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Permalink
https://escholarship.org/uc/item/6rw6f132

Journal
The Journal of cell biology, 181(3)

ISSN
0021-9525

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Publication Date
2008-05-01

DOI
10.1083/jcb.200802189

Peer reviewed
CENP-E combines a slow, processive motor and a flexible coiled coil to produce an essential motile kinetochore tether

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Introduction

Mitotic chromosome motions are driven by microtubule-based motors positioned at kinetochores as well as by the dynamic properties of spindle microtubules. Centromere protein E (CENP-E) and cytoplasmic dynein are the two microtubule motors in metazoans known to localize at kinetochores (Pfarr et al., 1990; Yen et al., 1992). Each uses chemical energy released from ATP hydrolysis to direct movement toward plus ends or minus ends of microtubules, respectively (Schroer et al., 1989; Wood et al., 1997). Mitotic centromere-associated kinesin, a nonmotile kinesin, is also found at the inner centromere, and its microtubule-depolymerizing activity plays a role in error correction as well as to generate proper kinetochore microtubule attachment (Desai et al., 1999a; Ohi et al., 2003). Besides the energy-consuming motors, however, energy stored in the microtubule lattice from cleavage of GTP during microtubule assembly can generate sufficient force to power aspects of chromosome movement (Koshland et al., 1988; Coue et al., 1991; Inoue and Salmon, 1995; Grishchuk et al., 2005). By simply maintaining a linkage with spindle microtubules, kinetochores can exploit the energy liberated from the microtubule lattice during disassembly to drive chromosome movement independent of a power stroke generated by motors and additional chemical energy (Howard and Hyman, 2003).

An important unsolved question is how the kinetochore holds onto both growing and shrinking microtubules. Using purified chromosomes attached to dynamic microtubules, either pan kinesin antibodies or those specific for CENP-E were shown to inhibit microtubule depolymerization-dependent motion of chromosomes in vitro (Lombillo et al., 1995a). Moreover, beads coated with kinesin were able to follow the end of disassembling microtubules (Lombillo et al., 1995b), suggesting a role of kinesin family members in coupling kinetochores to dynamic microtubules. Discovery of the ring structure of the yeast Dam1/DASH complex (Miranda et al., 2005; Westermann et al., 2005) has introduced such rings as plausible coupling devices for chromosome attachment and movement. In vitro, the ring complex can move processively on depolymerizing microtubule ends (Westermann et al., 2006), and harness microtubule dynamics to produce force (Asbury et al., 2006). However, the Dam1/DASH complex has not been found outside fungi.
Conversely, motor enzymes such as dynein, CENP-E, and mitotic centromere-associated kinesin are missing from yeast kinetochores (McIntosh, 2005).

Initially found to be a kinesin family member that localizes to kinetochores (Yen et al., 1992), CENP-E is a mitosis-specific kinesin with a cyclin-like accumulation and degradation, reaching its peak during G2 and early M phase followed by rapid degradation during the completion of mitosis (Brown et al., 1994). As CENP-E is an essential kinetochore component whose loss leads to high rates of whole chromosome missegregation (Weaver et al., 2003), CENP-E is one of the components directly responsible for the stable capture of spindle microtubules by kinetochores (Schaar et al., 1997; Wood et al., 1997; Yao et al., 2000; Putkey et al., 2002). Inhibition or removal of CENP-E not only leads to a failure of metaphase chromosome alignment, resulting in unattached chromosomes (Brown et al., 1997; Yao et al., 2000; Putkey et al., 2002). We also verified CE473-GFP to be dimeric by using total internal reflection fluorescence (TIRF) microscopy to measure the fluorescence intensity of single molecules immobilized on coverslips. Freshly prepared protein was nonspecifically adhered on a coverslip surface, and the fluorescence intensity of CE473-GFP spots was measured and compared with that of K560-GFP, the well-characterized dimeric motor head of human kinesin-1 (Case et al., 1997). The fluorescence intensity distributions between CE473-GFP (n = 432) and K560-GFP (n = 347) were indistinguishable (Fig. S1 C), confirming that most imaged CENP-E spots were dimers. When the fluorescent CENP-E molecules were followed over time to examine their photobleaching behavior, most CE473-GFP and K560-GFP spots disappeared in either two steps or a single step (Fig. S1 D), as expected if each spot consisted of either two GFPs or a single GFP. Furthermore, the initial fluorescence intensity of moving spots was measured to confirm that the majority of moving CENP-E was indeed dimeric (Fig. S1 E). Therefore, both the hydrodynamic measurements in solution and the fluorescence analysis of single molecules viewed microscopically demonstrate that CE473-GFP is dimeric.

We then tested the processivity of CENP-E in vitro by directly observing single molecules of CE473-GFP moving along microtubules using TIRF microscopy. X-rhodamine-labeled guanosine-5’-[(a,b)-methylene] triphosphate (GMP-CPP) microtubules were immobilized on a coverslip surface using an antitubulin antibody, and, subsequently, a low concentration (0.5–1 nM) of CE473-GFP was introduced into the flow chamber (Fig. 1 B). CENP-E single molecules moved processively and unidirectionally along microtubules (Fig. 1 C and Videos 1 and 2, available at http://www.jcb.org/cgi/content/full/jcb.200802189/DC1) with a median velocity of 8 nm/s (n = 320; four independent preparations; Fig. 1 E). Substantial variation in speed (ranging from 1.7 to 48 nm/s) was observed for CENP-E. However, this was not the result of heterogeneity in motor activity among different protein preparations, as large variation in the velocity was consistently observed in the assays performed on the same day. Furthermore, the velocity of movement often differed by a factor of two even between the molecules running on the same microtubule (Fig. 1 C, compare particle 1 with particle 2).

Processive movement of CENP-E motor single molecules was not smooth, and the CENP-E changed instantaneous velocity with multiple pauses within a single run (Fig. 1 D). Frequent changes in instantaneous velocity with large variations in the speed and even reversals in direction observed when frames were taken more rapidly in our single-molecule assays (Fig. S1, G and H; and Video 3, available at http://www.jcb.org/cgi/content/full/jcb.200802189/DC1) suggested that CENP-E motility has a one-dimensional diffusion component. A mean square displacement (MSD) plot of CENP-E was well fitted with biased Brownian movement (Fig. 1 F). Polynomial regression to MSD (p = \(v^{2} + 2Dt\)) produced a mean velocity of 9.7 ± 0.6 nm/s (n = 90) and a diffusion coefficient of 690 ± 94 nm^2/s (n = 90), which was larger than expected from the fluctuation of a kinesin with an 8-nm step size and ~10-nm/s speed (D = 80 nm^2/s; Svoboda et al., 1994).

Using purified recombinant CENP-E, we now find that CENP-E is a very slow, highly processive motor that maintains microtubule attachment for long periods with a highly flexible 230-nm-long contour length. These properties combine not only to power chromosome congression but also to provide an essential motile kinetochore tether.

**Results and discussion**

**CENP-E is a slow, processive motor that maintains microtubule attachment for long periods**

To characterize the motor properties of CENP-E, we purified the motor plus neck domain of *Xenopus laevis* CENP-E (aa 1–473) with GFP and a hexahistidine tag fused at the C terminus (CE473-GFP hereafter; Fig. 1 A). The presumed dimeric state of this CENP-E fragment was tested by measuring hydrodynamic parameters in solution. A sedimentation coefficient (6 S) and Stokes radius (5.2 nm) were obtained by sucrose gradient sedimentation and gel filtration chromatography, respectively (Fig. S1, A and B; available at http://www.jcb.org/cgi/content/full/jcb.200802189/DC1). This produced a calculated native molecular mass (Siegel and Monty, 1966) for CE473-GFP of ~130 kD, which is close to the predicted molecular mass of the dimer (160 kD), demonstrating that the preponderance of the CENP-E motor domain exists as a dimer in solution.
CENP-E is a slow, processive motor that maintains microtubule attachment for long periods.

The presence of a diffusional component in motility has been reported for other microtubule-dependent motors (Vale et al., 1989; Okada and Hirokawa, 1999; Kwok et al., 2006; Helenius et al., 2006; Furuta and Toyoshima, 2008) and suggests that CENP-E contains an additional electrostatic binding mode within the head-neck domain in addition to its force-generating strong microtubule-binding state. This weak binding of CENP-E to a microtubule might prevent the dissociation of CENP-E even when both motors are unbound, thus facilitating efficient reattachment. Consistent with this, CENP-E was able to maintain microtubule attachment for long periods, producing remarkably processive movement despite its slow velocity. The run time, which is defined as the time interval between the start and end of each processive run, was distributed exponentially, and the mean run time was determined by fitting the data into a cumulative distribution function. The mean duration of CENP-E processive runs was 195 ± 13 s (Fig. 1 G), a value 50 times longer than the reported association time of K560-GFP single molecules (Friedman and Vale, 1999). The longest run we observed (Fig. 1 D, kymograph 2) lasted >20 min. The slow velocity and longer association time of the CENP-E motor domain resulted in a mean run length of 1.5 ± 0.1 μm (Fig. 1 H), which is comparable with the processivity of conventional kinesin. Therefore, unlike fast-moving organelle transport kinesins, CENP-E is a slow motor, but its high processivity maintains interaction with a microtubule for long periods.

Similar mean velocities and diffusion coefficients for CENP-E were found in the experiments performed using more frequent frame rates and quantum dot-labeled CENP-E motor domain (Fig. S1 I).
CENP-E is a highly flexible, dimeric kinesin with a 230-nm discontinuous coiled coil

To assess the structural features of full-length CENP-E, we purified untagged 340-kD recombinant CENP-E close to homogeneity from insect cells infected with baculovirus-expressing full-length Xenopus CENP-E cDNA (Fig. 2 A). Immediately after purification, CENP-E was adsorbed to mica, and individual molecules were visualized by electron microscopy using the quick-freeze deep-etch technique and platinum replication (Heuser, 1989). Consistent with its predicted long α-helical coiled-coil domain, CENP-E was seen to be highly elongated. The two kinesin motor heads, as indicated by arrows, were found separated by a long coiled coil followed by globular tails (Fig. 2 B), directly demonstrating that CENP-E is a homodimer. The mean contour length of CENP-E was measured to be 230 ± 25 nm (n = 20; Fig. 2 C), which is almost three times longer than the overall length of conventional kinesin (Hirokawa et al., 1989).

A striking feature of CENP-E was the high flexibility of the α-helical coiled coil, as inferred from the wide variety of configurations that it adopted. Conventional kinesins mainly exist as either extended or folded conformations with a hinge located in the middle of the coiled coil (Hirokawa et al., 1989). In contrast, CENP-E molecules were never uniformly straight but rather displayed a myriad of diverse conformations, none of which could be categorized by frequency as preferred conformations. Throughout the long coiled-coil domain, CENP-E seemed to contain multiple hinges and local distortions that accommodated sharp bends. Analysis of the CENP-E sequence showed that its predicted coiled-coil structure is disrupted >20 times, almost always by segments containing proline or glycine residues (Fig. 2 D). Although it is unknown whether all of the predicted coiled-coil disruptors actually loop out from the coiled-coil axis, the discontinuities in the coiled coil are the mostly likely source of the high flexibility of CENP-E.

Measurement of hydrodynamic properties of full-length CENP-E in solution confirmed a highly elongated, flexible coiled-coil domain. A native molecular mass of 590 kD for recombinant Xenopus CENP-E was calculated from the measured values determined by sucrose gradient sedimentation and gel filtration of sedimentation coefficient (8.6 S) and Stokes radius (16 nm), respectively (Fig. S2, A and B; available at http://www.jcb.org/cgi/content/full/jcb.200802189/DC1). This value was close to the predicted molecular mass of a CENP-E homodimer (680 kD), which is consistent with our evidence from electron microscopy that CENP-E is dimeric. Calculating the frictional ratio (f/f₀) using the mass of the dimer (680 kD) and the measured Stokes radius produced a value of 2.8 (Fig. S2 C). If CENP-E were a rigid prolate ellipsoid with an axial ratio of ~100 (230-nm contour length/2-nm coiled-coil diameter), the f/f₀ would be >4. With the f/f₀ of 2.8, assuming no hydration and a prolate ellipsoid...
shape, the maximum possible axial ratio (a/b) of the CENP-E dimer was calculated (Harding and Colfen, 1995) to be 45 with an estimated length of 148 nm, which is obviously much shorter than CENP-E’s actual contour length as determined by electron microscopy. Therefore, CENP-E in solution is substantially elongated on average but is not rigidly extended, which is consistent with the high flexibility observed by electron microscopy.

**Full-length CENP-E is a slow plus end-directed kinetochore motor**

The characteristics of full-length CENP-E as a motor were determined using polarity-marked fluorescent microtubules and GFP-tagged full-length CENP-E (366 kD) tethered to a coverslip (Fig. 3 A). The directionality of CENP-E has been controversial over the years. Partial purification of native CENP-E was initially reported to contain minus end-directed motility (Thrower et al., 1995), whereas later the recombinant motor domain was shown to be a plus end-directed motor (Wood et al., 1997). It was also reported that native CENP-E purified from mitotic HeLa cells did not support any motility (DeLuca et al., 2001). In this study, we unambiguously show that full-length recombinant CENP-E powered motility toward plus ends of microtubules, with microtubule gliding seen solely with the bright minus end leading. The mean velocity of gliding was 30 ± 7.6 nm/s (n = 112; Fig. 3 B; Video 4, available at http://www.jcb.org/cgi/content/full/jcb.200802189/DC1). Although the speed difference between the motor domain and the full-length CENP-E could reflect autoinhibition of the C-terminal tail binding to the motor domain as recently proposed by Espeut et al. (2008), none of our electron micrographs (Fig. 2 B) offered support for motor binding to the tail of CENP-E.

**CENP-E motor activity is essential for metaphase chromosome alignment**

It has been well established that CENP-E is essential for stable microtubule capture at the kinetochore and is required for chromosome alignment (Schaar et al., 1997; Wood et al., 1997; Yao et al., 2000; Putkey et al., 2002; Weaver et al., 2003). To test whether it is the motor activity of CENP-E that is essential for chromosome alignment, we used the Xenopus egg extract system in which mitotic spindles can be assembled in vitro. When added to Xenopus extracts before spindle assembly (Fig. 3 C), CENP-E–GFP localized to kinetochores and remained kinetochore bound throughout metaphase (Fig. 3 D and Video 5, available at http://www.jcb.org/cgi/content/full/jcb.200802189/DC1) and at least until the early stage of anaphase (not depicted).

The contribution of CENP-E motor activity to chromosome alignment and its maintenance was tested by assembling spindles in CENP-E–depleted Xenopus extracts that were supplemented with either wild-type CENP-E or motor-dead CENP-E proteins (Fig. 4, A and B). To make a motor-dead CENP-E, we introduced a single point mutation in the ATP-binding P loop (T91N mutation) in the highly conserved kinesin motor domain of CENP-E (Nakata and Hirokawa, 1995).
This mutation provided a rigor binding to microtubules that could not be dissociated even in the presence of ATP, and the motor domain of CENP-E \( \text{T91N} \) failed to support microtubule gliding even in the presence of saturating ATP concentrations (unpublished data). As reported previously (Wood et al., 1997), chromosomes failed to align properly in the majority of bipolar spindles in CENP-E-depleted extracts (Fig. 4 C). When wild-type CENP-E was added back into CENP-E-depleted extracts before spindle assembly, chromosome alignment defects were rescued in part with the CENP-E localized to kinetochores (Fig. 4 C and D; and Fig. S3, available at http://www.jcb.org/cgi/content/full/jcb.200802189/DC1). Although still bound to kinetochores, rigor CENP-E \( \text{T91N} \) failed to rescue chromosome alignment, with the chromosomes instead spread along the spindle axis and located close to the spindle poles (Fig. 4, C and D). Accumulation at the broad region of spindle poles in CENP-E \( \text{T91N} \)-added extracts (Fig. S3) presumably reflected initial rigor binding all along spindle microtubules followed by poleward transport by flux.

Thus, CENP-E motor activity rather than simple microtubule binding is essential for accurate chromosome congression. This conclusion is tempered by the realization that in Xenopus extract spindles, chromosomes are prepositioned at the center (precongressed) during spindle assembly, and the process of chromosome alignment may differ from the typical prometaphase congression in somatic cells (Sawin and Mitchison, 1991). Nevertheless, it is clear that the plus end-directed motility of CENP-E, at a minimum, is required for maintenance of chromosome alignment even in a system with fewer required chromosome movements for establishing an initial alignment. We believe that the strong misalignment phenotype from CENP-E depletion in Xenopus extracts (Fig. 4), in which fast microtubule poleward flux dominates the forces acting on kinetochores (Maddox et al., 2003a), further supports CENP-E as an essential component for powering plus end-directed chromosome movement to counteract flux.

**CENP-E as a motile, flexible kinetochore tether**

Use of electron tomography has shown that the vertebrate kinetochore is composed of a fibrous protein network with multiple microtubule interactions (Dong et al., 2007). CENP-E is certainly part of the fibrous network for kinetochore attachment (as demonstrated with immunogold electron microscopy; Cooke et al., 1997; Yao et al., 1997), with its 230-nm length likely to be one of the longest fibers extending out from the kinetochore to capture microtubules. From our evidence, it is now clear that the long coiled coil of CENP-E could work advantageously for the...
initial capture of kinetochores to spindle microtubules by searching a large radial volume in cells (Fig. 5 E). After initial capture, the combination of slow processive movement along with one-dimensional diffusion would yield the translocation of CENP-E to the plus ends of microtubules, potentially damping with one-dimensional diffusion would yield the translocation of capture, the combination of slow processive movement along searching a large radial volume in cells (Fig. 5 E). After initial capture of kinetochores to spindle microtubules by congression in cells (Fig. 3 D; Skibbens et al., 1993). Furthermore, the high flexibility of CENP-E would allow multiple molecules to work together along the kinetochore fibers without forcing each other into unproductive conformations.

The CENP-E family kinesin (kinesin-7) has a longer family-specific neck domain than conventional kinesins (Endow, 1999). We hypothesize that the longer neck and highly flexible, discontinuous coiled-coil domain of CENP-E are specialized for three roles during mitosis: efficient microtubule capture by kinetochores, towing monooriented chromosomes through viscous cytoplasm (especially those initially far from the spindle equator), and, perhaps most importantly, as a motile kinetochore tether maintaining linkage to dynamic kinetochore microtubule plus ends. The last of these properties couples the slow, processive motility with one-dimensional diffusion that together offer a molecular explanation for how relatively few CENP-E molecules (≈50 dimers per human kinetochore; Brown et al., 1994) can sustain kinetochore attachment to individual dynamic (growing and shrinking) microtubules without losing connection. Combined with the highly flexible 230-nm-long coiled coil, slow plus end–directed motility, and ability of single molecules to remain microtubule bound for minutes, we propose that CENP-E is a part of the kinetochore slip clutch that is engaged on fluxing kinetochore microtubules (Maddox et al., 2003a). In effect, CENP-E combines a slow but processive motor, flexibility, reach, and stable microtubule binding to produce a motile molecular Velcro at kinetochores (Fig. 5).

Materials and methods

Cloning, expression, and purification of CENP-E
cDNA encoding Xenopus CENP-E residues 1–472 fused at the C terminus to a GFP-6His tag was cloned into pET23d (Novagen), and protein expression was induced at 13°C for ∼12 h with 10 nM IPTG in Rosetta (DE3). Bacterial pellets were suspended in lysis buffer (25 mM K-Pipes, pH 6.8, 300 mM KCl, 40 mM imidazole, 2 mM MgCl₂, 0.5 mM EGTA, 10 mM β-mercaptoethanol, 0.1 mM ATP, 1 mM PMSF, and protease inhibitors [from cocktail tablet; Roche]) and lysed by sonication after lysozyme treatment.

The high flexibility of CENP-E would allow multiple molecules to work together along the kinetochore fibers without forcing each other into unproductive conformations. (B) The 230-nm-long coiled coil of CENP-E functions as a safety catch for disassembling microtubules detached from the core kinetochore attachment components, thereby stabilizing the microtubule and enabling rescue. (C) CENP-E is likely to be a part of the kinetochore slip clutch that is engaged on fluxing kinetochore microtubules with its slow plus end–directed motility (Maddox et al., 2003a). CENP-E bound to the microtubule surface may affect kinetochore microtubule plus ends, thereby promoting growth and allowing recapture. (D) Unlike other shorter and more rigidly structured kinetochore capture components, multiple CENP-E molecules are likely to work together by allowing the simultaneous attachment at many different microtubule orientations relative to the kinetochore axis without forcing each other into unproductive conformations. (E) The highly flexible extended coiled coil of CENP-E mediates the initial capture of microtubules by searching a large volume in cells. (F) Its slow, processive motility powers monooriented chromosomes to congress using an adjacent kinetochore fiber (Kapoor et al., 2006).
(QIAGEN) for 1 h at 4 °C. The protein was further purified by HiTrap SP HP (GE Healthcare) using 100 mM–1 M KCl gradient elution in 25 mM Pipes, pH 6.8, 2 mM MgCl₂, 1 mM DTT, and 0.1 mM ATP. The peak fraction from the sulphopropyl column was then loaded onto Superose 6 10/300 (GE Healthcare) preequilibrated with the same buffer containing 100 mM KCl, and the peak fraction was used in the assay.

Full-length Xenopus CENP-E (340 kD) and CENP-E–GFP (366 kD) were expressed and purified from High Five cells (Invitrogen) as previously described (Abrieu et al., 2000) followed by a final purification with a Superose 6 column. To make rigor CENP-E, the Thr at residue 91 was mutated to Asn (ACG → AAC) using site-directed mutagenesis. Fresh proteins were used in all experiments with the exception of the Xenopus extracts experiment.

Sucrose gradient sedimentation and gel filtration chromatography

To measure the S value of CENP-E, 100 μl of protein was centrifuged through 2 ml of 5–40% sucrose gradient at 50,000 rpm for 6 h at 4 °C (TLS 55 rotor; Beckman Coulter). The sedimentation coefficient of CENP-E was determined by linear interolation of a standard curve (R² = 0.96) using proteins of known S values (BSA, 4.6 S; aldolase, 7.3 S; catalase, 11.3 S; thyroglobulin, 19 S). To measure the Stokes radius of CENP-E, Sephacryl S-500 HR (GE Healthcare) was packed into a column (XX 26/70; GE Healthcare) calibrated with proteins standards [20 nm myosin II, 12.5 S, IgM, 8.5 nm thyroglobulin, 5.2 nm catalase, and 4.8 nm aldolase; R² = 0.99]. 500 μl of partially purified protein was loaded, and CENP-E was detected by immunoblotting. The partial specific volume (v) of CENP-E was calculated to be 0.723 ± 0.002/ml from the mean partial specific volumes of the individual amino acids as described by Perkins (1986). Native molecular weight and frictional ratio (f/f₀) were calculated as described in Harding and Cölfen (1995) assuming no hydration and 50% weight and frictional ratio (f/f₀) were calculated as described previously (Abrieu et al., 2000) assuming no hydration and a prolate ellipsoid. The dimensions for CENP-E were calculated from the volume of a spherical mass of 680 kD and the partial specific volume, with V = a × M/A = 4/3, where M represents molecular weight, A represents Avogadro’s number (6.02 × 10²³), and a = 45b.

Single-molecule assays

CENP-E single-molecule assays were performed at room temperature using a digital TIRF imaging system described in detail in Adams et al. (2004). In brief, this consisted of a microscope (TE2000U; Nikon) equipped with a custom-modified TIRF epiluminoator and a 100× NA 1.45A TIRF objective (Nikon). Microscope automation, image acquisition, and processing were performed using MetaMorph software (MDS Analytical Technologies). TIRF images using an evanescent field illumination depth of 150 nm were acquired by the 488- and 568-nm lines of a 50-mW KrAr laser, with individual lines set by a custom-modified TIRF epiluminoator and a 100× NA 1.45A TIRF objective (Nikon). Microscope automation, image acquisition, and processing were performed using MetaMorph software (MDS Analytical Technologies). Flow cells were constructed with a slide and 22 × 22-mm square coverslip separated by two strips of double-stick tape. For measuring fluorescence intensity and photobleaching, −0.5 nM of motors was incubated for 10 min. Fluorescence intensity was measured by integrating the signal of a fluorescent spot per frame and subtracting the background (integrated area = 49 pixels). For motility assays, 50 μg/ml of a rat monoclonal tubulin antibody (Serotec) was incubated for 5 min followed by BRB80 (80 mM K-Pipes, pH 6.8, 1 mM MgCl₂, and 1 mM EGTA) wash, X-horodamine–labeled GMPPCP microtubules for 10 min, and 0.5 mg/ml casein-blocking solution for 10 min. Then, −0.5–1 nM CENP-E motors in motility buffer with 200 mg/ml scavenging system (BRB80, 1 mM DTT, 3 mM ATP, 4.5 mg/ml glucose, 0.2 mg/ml glucose oxidase, and 35 μg/ml catalase) was flowed into the chamber. Frames were captured every 5 s with 120 ms exposure (three frames with 40 ms exposure were averaged to increase the signal to noise ratio), and the duration of imaging was ~5–20 min, which results in a total exposure of ~7–28.8 s. Events that lasted more than two frames (>10 s) were tracked, and nonmoving molecules were excluded from our analysis. CENP-E processive runs were analyzed by drawing kymographs with MetaMorph (MDS Analytical Technologies), and for MSD analyses, the intensity centroid of moving CENP-E spots was tracked using the MetaMorph Track Object tool. Particle coordinates (xᵢ, yᵢ) and xᵢ = x(n+1) and yᵢ = y(n+1) were used to calculate MSD: 

$$\text{MSD}(n) = \sum_{i=0}^{N} \left[ (x_{i+1} - x_i)^2 / (N + 1) \right] + \sum_{j=0}^{N} \left[ (y_{i+1} - y_i)^2 / (N + 1) \right]$$

where ΔT is the data acquisition time interval (Qian et al., 1991). MSD was fitted with the equation for biased Brownian movement: MSD = a × v² + 2D (D, diffusion coefficient, v, mean velocity).

Microtubule gliding assay

Microtubule gliding assays were performed as described previously (Wood et al., 1997) with use of GFP antibody to coat CENP-E–GFP onto a coverslip. Time-lapse image acquisition was performed at room temperature using an inverted microscope (Eclipse TE 300; Nikon) with a 60× NA 1.4A objective, and the images were captured with a camera (CoolSNAP HQ; Photometrics) controlled by MetaMorph.

Electron microscopy

Freshly prepared CENP-E was maintained on ice overnight and processed for freeze drying the following morning. CENP-E proteins were adsorbed to a suspension of mica flakes followed by freeze drying and platinum replication (Heuser, 1989).

Spindle assembly, live imaging, and immunofluorescence in Xenopus extract

The preparation of Xenopus cytoskeletal factor–arrested egg extracts and the cyclic spindles assembly in vitro were performed as described previously (Desai et al., 1999b). For live imaging, ~5 μg/ml X-horodamine tubulin and 25 μM of recombinant CENP-E–GFP were added in the Xenopus extracts before spindle assembly, and squashed extract was imaged using a spinning disc confocal mounted on a microscope (TE2000e; Nikon) with a 100× NA 1.4A objective (Nikon) and 2 × 2 binning as described in Maddox et al. (2003b). CENP-E immunodepletion and immunofluorescence were performed as described previously (Abrieu et al., 2000). More than 95% of CENP-E was consistently depleted using this method. For rescue experiments, either wild-type or rigor CENP-E protein was added to CENP-E-depleted extract at endogenous level (~25 nM) after completion of DNA replication.

Online supplemental material

Fig. S1 shows that CE473-GFP forms a dimer. Fig. S2 shows hydrodynamic measurements demonstrating that purified full-length CENP-E in solution is dimeric with high flexibility. Fig. S3 shows CENP-E localization in the Xenopus extract spindles in Fig. 4. Video 1 shows processive movements of CE473-GFP, and Video 2 shows processive movements of Qdot525-labeled CENP-E (aa 1–53). Video 3 shows backward movement of Qdot525-labeled CENP-E (aa 1–473). Video 4 shows a microtubule gliding assay with CENP-E–GFP (366 kD) and polarity-marked microtubules. Video 5 shows that recombinant CENP-E–GFP is localized to kinetochores in Xenopus extract spindles. Online supplemental material is available at http://www.jcb.org/cgi/content/full/jcb.200802189/DC1.

We are grateful to Arshad Desai (University of California, San Diego, La Jolla, CA) for advice and reagents for assembling micromotors, Paul Maddox for encouragement, Ian Schroeder for help with TIRF imaging, Robyn Roht for help with EM, Ron Vale (University of California, San Francisco, San Francisco, CA) for providing K560-GFP construct, and members of the Cleveland laboratory for stimulating discussions.

This work was supported by a National Institutes of Health grant (GM29513) to D.W. Cleveland. Salary support for D.W. Cleveland is provided by the Ludwig Institute for Cancer Research.

Submitted: 28 February 2008
Accepted: 1 April 2008

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