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The microtubule cross-linker Feo controls the midzone stability, motor composition, and elongation of the anaphase B spindle in Drosophila embryos

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ABSTRACT Chromosome segregation during anaphase depends on chromosome-to-pole motility and pole-to-pole separation. We propose that in Drosophila embryos, the latter process (anaphase B) depends on a persistent kinesin-5–generated interpolar (ip) microtubule (MT) sliding filament mechanism that “engages” to push apart the spindle poles when poleward flux is turned off. Here we investigated the contribution of the midzonal, antiparallel MT-cross-linking nonmotor MAP, Feo, to this “slide-and-flux-or-elongate” mechanism. Whereas Feo homologues in other systems enhance the midzone localization of the MT- MT cross-linking motors kinesin-4, -5 and -6, the midzone localization of these motors is respectively enhanced, reduced, and unaffected by Feo. Strikingly, kinesin-5 localizes all along ipMTs of the anaphase B spindle in the presence of Feo, including at the midzone, but the antibody-induced dissociation of Feo increases kinesin-5 association with the midzone, which becomes abnormally narrow, leading to impaired anaphase B and incomplete chromosome segregation. Thus, although Feo and kinesin-5 both preferentially cross-link MTs into antiparallel polarity patterns, kinesin-5 cannot substitute for loss of Feo function. We propose that Feo controls the organization, stability, and motor composition of antiparallel ipMTs at the midzone, thereby facilitating the kinesin-5–driven sliding filament mechanism underlying proper anaphase B spindle elongation and chromosome segregation.

INTRODUCTION
During mitosis, chromosomes are separated by chromosome-to-pole motility (anaphase A) and spindle elongation (anaphase B; Gadde and Heald, 2004; Gerdes et al., 2010; Goshima and Scholey, 2010; Walczak et al., 2010; Zheng, 2010; McIntosh et al., 2012; Helmke et al., 2013; Civelekoglu-Scholey and Cimini, 2014). The Drosophila embryo mitotic spindle assembles by a centrosome-directed mechanism that can be augmented by chromatin- and augmin-directed microtubule (MT) assembly (Hayward et al., 2014) and then segregates chromosomes using both anaphase A and B (Brust-Mascher and Scholey, 2002; Maddox et al., 2002). Whereas anaphase A depends on a combined kinesin-13–dependent pacman-flux mechanism (Rogers et al., 2004), we propose that anaphase B depends on a slide-and-flux-or-elongate mechanism in which the persistent sliding apart of interpolar microtubules (ipMTs) driven by kinesin-5 “engages” to push apart the spindle poles when poleward flux is turned off (Cole et al., 1994; Kashina et al., 1996; Brust-Mascher and Scholey, 2002, 2011; Brust-Mascher et al., 2004, 2009; Cheerambathur et al., 2007; van den Wildenberg et al., 2008; Acar et al., 2013; Wang et al., 2013; Scholey et al., 2014). Thus, in pre–anaphase B spindles, the outward sliding of ipMTs is balanced by the kinesin-13 (KLP10A)-catalyzed depolymerization of their minus ends at the poles, producing poleward flux (Rogers et al., 2004), and the spindle maintains a steady length. After cyclin B degradation, however, the MT minus end–capping protein patronin (Goodwin and Vale, 2010) counteracts KLP10A activity at spindle poles to turn off ipMT minus end depolymerization so poleward flux ceases and the outwardly sliding ipMTs can now elongate the spindle.
(Wang et al., 2013). At the same time, ipMT plus ends display net growth and recruit MT-MT cross-linkers to assemble a more robust midzone, where the sliding motors act (Cheerambathur et al., 2007). However, although the patronin-mediated suppression of KLP10A activity is sufficient to convert MT poleward flux to anaphase B spindle elongation, the function of this spindle midzone reorganization is unclear (Wang et al., 2013).

The spindle midzone is a structure that assembles and self-organizes during anaphase and consists of a dense network of overlapping antiparallel ipMTs containing multiple motor proteins, regulatory molecules, and nonmotor microtubule-associated proteins (MAPs; Glotzer, 2009; Peterman and Scholey, 2009; Mitchison et al., 2013). The assembly, organization, and function of this structure depend critically on members of the Ase1p (also known as PRC1 and MAP65) family of homodimeric nonmotor MAPs, which bind and diffuse along single MTs, cross-linking antiparallel MTs into bundles and accumulating cooperatively between pairs of overlapping antiparallel MTs. Like kinesin-5 motors, members of the Ase1p protein family display a significant preference for cross-linking MTs into antiparallel versus parallel orientations (Kapitein et al., 2008; van den Wildenberg et al., 2008), and this preference for antiparallel MT binding is unique among nonmotor MAPs (Duellberg et al., 2013). Members of the Ase1p/PRC1 family specifically accumulate at the spindle midzone from anaphase to telophase and interact with motor proteins and other MAPs to facilitate their association with the midzone. Such motors and MAPs include kinesin-4 (Kurasawa et al., 2004) and CLASP1 (Liu et al., 2009) in mammalian cells and kinesin-5 (Khmelnitski et al., 2009) and kinesin-6 (Fu et al., 2009) in yeast. Thus Ase1p/PRC1 proteins play essential roles in spindle midzone organization and cytokinesis. Although the Ase1p-mediated recruitment of sliding motors has been shown to be crucial for anaphase B spindle elongation in yeast (Fu et al., 2009; Khmelnitski et al., 2009), in metazoan systems the interactions between Ase1p/PRC1 homologues and sliding motors and their roles in anaphase B spindle elongation are less well understood.

In this study, we took advantage of the Drosophila syncytial embryo system, which progresses through multiple mitoses without intervening cytokinesis, to investigate specifically the mitotic function of the Ase1p/PRC1 homologue, Feo. Drosophila contains two members of the Ase1p family, namely Feo, which plays essential roles in mitosis, and Sofe, which does not (and is not considered further here; Verni et al., 2004). In previous studies, based on the analysis of mutants and RNA interference in fixed spermatocytes, neuroblasts, and cultured S2 cells, it was found that Feo is required to organize a robust central spindle in telophase, to target Polo kinase to the midzone (D’Avino et al., 2007), and for contractile ring assembly and cytokinesis but it is dispensable for anaphase B (Verni et al., 2004). Here we reevaluated the mitotic roles of Feo in living embryo spindles, focusing on its possible role in anaphase B. We find that Feo controls the stability and motor composition of the spindle midzone and is required for proper anaphase B spindle elongation and chromosome segregation in Drosophila embryos.

RESULTS

Feo accumulates at the anaphase B and telophase spindle midzone downstream of cyclin B degradation and cdk1 inactivation

We created transgenic flies expressing Feo-green fluorescent protein (GFP) or Feo-mCherry and used spinning disk confocal microscopy to investigate its dynamics throughout the cell cycle (Figure 1 and Supplemental Video S1). Feo-GFP localized to the cytoplasm during interphase and only faintly to mitotic spindles from prophase to metaphase, but increasing amounts of Feo-GFP accumulated at the spindle midzone from anaphase B onset, appearing as a bright equatorial band of increasing intensity perpendicular to the spindle long axis (Figure 1, A and B, and Supplemental Video S1). To determine whether this accumulation is cell cycle regulated, we injected the cdk1 inhibitor purvalanol into metaphase embryos expressing Feo-mCherry and the chromosome-associated kinesin-4, KLP3A-GFP, which normally translocates to the midzone during anaphase B (Kwon et al., 2004). In wild-type embryo metaphase spindles, Feo localized faintly throughout the spindle, whereas KLP3A localized to chromosomes and weakly to spindles, as expected. After cdk1 inhibition, however, a premature metaphase spindle midzone is formed (Hu et al., 2012), and both Feo-mCherry and KLP3A-GFP accumulated on this structure, mimicking their midzone localization during anaphase B and telophase in wild-type spindles (Figure 1C and Supplemental Video S2). Thus, in wild-type spindles, cdk1 inactivation promotes the assembly of a spindle midzone containing both Feo and KLP3A. We further confirmed that Feo accumulation occurs downstream of cyclin B degradation, because the injection of excess nondegradable cyclin B into embryos containing anaphase B spindles caused Feo-GFP to rapidly dissociate from the spindle midzone (Figure 1D and Supplemental Video S3). Thus Feo accumulation at the anaphase/telophase midzone is regulated by cyclin B degradation and cdk1 inactivation, concordant with previous work showing that the localization and function of human PRC1 and yeast Ase1p are also regulated by cell cycle-dependent phosphorylation (Zhu et al., 2006; Fu et al., 2009; Khmelnitski et al., 2009).

Feo is essential for proper anaphase B spindle elongation and midzone organization

To examine the role of Feo in embryonic mitosis, we generated an antibody against Feo, which recognized a single band around 80 kDa by Western blotting against Drosophila embryonic extracts (Figure 1E), and tested its ability to inhibit Feo by microinjecting it into Drosophila syncytial embryos expressing fluorescent proteins. As expected (Brust-Mascher et al., 2009; Wang et al., 2013), injection of the anti-Feo antibody displaced Feo-GFP from spindles, forming fluorescent cytoplasmic immunoprecipitates, and caused a gradient of phenotypes with respect to spindle dynamics (Figure 1F and Supplemental Figure S1). To confirm the specificity of Feo inhibition, we also injected the anti-Feo antibody into flies expressing other GFP-tagged proteins, including kinesins-4, -5, -6, -8, -13, and -14, the MT tip tracker EB1, and the mitotic kinase polo, and observed that none were sequestered into cytoplasmic immunoprecipitates like those formed from Feo-GFP (unpublished data). This suggests that the anti-Feo antibody does not cross-react with and immunoprecipitate these other motors and microtubule-binding proteins.

To investigate the effect of Feo inhibition on spindle dynamics and chromosome segregation, we microinjected the anti-Feo antibody into flies expressing GFP-tubulin and red fluorescent protein (RFP)–histone (Figure 2). The anti-Feo–mediated inhibition of Feo function produced a striking gradient of mitotic defects from anaphase to telophase. In spindles proximal to the injection site, anaphase B spindle elongation was initiated but not sustained (e.g., s4 and s5 in Figure 2, A and B, right in E, and Supplemental Video S5), whereas distal spindles displayed normal anaphase B spindle elongation (e.g., s1 and s2 in Figure 2, A and B). In both cases, the midzone became progressively narrower until it disappeared in telophase (Figure 2, D and E).
FIGURE 1: Cell cycle–dependent localization of Feo to the spindle midzone. (A) Feo accumulates at the spindle midzone from anaphase B through telophase. Living embryos expressing Feo-GFP (green in merged image) were injected with tubulin (red). 0 s, anaphase B onset. See Supplemental Video S1. Line scans show relative intensity along interpolar (ipMT) bundles. To calculate the relative fluorescence intensity, the minimum and maximum fluorescence intensity in all images were set to 0 and 1, respectively. Tubulin line scan is red, and Feo line scan is light green. To illustrate better the midzone accumulation of Feo at anaphase B, we also calculated the relative fluorescence intensity of Feo by setting the maximum fluorescence intensity during anaphase B to 1.0 (dark green line scan). (B) Example plot of fluorescence intensity of Feo-GFP vs. time at the spindle midzone normalized to average metaphase fluorescence. Blue, maximum intensity; black, average intensity calculated in a box surrounding the spindle midzone; 0 s, anaphase B onset. (C) The cdk1 inhibition by injection of the cdk1 inhibitor purvalanol caused KLP3A-GFP (green in merged) and Feo-mCherry (red in merged) to accumulate prematurely at the spindle midzone. Control was injected with dimethyl sulfoxide. Tubulin is blue in merged. See Supplemental Video S2. (D) Excess nondegradable cyclin B injection inhibited Feo accumulation at the anaphase B spindle midzone. See Supplemental Video S3. (E) Western blot of affinity-purified
These effects of Feo inhibition were specific to anaphase/telophase, because ~95% of anti-Feo–injected spindles proceeded to anaphase normally (mistimed antibody microinjection during the previous anaphase/telophase led to aberrant metaphase spindle assembly in only 5.5% of spindles analyzed [219 spindles in 27 embryos]). Overall our results agree with those obtained in other Drosophila.
At the end of telophase, the distances between centrosomes and between paired nuclei were significantly decreased after Feo inhibition (Figure 3C), suggesting that incomplete pole-pole separation during anaphase B is correlated with the chromosome segregation defects observed after Feo inhibition. Previously we showed that patronin inhibition causes defects in anaphase B spindle elongation, which also led to defects in chromosome segregation (Wang et al., 2013). Both studies underscore the importance of anaphase B spindle elongation for chromosome segregation.

**Feo antagonizes the association of kinesin-5 with the anaphase B spindle midzone**

We tested the idea that the aforementioned defects in mitosis might reflect the effects of loss of Feo function on the localization of the three primary spindle midzone motors—kinesin-5 (KLP61F), kinesin-4 (KLP3A), and kinesin-6 (PavKLP). Previous work suggested that these motors play critical roles in anaphase B and cytokinesis. Kinesin-5 serves as the primary antiparallel MT-MT sliding motor driving poleward flux and spindle elongation (Sharp et al., 2000; Brust-Mascher et al., 2004, 2009); kinesin-4 somehow contributes to the down-regulation of poleward flux, leading to a corresponding enhancement of spindle elongation (Brust-Mascher et al., 2004; Kwon et al., 2004), and kinesin-6 bundles MTs of the late anaphase/telophase midzone and is required for cleavage furrow assembly and cytokinesis (Adams et al., 1998).

In the case of kinesin-5, as in previous immuno–electron microscopy and live-cell imaging (Sharp et al., 1999; Cheerambathur et al., 2008), in KLP61F-GFP–rescued flies (Cheerambathur et al., 2008), we observed that kinesin-5 colocalizes with tubulin throughout the spindle, is more concentrated around the poles during metaphase and anaphase, and displays a characteristic but previously unexplained narrow dip in fluorescence intensity at the equator during late anaphase B and telophase (Figure 4, A and B). We emphasize, however, that KLP61F is localized all along ipMT bundles through anaphase B, including ipMTs crossing the midzone, where this decrease in intensity occurs and therefore it is appropriately positioned to slide apart antiparallel MTs to drive poleward flux and pole–pole separation in wild-type embryo spindles (Figure 4, A and B). We observed that the dip in KLP61F fluorescence intensity during anaphase B and telophase coincides with the site of Feo accumulation in the center of the midzone (Figure 4A and Supplemental Video S6). We also note that KLP61F is less concentrated at the equator during metaphase, but this reflects steric exclusion by chromosomes at the equator (Figure 4B, bottom).

Surprisingly, after Feo inhibition in KLP61F-GFP rescued flies (Cheerambathur et al., 2008), the distribution of KLP61F is significantly altered from anaphase B on (Figure 4C and Supplemental Videos S7 and S8). KLP61F concentrates at the spindle midzone, and the characteristic dip does not form. Feo inhibition does not affect the distribution of KLP61F during metaphase, suggesting that Feo specifically regulates KLP61F during anaphase B by limiting the amount of KLP61F associated with the center of the antiparallel overlap zone. To explore this further, we compared the KLP61F-to-tubulin ratio in untreated and Feo-inhibited embryos. In the inhibited embryo spindles, this ratio is not affected during metaphase, but it is higher at the midzone from anaphase B on (Figure 4C). Quantification of the KLP61F-to-tubulin ratio at the spindle midzone during metaphase and anaphase B shows that in untreated embryos, the anaphase B ratio is similar to the metaphase ratio, but after Feo inhibition the ratio during anaphase B is significantly increased (Figure 4D). Ase1p and kinesin-5 have a similar preference for cross-linking MTs into antiparallel versus parallel overlap zone.

**Feo is essential for proper chromosome segregation in Drosophila embryos**

After Feo inhibition, we also observed chromosome segregation defects characterized by the presence of chromosome bridges (arrowheads) between the two daughter nuclei even after nuclear envelope re-formation (region I). Away from the injection site, daughter nuclei appear to be properly segregated, but midbodies were not present between them (region II). The arrow shows the injection site. Bar, 10 μm. (B) Chromosome-to-pole distance vs. time in untreated and anti-Feo–injected embryos. 0 s, anaphase A onset. Averages of ≥6 spindles. (C) Distance between the inner edges of daughter nuclei (inner) and between centrosomes (centrosomes) at the end of telophase in untreated and anti-Feo–injected embryos. Twenty-six spindles in two embryos were measured for control (untreated), and 34 spindles in three embryos were measured after Feo inhibition.
Feo controls the anaphase B midzone

**FIGURE 4:** Influence of Feo on the spindle midzone localization of the MT sliding motor, kinesin-5. (A) Dynamics of KLP61F (kinesin-5)-GFP and Feo-mCherry during Drosophila embryo mitosis. In the merged images and line scans, Feo is shown in red and KLP61F in green. See Supplemental Video S6. (B) KLP61F localizes all along spindle MTs. Top, time series of a single plane through the center of the spindle; KLP61F-GFP decorates all spindle MTs, the intensity in the center of the midzone decreases during anaphase B, and a gap is clearly visible at telophase. Middle, Z-series through a spindle at the onset and the second half of anaphase B. KLP61F clearly labels MTs from pole to pole. Bottom, comparison of KLP61F and chromosome localization. Time 0, anaphase B onset. Bar, 5 μm. (C) Effect of Feo inhibition on the dynamics of KLP61F in Drosophila embryo mitosis. The midzone association of kinesin-5 is enhanced during anaphase B after Feo inhibition. Bottom, ratio of KLP61F:tubulin fluorescence intensity. The maximum ratio in all images is set to 1.0 to normalize the data. Repeated in >20 embryos. See Supplemental Videos S7 and S8. (D) KLP61F/tubulin ratio at the spindle midzone during metaphase and anaphase B in untreated and Feo-inhibited embryos. The KLP61F/tubulin ratio during anaphase was normalized with the ratio during metaphase. The number of spindles/embryos analyzed is shown at the bottom. p < 2.2e-16. Bars, 5 μm.

Feo is required for kinesin-4 accumulation to the spindle midzone during anaphase B and telophase

We next examined the effect of loss of Feo function on the localization of kinesin-4. In mammalian parallel orientations, and therefore we propose that in wild-type embryos, Feo and KLP61F competitively bind to antiparallel MTs, and thus KLP61F replaces antibody-dissociated Feo in bridging antiparallel MTs at the center of the spindle midzone when Feo is absent.

However, kinesin-5 cannot functionally substitute for the loss of Feo from the spindle midzone because, as shown, defects in anaphase B spindle elongation clearly occur (Figure 2). Thus it is possible that in wild-type embryos, Feo accumulation at the spindle midzone limits the number of KLP61F motors that associate with midzonal antiparallel ipMTs to somehow optimize their MT-MT sliding activity and also helps to organize antiparallel MTs into an ordered structure that facilitates kinesin-5–driven anaphase B spindle elongation. This effect of Feo in regulating kinesin-5 localization differs from the situation in Saccharomyces cerevisiae, in which Ase1p appears to positively regulate the spindle localization of the sliding motors. In the latter case, the kinesin-5, CIN8, must bind to Ase1p in order to associate with the midzone and drive spindle elongation (Khmelinskii et al., 2009).

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cells, PRC1 is required for localization of kinesin-4, and direct interaction between Feo and kinesin-4 has been reported in Drosophila S2 cells (D’Avino et al., 2007). To test whether Feo is required to target the Drosophila kinesin-4 KLP3A to the spindle midzone, we compared the dynamics of both proteins (Figure 5A) and examined how the loss of Feo function affects the dynamics of KLP3A-GFP during mitosis (Figure 5B). Consistent with previous immunostaining (Kwon et al., 2004), in wild-type embryos, KLP3A-GFP concentrated on chromosomes and only weakly localized to spindle MTs during metaphase, but during anaphase and telophase, KLP3A-GFP dissociated from the chromosomes and concentrated on the spindle midzone, displaying strong colocalization with Feo (Figure 5A and Supplemental Videos S9 and S10). After Feo inhibition, however, KLP3A localized normally to metaphase chromosomes but did not translocate to the midzone, instead remaining associated with chromosomes throughout anaphase B and telophase (Figure 5B). This suggests that Feo targets KLP3A to the anaphase spindle midzone, in agreement with previous data showing that the midzone localizations of PRC1 and KIF4 are interdependent in mammalian cells (Kurasawa et al., 2004; Zhu and Jiang, 2005). The midzone defects resulting from Feo inhibition (Figure 2) are not solely due to failures in KLP3A targeting, because in KLP3A-inhibited embryos, midzones splay out into broader structures than those in controls (Kwon et al., 2004), whereas in Feo-inhibited embryos, midzones become narrower. Thus the midzone defects observed after Feo inhibition are a direct result of loss of Feo-dependent midzonal MT bundling.

**Feo inhibition does not influence kinesin-6 localization to the spindle midzone**

Feo does not seem to influence the association of PavKLP with the spindle midzone; in wild-type embryos, PavKLP accumulated to a narrower region of the midzone than Feo (Figure 5C and Supplemental Videos S11 and S12) and retained

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**FIGURE 5:** Influence of Feo on the spindle midzone localization of the MT motors, kinesin-4 and kinesin-6. (A) Dynamics of KLP3A (kinesin-4)-GFP and Feo-mCherry during Drosophila embryo mitosis. In the merged images and line scans, Feo is shown in red, KLP3A in green, and tubulin in blue. See Supplemental Videos S9 and S10. (B) Effect of Feo inhibition on the dynamics of KLP3A during anaphase B and telophase (compare with A). Repeated in 12 embryos. (C) Dynamics of Pav (kinesin-6)-GFP and Feo-mCherry during mitosis. In the merged images and line scans, Feo is shown in red, Pav in green, and tubulin in blue. See Supplemental Videos S11 and S12. (D) Effect of Feo inhibition on the dynamics of Pav-GFP during anaphase B and telophase (compare with C). Repeated in six embryos. Bars, 5 μm. 0 s, anaphase B onset.
this localization after Feo inhibition (Figure 5D). This is consistent with previous work in S2 cells (D’Avino et al., 2007), but differs from results showing that loss of Feo activity in spindles of other Drosophila cell types leads to the mislocalization of this kinesin-6 during telophase and to defects in cytokinesis (Verni et al., 2004). In addition, in Schizosaccharomyces pombe, Ase1p recruits the kinesin-6, klp9p, to the spindle midzone, where this motor drives anaphase B spindle elongation (Fu et al., 2009).

**Feo inhibition has no effect on the redistribution of ipMT plus ends at anaphase B onset**

Ase1p/PRC1 family members and their associated proteins (e.g., kinesin-4, Bieling et al., 2010; CLASP1, Liu et al., 2009) have been reported to be involved in regulating MT plus end dynamics. We tested whether Feo has a role in the redistribution of MT plus ends observed at anaphase B onset (Cheerambathur et al., 2007) by injecting Feo antibody into flies expressing EB1-GFP, a marker of growing MT plus ends (Figure 6A). After Feo inhibition, spindles were narrower, but EB1-GFP still redistributed to the spindle midzone at anaphase B onset (Figure 6, B vs. C). Given that the midzone localization of kinesin-4 was abolished after Feo inhibition, our observation that Feo inhibition has no effect on EB1 concentration at anaphase B onset suggests that neither Feo nor kinesin-4 at the midzone is responsible for the redistribution of MT plus ends at anaphase B onset.

**Feo and kinesin-5 display different patterns of localization along antiparallel MTs in the anaphase B spindle midzone**

Our functional data suggest that kinesin-5 and Feo cooperate at the spindle midzone to produce the robust anaphase B spindle elongation and faithful chromosome segregation observed in wild-type embryo spindles. To estimate their relative distribution on antiparallel ipMTs at the midzone, we evaluated the maximal size of the zone of overlapping MTs by tracking EB1-GFP in living transgenic embryos and measuring the excursions of growing MT plus ends that move across the equator from one half-spindle to the opposite half-spindle. These measurements suggest a midzone length of ~3–4 μm (Figures 6A and 7A). During anaphase B, Feo is localized in a tight band at the center of the overlap zone marked by EB1 (Figures 6B and 7A), where it could organize adjacent antiparallel MTs into bundles with optimal spacing. Kinesin-5 is distributed mainly along midzonal antiparallel ipMTs adjacent to this Feo zone (Figure 7A), where it could slide apart optimally spaced MTs, supporting the hypothesis that Feo partially restricts the association of kinesin-5 with the center of the midzone, possibly by steric inhibition as a result of its being assembled into a dense, multimeric midzone matrix (Schuyler et al., 2003; Kapitein et al., 2008; Hutterer et al., 2009). An accurate picture of the distribution of overlapping MTs and associated proteins at the midzone would require high-resolution studies (e.g., multiple-label immuno–electron microscopy; Sharp et al., 1999), but our light microscopy data (Figure 7A) support the model shown in the cartoon in Figure 7B.

**DISCUSSION**

The spindle midzone appears to be a conserved structure overall, containing common molecular components that associate with its dense network of antiparallel MTs (Glotzer, 2009; Peterman and Scholey, 2009; Mitchison et al., 2013). However, current and previous studies suggest that these components display system-specific variations in their interactions, targeting, and functions, conforming to the general idea that the mitotic spindle consists of conserved...
molecules and biochemical modules that are deployed in a system-specific, combinatorial manner (Goshima and Scholey, 2010). The role of the key Ase1p family member, PRC1/Feo, appears to differ in different systems. In this study, we focused on its role in Drosophila syncytial embryo mitosis. Previous work on Feo in Drosophila was based on fixed immunostaining data and focused on the role of Feo in organizing the spindle midzone during telophase and cytokinesis. Here we characterized the real-time dynamics of Feo and its interactions with key MT-MT cross-linking motors and uncovered an essential role for Feo in anaphase B spindle elongation and chromosome segregation. Using the Drosophila syncytial embryo, in which rounds of mitosis occur very rapidly without intervening cytokinesis, our work revealed the roles of Feo during late mitosis. Thus, in contrast to previous studies (Verni et al., 2004), we show that Feo inhibition caused significant defects in anaphase B spindle elongation and midzone assembly, which lead to defects in chromosome segregation and telophase midbody structure.

To our knowledge, this is the first time that an Ase1p family member has been shown to contribute to anaphase B spindle elongation in metazoan systems. Of interest, the mechanism by which Feo contributes to anaphase B spindle elongation seems to differ from the role of Ase1p in yeast. Whereas Ase1p recruits sliding motors (kinesin-5 or kinesin-6) to the spindle midzone to initiate anaphase B spindle elongation in yeast (Khmelinskii et al., 2009), in Drosophila embryos, Feo appears to restrict the association of the sliding motor, kinesin-5, to the spindle midzone. Drosophila embryo spindle midzones contain a single type of kinesin-5, kinesin-4, and kinesin-6, whereas S. cerevisiae spindles use two distinct types of kinesin-5 motors, but no kinesin-4 or kinesin-6 is present (Saunders and Hoyt, 1992; Straight et al., 1998). Whereas kinesin-5 appears to be the primary motor at the spindle midzone driving anaphase B in Drosophila embryo and S. cerevisiae mitotic spindles (Straight et al., 1998; Brust-Mascher et al., 2009), kinesin-6 fulfills this role in S. pombe (Fu et al., 2009). We find that the midzone association of Feo is not required for the association of kinesin-6 with the Drosophila embryo mitotic spindle midzone, consistent with results in mammalian cells but different from the situation in S. pombe, where Ase1p is required for kinesin-6 localization (Kurasawa et al., 2004; Fu et al., 2009).

In many systems, including Drosophila S2 cells, members of the Ase1p family have been shown to directly interact with kinesin-4 during late mitosis (Kurasawa et al., 2004; Zhu and Jiang, 2005; D’Avino et al., 2007). In the present study, we show that Feo is required for the translocation of Drosophila kinesin-4, KLP3A, from chromosomes to the spindle midzone during anaphase B and telophase and that accumulation and colocalization of Feo and KLP3A to the anaphase B spindle midzone are regulated by cyclin B degradation and cdk1 inactivation, as shown in mammalian cells (Kurasawa et al., 2004; Zhu et al., 2006). In S2 cells, the interaction between Drosophila Feo and KLP3A depends on their phosphorylation states (D’Avino et al., 2007), and it has also been shown that PRC1 is directly phosphorylated by cdk1 in early mitosis and that its dephosphorylation may increase its affinity for MTs (Zhu and Jiang, 2005; Zhu et al., 2006; Subramanian et al., 2013). It is therefore possible that in Drosophila embryos, in response to cyclin B degradation and cdk1 inactivation, dephosphorylated Feo increases its binding affinity to MTs and KLP3A and thus recruits KLP3A to the antiparallel overlap to organize the anaphase B spindle midzone. Of note, both Feo and KLP3A inhibition destabilize IPMTs and cause defects in anaphase B spindle elongation; however, KLP3A inhibition causes disorganized “splayed” MTs with an increased spindle width-to-length ratio during anaphase B (Kwon et al., 2004), whereas Feo inhibition causes a narrower and denser spindle midzone, suggesting that in addition to recruiting KLP3A, Feo has independent roles in the organization of the anaphase B spindle midzone.

The present study illuminates the key role of Feo in assembling and organizing the spindle midzone. Our results suggest that Feo localization to the spindle midzone from early anaphase on is required to allow the key MT sliding motor, kinesin-5, to drive normal anaphase B spindle elongation. Consequently, after its antibody-induced dissociation, only a narrow midzone can assemble, leading to a reduced rate and extent of spindle elongation. However, while Feo is required for kinesin-5–driven anaphase B spindle elongation, it does not appear to be required for kinesin-5–driven poleward flux during pre–anaphase B because it does not start to accumulate on the spindle midzone until anaphase B onset. Members of the kinesin-5 family of motors and the Ase1p family of nonmotor MAPs both localize to antiparallel IPMTs of the anaphase B spindle midzone in fly embryo spindles, and they share a similar preference for cross-linking MTs into antiparallel versus parallel polarity patterns in vitro. It is striking that kinesin-5 localization to the embryo spindle midzone increases when Feo is depleted, yet the additional kinesin-5 cannot substitute for the decrease in Feo function despite sharing its antiparallel MT cross-linking activity. This suggests that Feo-dependent assembly and organization of the midzone is required for proper anaphase B spindle elongation, probably by organizing antiparallel

FIGURE 7: Model for the functional interaction between Feo and kinesin-5 on antiparallel ipMTs at the spindle midzone during anaphase B in Drosophila embryos. (A) Diagrammatic representation of the relative distributions of Feo, EB1, and KLP61F along the anaphase B spindle. Center (0), spindle equator. Vertical axis shows relative fluorescence intensity (1 is maximum for each protein). (B) Model of the anaphase B spindle midzone showing the spatial relationship between Feo, EB1, and KLP61F. Feo preferentially bundles antiparallel ipMTs and also localizes weakly to parallel MTs. EB1, a marker of growing MT plus ends, is concentrated at the midzone. KLP61F cross-links parallel and antiparallel MTs and slides apart antiparallel ipMTs. The kinesin-5–driven antiparallel ipMT sliding requires Feo-mediated midzone organization to produce normal anaphase B spindle elongation.
ipMTs into properly spaced networks containing the appropriate number of kinesin-5 motors consistent with optimal outward sliding of the ipMTs, which drive pole–pole separation (Lansky et al., 2015). We propose that these cell cycle–regulated Feo and kinesin-5–dependent midzone-associated processes (Cheerambathur et al., 2007; Bastos et al., 2014) cooperate with events at the spindle pole, where, after cyclin B degradation, patronin-mediated suppression of kinesin-13 activity converts MT poleward flux into anaphase B spindle elongation (Wang et al., 2013). Finally, we note that the frequency of anaphase B errors that accompany aging in humans has been associated with an increased risk of cancer (Ford, 2013), suggesting that these processes may be medically relevant.

**MATERIALS AND METHODS**

**Generation of transgenic flies**

For Feo-mCherry transgenic flies, a full-length Feo (fascetto) cDNA was subcloned into an mCherry coding sequence in the pWR-pUbq vector (provided by N. Brown, University of Cambridge, Cambridge, United Kingdom) downstream of the polyubiquitin promoter (Wang et al., 2013), and the resulting transgenic flies with a single insertion in the third chromosome were generated by BestGene (Chino Hills, CA). For KLP3A-GFP transgenic flies, a cDNA encoding full-length KLP3A (provided by Michael Goldberg, Cornell University, Ithaca, NY; Kwon et al., 2004) was subcloned with a GFP coding sequence into the pWR-pUbq vector in a similar way, and the resulting transgenic flies with a single insertion in the X chromosome were generated by BestGene.

**Drosophila stocks and embryo preparation**

All flies were maintained at 25°C. Transgenic flies coexpressing GFP-tubulin and RFP-histone were generated as described (Wang et al., 2013). Flies with an EB1-GFP (Cheerambathur et al., 2007), KLP61F-GFP (Cheerambathur et al., 2008), PavKLP-GFP (Sommi et al., 2011), or Feo-GFP transgene (Wang et al., 2013) were expressed as previously described. Flies expressing Polo-GFP were kindly provided by Claudio Sunkel (Institute for Molecular and Cell Biology, University of Porto, Porto, Portugal). Flies coexpressing Feo-mCherry and KLP3A-GFP or KLP61F-GFP or PavKLP-GFP or EB1-GFP were generated by standard crossing. Embryos were prepared as described (Wang et al., 2013).

**Protein purification, antibody generation, and microinjection**

Embryos were premicroinjected with rhodamine-labeled tubulin or HiLyte Fluor 647–labeled tubulin (Cytoskeleton, Denver, CO) when necessary to visualize tubulin. GST-cyclin B was purified as described (Cheerambathur et al., 2007). Anti-Feo antibody was affinity purified with GST-tagged Feo N-terminal fragments (amino acids 1–250 and 1–480) from a rabbit anti-Feo serum against histidine-tagged full-length Feo protein. To purify wild-type full-length Feo protein, Feo cDNA was cloned into pFastBac HTc vector (Invitrogen, Carlsbad, CA) and then expressed in baculovirus system and purified with Ni-nitriloacetic acid agarose (Qiagen, Valencia, CA). All injected proteins were dialyzed in phosphate-buffered saline (PBS) and concentrated as protein concentration times the estimated injection volume/injection site volume.

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