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Neil J. Ganem
Dartmouth College

Duane A. Compton
Dartmouth College

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The KinI kinesin Kif2a is required for bipolar spindle assembly through a functional relationship with MCAK

Neil J. Ganem and Duane A. Compton

Department of Biochemistry, Dartmouth Medical School, Hanover, NH 03755

Although the microtubule-depolymerizing KinI motor Kif2a is abundantly expressed in neuronal cells, we now show it localizes to centrosomes and spindle poles during mitosis in cultured cells. RNAi-induced knock-down of Kif2a expression inhibited cell cycle progression because cells assembled monopolar spindles. Bipolar spindle assembly was restored in cells lacking Kif2a by treatments that altered microtubule assembly (nocodazole), eliminated kinetochore–microtubule attachment (loss of Nuf2), or stabilized microtubule plus ends at kineto-

chores (loss of MCAK). Thus, two KinI motors, MCAK and Kif2a, play distinct roles in mitosis, and MCAK activity at kinetochores must be balanced by Kif2a activity at poles for spindle bipolarity. These treatments failed to restore bipolarity to cells lacking the activity of the kinesin Eg5. Thus, two independent pathways contribute to spindle bipolarity, with the Eg5-dependent pathway using motor force to drive spindle bipolarity and the Kif2a-dependent pathway relying on microtubule polymer dynamics to generate force for spindle bipolarity.

Introduction

The spindle is a microtubule-based structure responsible for accurate chromosome segregation during mitosis and meiosis (McIntosh and Koonce, 1989; Hyman and Karsenti, 1996; Compton, 2000). Organization of microtubules into a fusiform, bipolar spindle requires numerous nontubulin accessory proteins including several members of the kinesin gene family (Vale and Fletterick, 1997; Sharp et al., 2000). Kinesin-related proteins with the conserved motor domain positioned at the NH₂ terminus (KinN kinesins) display plus-end-directed motility and perform diverse functions during spindle morphogenesis including kinetochore–microtubule interactions (CENP-E; Yao et al., 1997), generation of polar ejection force (Kid; Levesque and Compton, 2001), and spindle bipolarity (Eg5; Sawin et al., 1992). Kinesin-related proteins with the conserved motor domain positioned at the COOH terminus (KinC kinesins) display minus-end-directed motility and participate in focusing microtubule minus ends at spindle poles (Endow et al., 1994; Mountain et al., 1999). Finally, kinesin-related proteins with the conserved motor domain positioned in the middle of the protein (KinI kinesins) lack motility and use their catalytic motor domains to induce

conformation changes in protofilament structure at microtubule ends, which stimulates microtubule depolymerization by promoting catastrophe (Desai et al., 1999; Moores et al., 2002; Ogawa et al., 2004). MCAK/XKCM1/Kif2c (hereafter referred to as MCAK) is the best characterized member of the KinI subgroup. MCAK localizes to kinetochores and spindle microtubules (Wordeman and Mitchison, 1995; Walczak et al., 1996) where it regulates microtubule dynamics essential for proper chromosome attachment to spindle microtubules, chromosome movement, and correction of attachment errors (Maney et al., 1998, 2001; Kline-Smith and Walczak, 2002; Walczak et al., 2002; Hunter et al., 2003; Ohi et al., 2003; Kline-Smith et al., 2004). However, MCAK is not the only member of KinI kinesin-related proteins. Kif2a is a closely related KinI kinesin-related protein that plays a role in neurogenesis (Noda et al., 1995; Debernardi et al., 1997; Morfini et al., 1997; Homma et al., 2003). In this paper we demonstrate that Kif2a is essential for bipolar spindle assembly during mitosis in cultured vertebrate cells.

Results and discussion

To build Kif2a-specific antibodies we immunized rabbits with the unique 119 amino acids of the NH₂-terminal end of human Kif2a. A single protein species of 110 kD was detected by immunoblot analysis with these antibodies in total cell extracts prepared from human HeLa, hamster CHO, and frog A6 cells (Fig. 1 A). The molecular mass of this protein is consistent with the predicted size of Kif2a, indicating

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Address correspondence to Duane A. Compton, Dept. of Biochemistry, Dartmouth Medical School, 410 Remsen Bldg., Hanover, NH 03755. Tel.: (603) 650-1990. Fax: (603) 650-1128. email: duane.a.compton@dartmouth.edu

Key words: mitotic spindle; kinetochore; microtubule; centrosome; poleward flux

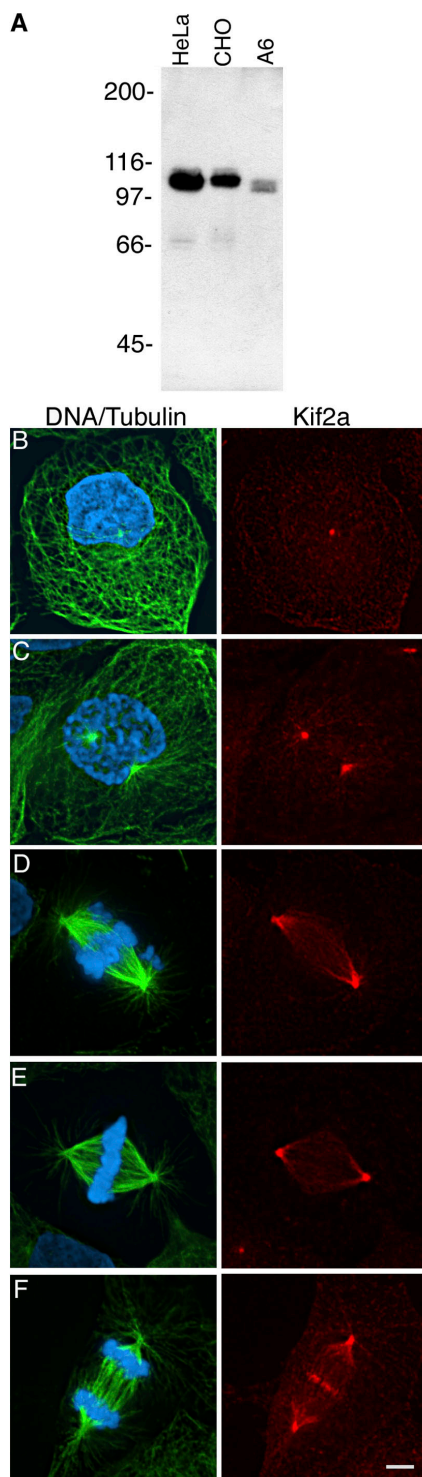


Figure 1. Characterization of Kif2a in cultured vertebrate cells. (A) Total cell protein from ~100,000 human HeLa, frog A6, or hamster CHO cells was separated by SDS-PAGE and blotted with Kif2a polyclonal antibodies. Positions of myosin (200), β -galactosidase (116), phosphorylase B (97), albumin (66), and ovalbumin (45) are shown in kilodaltons. (B–F) Human CFPAC-1 cells were processed for indirect immunofluorescence microscopy using Kif2a- (red) and tubulin-specific antibodies (green), as well as the DNA-specific dye DAPI (blue). Cells are in interphase (B), prophase (C), prometaphase (D), metaphase (E), and anaphase (F). Bar, 10 μ m.

that the antibodies are specific for Kif2a, they cross react to Kif2a from several vertebrate species, and Kif2a is expressed in cultured vertebrate cells. Kif2a localized to centrosomes in interphase and mitotic human CFPAC-1 cells (Fig. 1, B–F), but also to spindle microtubules being concentrated at poles (Fig. 1, D–F). In anaphase and telophase, Kif2a localized to the spindle midzone and midbody in addition to spindle poles (Fig. 1 F). We detected Kif2a at kinetochores, but the signal was just slightly above background levels and was only observed after depolymerization of microtubules with nocodazole (unpublished data). This localization pattern for Kif2a is different from that reported for the related KinI motor MCAK (Wordeman and Mitchison, 1995; Walczak et al., 1996).

To determine if Kif2a is required for mitotic spindle assembly or function, we used siRNA to knock down Kif2a protein levels in human U2OS cells (Fig. 2). 48 h after transfection, immunoblot analysis demonstrated that Kif2a levels were reduced >90% in cells transfected with Kif2a-specific RNA relative to cells transfected with nonspecific RNA (Fig. 2 A). At this time point, cells transfected with

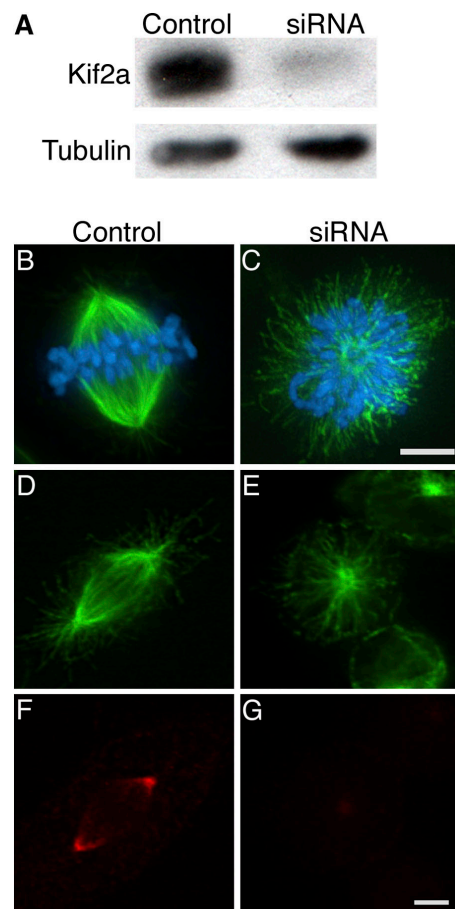


Figure 2. Kif2a is essential for spindle bipolarity. (A) Human U2OS cells were transfected with either nonspecific (control) or Kif2a-specific (siRNA) RNAs. Total cell protein was separated by size using SDS-PAGE and blotted with Kif2a-specific antibodies and tubulin antibodies. (B–G) Human U2OS cells were transfected with either nonspecific (control) or Kif2a-specific (siRNA) RNAs, fixed with either glutaraldehyde (B and C) or methanol (D–G), and stained for tubulin (green), DNA (blue), and/or Kif2a (red). Bars, 10 μ m.

Kif2a-specific RNA had an average mitotic index of $\sim 36\%$ in contrast to the mitotic index of $\sim 3\%$ observed after transfection with nonspecific RNA. Most ($>99\%$) mitotic cells transfected with nonspecific RNA had bipolar spindles with Kif2a localized at spindle poles (Fig. 2, B, D, and F). In contrast, $>90\%$ of mitotic cells transfected with Kif2a-specific RNA had monopolar spindles (Fig. 2, C and E; and see Fig. 5 H), and microtubule attachment to kinetochores was detected by electron microscopy (not depicted). Kif2a was undetectable on these monopolar spindles by indirect immunofluorescence, which is consistent with the results of immunoblot analysis (Fig. 2 G). Centrosome staining demonstrated that initial separation of centrosomes at prophase in cells lacking Kif2a was not maintained when cells entered prometa/metaphase (Fig. S1, available at <http://www.jcb.org/cgi/content/full/jcb.200404012/DC1>).

Kif2a has been shown to catalyze microtubule depolymerization in the absence of motile function (Desai et al., 1999). To determine if altered microtubule dynamics might contribute to the failure to maintain centrosome separation in mitotic cells lacking Kif2a, we suppressed microtubule dynamics by addition of 100 nM nocodazole (Fig. 3). Whereas only $\sim 8\%$ of mitotic cells transfected with Kif2a-specific RNA alone formed bipolar spindles, $\sim 58.6\%$ of mitotic cells transfected with Kif2a-specific RNA and treated with nocodazole for 5 h before fixation assembled bipolar spindles (Fig. 3 C and see Fig. 5 H, $n = 232$, five independent experiments). Chromosomes did not align efficiently on these bipolar spindles, which is consistent with disruption of microtubule dynamics by nocodazole treatment. Restoration of spindle bipolarity by nocodazole treatment was specific to cells lacking Kif2a activity because nocodazole treatment did not alter the percentage of cells with monopolar spindles induced with 100 μM monastrol (Fig. 3 D and see Fig. 5 I, $n = 282$, six independent experiments), a cell-permeable inhibitor of Eg5 function (Mayer et al., 1999). Microtubules in monopolar spindles induced by Eg5 inhibition end abruptly near chromosomes in contrast to microtubules in monopolar spindles induced by the lack of Kif2a, which extend beyond chromosomes to the cell cortex (Fig. 3, A and B). A similar difference between cells lacking Kif2a or Eg5 activity was observed with low doses of taxol, as 53.6% ($n = 41$) of mitotic cells lacking Kif2a and treated with 15 nM taxol had either multipolar or bipolar spindles, whereas 94.3% ($n = 35$) of mitotic cells treated with both 100 μM monastrol and 15 nM taxol had monopolar spindles.

To alter microtubule dynamics in mitotic cells through a mechanism independent of microtubule drugs, we used siRNA to knock down expression of the microtubule destabilizing protein MCAK (Fig. 4). 48 h after transfection, immunoblot analysis indicated that both MCAK and Kif2a were reduced to levels equivalent to when each protein was knocked down alone (Fig. 4 A). In contrast to the monopolar spindles induced by knockdown of Kif2a alone (Fig. 4 B), most cells ($\sim 91\%$) lacking MCAK alone had bipolar spindles (Cassimeris and Morabito, 2004; Holmfeldt et al., 2004). Simultaneous knockdown of MCAK and Kif2a resulted in bipolar spindles with aligned chromosomes in a majority of cells (70%, $n = 200$, four independent experiments; Fig. 4 D and see Fig. 5 H). Astral microtubules were

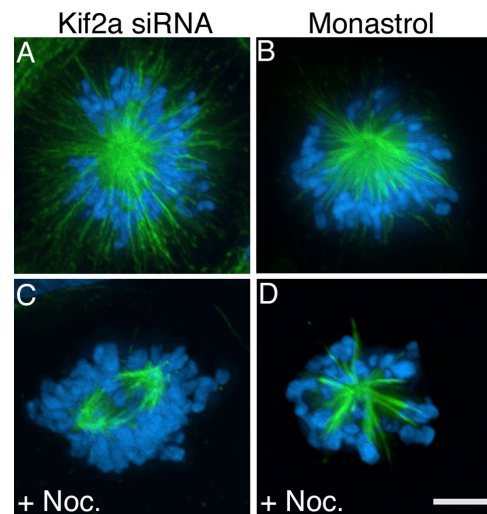


Figure 3. Nocodazole treatment restores spindle bipolarity to cells lacking Kif2a. U2OS cells stained for tubulin (green) and DNA (blue) after transfection with Kif2a-specific RNA (A and C) or treatment with 100 μM monastrol (B and D) in the presence (C and D) or absence (A and B) of 100 nM nocodazole. Bar, 10 μm .

extremely long in these spindles and often extended from one spindle pole to the cell cortex beyond the opposite spindle pole. Anaphase cells were observed and the mitotic index was $\sim 6\%$, a significant decrease from that observed after diminution of Kif2a alone. Reduction of MCAK levels failed to prevent monopolarity in monastrol-treated cells, as only $\sim 5\%$ had bipolar spindles (Fig. 4 E and see Fig. 5 I; $n = 84$, two independent experiments).

These results demonstrate that MCAK and Kif2a play functionally different roles during mitosis in cultured cells, a conclusion paralleled by analyses of Klp59C and Klp10A in fruit flies (Goshima and Vale, 2003; Rogers et al., 2003). MCAK is necessary for correction of chromosome attachment errors on spindles (Maney et al., 1998; Walczak et al., 2002; Kline-Smith et al., 2004), but not bipolar spindle assembly (Cassimeris and Morabito, 2004; Holmfeldt et al., 2004). In contrast, Kif2a is essential for bipolar spindle assembly. Also, the relative contributions made by MCAK and Kif2a to spindle assembly differ between mitotic and meiotic systems. MCAK plays a dominant role in regulating microtubule plus end dynamic instability in meiotic systems through antagonism of TOGp/XMAP215 activity (Tournabize et al., 2000; Kinoshita et al., 2001) such that massive monopolar arrays form if it is inhibited (Walczak et al., 1996). Perturbation of Kif2a also leads to monopolar arrays in meiotic extracts (Gaetz and Kapoor, 2004). Meiotic systems have far greater pools of tubulin subunits relative to mitotic cells, and perturbation of either MCAK or Kif2a most likely result in monopolar arrays in meiotic systems because excess microtubule polymer overwhelms the machinery that organizes microtubules into bipolar arrays. Finally, MCAK activity must be appropriately balanced by Kif2a activity for bipolar spindle assembly in cultured cells. This unexpected functional relationship between two KinI kinesin-related proteins indicates that MCAK and Kif2a activities are spatially restricted in spindles with MCAK regulating

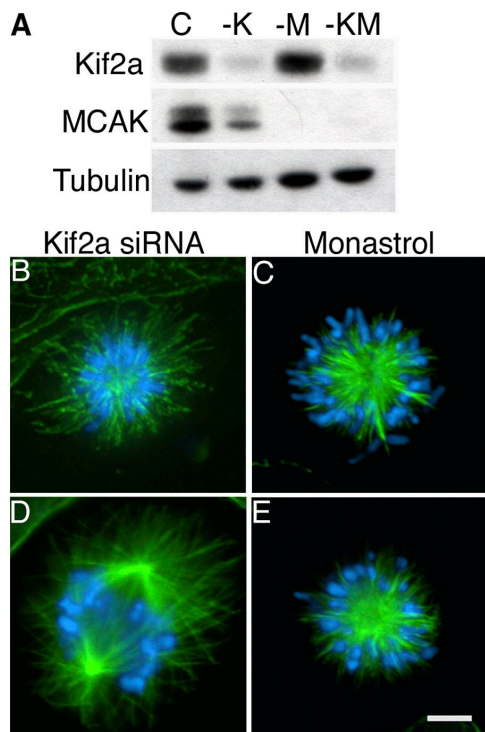


Figure 4. MCAK knockdown restores spindle bipolarity to cells lacking Kif2a. (A) U2OS cells were transfected with nonspecific (C), Kif2a-specific (-K), MCAK-specific (-M), or both Kif2a- and MCAK-specific (-KM) RNAs. Total cell protein was separated by size using SDS-PAGE and blotted with Kif2a-, MCAK-, and tubulin-specific antibodies. (B–E) Human U2OS cells were stained for tubulin (green) or DNA (blue) after transfection with Kif2a-specific RNA alone (B), transfection with both Kif2a- and MCAK-specific RNA (D), treatment with 100 μ M monastrol alone (C), or transfection with MCAK-specific RNA and treatment with 100 μ M monastrol (E). Bar, 10 μ M.

microtubule plus ends at kinetochores and Kif2a regulating microtubule minus ends at poles.

To test if kinetochore-induced microtubule depolymerization is necessary for monopolar spindle assembly in the absence of Kif2a, we eliminated kinetochore–microtubule attachment using siRNA to knock down expression of Nuf2, a component of a conserved, essential kinetochore complex containing Nuf2 and Hec1 (Fig. 5; DeLuca et al., 2003). We used Hec1-specific antibodies to monitor the efficiency of Nuf2 knockdown because the stability of Hec1 relies on the presence of Nuf2 (DeLuca et al., 2003), and Nuf2-specific antibodies were not available. 48 h after transfection, immunoblot analysis indicated that Kif2a and Hec1 were reduced to levels equivalent to when each protein was knocked down alone (Fig. 5 A). Most (~98%; $n = 77$, three independent experiments) cells lacking Nuf2 alone had bipolar spindles with unattached chromosomes (Fig. 5 C), which is consistent with previous findings (DeLuca et al., 2003). In contrast to the monopolar spindles formed in cells lacking Kif2a alone (Fig. 5 D), ~94% ($n = 213$, three independent experiments) of cells lacking both Kif2a and Nuf2 assembled bipolar spindles with unattached chromosomes (Fig. 5, E and H). Knockdown of Nuf2 failed to restore bipolarity to cells lacking Eg5 activity (Fig. 5, F, G, and I; $n = 83$, three

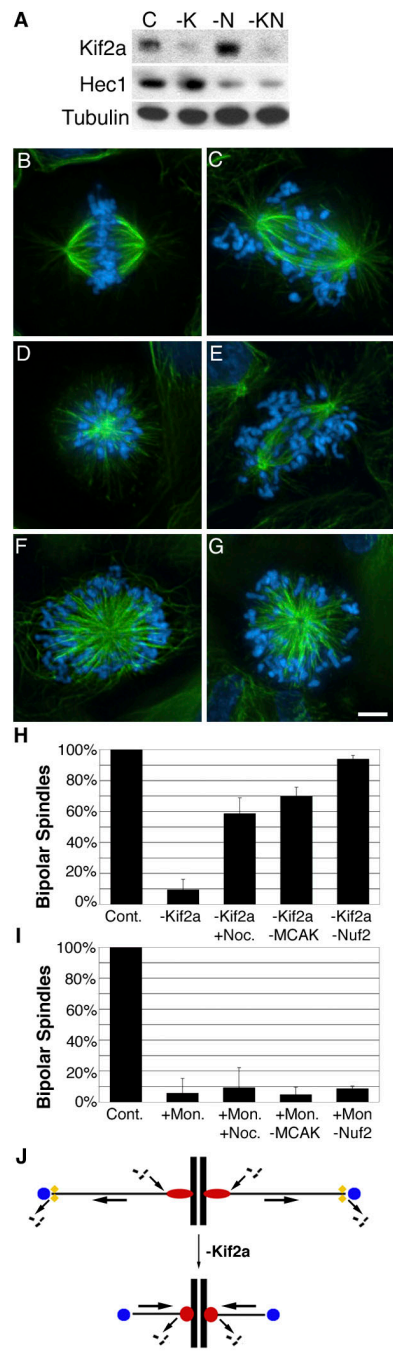


Figure 5. Nuf2 knockdown restores spindle bipolarity to cells lacking Kif2a. (A) U2OS cells were transfected with nonspecific (C), Kif2a-specific (-K), Nuf2-specific (-N), or both Kif2a- and Nuf2-specific (-KN) RNAs. Total cell protein was separated by size using SDS-PAGE and blotted with Kif2a-, Hec1-, and tubulin-specific antibodies. (B–E) Human U2OS cells were stained for tubulin (green) or DNA (blue) after transfection with control RNA (B), Kif2a-specific RNA (D and E), Nuf2-specific RNA (C, E, and G), or treatment with 100 μ M monastrol (F and G). Bar, 10 μ M. (H and I) Quantification of the percentage of mitotic U2OS cells with bipolar spindles under various conditions after Kif2a knockdown (H) or monastrol treatment (I). Error bars represent SD. (J) Kif2a depolymerizes microtubule minus ends associated with poleward microtubule flux. Flux generates tension that induces kinetochores to add tubulin subunits. When flux is perturbed in the absence of Kif2a, tension at kinetochores is lost and kinetochores depolymerize microtubules drawing centrosomes together into monopolar arrays.

independent experiments), and knockdown of both MCAK and Nuf2 showed no detectable difference from knockdown of Nuf2 alone (not depicted).

These results demonstrate that Kif2a and Eg5 act through independent pathways to promote spindle bipolarity, and that each pathway is necessary but not sufficient for bipolar spindle assembly in cultured cells. Eg5 uses motor force to slide antiparallel microtubules relative to one another to promote spindle bipolarity (Sharp et al., 2000) in a mechanism that is insensitive to treatments that alter microtubule dynamics and independent of kinetochore–microtubule interactions. In contrast, Kif2a is a microtubule-depolymerizing enzyme that promotes spindle bipolarity in a mechanism that is sensitive to treatments that alter microtubule dynamics. Similar differences have been noted for the Cin8 and Kip3 (and Kar3) kinesin-related proteins in yeast (Cottingham et al., 1999). It is unlikely that loss of Kif2a leads to monopolar spindles due to microtubules being excessively long because cells lacking both Kif2a and MCAK have unusually long microtubules but still assemble bipolar spindles (Fig. 4 D). Instead, kinetochore-derived forces draw centrosomes together in the absence of Kif2a. This finding explains why centrosome separation in the absence of Kif2a is normal until nuclear envelope break down (Fig. S1), and why treatments that stabilize microtubule plus ends, like low doses of nocodazole or loss of MCAK (Jordan et al., 1992; Vasquez et al., 1997; Kinoshita et al., 2001; Kline-Smith and Walczak, 2002; Ohi et al., 2003), or eliminate kinetochore–microtubule attachment promote spindle bipolarity in the absence of Kif2a. We cannot discount the possibility that Kif2a acts directly at kinetochores, but that is inconsistent with Kif2a localization to poles. Alternatively, recent data shows that disassembly of microtubule minus ends associated with poleward microtubule flux requires the targeting of Klp10A (Rogers et al., 2003) and Kif2a (Gaetz and Kapoor, 2004) to spindle poles. Force from poleward flux generates tension that induces kinetochores to add tubulin subunits to microtubule plus ends (Maddox et al., 2003). In that context, disruption of flux by the loss of Kif2a would relax tension at kinetochores. Consequently, kinetochores would switch to the depolymerization state and remove tubulin subunits from microtubule plus ends, drawing centrosomes together into monopolar spindles (Fig. 5 J). Although stabilization of microtubule plus ends at kinetochores through loss of MCAK is sufficient to restore kinetochore tension and prevent monopolarity, other kinetochore proteins execute microtubule depolymerization during chromosome movement because chromosome velocity is unaffected by depletion of MCAK from centromeres (Kline-Smith et al., 2004). Together, these data lead us to speculate that force from microtubule polymer dynamics (for review see Inoue and Salmon, 1995) contributes to bipolar spindle assembly.

Materials and methods

Cell culture

Human HeLa, CFPAC-1, and U2OS cells were maintained in Dulbecco's modified medium, Iscove's modified Dulbecco's medium, and McCoy's medium, respectively, containing 10% FBS, 50 IU/ml penicillin, and 50 µg/ml streptomycin.

Antibodies

The region of the cDNA encoding the NH₂-terminal 119 aa of human Kif2a (GenBank/EMBL/DBJ accession no. BG721281) was amplified using forward (5'-ATCGGATCCTGCTGCTCCAGATGAGGT-3') and reverse (5'-CTTCTCTGGGGCACGGGAAATCTTAAG-3') primers and inserted into the BamHI–EcoRI sites of pRSET-B plasmid (Invitrogen). The His-tagged recombinant protein was purified by affinity chromatography and used to immunize two rabbits (Covance Research Products).

Other antibodies used in this work included MCAK (Mack and Compton, 2001), DM1α (Sigma-Aldrich), Hec1 (Novus Biologicals), and human anticentrosome antibody (provided by J.B. Rattner, University of Calgary, Alberta, Canada).

Indirect immunofluorescence microscopy

Cells were extracted in microtubule-stabilizing buffer (4 M glycerol, 100 mM Pipes, pH 6.9, 1 mM EGTA, 5 mM MgCl₂, and 0.5% Triton X-100) followed by fixation in methanol (Kif2a staining) or 1% glutaraldehyde (microtubule staining). Subsequent antibody incubations and washes were done in TBS-BSA (10 mM Tris, pH 7.5, 150 mM NaCl, and 1% BSA). Primary antibodies were detected using species-specific fluorescein- or Texas red-conjugated secondary antibodies (Vector Laboratories). DNA was detected with 0.2 µg/ml DAPI (Sigma-Aldrich). Coverslips were mounted with ProLong Antifade mounting medium (Molecular Probes).

Fluorescent images were captured with a cooled CCD camera (model Orca II; Hamamatsu) mounted on a microscope (model Axioplan 2; Carl Zeiss Microimaging, Inc.). A series of 0.5-µm optical sections were collected in the z plane for each channel (DAPI, fluorescein, and/or Texas red) using a 63× 1.4NA objective and deconvolved using Openlab software (Improvision Inc.) to eliminate extraneous fluorescence background.

RNA interference

Kif2a levels were reduced using the sequence 5'-GGCAAAGAGAU-UGACCUGG-3'. Nuf2 and MCAK levels were reduced using published sequences (DeLuca et al., 2003; Cassimeris and Morabito, 2004). All siRNA duplexes were synthesized with 3' dTdT overhangs and were annealed (Ambion). Approximately 50,000 U2OS cells were plated on coverslips in 35-mm dishes the day before transfection and grown without antibiotics. Double stranded RNAs were transfected into cells using OligofectamineTM reagent (Invitrogen) as described previously (Elbashir et al., 2001). Samples were analyzed 48 h after transfection by either indirect immunofluorescence or immunoblot analysis.

Online supplemental material

Fig. S1 shows centrosome position during both prophase and metaphase in control cells and cells lacking Kif2a. Online supplemental material is available at <http://www.jcb.org/cgi/content/full/jcb.200404012/DC1>.

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