; Glotzer, 2004). At the metaphase-to-anaphase transition the spindle midzone is displaced toward the posterior end of the zygote and is believed to positively influence furrow formations, whereas astral microtubules repress furrowing at the poles (Albertson, 1984; Keating and White, 1998; Glotzer, 2004). Thus asymmetric positioning of the spindle before anaphase ultimately leads to a displaced furrow and unequal-sized daughter cells. In neuroblasts, however, the spindle is symmetrically positioned before anaphase, suggesting that other mechanisms are responsible for neuroblast asymmetric division (Cai *et al.*, 2003; Kaltschmidt *et al.*, 2000; Siller and Doe, 2009). Recently, components normally restricted to the cleavage furrow were found to be localized to the neuroblast basal cortex (Cabernard *et al.*, 2010). We found that this “basal furrow domain” restricts cortical extension that normally happens at both poles during the division of many small, symmetrically dividing cells. This restriction of cortical extension limits the size of the future basal cell while allowing the future apical cell to expand during anaphase (Figure 4C).

The contribution of asymmetric cortical extension to daughter cell size explains why *asl* and *cnn* mutants, which lack astral microtubules, divide asymmetrically although they have a symmetric spindle (Bonaccorsi *et al.*, 1998; Basto *et al.*, 2006). We argue that the asymmetric spindle poles (large apical pole, small basal pole) observed in wild-type neuroblasts late in the cell cycle (Fuse *et al.*, 2003) are a consequence, not a cause, of asymmetric cortical extension. Lack of basal cortical extension prevents the spindle from growing at this pole, whereas it is free to grow at the apical pole. In

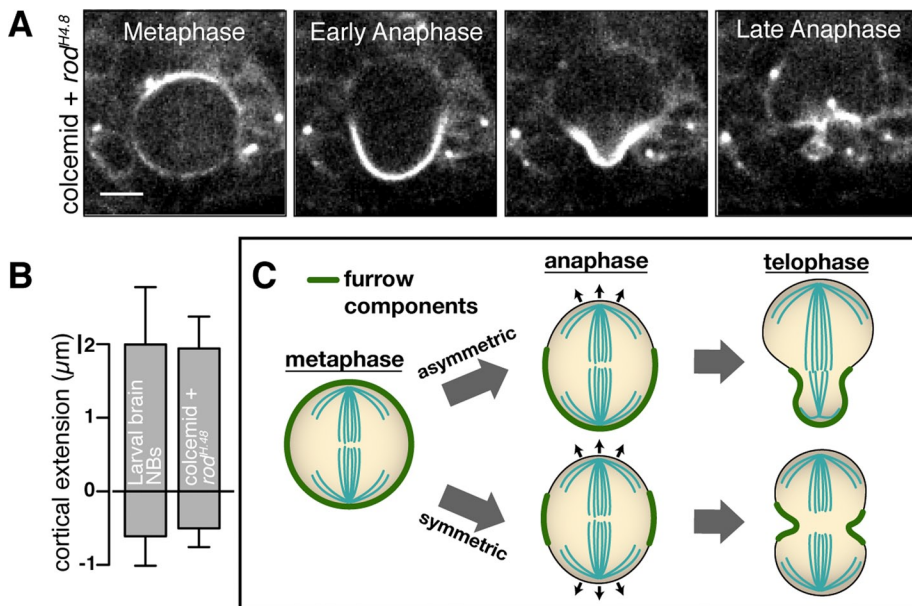


FIGURE 4: Asymmetric cortical extension does not require spindle-induced equatorial contraction. (A) Sequence of a Colcemid treated *rod* mutant neuroblast expressing Jupiter-cherry to ensure the absence of a spindle. Scale bar, 10 μm . (B) Quantification of cortical extension in Colcemid-treated *rod* mutants. Error bars, 1 SD. (C) Model for the role of the basal furrow domain in daughter cell size asymmetry.

fact, in S2 cells where cortical extension has been inhibited at both poles by RNA interference knockdown of Rho kinase (Hickson *et al.*, 2006), spindles become bent and distorted during anaphase. We propose that asymmetric cortical extension in neuroblasts channels spindle growth into the apical pole, resulting in a highly symmetric spindle. The basal myosin domain does not passively halt the extension at the basal pole by acting against the forces produced by spindle-directed equatorial contraction, but instead is directly responsible for asymmetric cortical extension. Nevertheless, the basal and spindle-directed myosin domains must work together for asymmetric cortical extension to occur (Figure 3F).

How is the basal furrow domain regulated? Several components that function upstream of myosin II in the canonical furrow (e.g., Pav and Anillin) are also present in the basal furrow domain (Cabernard *et al.*, 2010). Previous work showed that the polarity protein Pins is required to establish a basal myosin domain, whereas *Gai* mutants do not perturb the domain (Cabernard *et al.*, 2010). A linear model in which $G\beta\gamma$ promotes the basal myosin domain but is inhibited by *Gai*, which is in turn inhibited by Pins, is consistent with these observations, and our data provide further support. Pins binds *Gai* such that it is dissociated from $G\beta\gamma$, so in cells that lack Pins, heterotrimeric G-protein complex assembly is favored and both $G\alpha$ and $G\beta\gamma$ activity are reduced. As *Gai* mutants have normal basal furrow domains (Cabernard *et al.*, 2010), *Gai* is not required for the basal furrow domain, suggesting that $G\beta\gamma$ is the relevant furrow domain regulator. The *Gai* overexpression results presented here are also consistent with this model. We observed that expression of *Gai*, which would inactivate any free $G\beta\gamma$, leads to loss of the basal furrow domain. However, expression of constitutively active *Gai*, which does not bind $G\beta\gamma$ but does bind downstream effectors, has no effect on the furrow domain. Thus we propose that $G\beta\gamma$ activity is essential for establishing the basal myosin domain, asymmetric cortical extension, and unequal daughter cell size. Future work will be directed toward rigorously testing this model and identifying the link between polarity proteins and myosin.

MATERIALS AND METHODS

Fly stocks

All mutant chromosomes were balanced over CyO actin:GFP, TM3 actin:GFP, Ser, e, or TM6B Tb. We used Oregon R as wild type and the following mutant chromosomes and fly strains: *Sqh:GFP* (Royou *et al.*, 2002), *worGal4* (Albertson and Doe, 2003), *UAS-Cherry:Jupiter* (Cabernard and Doe, 2009), *worGal4* and *UAS-Dlg:eGFP* (Koh *et al.*, 1999), *UAS-Gai* and *UAS-GaiQ205L* (Schaefer *et al.*, 2001; kindly provided by J.A. Knoblich), *mod⁴* (Guan *et al.*, 2000), *pins^{P89}* (Yu *et al.*, 2000), *FRT82B Sas-4^M* (Basto *et al.*, 2006), *rod^{H4.8}* (Basto *et al.*, 2000), *UAS-His2A-mRFP* (Emery *et al.*, 2005), *UAS-moe-RFP* (Schwabe *et al.*, 2005), and *UAS-Jupiter-GFP* (Karpova *et al.*, 2006). All crosses were performed at 25°C except *UAS-Gai* crosses, which were performed at room temperature and transferred to 30°C approximately 8 h prior to imaging.

Cell culture

Drosophila S2 cells were maintained at room temperature in Schneider's medium (Sigma-Aldrich, St. Louis, MO) supplemented with 10% fetal bovine serum (Sigma-Aldrich). Cells were transiently transfected with pMT-Zeus-Cherry with Effectene (Qiagen, Valencia, CA) according to the manufacturer's instructions, and expression was induced with 0.5 M copper sulfate 20 h prior to imaging. Immediately before imaging, cells were resuspended in Schneider's medium supplemented with 10% fetal bovine serum.

In vitro neuroblast culture

Primary cell cultures were made from embryos aged 6–8 h as previously described (Grosskortenhaus *et al.*, 2005). They were then prepared for live imaging by resuspension in Chan and Gehrings balanced saline solution supplemented with 2% FBS.

Live imaging

Second or third larval brains were prepared for imaging as previously described (Siller *et al.*, 2005). Five to nine Z steps were collected at 1- to 2- μm intervals every 6–12 s. Live imaging was performed using a spinning disk confocal microscope equipped with a Hamamatsu EM-CCD camera (Hamamatsu, Japan) using a 63 \times 1.4 numerical aperture oil immersion objective. Pixel intensity measurements were performed using ImageJ (National Institutes of Health, Bethesda, MD). A linear photobleaching correction was added, using the intensity of a region outside the cell as a reference. Cortical extension was determined by measuring the position of the cell edges at the poles with a section from the middle of the cell as determined by examination of sections throughout the cell.

Colcemid treatment was performed on the strain +; *worGal4*, *UAS-Cherry:Jupiter*, *Sqh:GFP*; *rod^{H4.8}* (Cabernard *et al.*, 2010), using a final concentration of 0.1 $\mu\text{m}/\text{ml}$, with live imaging beginning immediately after treatment.

Three-dimensional reconstructions for surface area analysis were done using the BoneJ plugin for ImageJ with a sampling value of 2 (Doube *et al.*, 2010).

ACKNOWLEDGMENTS

We thank Brad Nolen and members of the Prehoda lab for helpful comments. C.Q.D. is an Investigator of the Howard Hughes Medical Institute. This work was supported by National Institutes of Health Grant 068032 (K.E.P.) and the American Heart Association (C.C.).

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