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CENP-32 is required to maintain centrosomal dominance in bipolar spindle assembly

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Abbreviations used: CENP-32, centromere protein 32; CG-NAP, centrosome and Golgi–localized, PKN-associated protein; Dgt2-9, dim γ-tubulin 2-9; PCM, pericentriolar material; γ-TuRC, γ-tubulin ring complex.

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ABSTRACT

Centrosomes nucleate spindle formation, direct spindle pole positioning, and are important for proper chromosome segregation during mitosis in most animal cells. We previously reported that centromere protein 32 (CENP-32) is required for centrosome association with spindle poles during metaphase. In this study, we show that CENP-32 depletion seems to release centrosomes from bipolar spindles whose assembly they had previously initiated. Remarkably, the resulting anastral spindles function normally, aligning the chromosomes to a metaphase plate and entering anaphase without detectable interference from the free centrosomes, which appear to behave as free asters in these cells. The free asters, which contain reduced but significant levels of CDK5RAP2, show weak interactions with spindle microtubules but do not seem to make productive attachments to kinetochores. Thus CENP-32 appears to be required for centrosomes to integrate into a fully functional spindle that not only nucleates astral microtubules, but also is able to nucleate and bind to kinetochore and central spindle microtubules. Additional data suggest that NuMA tethers microtubules at the anastral spindle poles and that augmin is required for centrosome detachment after CENP-32 depletion, possibly due to an imbalance of forces within the spindle.

INTRODUCTION

Accurate spindle formation is essential for the proper segregation of chromosomes during cell division. The bipolar mitotic spindle is composed of polar microtubules organized with their minus ends anchored to centrosomes and plus ends projecting outward to form the spindle and asters. In Saccharomyces cerevisiae, the spindle pole body, which is functionally equivalent to the centrosome, serves as the microtubule-organizing center and contributes to bipolar spindle formation in the mitotic nucleus (Byers and Goetsch, 1974).

In animal cells, spindles usually form as the result of the separation and activation of centrosomes during the G2 phase of the cell cycle (Kimura et al., 1997; Wakefield et al., 2000; Jackman et al., 2003; Dossay et al., 2005; Hachet et al., 2007; Portier et al., 2007).

The pericentriolar material (PCM) is created by pericentrin/kendrin and centrosome and Golgi–localized, PKN-associated protein (CG-NAP), which are stabilized by Kizuna (Oshimori et al., 2006, 2009). CG-NAP and kendrin help to organize microtubule nucleation in the mammalian centrosome by anchoring the γ-tubulin ring complex (γ-TuRC) together with several other proteins (Dietenberg et al., 1998; Hung et al., 2000; Takahashi et al., 2000, 2002; Casenghi et al., 2003). Centrosomal localization of CG-NAP and Kizuna is regulated by Cep72, which is also involved in γTuRC recruitment to the centrosome (Oshimori et al., 2009).

Centrosomes nucleate the assembly of highly dynamic microtubules that shift between states of polymerization and depolymerization, effectively probing the cytoplasm and enabling them to be captured by kinetochores on mitotic chromosomes (Mitchison and Kirschner, 1985; Kirschner and Mitchison, 1986). A subset of microtubule plus ends from opposite poles also takes part in antiparallel interactions in the midzone and stabilizes the central spindle.
Correct centrosome positioning and microtubule nucleation in the mitotic cell are thus extremely important for establishing and maintaining normal spindle shape and structure (O’Connell and Wang, 2000; Lüders and Stearns, 2007; Dunsch et al., 2012; Laan et al., 2012; Kiomitsu and Cheeseman, 2013).

Despite the obvious importance of centrosomes in nucleating formation of bipolar spindles, spindles can form in the absence of centrosomes (e.g., in oocytes) and after centrosome inactivation (Uzbekov et al., 1995; Khodjakov et al., 2000). Centrosome-independent mechanisms for mitotic spindle formation appear to involve the Ran-GTP-mediated polymerization of microtubules near chromosomes (Karsenti et al., 1984; Heald et al., 1996; Wilde and Zheng, 1999; Wilde et al., 2001; Kalab et al., 2002; O’Connell and Khodjakov, 2007), capture of the microtubule plus ends by kinetochores, and clustering of the microtubule minus ends by microtubule motor proteins to form the spindle poles (Sawin et al., 1992; Merdes et al., 1996; Walczak et al., 1998; Sharp et al., 2000). Ran is also required for the correct cortical localization of nuclear mitotic apparatus protein (NuMA; Wilde and Zheng, 1999; Zhang et al., 1999; Wiese et al., 2001). NuMA interacts with cytoplasmic dynein and dynactin, and they transport it toward microtubule minus ends (Merdes et al., 1996, 2000). These motor proteins accumulate at the pericentrosomal region and play a role in generating pulling forces on astral microtubules (Gagliò et al., 1995; Haren et al., 2009).

De novo microtubule generation can occur within mitotic spindles of vertebrate cells in a centrosome-independent manner (Goshima et al., 2008). The dim γ-tubulin 2-9 (Dgt2-9) octamer complex, or augmin, was reported to nucleate microtubule formation within mitotic spindles in a γ-tubulin-dependent process (Uehara et al., 2009). Thus there may exist more than one centrosome-independent spindle formation mechanism that maintains acentrosomal spindles in mitosis across a wide range of species, including plants (Masoud et al., 2013).

Previously our mitotic chromosome proteomics analysis predicted mitotic chromosome association of centromere protein 32 (CENP-32)/C9orf119. Streptavidin-binding peptide–tagged CENP-32 accumulated on mitotic spindles and at kinetochores (Supplementary Figure S1A; Ohta et al., 2010). Our proteomics-based bioinformatics MCCP (multi-classifier combinatorial proteomics) analysis predicted that CENP-32 function is related to CLASP1 and 2, whose association with microtubules, kinetochores, and centrosomes is well known. CLASPs are involved in multiple microtubule-dependent processes in mitosis, including kinetochore fiber dynamics, mitotic spindle assembly, and the stabilization of microtubule plus ends near the cortex (Maiato et al., 2005; Mimori-Kiyosue et al., 2005; Miller et al., 2009; Reis et al., 2009; Patel et al., 2012; Bird et al., 2013). Recent work identified CLASP1 as being required for proper positioning of the mitotic spindle (Samora et al., 2011; Espiritu et al., 2012; Kiomitsu and Cheeseman, 2012).

We previously reported that centrosomes detach from mitotic spindles in CENP-32–depleted cells (Ohta et al., 2010). A second pericentriolar protein, CDK5RAP2, is also required to maintain centrosome engagement and cohesion (Fong et al., 2008; Barrera et al., 2010). CDK5RAP2 was reported to be recruited to centrosomes by dynein and be required to maintain attachment of centrosomes to spindle poles in mitosis (Barr et al., 2010; Jia et al., 2013).

It is unclear how the link between centrosomes and the mitotic spindle poles is maintained. Here we study whether CENP-32–depleted spindles, which apparently do not form a stable connection with centrosomes, could facilitate normal chromosome arrangements to a metaphase plate and segregation during anaphase. We find that CENP-32 is necessary for CG-NAP recruitment to centrosomes and for maintaining interactions between the mitotic spindle and centrosomes. Moreover, augmin is necessary for the centrosome detachment and subsequent anastral mitosis that occurs in the absence of CENP-32. Our findings demonstrate that CENP-32 is required for centrosome integration into the spindle and that its absence causes abnormal microtubule organization, possibly by interfering with CG-NAP localization and/or function.

RESULTS

CENP-32 depletion leads to centrosome mislocalization from spindle poles

We previously reported that CENP-32 depletion resulted in the detachment of centrosomes from spindle poles during metaphase (Figure 1B; Ohta et al., 2010). To investigate the timing of this centrosomal dissociation, we used immunofluorescence staining against γ-tubulin and α-tubulin to visualize centrosomes and spindle microtubules, respectively (Figure 1A). Compared to cells treated with control small interfering RNA (siRNA), centrosomes of CENP-32 siRNA cells exhibited premature movement away from the nuclear surface before nuclear envelope breakdown (Figure 1A; prophase). Of interest, subsequent bipolar spindle formation was not abolished upon CENP-32 depletion, even though centrosomal detachment persisted after anaphase onset (Figure 1A; white arrowheads). Overexpression of enhanced green fluorescent protein (EGFP);CENP-32 reduced the percentage of centrosomal dissociation after CENP-32 siRNA treatment (Figure 1C). This rescue suggests that the centrosomal dissociation was caused by CENP-32 depletion.

Time-lapse microscopy was also used to analyze spindle formation in CENP-32–depleted U2OS cells overexpressing GFP-α-tubulin and histone H2B:monomeric red fluorescent protein fusion proteins (Figure 1D). In CENP-32–depleted cells, detached centrosomes could be distinguished from the remainder of the bipolar spindle as isolated spots of microtubule staining (Figure 1D; arrow heads). Centrosomes appeared to dissociate from the mitotic spindle immediately after bipolar spindle formation in CENP-32 siRNA cells (Figure 1D). Despite this detachment, CENP-32 depletion did not have a dramatic effect on spindle morphology or chromosome alignment during prometaphase (Figure 1D).

These results reveal that CENP-32 is required for centrosome tethering at spindle poles and suggest that the protein is required for centrosomes to act as dominant polarity determinants of spindle pole formation.

Dissociated centrosomes have an altered protein composition

It was shown previously that dissociated centrosomes in CENP-32–depleted cells maintain γ-TuRC complex and pericentrin localization and that those proteins are absent from spindle poles after centrosome detachment (Ohta et al., 2010). Here we confirmed that γ-tubulin staining was present on detached centrosomes in CENP-32–depleted cells (Figure 1), albeit at decreased levels after their disassociation from the spindle poles (quantification in Figure 2I). In both control and CENP-32–depleted cells, weak γ-tubulin signals were also found on mitotic spindles. Double immunofluorescence labeling against the γ-TuRC complex and various centrosomal proteins was then used to assess the composition of the detached centrosomes in the depleted cells.

Aurora A is required for centrosome maturation and bipolar spindle formation (Hannak et al., 2001). Compared to control cells, CENP-32 siRNA cells exhibited a weaker (but still positive) staining of Aurora A on the detached centrosomes and a stronger diffuse...
signal on spindle microtubules (Figure 2, A and E). In contrast, NuMA did not accumulate on the dissociated centrosomes but did continue to localize at the poles of the bipolar spindles (Figure 2, B and F). CG-NAP signals were also substantially decreased on detached centrosomes in CENP-32-depleted cells (Figure 2, C and G). It was previously reported that lack of CDK5RAP2 is able to cause centrosomal release from the spindle (Barr et al., 2010). However, CDK5RAP2 could still be detected on the detached centrosomes, albeit at decreased levels, in CENP-32–depleted cells (Figure 2, D and H).

FIGURE 1: (A) Mitotic centrosome position in control and CENP-32 siRNA–transfected cells at prophase, prometaphase (PM), metaphase (or prometaphase with metaphase plate: PM + MP), and anaphase. Cells were immunolabeled with γ-tubulin (green) and α-tubulin (red) antibodies. DAPI staining was used to detect DNA (blue). (B) Quantification of phenotypes in CENP-32–depleted cells with metaphase plate. (C) Quantification of phenotypes in CENP-32–depleted cells with overexpression of EGFP (top) or EGFP:CENP32 (bottom). (D) Time-lapse analysis of control and CENP-32 siRNA–transfected cells. The number shows time (minutes) from nuclear envelope breakdown. White arrowheads indicate centrosomes dissociated from the mitotic spindle in A and D.
FIGURE 2: Pericentrosomal protein localization (A, Aurora A; B, NuMA; C, CG-NAP; and D, CDK5RAP2) in control and CENP-32–depleted cells with DAPI and α-tubulin or γ-tubulin. (E–I) Quantification of immunofluorescence intensities of control and CENP-32 siRNA–transfected cells. **p < 0.05, ***p < 0.01.
CG-NAP knockdown does not affect centrosome dissociation from mitotic spindles

We next tested the hypothesis that CG-NAP mislocalization caused by CENP-32 depletion might serve as a trigger for centrosome dissociation from mitotic spindles. U2OS cells were depleted of CG-NAP by two different siRNA transient transfections (Supplemental Figure S1B). Immunofluorescence labeling against α-tubulin and pericentrin revealed that CG-NAP loss could occasionally result in a centrosome-dissociation phenotype in mitotic cells (Supplemental Figure S1C). However, this phenotype was observed at a much lower frequency than that found in CENP-32–depleted cells (Supplemental Figure S1, D and E; ~6% in C-GAP-depleted cells vs. ~33% in CENP-32–depleted cells). CG-NAP depletion also resulted in premature centrosome movement away from the nuclear surface in ~50% of prophase cells, similar to that observed in cells depleted of CENP-32 (unpublished data).

Thus, although loss of CG-NAP can explain certain aspects of the CENP-32–depletion phenotype, it cannot fully explain the dissociation of centrosomes from the spindle or their loss of polarity dominance.

Microtubule regrowth assay with CENP-32–depleted cells

Next we examined the spindle formation ability of dissociated centrosomes, using a microtubule regrowth assay. Microtubules in control and CENP-32–depleted cells were disrupted by cold treatment for 30 min (Figure 3, A and D). In the control, microtubule initiation from centrosomes was observed by 3 min after the transfer to prewarmed medium (Figure 3B). After 25 min, 74% of cells showed bipolar spindles (Figure 3, C and G). In CENP-32–depleted cells, we observed significantly lower levels of microtubule polymerization from centrosomes by 3 min after the transfer to prewarmed medium (Figure 3E). Furthermore, 17% of cells showed anastral spindle after 25 min (Figure 3, F and G). These results suggest that the anastral spindle poles remain capable of nucleating microtubules after centrosome detachment, whereas nucleation by the detached centrosomes is significantly impaired.

The ultrastructure of dissociated centrosomes

Transmission electron microscopy was used to visualize the ultrastructure of dissociated centrosomes in CENP-32–depleted cells. Centrosomes in control and CENP-32–depleted cells contained centrioles with nine triplet microtubules assembled in a normal cartwheel structure (Figure 3, H and I). No major differences were observed in the structure of the centrioles or PCM surrounding dissociated centrosomes in comparison to normal, spindle pole–associated centrosomes (Figure 3J). However, we did observe an apparent decrease in the density of microtubules surrounding the dissociated centrosomes in CENP-32–depleted cells (Figure 3K). Furthermore, most microtubules associated with the detached centrosomes appeared to be shorter than those at control centrosomes. This observation is consistent with the impaired astral microtubule assembly observed in CENP-32–depleted cells in the microtubule regrowth assay.

Microtubule instability at spindle poles without centrosomes

We next examined the stability of the spindle microtubules in cells with dissociated centrosomes induced by CENP-32 depletion. We did this by using a cold-stability assay. Under the conditions used, only stable spindle microtubules associated with kinetochores (K-fibers) withstand the cold treatment at 4°C for 10 min. After the cold treatment, cells were immunostained for the kinetochore marker, CENP-C, and α-tubulin to detect spindle morphology.

CENP-32 siRNA cells exhibited two distinct phenotypes after cold treatment. The first, observed in 18% of CENP-32–depleted cells, consisted of fragmented acentrosomal spindle poles (Figure 4, A and B, arrowheads, and C, unfocused). The second phenotype, observed in 22% of CENP-32–depleted cells after cold treatment, consisted of intact focused anastral spindle poles. These spindles contained robust microtubule bundles (Figure 4, A and B, white arrows, and C, robust).

This experiment suggested that kinetochore–microtubule interactions were apparently stable after cold treatment in both control and CENP-32–depleted cells (Figure 4A). However, we were unable to observe any microtubule signal associated with the dissociated centrosomes in CENP-32–depleted cells, whereas a faint signal was seen in the cells exposed to control RNA interference (RNAi; Figure 4, A and B). Therefore the microtubules nucleated by the detached spindle poles apparently do not interact with kinetochores and also apparently interact more weakly with the detached centrosomes.

Taken together, the experiments in this and the preceding section reveal that CENP-32 is required for either the nucleation or stability of astral microtubules.

Augmin knockdown partially suppresses centrosome detachment in CENP-32–depleted cells

The augmin complex is required for the recruitment of the γ-TuRC to microtubules and promotes centrosome-independent microtubule formation (Uehara et al., 2009). To investigate the possibility that augmin is required for anastral spindle formation in CENP-32–depleted cells, we codepleted the augmin subunit Dgt6 using siRNA. As expected, spindles contained a reduced microtubule density after Dgt6 RNAi depletion (Figure 5A). Double staining for α-tubulin and pericentrin revealed that in single-target siRNA experiments, 33% of mitotic cells had more than one dissociated centrosome in CENP-32–depleted cells, whereas only 2% of mitotic cells exhibited this phenotype when Dgt6 levels were reduced (Figure 5B). We found that 5.6% of mitotic cells contained more than one dissociated centrosome in CENP-32 and Dgt6 double-depleted cells (Figure 5B). This result indicates that siRNA targeting of the augmin complex partly suppressed the centrosome-detachment phenotype in CENP-32–depleted cells. CENP-32 and Dgt6 double-depleted cells also exhibited a multipolar phenotype and prometaphase accumulation (Figure 5B; 37% of cells).

CENP-32 depletion causes a decrease in monopolar spindle formation accompanied by centrosome detachment

Bipolar spindles form as a result as balanced interactions between motor proteins, and imbalances in those forces can cause the spindles to collapse (Saunders and Hoyt, 1992; Saunders et al., 1997; Straight et al., 1998; Nazarova et al., 2013). A widely accepted approach used to look at the effect of disrupting this balance of forces involves the use of monastrol, an inhibitor of the bipolar kinesin-5 Eg5, during spindle assembly (Mayer et al., 1999). This inhibition results in efficient collapse of the spindle.

We found that although monopolar spindle formation still occurs with high efficiency in monastrol-treated, CENP-32–depleted cells, the process is less efficient than it is in control depleted cells (Figure 5C; 90% in control and 66% in CENP-32 depleted). The background of normal bipolar spindles observed after monastrol treatment was unchanged by CENP-32 depletion (Figure 5C; 10% in control and 13% in CENP-32 depleted).

Of interest, the decrease in monopolar spindles observed in the CENP-32–depleted cells was balanced primarily by an increase in bipolar, monoastral spindles (Figure 5C; not observed in controls,
FIGURE 3: Immunostaining of α-tubulin (red) and γ-tubulin (green) after microtubule regrowth for 0 (A, D), 3 (B, E), and (C, F) 25 min. (G) Quantification of reformation of mitotic spindles in this assay. Observation of centrosomes under a transmission electron microscope of centrioles in control (H) and CENP-32 siRNA (I) cells. The number indicates the number of the triplet microtubules. Scale bar, 500 nm. (J) Microtubules around centrosomes in control and (K) CENP-32 siRNA cells. Scale bar, 1 μm. Red lines indicate observed microtubules.
and that if NuMA is depleted in addition to CENP-32, the integrity of the anastral spindles might not be maintained.

Single siRNA treatment targeting NuMA depleted 75% of endogenous NuMA (Figure 6, A and B). Double depletion of NuMA and CENP-32 did not suppress the centrosome detachment seen in metaphase cells after CENP-32 depletion (Supplemental Figure S2A; CENP-32 depletion, 48%; NuMA plus CENP-32 double depletion, 54%). To investigate whether double-depleted cells are able to construct functional bipolar spindles and align chromosomes at a metaphase plate, we determined the frequency of metaphase cells in cultures after double depletion of NuMA and CENP-32 or single depletion of CENP-32 (Figure 6D). This analysis showed that double depletion of NuMA and CENP-32 caused a marked decrease in the frequency of cells with a metaphase plate after CENP-32 depletion (Figure 6C and Supplemental Figure S2B; CENP-32 depletion, 46.9%; NuMA plus CENP-32 double depletion, 21.4%).

CENP-32 is also important for spindle organization in chicken mitotic cells

We generated a CENP-32 deletion mutant in chicken DT40 cells (Supplemental Figure S3). Of interest, this mutation was not lethal to chicken cells but did induce a mild growth defect. Compared to wild-type cells (doubling time of 10.7 h), loss of CENP-32 caused the doubling time to increase to 14.5 h (Figure 7A). CENP-32 depletion also caused a small increase in the percentage of cells in the G2/M phase (19.2% in the CENP-32 knockout compared with 11.9% for wild-type cells) as measured by fluorescence-activated cell sorting (FACS) analysis (Figure 7B). However, CENP-32 depletion did not cause a significant increase in the percentage of cells phosphorylated at serine 10 in histone H3 (1.9% in the CENP-32 knockout and 1.7% in wild-type cells; Supplemental Figure S4) as measured by FACS analysis. The increase in G2/M cells may have been partly explained by the fact that CENP-32–knockout cells exhibited an increase in number of cells with multipolar spindles (from 3.5% in controls to 18.8%; Figure 7, C and D).

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**DISCUSSION**

Time-lapse microscopy analysis revealed that CENP-32 depletion resulted in centrosome dissociation from mitotic spindles during early prometaphase. After centrosome dissociation, the mitotic spindles remained active and promoted chromosome alignment to a metaphase plate. Moreover, these mitotic spindles without centrosomes ("anastral" spindles) were able to advance to anaphase. Remarkably, the detached centrosomes appeared to have lost the ability to act as dominant determinants of spindle polarity. Although they could still nucleate microtubules, the efficiency of microtubule
Augmin depletion dissolves the centrosome-dissociation phenotype. (A) Dgt6 or Dgt6- and CENP-32–depleted cells show reduced mitotic spindle microtubule density. The most frequently observed spindles in Dgt6- and CENP-32–depleted cells are indicated with α-tubulin (green), pericentrin (red), and DAPI (blue). (B) Quantification of phenotypes in CENP-32– and Dgt6-depleted cells (bottom table). (C) Quantification of phenotypes in monastrol-treated CENP-32–depleted cells (bottom table). (D) The most frequently observed spindles in monastrol-treated CENP-32–depleted cells are indicated with α-Tubulin, γ-Tubulin, and DAPI.
CG-NAP is important for centrosome maturation (Takahashi et al., 1999, 2002). We speculate that CENP-32 may promote centrosome maturation by controlling CG-NAP localization to centrosomes in early mitosis. Our data suggest that not only centrosome targeting of CG-NAP but also other structural changes resulting from CENP-32 depletion must contribute to centrosome dissociation from mitotic spindle poles.

Mitotic spindle poles without centrosomes are tethered by NuMA

The fact that chromosomes were still able to align at a metaphase plate after centrosome detachment in CENP-32–depleted cells suggests that the basic functionality of the anastral mitotic spindle is maintained even in the absence of centrosomes. Of importance, both a microtubule regrowth assay and electron microscope observations suggested that microtubule elongation from detached centrosomes is impaired in cells with anastral spindles. In contrast, a
our indirect immunofluorescence microscopy analysis showed CDK5RAP2 remained on disassociated centrosomes in CENP-32–depleted cells, albeit at lower levels (Figure 1D). This result suggests that CENP-32 might anchor centrosomes at mitotic spindle poles via a mechanism different from CDK5RAP2, although it is possible that the decrease in centrosome-associated CDKRAP2 levels after CENP-32 depletion is sufficient to promote centrosome detachment. Augmin knockdown was able to suppress this centrosomal-dissociation phenotype (Moutinho-Pereira et al., 2013). Moreover, double knockdown of CENP-32 and Dgt6 (augmin) partly suppressed the centrosome detachment phenotype seen after CENP-32 depletion. Augmin recruits γ-TuRC to microtubule walls and contributes to centrosome-independent microtubule generation (Goshima et al., 2008; Kamasaki et al., 2013). It was recently reported that augmin is involved in centrosome-independent mitotic spindle assembly in Drosophila S3 cells and plants (Ho et al., 2011; Hotta et al., 2012; Hashimoto, 2013; Moutinho-Pereira et al., 2013). One role of augmin is apparently to make the central portion of the spindle more robust. This could create a functional junction between the aster and the central region of the spindle. Spindle forces acting after loss of CENP-32 might cause a weakened junction to break, thereby releasing the centrosome and leaving behind an anastral spindle. In the absence of augmin, the less robust central spindle might be more flexible, so that the centrosome does not detach when the same forces act. Alternatively, γ-TuRC levels might potentially increase at centrosomes after augmin depletion. This could potentially increase their microtubule nucleation ability, thereby integrating them more robustly into the body of the spindle. However, note that under the conditions used here, Dgt6 knockdown did not cause an increase in γ-TuRC levels at spindle poles as measured by indirect immunofluorescence. It is also possible that the presence of augmin is somehow required in order for the forces that occur after depletion of CENP-32 to cause spindle detachment. The nature of those forces is unknown, but it has been shown that depletion of dynein subunits in Drosophila cells can result in centrosome detachment from spindles (Morales-Mulia and Scholey, 2005). Of note, those spindle poles became unfocused, and the spindles were significantly longer than wild type, phenotypes that are occasionally seen in the case of CENP-32 depletion. An alternative explanation is that some factor downstream of the large scaffolding protein CG-NAP is involved in linking centrosomes to the central region of the spindle.

That forces within the spindle cause the centrosome detachment is also suggested by the observation that CENP-32 depletion results in a small but significant decrease in the formation of monopolar cold-stable microtubule depolymerization assay showed that kine-tochore–microtubule interactions remained robust in CENP-32–depleted cells. This therefore raises the question of how the anastal spindles in CENP-32–knockdown cells are maintained.

NuMA was previously shown to interact with dynein to promote spindle pole formation in Xenopus egg extracts by cooperating with cytoplasmic dynein to cross-link parallel microtubules (Merdes et al., 1996, 2000). Furthermore, after CENP-32 depletion, NuMA remained at the poles of the anastal spindles and did not associate with the dissociated centrosomes. Bipolar spindle morphology was severely compromised after double depletion of CENP-32 and NuMA. Thus NuMA is one of the factors that maintain the anastal spindles in CENP-32–depleted cells.

Factors that form and maintain anastal mitotic spindles

Centrosomal dissociation from the mitotic spindle was previously reported in CDK5RAP2-knockdown cells (Barr et al., 2010). In fact,
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spindles in monastrol-treated cells. This decrease is balanced by an increase in the number of bipolar, monoastral spindles, suggesting that in those cells, the spindle forces normally counteracted by Eg5 cause the detachment of a centrosome from one pole instead of pulling the two poles together.

**CENP-32 and centrosomal dominance in spindle assembly**

Even though spindles can form perfectly well in the absence of centrosomes (e.g., in eggs and higher plants, as well as in vertebrate cultured cells after experimental manipulations), it is generally believed that where they are present, centrosomes act as dominant determinants of spindle pole formation. Remarkably, the microtubules associated with CENP-32-detached centrosomes do not make stable attachments to kinetochores and appear not to insinuate themselves into the body of the spindle. These results suggest that microtubules associated with detached centrosomes in CENP-32–depleted cells behave like astral microtubules rather than like components of the central body of the spindle. This is despite the fact that the detached centrosomes retain both Aurora A and γ-tubulin. Thus CENP-32 is apparently required for centrosomes to behave like spindle poles capable of nucleating both spindle and astral microtubules.

Future experiments with CENP-32–depleted cells may allow identification of the determinants that enable centrosomes to act as dominant determinants of spindle pole formation.

**MATERIALS AND METHODS**

**Cell culture**

U2OS cells in exponential growth were seeded onto coverslips and grown overnight at 37°C in RPMI/10% fetal bovine serum (FBS) at 5% CO₂. DT40 cells with the CENP-32 mutant were maintained in RPMI 1640 medium supplemented with 1% (vol/vol) FBS, 1% chicken serum (Life Technologies, Grand Island, NY), 100 U/ml penicillin, 100 μg/ml streptomycin (Wako, Osaka, Japan) at 39°C, and 5% CO₂ in a humid incubator.

**Drug treatment**

Monastrol (Sigma-Aldrich, St. Louis, MO) was dissolved in dimethyl sulfoxide (DMSO) and used at a final concentration of 68 μM for 12 h. DMSO was added to mock-treated controls.

**Transfection and indirect immunofluorescence microscopy**

A 100-pmol amount of siRNA (control, AACUGACCGGAUAC- UUCGAdTdT; CENP-32, GCAGACCCUCGGACACAAAdTdT; Ohta et al., 2010; CG-NAP-si1, GCUUCUAUUUUGUCAGAAdTdT, Ozaki et al., 2012; CG-NAP-si2, GCAUGGAUGCUUCGACAdTdT; Dgt6, CAGUAAAAGCUACGGAAdTdT, Uehara and Goshima, 2010); NuMA, CUAGCUAGCCAGUACGAdTdT, Haren et al., 2009) was administered to U2OS cells at 30–40% confluence by transfection with Oligofectamine or Lipofectamine RNAi MAX (Life Technologies) in complete medium without antibiotics. To rescue siRNA treatment. Cells maintained in this medium for 48–72 h were transfected to U2OS cells at 30–40% confluence by Transfection and indirect immunofluorescence microscopy

**Immunofluorescence**

Immunofluorescence was done as described above.

**Electron microscopy**

The culture cells were fixed with 1% glutaraldehyde in PBS, pH 7.3, for 1 h at 4°C. They were then postfixed with 1.5% osmium tetroxide (or osmic acid) in 0.1 M phosphate buffer, pH 7.3, for 1 h at 4°C and dehydrated in a graded series of ethanol. After dehydration, they were embedded in Epon 812 (TAAB Laboratories Equipment, Berkshire, United Kingdom). They were observed with a JEM-1400Plus electron microscope (JEOL, Tokyo, Japan).

**FACS**

FACS analysis was carried out on DT40 cells maintained in media containing bromodeoxyuridine (BrdU) for 30 min before harvest. Cells were washed once in PBS with 1% BSA and fixed in 70% ethanol. The fixed cells were washed again once with PBS and probed with anti-BrdU fluorescein isothiocyanate (1:100, 1:10,000 and anti-mouse IgG at 1:2000. The secondary antibodies were IRDye 800CW donkey anti–a-rabbit immunoglobulin IgG at 1:10,000 and anti-mouse IgG at 1:10,000 (926-32213, 926-32210; LI-COR Biosciences, Lincoln, NE).

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