C-Nap1, a Novel Centrosomal Coiled-Coil Protein and Candidate Substrate of the Cell Cycle-regulated Protein Kinase Nek2

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Abstract. Nek2 (for NIMA-related kinase 2) is a mammalian cell cycle–regulated kinase structurally related to the mitotic regulator NIMA of Aspergillus nidulans. In human cells, Nek2 associates with centrosomes, and overexpression of active Nek2 has drastic consequences for centrosome structure. Here, we describe the molecular characterization of a novel human centrosomal protein, C-Nap1 (for centrosomal Nek2-associated protein 1), first identified as a Nek2-interacting protein in a yeast two-hybrid screen. Antibodies raised against recombinant C-Nap1 produced strong labeling of centrosomes by immunofluorescence, and immunoelectron microscopy revealed that C-Nap1 is associated specifically with the proximal ends of both mother and daughter centrioles. On Western blots, anti–C-Nap1 antibodies recognized a large protein (>250 kD) that was highly enriched in centrosome preparations. Sequencing of overlapping cDNAs showed that C-Nap1 has a calculated molecular mass of 281 kD and comprises extended domains of predicted coiled-coil structure. Whereas C-Nap1 was concentrated at centrosomes in all interphase cells, immunoreactivity at mitotic spindle poles was strongly diminished. Finally, the COOH-terminal domain of C-Nap1 could readily be phosphorylated by Nek2 in vitro, as well as after coexpression of the two proteins in vivo. Based on these findings, we propose a model implicating both Nek2 and C-Nap1 in the regulation of centriole–centriole cohesion during the cell cycle.

The closest known mammalian relative to NIMA is a kinase termed Nek2 (for NIMA-related kinase 2)1 (Fry and Nigg, 1997). This kinase undergoes cell cycle–dependent changes in abundance and activity, reminiscent of NIMA (Schultz et al., 1994; Fry et al., 1995). It is highly expressed in male germ cells (Rhee and Wolgemuth, 1997; Tanaka et al., 1997), and data have been reported consistent with a role for Nek2 in meiotic chromosome condensation (Rhee and Wolgemuth, 1997). However, overexpression of active Nek2 in somatic cells has no obvious effect on chromosome condensation; instead, it induces striking alterations in the structure of the centrosome, the principal microtubule-organizing center of mammalian cells (Fry et al., 1998). Furthermore, immunofluorescence microscopy and subcellular fractionation concur to demonstrate that endogenous Nek2 associates with centrosomes, strongly suggesting that one physiological function of this kinase may relate to the centrosome cycle (Fry et al., 1998).

1. Abbreviations used in this paper: CDK, cyclin-dependent kinase; C-Nap1, centrosomal Nek2-associated protein 1; IEM, immunoelectron microscopy; INMP, intranuclear matrix protein; Nek2, NIMA-related kinase 2; PCM, pericentriolar material; SPB, spindle pole body; γ-TuRCs, γ-tubulin-containing ring complexes.
The mammalian centrosome is an organelle of about 1 μm in diameter. It comprises two barrel-shaped centrioles that are made of nine short triplet microtubules and are surrounded by an amorphous matrix known as the pericentriolar material (PCM) (for review see Brinkley, 1985; Vorobjev and Nadezhdina, 1987; Kimble and Kuriyama, 1992; Kalt and Schliwa, 1993; Kellogg et al., 1994; Lange and Gull, 1996). Major progress has recently been made with the demonstration that microtubules are nucleated from γ-tubulin-containing ring complexes (γ-TuRCs), which are concentrated within the PCM (Moritz et al., 1995; Zheng et al., 1995). γ-Tubulin forms complexes with Spc97/98, two evolutionarily conserved proteins first identified in budding yeast spindle pole bodies (Geissler et al., 1992; Kalt and Schliwa, 1993; Stearns and Wines, 1997), and there is also evidence for an important role of pericentrin and other coiled-coil proteins in organizing γ-TuRCs into higher order lattice structures (Doxsey et al., 1994; Dictenberg et al., 1998). However, in spite of this recent progress, it is clear that the inventory of centrosome components is far from complete.

Centrosome structure and function is regulated in a cell cycle–dependent manner (for reviews see Mazia, 1987; Kellogg et al., 1994; Tournier and Bornens, 1994). Once in every cell cycle, and beginning around the G1/S transition, centrioles are duplicated (e.g., Kuriyama and Borisy, 1981a; Vorobjev and Chentsov, 1982; Kohanski and Borisy, 1990; Chrétien et al., 1997). Late in G2, centrosomes then grow in size (a process referred to as maturation) through the recruitment of additional PCM proteins (Rieder and Borisy, 1982; Kalt and Schliwa, 1993; Lange and Gull, 1995). At the G2/M transition, the duplicated centrosomes separate and migrate to opposite ends of the nucleus. Concomitantly, their microtubule-nucleating activities increase dramatically in preparation for spindle formation (McGill and Brinkley, 1975; Snyder and McIntosh, 1975; Gould and Borisy, 1977; Kuriyama and Borisy, 1981b; for reviews see Brinkley, 1985; Vorobjev and Nadezhdina, 1987; Karsenti, 1991). By what mechanisms these events are controlled remains largely unknown, but data obtained using phosphoepitope-specific antibodies strongly suggest that phosphorylation of centrosomal proteins plays a major role (Vandré et al., 1984, 1986; Centonze and Borisy, 1990). More direct support for this view stems from the observation that cyclin-dependent kinases (CDKs) enhance the microtubule-nucleating activity of centrosomes at the G2/M transition (Verde et al., 1990, 1992; Buendia et al., 1992) and are involved in promoting centrosome separation (Blangy et al., 1995; Sawin and Mitchison, 1995). Similarly, polo-like kinase 1, a cell cycle regulatory kinase structurally distinct from CDKs, has recently been implicated in centrosome maturation (Lane and Nigg, 1996).

The precise role of Nek2 at the centrosome remains to be determined, but it is intriguing that overexpression of this kinase in human cells causes a pronounced splitting of centrosomes. This led us to propose that Nek2-dependent phosphorylation of previously unidentified proteins may cause a loss of centriole–centriole cohesion, and that this event might represent an early step in centrosome separation at the G2/M transition (Fry et al., 1998). With the aim of identifying potential substrates (or regulators) of Nek2, we have now performed a yeast two-hybrid screen, using full-length Nek2 as a bait. We report here the molecular characterization of a novel coiled-coil protein that we call C-Nap1 (for centrosomal Nek2-associated protein 1). C-Nap1 represents a core component of the mammalian centrosome and the first candidate substrate for a member of the NIMA protein kinase family to be identified.

Materials and Methods

Two Hybrid Interaction Screening

A yeast two-hybrid screen was performed essentially as described previously (Durfee et al., 1993; Harper et al., 1993; Blangy et al., 1997). In brief, yeast strain Y190 expressing the fusion protein GAL4 DBD-Nek2 was transformed with a human B cell cDNA library constructed in plasmid pACT (Durfee et al., 1993) and plated on selective medium (lacking tryptophan, leucine, and histidine) supplemented with 50 mM 3-amino triazole. From 4 × 10⁶ transformants, 43 His⁺ colonies were obtained, of which 18 were positive for LacZ expression. Bait loss through cycloheximide selection was only successful for 11 out of 18. Of these, eight remained LacZ positive when mated with Y187 carrying DBD-Nek2, but were negative when mated with Y187 carrying DBD-p53, -lamin, -cdk2, -SNF1, or -Eg5. The library cDNAs in these eight positives were amplified by PCR, using pACT2-specific primers (AAAT CACTACAATTGATGATG and GCTCTAGAAGTGAATGATCCGCGG), and the PCR products were digested with the frequently cutting restriction endonuclease, Aul, to determine the number of different clones. Three clones (2-6, 2-27, and 2-31) contained an identical 1.7 kb insert, while a fourth clone (2-2) was slightly larger (1.9 kb) and yielded a very similar digestion pattern. For subsequent analyses, the PCR products from clones 2-6 and 2-2 were digested with EcoRI and XbaI and subcloned into the EcoRI-XbaI sites of pBlueScript-KS vector carrying the myc-epitope tag (Schmidt-Zachmann and Nigg, 1993) to generate the plasmids pBSmyc2-6 and pBSmyc2-2. To construct the eukaryotic expression plasmid PCMVmyc2-6, the myc2-6 fusion was excised from pBSmyc2-6 on a HindIII-XbaI fragment and subcloned into the HindIII-XbaI sites of pRCMV (Invitrogen Corp., Carlsbad, CA).

Antibody Production

To generate antibodies, a (His)₉-tagged fusion protein of clone 2-6 was made using the QIAexpress bacterial expression system (QIAGEN, Inc., Valencia, CA). The plasmid pQE102-6 was constructed by subcloning a BamH1-XbaI (blunt end with Klenow) fragment from pBSmyc2-6 into the pQE10 vector (QIAGEN) digested with BamHI and HindIII (blunt ended). Overexpression was found to be optimal in the Escherichia coli strain M15[pREP4] using Super medium (25 g bacto-tryptone, 15 g bacto-extract yeast, and 5 g NaCl per liter). Recombinant protein was expressed and purified under denaturing conditions as described by the manufacturer (QIAGEN), before concentrating on a Centricron-30 column (Amicon, Inc., Beverly, MA) and further purification on a preparative 12% SDS–polyacrylamide gel as described in Fry et al. (1998). For immunizations, 500 μg of the purified (His)₉-2-6 protein was injected subcutaneously into New Zealand white rabbits (Elevage Scientifique des Dombes, Chatillon sur Chalaronne, France) on days 1, 28, 56, and 84. Immune serum was obtained on days 70 and 98. For affinity purification of antibodies, two different methods were used. In the first, (His)₉-2-6 protein was expressed in E. coli, purified under native conditions and bound to a Ni²⁺-resin column as described by the manufacturer (QIAGEN), before concentrating on a Centricron-30 column (Amicon, Inc., Beverly, MA) and further purification on a preparative 12% SDS–polyacrylamide gel as described in Fry et al. (1998). The purified antibody was dialyzed extensively against PBS before use. In the second approach, 1.0 mg of purified (His)₉-2-6 protein was covalently coupled to 400 mg CNBr-activated Sepharose 4B (Pharmacia Biotech, Piscataway, NJ) as recommended by the manufacturer, and this affinity matrix was used to purify antibodies from immune serum as described by Harlow and Lane (1988).

Cell Culture and Transfections

Human U2OS osteosarcoma and KE37 T-lymphoblastoid cells were grown at 37°C in a 7% CO₂ atmosphere in DME supplemented with 10% heat-inactivated FCS and penicillin-streptomycin (100 IU/ml and 100 μg/ml,
U2OS cells were seeded onto HCl-treated glass coverslips at a density of 1 × 10^5 cells per 35-mm dish and transfected with 10 μg of plasmid DNA, using calcium phosphate precipitates as previously described (Krek and Nigg, 1991).

**Immunofluorescence and Immunoelectron Microscopy**

Immunofluorescence microscopy on U2OS cells was performed as described in Fry et al. (1998), using a Zeiss Axioskop II microscope (Thornwood, NY) and 40 or 63 × oil immersion objectives. Affinity-purified anti-C-Nap1 antibodies (R62 and R63) were used at 1.0 μg/ml. Photographs were taken using a Quantix 1400 CCD camera (Photometrics, Inc., Tuscon, AZ) and IP-Lab software, and the images were processed using Adobe Photoshop (San Jose, CA).

Negative staining immunoelectron microscopy of isolated centrosomes was performed by diluting centrosomes 1:10 in PBS before sedimentation onto Pioloform and carbon-coated grids using an airfuge (65,000 g-av, 15 min; Beckman Instruments, Inc., Fullerton, CA). Grids were blocked with 0.1% gelatin in PBS, incubated with affinity-purified anti-C-Nap1 IgGs (R63, 1 μg/ml in blocking buffer, 45 min) and protein A–6-nm gold (60 min; Slot and Geuze, 1985). Grids were negatively stained with Nano-W (Nanoprobes, Stony Brook, NY). For preembedding electron microscopy of whole cells, U2OS cells, grown on coverslips, were fixed with 3% paraformaldehyde/2% sucrose for 10 min, permeabilized with PBS (0.1% Tween-20). All antibody incubations were carried out in blocking buffer, and bound IgGs were visualized using alkaline phosphatase–conjugated anti–rabbit IgG secondary antibodies (Promega Corp., Madison, WI).

To prepare cell extracts for immunoblotting, cells were harvested and washed once in PBS + 1 mM PMSF before resuspending in extraction buffer (50 mM Tris-HCl, pH 7.5, 1% Triton X-100, 150 mM NaCl, 5 mM EDTA, 1 mM PMSF, 1 mg/ml aprotinin, 1 μg/ml leupeptin, 1 μg/ml pepstatin A, 20 mM β-glycerophosphate) to give 10^6 cells/μl. Extracts were left 30 min on ice, passed 10 times through a 27-g needle, and centrifuged at full speed in a microfuge at 4°C for 3 min before analysis by SDS-PAGE. Immunoblotting was performed by electrophoretic transfer onto nitrocellulose membranes using a T3 polymerase with the TNT Coupled Reticulocyte Lysate System (Promega Corp.) in the presence of [35S]methionine (Amersham Corp., Arlington Heights, IL). 10 μl of the in vitro translation reaction was then mixed with 50 μl of NeK2 kinase buffer (Fry and Nigg, 1997), in the presence or absence of [32P-γ]ATP, and added to baculovirus-expressed wild-type or catalytically inactive NeK2 immunoprecipitates prepared as described in Fry and Nigg (1997). Samples were incubated at 30°C for 20 min, diluted 1:5 with NEB buffer (Fry and Nigg, 1997), and spun briefly in a microfuge, and the supernatant was recovered. For immunoprecipitation of myc-tagged proteins, samples were preclarified for 1 h with protein G-Sepharose beads (Pharmacia Biotech) and then incubated for 1 h at 4°C with protein G-Sepharose beads previously coupled with anti-myc monoclonal antibody 9E10. Immunoprecipitates were finally washed four times in NEB buffer, resuspended in 50 μl of protein sample buffer, and analyzed by SDS-PAGE.

**cDNA Library Screening and Sequence Analysis**

Overlapping cDNA clones spanning the entire coding sequence of C-Nap1 were isolated from a human placenta 5'-STRETCH PLUS cDNA library (CLONTECH Laboratories, Palo Alto, CA). DNA probes were labeled with 32P using either the random-primed DNA labeling kit (Boehringer Mannheim, Mannheim, Germany) for larger DNA fragments or polynucleotide kinase (Biofinaex, Praroman, Switzerland) for oligonucleotide probes. Positive phages were purified, lambda DNA was prepared from agarose plate lysates using the QIAGEN Lambda Kit, and inserts were excised and subcloned into the NotI site of a pBlueScript-KS vector. DNA sequencing in both directions was carried out using a 377 DNA Automated Sequencer (Perkin-Elmer Corp., Norwalk, CT).

**Miscellaneous Techniques**

Isolation of human centrosomes from the human T-lymphoblastoid cell line KE37 was carried out as described previously (Moudjou and Bornens, 1994). Treatment of asynchronously growing U2OS cells with nocodazole or taxol was as described in Fry et al. (1998), while cold treatment was performed by placing the tissue culture dishes on ice for 30 min. RNA purification from mouse organs and RNase protection assays were performed as described in Tanaka et al. (1997). To prepare an antisense RNA probe for mouse C-Nap1, the transcript complementary to the intranuclear matrix protein (INMP) cDNA sequence (sequence data available from GenBank/EMBL/DDBJ under accession number U33198, nucleotides 1105–1733) was synthesized on an INMP cDNA fragment with T7 RNA polymerase (Promega Corp.) in the presence of 11.25 μM [α-32P]UTP (90 Ci/ mmol, DuPont-NEN, Boston, MA). The template cDNA fragment was isolated by performing a reverse-transcriptase reaction on mouse testis RNA, followed by PCR amplification with specific primers corresponding to the 3' and 5' region of the published INMP sequence (Menz et al., 1996). For normalization, a transcript complementary to the mouse NF-Ya mRNA sequence was used (Tanaka et al., 1997).

For chromosomal mapping, two specific primers (AGCCACACGCAGAGGCACACAGAC and AGAGGAGTACAGTGCCTACA-GCC) were used to amplify a 265-bp PCR fragment in the 3' UTR of C-Nap1 using the Genbridge 4 Radiation Hybrid DNA panel (HGMP Resource Centre, UK). PCR products were analyzed by Southern blotting with a hybridization probe generated by PCR on the C-Nap1 cDNA, using the above oligonucleotides. The results were analyzed at http://www.genome.wi.mit.edu.

**Results**

**Identification of a Centrosomal Protein through Two-Hybrid Interaction with NeK2**

To identify cellular proteins that might interact with NeK2, a yeast two-hybrid screen was performed (Fields and Song, 1989; Durfee et al., 1993) using the full-length NeK2 as a bait. A human cDNA library was screened, and library clones able to activate both the auxotrophic selection marker His3 and the lacZ reporter gene specifically in the presence of the bait were selected. Among the clones showing a specific interaction with NeK2, three (designated 2-6, 2-27, and 2-31) were found to contain identical inserts of 1.7 kb, while a fourth (2-2) encoded a slightly longer version (1.9 kb) of the same cDNA. To determine which domains of NeK2 were required for the observed interactions, clones 2-6 and 2-2 were tested in the yeast two-hybrid assay against different NeK2 constructs (data not shown): both clones interacted strongly with full-length wild-type NeK2, a catalytically inactive mutant of NeK2 (NeK2-K37R; Fry et al., 1995), and the COOH-terminal noncatalytic domain of NeK2. There was also a significant interaction, albeit slightly less strong, with the NH2-termi-
Figure 1. Localization of transfected and endogenous C-Nap1 to the centrosome. (A) U2OS osteosarcoma cells were transiently transfected with myc-epitope-tagged clone 2-6 and fixed with methanol after 24 h. Cells were double-stained with anti-myc (a) and anti-γ-tubulin (b) antibodies. In b, nuclei can be seen due to staining of the DNA with Hoechst dye 33258. (B) Asynchronous U2OS cells were fixed with methanol and stained with anti-C-Nap1 antibodies (a, preimmune R63; b, immune R63; c, immune R62). Cells stained for C-Nap1 (c) were also double-stained for the centrosomal component p150Glued (d), demonstrating colocalization at the centrosome (c and d, arrowheads). Bar, 10 μm.

The above data strongly suggested that the C-Nap1 protein isolated originally by two-hybrid interaction with Nek2 represents a COOH-terminal fragment of a much larger protein. To obtain the full-length sequence, we

The COOH-terminal noncatalytic domain contains a leucine zipper motif that is implicated in Nek2 homodimerization (Fry, A.M., L. Arnaud, and E.A. Nigg, manuscript in preparation). Deletion of this leucine zipper did not reduce the strength of the association between Nek2 and clones 2-6 or 2-2, indicating that this motif was not by itself responsible for the observed interaction.

As a first approach to assess the significance of this interaction, the subcellular localization of the protein encoded by clone 2-6 was examined by transfecting a myc-tagged version of this cDNA into human tissue culture cells. Immunofluorescence microscopy revealed that ectopically expressed myc:2-6 protein localized primarily to the centrosome, as demonstrated by colocalization of antimyc antibodies with antibodies for the centrosomal marker γ-tubulin (Fig. 1 A, a and b). In weakly expressing cells, the myc:2-6 protein appeared to be exclusively associated with the centrosome, but in cells expressing higher levels, myc:2-6 could also be seen in the cytoplasm (Fig. 1 A, a).

To verify this localization, two rabbit antisera (R62 and R63) were raised against bacterially expressed, polyhistidine-tagged 2-6 protein. When used for immunofluorescent staining of human tissue culture cells, both antibodies produced highly specific staining of the centrosome (Fig. 1 B, b and c), whereas no such staining was observed with the corresponding preimmune sera (Fig. 1 B, a and data not shown). The identity of the stained organelles was confirmed by performing double labeling experiments, using R62 in combination with a monoclonal antibody against p150Glued, a component of the dynactin complex that is known to concentrate at the centrosome (e.g., Blangy et al., 1997; Fig. 1 B, d). From these results, we conclude that the protein encoded by cDNA 2-6 is a novel component of the centrosome. Hence, in the following we refer to this protein as C-Nap1.

C-Nap1 Is a High Molecular Mass Coiled-Coil Protein

To determine the size of the endogenous C-Nap1 protein, the anti–C-Nap1 antibodies (R62 and R63) were used for Western blotting. On total extracts of U2OS cells, both anti–C-Nap1 antibodies consistently identified a protein migrating at ~250 kD that was not detected by the corresponding preimmune sera (Fig. 2, lanes 2 and 1, respectively, and data not shown). When similar immunoblots were performed on total extracts prepared from the KE37 T-lymphoblastic cell line, both anti–C-Nap1 antibodies again recognized a protein migrating at 250 kD (Fig. 2, lane 3 and data not shown), and importantly, this protein was substantially enriched in centrosome preparations isolated from these cells (Fig. 2, lane 4). Indeed, the level of enrichment of C-Nap1 in purified centrosomes was very similar to that observed for Nek2, previously determined to be in the order of 100-fold (Fig. 2, lanes 5 and 6; and see Fry et al., 1998). Extrapolating from Nek2, this result also implies that the existence of a noncentrosomal pool of C-Nap1 is to be expected. Interestingly, both C-Nap1 antibodies also recognized a band at ~190 kD in KE37 total cell extracts and centrosomal preparations (Fig. 2, lanes 3 and 4). As this band showed a similar degree of enrichment in centrosome preparations as the full-length C-Nap1 protein, we believe that it may be structurally related (e.g., either represent a breakdown product of C-Nap1 or an alternatively spliced form; see legend to Fig. 3).

The above data strongly suggested that the C-Nap1 protein isolated originally by two-hybrid interaction with Nek2 represents a COOH-terminal fragment of a much larger protein. To obtain the full-length sequence, we
screened a human placenta cDNA library with the available C-Nap1 probe, and after repeated rounds of screening, isolated several overlapping cDNAs spanning a total of 8,347 nucleotides (Fig. 3 A). Conceptual translation of these cDNAs revealed the presence of a continuous open reading frame coding for a protein of 2,442 amino acids, with a calculated molecular mass of 281 kD (Fig. 3 B). Although we have been unable to determine the size of the C-Nap1 mRNA by Northern blot analysis, presumably because of the low abundance of this transcript (see below), we are confident that the isolated cDNAs encompass the complete protein. In fact, the putative initiator methionine is preceded by several stop codons, and the 3' ends of some cDNAs include a polyadenylation signal and a poly(A) tail. Taken together with the results of our immunochemical data, these cDNA sequence analyses indicate that C-Nap1 is a novel, high molecular mass component of the mammalian centrosome. The C-Nap1 gene was mapped by radiation hybrid mapping close to the centromeric region of human chromosome 20, between the framework markers WI-6874 and AFMA152YG1. This places C-Nap1 between the genetic markers D20S195 and D20S908 at approximately 20q11.2.

Examination of the predicted C-Nap1 protein sequence indicates that this protein is likely to form extended coiled-coil domains, with a possible kink or hinge region near the center (Fig. 3 C). Database searches revealed that the COOH terminus of C-Nap1 is almost identical to a previously described mouse protein, termed INMP (see Discussion). Furthermore, while this manuscript was in preparation, a human cDNA corresponding to the first 5 kb of the C-Nap1 sequence reported here was deposited in the Genbank database (under accession number AF022655). In keeping with our results, this independently isolated cDNA was identified with the aid of human autoimmun sera reactive against centrosomal antigens (Mack et al., 1998). Beyond these matches, however, no obvious close relatives of C-Nap1 could be identified in any species, apart from weak similarities to many coiled-coil proteins.

**C-Nap1 Is a Low Abundance Ubiquitous Protein**

Recent studies on the expression of Nek2 have shown that this kinase is highly abundant in testis, and detailed in situ hybridization analyses revealed a stage-specific expression pattern during spermatogenesis (Rhee and Wolgemuth, 1997; Tanaka et al., 1997). To investigate whether C-Nap1 might show a similar tissue-specific expression, we have isolated a cDNA coding for murine C-Nap1 and performed RNase protection experiments. As shown in Fig. 4, C-Nap1 was expressed ubiquitously and at comparable levels in all tissues analyzed, including testis. Thus, there is presently no evidence for a specialized function of C-Nap1 in meiotic processes. It is noteworthy, however, that the intensities of the signals observed with the C-Nap1 probe were similar to those obtained with a control probe for the ubiquitous transcription factor NF-Ya, labeled to a comparable specific activity (Fig. 4). As it has been calculated previously that there are only about 10 copies of the NF-Ya mRNA per cell (Schmidt and Schibler, 1995), our results would indicate that the level of C-Nap1 mRNA may be similarly low.

**C-Nap1 Is a Centriole-associated Core Component of the Centrosome**

To determine whether the centrosomal association of C-Nap1 was dependent upon the presence of microtubules, immunostaining on cells was performed under conditions that perturb the microtubule network. In cells incubated at 4°C for 30 min or treated with nocodazole for 4 h, interphase microtubules were completely depolymerized (Fig. 5 b and c). Yet, in comparison to untreated cells (Fig. 5 a and e), there was no observable decrease in the concentration of C-Nap1 at the centrosome after cold or nocodazole treatment (Fig. 5 f and g). In cells incubated with taxol, on the other hand, the microtubules appeared as dense bundles, which in many interphase cells were detached from the centrosome (Fig. 5 d). In such cells, C-Nap1 was clearly still associated with the centrosome and not with the microtubule bundles (Fig. 5 h). Thus, C-Nap1, like Nek2, associates with centrosomes independently of microtubules, and by this criterion, should be considered as a bona fide component of the core centrosome (Oegema et al., 1995).

To examine the localization of C-Nap1 at the ultrastructural level, immunoelectron microscopy (IEM) was performed, initially using anti-C-Nap1 antibodies on isolated centrosomes. After negative staining IEM, gold particles were specifically found at one end of the centriole (Fig. 6...
Whenever a pair of centrioles was visible, gold particles were consistently present at the proximal ends of each centriole (data not shown). Next, preembedding IEM was carried out, using anti–C-Nap1 antibodies on U2OS human osteosarcoma cells (Fig. 6, b–g). Again, the bulk of gold particles was tightly associated with the proximal ends of the centrioles. There were very few gold particles located elsewhere in the PCM, and little penetration into the lumen of the centriole barrel could be observed. In many sections, one centriole was more heavily labeled than the other; however, when serial sections through a pair of centrioles were compared, both centrioles showed proximal end labeling, suggesting that mother and daughter centrioles were labeled to a similar extent (Fig. 6, b and c). We have also performed preembedding IEM, using anti-Nek2 antibodies on U2OS cells. In this case, we observed a pattern of staining remarkably similar to that seen with anti–C-Nap1 antibodies, with a concentration of Nek2 also at the proximal ends of centrioles (Fig. 6, h–k).

Hence, C-Nap1 and Nek2 demonstrate a striking colocalization within the substructure of the centrosome. Attesting to the specificity of these results, antibodies against γ-tubulin produced a much more diffuse labeling throughout the PCM (data not shown), in agreement with previously published results (Fuller et al., 1995; Moudjou et al., 1996). Also, no centrosome-specific staining was observed when using the immunogold-labeled secondary antibodies alone (data not shown).

**Loss of C-Nap1 Staining on Mitotic Spindle Poles**

The intensity of centrosome staining by anti–C-Nap1 antibodies showed little if any variation during interphase stages of the cell cycle, and fairly uniform staining was also observed when anti–C-Nap1 antibodies were used to stain centrosomes isolated from exponentially growing KE37 cells (data not shown). In striking contrast, however, mitotic spindle poles were poorly labeled by anti–C-Nap1 antibodies, and in some mitotic cells staining of spindle poles was undetectable altogether (Fig. 7). Diminished C-Nap1 staining could be detected at the onset of prophase, concomitant with centrosome separation, increased microtubule-nucleating activity, and chromosome condensation (Fig. 7 A, a–c), and it persisted throughout the remainder of mitosis (e.g., anaphase cell in Fig. 7 A, d and e). The apparent loss of C-Nap1 from mitotic spindle poles is in

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marked contrast to what is seen with many other centrosomal components, such as γ-tubulin, which show an increased accumulation at the spindle poles as a result of centrosome maturation in late G2 (Fig. 7B, compare a–d with e–h). The most straightforward interpretation of these results is that the observed loss of C-Nap1 staining during mitosis reflects a significant reduction in the amount of C-Nap1 protein at mitotic spindle poles. However, we cannot rigorously exclude cell cycle–dependent epitope masking as an alternative explanation.

**Phosphorylation of C-Nap1 by Nek2**

The observed interaction between the COOH-terminal fragment of C-Nap1 and Nek2 raised the possibility that C-Nap1 might be a substrate of Nek2. To examine this possibility, the myc-tagged C-Nap1 clone 2-6 (i.e., the COOH-terminal 53 kD of C-Nap1) was transcribed and translated in a rabbit reticulocyte lysate and incubated in the presence of recombinant Nek2 proteins immunoprecipitated from baculovirus-infected insect cells. In the presence of active Nek2, the myc:2-6 protein showed a drastic reduction in electrophoretic mobility (Fig. 8A, lane 1) and could be radiolabeled by incubation with [32P]-γATP (Fig. 8A, lane 3), whereas neither of these results was observed in the presence of the catalytically inactive Nek2 mutant (Fig. 8A, lanes 2 and 4). Hence, Nek2 could readily phosphorylate the COOH-terminal fragment of C-Nap1 in vitro. To determine whether this was also the case in vivo, the C-Nap1 clone myc:2-6 was cotransfected with either active or inactive Nek2 into U2OS cells. 24 h after transfection, cells were lysed directly into hot sample buffer to prevent reactions proceeding during extraction, and the electrophoretic mobility of the myc:2-6 protein was examined by immunoblotting with anti-myc antibodies. Consistent with the data obtained in vitro, myc:2-6 showed a conspicuously reduced gel mobility when coexpressed with active Nek2 (Fig. 8B, lane 3), but not when expressed alone (Fig. 8B, lane 2) or in combination with the catalytically inactive Nek2 (Fig. 8B, lane 4). Thus, under the conditions used here, the COOH-terminal fragment of C-Nap1 behaves as a good substrate for Nek2 both in vitro and in living cells.

**Discussion**

Studies on the mammalian centrosome began more than a hundred years ago, yet many of its components still await identification, and the mechanisms regulating centrosome function during the cell cycle remain largely unknown. By combining genomic sequence information with analytical mass spectrometry, rapid progress is currently being made...
on the identification of proteins present in spindle pole bodies (SPBs) purified from the yeast Saccharomyces cerevisiae (Rout and Kilmartin, 1990; Wigge et al., 1998). However, although many yeast SPB components may have been conserved during evolution, the substantial morphological differences between fungal SPBs and higher eukaryotic centrosomes suggest that this approach is unlikely to yield a complete inventory of the mammalian centrosome. Thus, complementary approaches for identifying novel centrosome components are needed. One straightforward strategy consists of the search for proteins that are able to interact with already known centrosomal proteins. Here, we have applied this strategy, using the centrosome-associated protein kinase Nek2 as a bait in a yeast two-hybrid screen. Through this approach, we have identified C-Nap1, a novel high molecular mass component of the centrosome. C-Nap1 not only colocalizes with Nek2 at the centrosome, but it also represents the first candidate physiological substrate of this cell cycle-regulated protein kinase.

Both immunocytological studies and biochemical analyses of purified centrosomes demonstrate that C-Nap1 is a bona fide component of the core centrosome. Furthermore, ultrastructural analyses using antibodies raised against the COOH terminus of C-Nap1 revealed a highly concentrated localization of the corresponding epitopes to the proximal ends of both mother and daughter centrioles. Although antigen accessibility is a general concern with preembedding IEM, this strikingly restricted localization is remarkable, as other centrosomal proteins show very different distributions within the centrosome: centrin was reported to be concentrated at the distal ends of centrioles, as well as within their lumen (Paolelli et al., 1996), whereas γ-tubulin has a broader distribution, localizing to the periphery of the PCM, to pericentriolar satellites and to the ends of centriolar and procentriolar microtubules (Fuller et al., 1995; Moudjou et al., 1996). Of those centrosomal proteins whose localization has been determined by electron microscopy, only a vertebrate Spc110p-related protein shows a localization somewhat similar to that described here for C-Nap1 (Tassin et al., 1997). However, while the Spc110p-related protein was reported to be associated preferentially with the mother centriole (Tassin et al., 1997), no biased distribution between mother and daughter centrioles could be detected in the case of C-Nap1, regardless of whether isolated centrosomes were analyzed or centrosomes in situ.

Cloning and sequencing of several overlapping cDNAs revealed that human C-Nap1 is a protein with a predicted molecular mass of 281 kD. The COOH terminus of C-Nap1 is very closely related to a previously described putative nuclear matrix protein, termed INMP, that was isolated by expression cloning using a monoclonal antibody raised to a nuclear matrix preparation (Menz et al., 1996). Based on cDNA sequencing, mouse INMP was reported to comprise 446 residues. However, our sequencing of C-Nap1 cDNAs isolated from both human placenta and mouse testis (Tanaka, K., T. Mayor, and E.A. Nigg, unpublished re...
results) libraries revealed that an open reading frame extends throughout the alleged 5’ untranslated region of INMP. Thus, we believe that INMP most likely represents a COOH-terminal fragment of a mouse C-Nap1 homologue. It would be premature to exclude the existence of C-Nap1–related proteins within the nucleus, but our localization studies provide no evidence for a nuclear localization of the C-Nap1 protein described here. Additional support that C-Nap1 is a bona fide centrosomal protein stems from the independent identification of the same protein in a study on the characterization of human autoimmune sera with centrosomal reactivity (Mack et al., 1998).

On the basis of primary structure analyses, C-Nap1 is predicted to consist almost entirely of coiled-coil domains (Lupas, 1996). Several other recently described centrosomal proteins also contain predicted coiled-coils (for review see Stearns and Winey, 1997), suggesting that protein–protein interactions based on such domains may be ideally suited for the maintenance of a dynamic centrosome structure. The only regions within C-Nap1 that contain clusters of helix-breaking proline residues are located towards the extreme NH2 and COOH termini, and intriguingly, near the center of the protein. Thus, it is possible that C-Nap1 might contain a hinge or even fold back upon itself. Provided that recombinant C-Nap1 can be produced in a soluble form, it will be interesting to determine its shape. If this 281-kD coiled-coil protein were to form a straight rod, then its predicted length would be around 300 nm (comparable to the average intercentriolar distance; e.g., Chrétien et al., 1997). At present, we have no information about the oligomerization state of C-Nap1 or its ability to interact with other coiled-coil proteins. However, electron-dense filaments have been shown to extend between the proximal ends of the two centrioles (Bornens et al., 1987; Paintrand et al., 1992; and references therein), and it is attractive to speculate that C-Nap1 could be a component of a
Figure 8. Phosphorylation of the COOH-terminal fragment of C-Nap1 by Nek2 in vitro and in vivo. (A) The C-Nap1 COOH-terminal construct myc:2-6 was in vitro translated in the presence of [35S]methionine and then mixed with immunoprecipitates of recombinant wild-type Nek2 (lanes 1 and 3) or catalytically inactive Nek2 (lanes 2 and 4) for 20 min at 30°C in the absence (lanes 1 and 2) or presence (lanes 3 and 4) of [35S]ATP. The Nek2 immunoprecipitates were then removed, and the myc:2-6 proteins were immunoprecipitated with anti-myc antibodies. Proteins were resolved on 12% SDS–polyacrylamide gels and subjected to autoradiography. To selectively enhance the signal due to 32P (lanes 3 and 4), an additional piece of x-ray film was used to block the signal caused by 35S; the faint band corresponding to the size of myc:2-6 in lane 4 is most likely due to weak penetration of the 35S signal. Molecular mass markers (kD) are indicated on the left. (B) 10-cm dishes of exponentially growing U2OS cells were transiently transfected with no DNA (lane 1), 25 μg pCMV-myc:2-6 (lane 2), 12.5 μg pCMV-myc:2-6 + 12.5 μg pCMV-Nek2 (lane 3), or 12.5 μg pCMV-myc:2-6 + 12.5 μg pCMV-Nek2-K37R (lane 4). After 24 h, cell extracts were resolved on 12% SDS–polyacrylamide gels and immunoblotted with anti-myc antibodies (top) or anti-Nek2 antibodies (bottom). The C-Nap1 myc:2-6 protein exhibits a significant reduction in gel mobility specifically in the presence of active Nek2 both in vitro and in vivo.

structure connecting the proximal ends of centrioles to each other (Fig. 9).

In this context, it is intriguing that Nek2 is able to phosphorylate the COOH-terminal domain of C-Nap1 both in vitro and in vivo, suggesting that C-Nap1 may be a physiological substrate of Nek2. This invites speculations as to the mechanisms by which overexpression of Nek2 disrupts centrosome structure. In a previous study, we have shown that overexpression of Nek2 in cultured cells produces a striking splitting of centrosomes (Fry et al., 1998). Since this effect was strictly dependent on Nek2 activity, we have proposed that increased Nek2 activity may lead to the degradation or disassembly of a molecular “glue” that holds the two centrosomes in close apposition during most stages of the cell cycle. In the simplest interpretation of this model, the hypothetical glue would be composed of structural proteins, at least one of which would be a direct substrate of Nek2. Phosphorylation of this protein at the appropriate stage of the cell cycle could then either target it for degradation or cause it to dissociate from centrosomes, with both processes leading to a weakening of centriole–centriole cohesion. As C-Nap1 most likely fulfills a structural role and clearly can be phosphorylated by Nek2, it would seem to qualify as a candidate component of this hypothetical centrosomal glue (Fig. 9). We have observed that C-Nap1 immunoreactivity is severely diminished at the spindle poles of mitotic cells, consistent with the notion that the association of C-Nap1 with the centrosome may be cell cycle regulated. However, as yet we have no information on either the timing or the molecular consequences of C-Nap1 phosphorylation by Nek2, and the model shown in Fig. 9 should therefore be considered as a working hypothesis. Testing of this model will require a detailed analysis of C-Nap1 phosphorylation during the cell cycle, and the mapping of Nek2-dependent phosphorylation sites. Such information may then set the stage for directly testing the consequences of overexpressing wild-type and phosphorylation-site mutant versions of C-Nap1 in living cells.

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