N-Acetyl-D-Glucosamine Kinase Interacts with Dynein-Lis1-NudE1 Complex and Regulates Cell Division

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N-acetyl-D-glucosamine kinase (GlcNAc kinase or NAGK) primarily catalyzes phosphoryl transfer to GlcNAc during amino sugar metabolism. Recently, it was shown NAGK interacts with dynein light chain roadblock type 1 (DYNLRB1) and upregulates axo-dendritic growth, which is an enzyme activity-independent, non-canonical structural role. The authors examined the distributions of NAGK and NAGK-dynein complexes during the cell cycle in HEK293T cells. NAGK was expressed throughout different stages of cell division and immunocytochemistry (ICC) showed NAGK was localized at nuclear envelope, spindle microtubules (MTs), and kinetochores (KTs). A proximity ligation assay (PLA) for NAGK and DYNLRB1 revealed NAGK-dynein complex on nuclear envelopes in prophase cells and on chromosomes in metaphase cells. NAGK-DYNLRB1 PLA followed by Lis1/NudE1 immunostaining showed NAGK-dynein complexes were colocalized with Lis1 and NudE1 signals, and PLA for NAGK-Lis1 showed similar signal patterns, suggesting a functional link between NAGK and dynein-Lis1 complex. Subsequently, NAGK-dynein complexes were found in KT and nuclear membranes where KT were marked with CENP-B ICC and nuclear membrane with lamin ICC. Furthermore, knockdown of NAGK by small hairpin (sh) RNA was found to delay cell division. These results indicate that the NAGK-dynein interaction with the involvements of Lis1 and NudE1 plays an important role in prophase nuclear envelope breakdown (NEB) and metaphase MT-KT attachment during eukaryotic cell division.

INTRODUCTION

N-acetylglucosamine kinase (GlcNAc kinase or NAGK; E.C. 2.7.1.59) was first discovered in 1970 (Datta, 1970), and it’s mRNA and proteins were later found to be expressed in a variety of cell lines and tissues (Hinderlich et al., 2000, Ligos et al., 2002). NAGK belongs to the sugar kinase/heat shock protein 70/actin superfamily (Hurley, 1996), which is characterized by a V-shaped fold consisting of two domains with the βββκκαα configuration. Members of this super family bind substrates and undergo a large conformational change involving a reduction of gap angle between these two domains (Hurley, 1996). The canonical function of NAGK is to phosphorylate GlcNAc to produce GlcNAc-6-phosphate in the GlcNAc recycling salvage pathway. The GlcNAc-6-phosphate is subsequently used to produce UDP-GlcNAc that is utilized to synthesize N-O-glycans and glycolipids (Hakomori, 2000; Schachter, 2000; Van den Steen et al., 1998), and glycosaminoglycans (Eska and Lindahl, 2001).

In addition to its enzymatic role as a kinase, our laboratory reported a non-canonical, i.e., a non-enzymatic function for NAGK, which was first implicated in the differential distributions of the protein in different types of brain cells. In the mature rat brain, NAGK is present predominantly in neuronal dendrites and its levels are very low in neuronal axons and in the processes of neuroglial cells, such as, astrocytes and oligodendrocytes (Lee et al., 2014a), suggesting it has a specific role in neuronal dendrites. Furthermore, exogenous overexpression of NAGK promoted dendritic complexity in cultured rat hippocampal neurons, whereas its knockdown caused dendritic degeneration (Lee et al., 2014a). Surprisingly, the overexpressions of kinase-deficient mutant NAGKs also upregulated dendritic arborization and overexpression of the NAGK small domain, but not of the large domain, resulted in dendritic degeneration (Lee et al., 2014b), confirming a non-canonical structural function of NAGK. A similar non-canonical function associated with axonal arborization was observed in the developing axons of hippocampal neurons (Islam et al., 2015b). In order to elucidate the structural role of NAGK, Islam et al. (2015a) performed yeast two-hybrid screening and found NAGK interacted with dynein light chain roadblock type 1 (DYNLRB1). Interaction between NAGK and dynein molecular motor was further confirmed by immunocytochemistry (ICC) and proximity ligation assay (PLA).

By transfecting a peptide derived from the C-terminal DYNLRB1, which interacted with NAGK, interfered with NAGK...
function during neuronal development and resulted in dendrite degeneration and delayed axonal growth (Islam et al., 2015a; 2015b). Interestingly, NAGK-dynein interactions occur in the branch points of developing neuronal dendrites and axons where NAGK-dynein complexes are associated with Golgi outposts (Islam et al., 2015a, 2015b). Since these outposts serve as local microtubule-organizing centers (Or-McKenney et al., 2012), evidence to date suggests that NAGK regulates dynein motor complex during the directional transport of Golgi outposts to branching sites, and/or during the microtubule-organizing function of Golgi outposts in dendritic and axonal branch points.

In contrast to cytoplasmic expression, NAGK is ubiquitously present in nuclei. Yeast two-hybrid selections have shown that NAGK interacts with small nuclear ribonucleoprotein (snRNP)-associated polypeptide N (snRPN), and general transcription factor IIH polypeptide 5 (GTF2H5) and an ICC study showed the colocalization of NAGK signals with speckle and paraspeckles, suggesting its participation in gene expression (Sharif et al., 2015). Furthermore, NAGK ICC signals were reported to surround the cytoplasmic face of the outer nuclear membrane (Sharif et al., 2015). In our previous studies, we found NAGK interacts with cytoplasmic dynein motor complex that is also present on nuclear envelopes (Hu et al., 2013). Others have implicated dynein in prophase nuclear envelope breakdown (NEB) and mitotic chromosome separation based on its microtubule-anchoring activities during cell division (Sharp et al., 2000; Salina et al., 2002). Although the molecular mechanisms responsible for cell division have been substantially elucidated, the ways in which they are organized and controlled to create an effective, accurate cellular machine are yet to be explored. In the present study, we focused on the modulatory effect of NAGK on dynein function during different phases of cell division.

MATERIALS AND METHODS

Antibodies and plasmids

The following antibodies were used at the indicated dilutions: chicken polyclonal NAGK (1:1000 for ICC; GW22347, Sigma, USA); mouse monoclonal NAGK (1:10 for PLA; Santa Cruz Biotechnology, USA); rabbit polyclonal NAGK (1:50 for PLA; GenTex, USA); rabbit polyclonal DYNLRB1/LC7 (1:50 for ICC and 1:25 for PLA; Proteintech Group, USA); mouse polyclonal dynein IC1 (1:25 for PLA; Santa Cruz Biotechnology); rabbit polyclonal Lis1 (Santa Cruz); rabbit polyclonal NudE1 (1:50 for ICC, Proteintech Group); rabbit polyclonal CENP-B (1:500; Abcam, UK); mouse monoclonal alpha-tubulin (1:10; brth preparation, Developmental Studies Hybridoma Bank, University of Iowa, USA); rabbit polyclonal lamin B (1:1000; Young In Frontier Inc., Korea). The fluorescently labeled secondary antibodies Alexa Fluor 488 and 568 (Invitrogen, USA) were used to detect primary antibodies. The plasmids used for transfection were pDsRed2 vector and NAGK short hairpin (sh) RNA, as previously described (Lee et al., 2014a; 2014b).

Cell culture, nocodazole treatment, and transfection

GT1-7 and HEK293T cells were grown on polylysine coated glass coverslips in DMEM (Invitrogen) supplemented with 10% fetal bovine serum and 1% Penicillin Streptomycin. Cells were treated with 70 ng/ml nocodazole (Sigma, USA) for 18.5 h in the same medium. HEK293T cells were transfected with indicated plasmids using Lipofectamine™ 2000 reagent (Invitrogen) according to the manufacturer’s instructions.

Immunocytochemistry (ICC)

Cells were fixed by sequential paraformaldehyde/methanol fixation (Moon et al., 2007). Briefly, cells were incubated in 4% paraformaldehyde in PBS (20 mM sodium phosphate buffer, pH 7.4, 0.9% NaCl) at RT for 10 min, and then incubated in methanol at -20°C for 20 min. Fixed cells were first incubated in preblocking buffer (5% normal goat serum, 0.08% Triton X-100 in PBS, pH 7.4) at 4°C for more than 6 h, then treated with indicated antibodies, and mounted on glass slides using mounting medium containing DAPI.

Proximity Ligation Assay (PLA)

Generic in situ PLA was performed using a Duolink assay kit (Olink Bioscience, Sweden), according to the manufacturer’s instructions with minor modifications. Briefly, cells were incubated with primary antibodies in preblocking buffer (5% normal goat serum, 0.05% Triton X-100 in PBS, pH 7.4) at 4°C overnight, and then washed three times for 20 min in preblocking buffer at RT. For secondary antibodies conjugated with oligonucleotides, PLA probe anti-mouse MINUS and PLA probe anti-rabbit PLUS were diluted in preblocking buffer and applied for 2 h at 37°C in a humidity chamber. Other aspects of the assay were performed according to the manufacturer’s instructions. For the combination of PLAs and ICC, the PLA steps were performed first and then primary antibodies were added to cells overnight at 4°C. Cells were then incubated with fluorophore conjugated secondary antibodies as per the above-mentioned ICC procedure.

Image acquisition

An Olympus microscope BX53 equipped with UNA, BNA, and GW filter sets was used to take epifluorescence images. Images were acquired at high-resolution using a digital camera DP72 (Olympus, Japan) using CellSens Imaging Software (Olympus), and processed using Adobe Systems Photoshop 7.0 software (Adobe, USA).

Statistics

Cells (n = 500) from 2-3 different positions on each coverslip were counted at 48 h and 72 h after transfection and numbers of transfected cells were then expressed as percentages of all cells.

RESULTS

NAGK was expressed throughout the different stages of cell division

In interphase HEK293T cells, NAGK was present in cytoplasm, in nuclear speckles and paraspeckles, and on nuclear envelopes (Sharif et al., 2015). To study NAGK expression patterns during cell division we performed ICC using anti-NAGK and anti-tubulin antibodies in asynchronous HEK293T cells and followed by DAPI staining. NAGK expression in interphase cells was in line with our previous findings (Sharif et al., 2015). Cytoplasmic NAGK signals were mostly localized on microtubules (MTs) (Fig. 1A, Interphase, arrowheads) as was in previous observations in hippocampal neurons (Islam et al., 2015a). At prophase when chromosomes started to condense, some NAGK signals colocalized with MT on nuclear envelopes (Fig. 1B, Prophase, inset, arrowheads). During metaphase, condensed chromosomes are sequentially attached to spindle microtubules at their kinetochores, and during this stage, NAGK signals localized on thin MT spindle shafts and at the plus ends of spindles attached to chromosomes (Fig. 1C, Metaphase,
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A similar distribution pattern was observed during anaphase (Fig. 1D, Anaphase) and telophase (Fig. 1E, Telophase). Finally, as the cytokinesis phase approached, NAGK started to redistribute throughout cells, including nuclei (Fig. 1F, Cytokinesis). These observations suggest that NAGK contributes to the anchoring of MT spindles to the nuclear envelope during prophase and to chromosomes during nucleokinesis.

NAGK interacted with dynein light chain roadblock type 1 (DYNLRB1) during cell division

Previously, we reported that NAGK interacted with DYNLRB1 subunits of dynein complex (Islam et al., 2015a). Dynein localizes to a variety of subcellular structures during G2 and mitosis, including the nuclear envelope (NE), centrosomes, kinetochores (KTs), mitotic spindles, and the cell cortex (Dujardin and Vallee, 2002; Kiyomitsu and Cheeseman, 2012; Pfarr et al., 1990; Steuer et al., 1990; Tanenbaum et al., 2010). In particular, dynein is involved in chromosome movements, spindle organization and positioning, and checkpoint silencing during mitosis (Howell et al., 2001; Sharp et al., 2000; Varma et al., 2008). Therefore, we investigated the in situ localization of NAGK-dynein interactions through different stages of cell division by performing PLA on HEK293T cells for NAGK and DYNLRB1 and then followed with tubulin ICC and DAPI counter-staining (Fig. 2). PLA signals were observed at different z-axis layers of cells but were mostly studied at the equatorial nuclear layer. In interphase cells, NAGK-DYNLRB1 interactions (PLA assay) were found in cytoplasmic areas with many PLA signals on and around the nuclear envelope (Fig. 2A, arrowheads). We also observed NAGK-dynein interactions on the prophase nuclear envelope when chromosomes began to condensate (Fig. 2B, arrowheads). Dynein plays a critical role in nuclear envelope invagination during early prophase (Salina et al., 2002), and thus, NAGK-DYNLRB1 interactions may be a part of the dynein-adaptor protein complexes that guide prophase nuclear invagination prior to NEB. At metaphase, chromosomes aligned themselves at the equatorial plane and NAGK-dynein interactions (PLA) occurred on the mitotic spindle shafts and on the chromosomes (Fig. 2C, red arrowheads), and in peripheral cytoplasmic areas (Fig. 2C, white arrowheads). These PLA signals in peripheral cytoplasm may be on astral MTs, because dynein links these MTs to the cell cortex during cell division. Similar localizations of NAGK-DYNLRB1 PLA signals were observed during anaphase, telophase, and cytokinesis (Figs. 2D-2F, red arrowheads-on chromosomes and/or on polar MTs, white arrowheads-others). These observations indicate NAGK might regulate dynein function on mitotic MT spindles and at kinetochores (KTs) on chromosomes.

NAGK interacted with Lis1 during cell division on nuclear envelopes

Previous studies suggest dynein plays its role in cell division with the cooperation of Lis1 and NudE1/NudEL1 (Shu et al., 2004; Yan et al., 2003). Raaijmakers et al. (2013) found that dynein must be complexed with Lis1 and NudE1/NudEL1 to establish appropriate spindle orientations, and later Moon et al. (2014) suggested LIS1 acted via the LIS1–NDEL1–dynein complex to regulate astral MT plus-ends dynamics to establish proper spindle organization. To investigate the possibility of an interaction between NAGK and dynein-Lis1 complex, we conducted NAGK-Lis1 PLA on asynchronous HEK293T cells and then performed tubulin ICC and DAPI counter-staining. As shown in Fig. 3A (arrowheads), NAGK interacted with Lis1 in
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NAGK-DYNLRB1 interaction during the cell cycle. Proximity ligation assays (PLAs) were performed on HEK293T cells using anti-NAGK and anti-DYNLRB1 antibodies and were followed by ICC using a mouse anti-tubulin antibody (green). Cells were then stained with DAPI to visualize chromosomes. PLA/DAPI, PLA/ICC (tubulin), and merge images are shown. (A) In interphase, NAGK-DYNLRB1 PLA dots around nuclear membrane are indicated by arrowheads. (B) In prophase, chromosomes were condensed (DAPI stained) and NAGK-DYNLRB1 interactions on nuclear envelopes are shown by arrowheads. (C) In metaphase, NAGK-dynein complexes were also observed on chromosomes (Fig. 3F; red arrowheads), and in other cytoplasmic areas (Fig. 3F; white arrowheads). Based on these observations, the localization of NAGK-Lis1 interactions (PLA signals) and of NAGK-dynein complexes during cell division were notably similar, particularly in prophase and metaphase cells, which suggested NAGK interacts with dynein-Lis1 complex.

NAGK interacted with dynein-Lis1-NudE1 complex on nuclear envelopes during prophase nuclear invagination
Salina et al. (2002) reported that dynein facilitates prophase nuclear envelope invagination leading to NEB by pulling nuclear membranes. To investigate NAGK-dynein interactions during prophase NEB, we performed NAGK-DYNLRB1 PLA followed by lamin-B ICC and DAPI counter-staining in prophase hypothalamic GT1-7 cells (Fig. 4A) (this cell line was utilized to support our previous findings in HEK293T cells; Fig. 2B). Here, NAGK-dynein PLA signals were also observed around prophase nuclear envelopes (Fig. 4A, arrowheads) and invagination was marked by stronger in situ lamin B signals (Fig. 4A, green arrow) as nuclear lamina disassembles and concentrates at invagination areas during the initiation of prophase NEB (Beaudouin et al., 2002; Georgatos et al., 1997). It has been established that the recruitment or targeting of dynein to nuclear envelopes are regulated by Lis1 (Egan et al., 2012; Splinter et al., 2012) and NudE1/NudEL1 (Bolhy et al., 2011), and dynein-Lis1-NudE1 interactions contribute to dynein-dependent nuclear envelope breakdown (Hebbar et al., 2008). We next examined NAGK’s role in NEB by studying its interaction with dynein-Lis1-NudE1 complex. First, we conducted NAGK-DYNLRB1 PLA followed by Lis1 ICC and DAPI counter-staining in HEK293T cells, and found NAGK-dynein complex (Fig. 4B, original image, inset, red arrowheads) colocalized with Lis1 ICC puncta (Fig. 4B, original image, inset, green arrowheads) at the nuclear envelope during prophase. In a similar experiment on HEK293T cells, NAGK-DYNLRB1 PLA was followed by NudE1 ICC and DAPI counter-staining, and NAGK-dynein-NudE1 interactions were observed on nuclear envelopes during NEB. Furthermore, NAGK-dynein interactions (Fig. 4C, original image, inset, red arrowheads) colocalized with these NudE1 signals (Fig. 4C, original image, inset, green arrowheads). These observations strongly indicate that NAGK interacts with dynein-Lis1-NudE1 complex and is involved in prophase NEB.

Colocalization of NAGK with Lis1 and with NudE1 complex on NEs observed after microtubule disruption
Nocodazole, a MT disrupting agent, delays NEB (Salina et al., 2002) and increases Lis1-NudE1/NudEL1-dynein complex accumulation at nuclear envelopes (Hebbar et al., 2008). We reasoned that nocodazole would disrupt both MT and MT-associated NAGK signals but would not change NAGK expression at nuclear envelopes if it interacts with Lis1-NudE1 during NEB. To investigate this rationale, we treated HEK293T cells with nocodazole, and found NAGK-dynein complexes were also observed on chromosomes of prophase cells (Fig. 3B; arrowheads). In metaphase, NAGK-Lis1 PLA signals were present in chromosomal alignment and colocalized with MT spindles (Fig. 3C; red arrowheads), and some signals were also observed in astral sites (Fig. 3C; white arrowheads). Similar NAGK-Lis1 interactions were found during anaphase and telophase (Figs. 3D and 3E), on mitotic spindles and on chromosomes (red arrowheads), and in astral cytoplasmic areas (white arrowheads). During cytokinesis, NAGK-Lis1 PLA signals were also observed in chromosomes areas (Fig. 3F; red arrowheads) and in other cytoplasmic areas (Fig. 3F; white arrowheads). Based on these observations, the localization of NAGK-Lis1 interactions (PLA signals) and of NAGK-dynein complexes during cell division were notably similar, particularly in prophase and metaphase cells, which suggested NAGK interacts with dynein-Lis1 complex.
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Fig. 3. NAGK-Lis1 interaction during the cell cycle. PLA was carried out on fixed HEK293T cells using primary antibodies against NAGK and Lis1, subsequent ICC was performed using anti-tubulin antibody and then DAPI staining was done. In interphase (A) and prophase cells (B), NAGK-Lis1 interactions were found on nuclear membranes (arrowheads). In metaphase (C), anaphase (D), telophase (E) and cytokinesis (F) stages, PLA signals were observed on chromosomes and/or on MTs (red arrowheads), and in other cytoplasmic areas (white arrowheads). Scale bar; 5 μm.

with nocodazole to arrest cell division at G2/M transition, at which NEB is a key prerequisite for commitment to M phase. After nocodazole treatment, the proportion of prophase cells increased, and when we double-stained nocodazole treated cells with anti-NAGK and -tubulin antibodies, it was found MTs were disrupted and tubulin staining was mostly present at one location, which was assumed to be the centrosome (Fig. 5A, red arrowhead). NAGK signals were also concentrated at this location and overlapped those of tubulin (Fig. 5A, green arrowheads). Furthermore, double-staining in the same manner showed NAGK expression around the nuclear envelope and a significant number of NAGK/Lis1 (Fig. 5B, inset, arrowheads) and NAGK/NudE1 colocalization (Fig. 5C, inset, arrowheads). In addition, cytoplasmic signals for NAGK and Lis1 were weaker than the dispersed cytoplasmic signals observed in the absence of nocodazole (Fig. 5D, arrows). These findings confirm interaction between NAGK and dynein-Lis1-NudE1 complex on nuclear envelopes during the G2/M phase and suggest the participation of NAGK in dynein-mediated NEB.

NAGK interacted with dynein-Lis1-NudE1 complex on chromosomes during metaphase
When we analyzed the positioning of NAGK-dynein (Fig. 6A) and NAGK-Lis1 (Fig. 6B) PLA signals in metaphase cells, both were observed on MT spindles and chromosomes (red arrowheads), suggesting their involvements in chromosome movement and pole focusing, and in other cytoplasmic areas assumed to be related in movement of astral microtubules towards the cortex. Interestingly, majority of NAGK-dynein and
Fig. 5. Colocalization of NAGK with Lis1 and NudE1 on nuclear envelopes after microtubule disruption in prophase cells. HEK293T cells were treated with nocodazole, double labeled with indicated antibodies and then stained with DAPI. (A) Tubulin signals disrupted by nocodazole and concentrated around centrosome are shown by red arrowheads (Tub/DAPI). In addition, colocalization of NAGK signal clusters (green arrowheads, NAGK) with tubulin are indicated by yellow arrowheads (merge). (B) ICCs with anti-NAGK (green) and anti-Lis1 (red) primary antibodies were performed on nocodazole treated prophase HEK293T cells, which were then stained with DAPI (blue). DAPI staining revealed invagination of prophase nuclear envelopes. Colocalizations of NAGK signals with Lis1 signals on nuclear envelopes are shown by arrowheads (boxed area enlarged inset) during nuclear invagination. (C) Nocodazole treated prophase HEK293T cells were double stained with anti-NAGK (green) and anti-NudE1 (red) antibodies in a similar method. Colocalization between NAGK and NudE1 signals are shown by arrowheads on a nuclear envelope (arrowheads, inset). (D) ICC doublelabeling with anti-NAGK (green) and anti-Lis1 (red) primary antibodies of control (not treated with nocodazole) HEK293T cells showed dispersed cytoplasmic signals (arrows). Scale bar; 5 μm.

Fig. 6. NAGK interacted with dynein-Lis1-NudE1 complex on chromosomes in metaphase. (A) (a) NAGK-DYNLRB1 PLA and subsequent tubulin ICC and DAPI staining were performed in HEK293T cells. NAGK-dynein interactions (PLA) on chromosomes are indicated by red arrowheads, and in other cytoplasmic areas by white arrowheads. (b) Statistics: Majority of NAGK-dynein PLA signals were on chromosomes (53.84%) in metaphase. (B) (a) NAGK-Lis1 PLA was performed on HEK293T cells and followed by tubulin ICC and DAPI staining. NAGK-Lis1 interactions (PLA) were localized to chromosomes (red arrowheads) and in other cytoplasmic areas (white arrowheads). (b) NAGK-Lis1 interactions in metaphase were occurred on chromosomes (64.28%) at a higher frequency compared to other cytoplasmic areas. NAGK-DYNLRB1 PLA followed by Lis1 (C) or NudE1 (D) immunostaining, and DAPI staining revealed colocalizations between NAGK-dynein complex (original image, inset, red arrowheads) and Lis1 (C, original image, inset, green arrowheads), and NudE1 (D, original image, inset, green arrowheads) in metaphase. Merged images show colocalization on chromosomes (C, D, original image, inset, yellow arrowheads). Scale bar; 5 μm.

NAGK-Lis1 PLA signals (Figs. 6A-b, 53.84% and Figs. 6B-b, 64.28%, respectively) were in chromosomal areas. These findings imply that NAGK works with dynein and Lis1 to move chromosomes in metaphase cells. Indeed, NAGK-DYNLRB1
A NAGK CENP-B/DAPI
merge

5 μm

B NAGK/DYNLRB1 /DAPI

NAGK/DYNLRB1 /CENP-B

CENP-B/DAPI
merge

5 μm

PLA followed by Lis1 or NudE1 immunostaining and DAPI counterstaining for chromosomes showed that NAGK-dynein complex colocalized with Lis1 (Fig. 6C, original image, inset, arrowheads). These results show that NAGK colocalizes with DYNLRB1-Lis1-NudE1 complex on chromosomes in metaphase cells.

**Presence of NAGK-dynein complex at kinetochores during metaphase**

Kinetochores (KTs) connect chromosomes to the spindle apparatus and pull chromosomes along spindles (Meraldi et al., 2006; Musacchio and Salmon, 2007). It has been reported that Lis1 and NudE1/NudEL1 link dynein to KTs (Liang et al., 2007; Mesngon et al., 2006; Stehman et al., 2007; Vergnolle and Taylor, 2007). To study the presence of NAGK on KTs, we double-stained HEK293T cells with antibodies against NAGK and KT marker protein CENP-B, and found NAGK colocalized with CENP-B at KTs in metaphase cells (Fig. 7A, inset, arrowheads). NAGK-dynein PLA followed by CENP-B immunostaining and DAPI counter-staining was then used to locate NAGK-dynein complex on KTs (Fig. 7B). NAGK-dynein complex (red arrowheads) colocalized with CENP-B (green arrowheads) on chromosomes (DAPI, blue) in metaphase. In addition, CENP-B signals were observed on centrosomes (green arrows). These results confirm that NAGK and dynein interact at KTs in metaphase.

**Knockdown of NAGK delayed cell division**

Because the above experiments showed interactions between NAGK and dynein-Lis1-NudE1 complex during different stages of mitotic cell division, we investigated the role of NAGK in cell division. NAGK was knocked-down by transfecting HEK293T cells with a plasmid construct expressing NAGK small hairpin (sh) RNA. Cells were transfected with a red fluorescent protein tagged pDsRed2 control vector (Fig. 8A-a, pDsRed2) or co-transfected with this control vector and NAGK shRNA vector (Lee et al., 2014a) (Fig. 8A-b, sh-NAGK + pDsRed2). After incubation for 48 h to allow protein expression, live images of transfected cells were captured, and same culture areas were imaged 24 h later. During this 24 h (between 48 h and 72 h in culture), most cells transfected with control pDsRed2 plasmid divided (Fig. 8A-a, red arrows show the same mark in two different time-lapse images). In contrast, a number of cells transfected with NAGK shRNA vector did not divide (Fig. 8A-b, arrow and arrowheads indicating corresponding cells). Statistical analysis (Fig. 8B, n = 500 cells) showed during this 24 hr period, control cells multiplied at the same rate as non-transfected cells, and that the transfected to non-transfected cell ratio remained almost the same after 72 h in culture (Fig. 8B-a). On the other hand cells transfected with NAGK shRNA vector did not divide (Fig. 8A-b, arrow and arrowheads indicating corresponding cells). These results show that NAGK knockdown delayed cell division process because of lack of interaction between NAGK and dynein-Lis1-NudE1 complex.

**DISCUSSION**

This study was undertaken to investigate the role of NAGK-dynein complexes during cell division. The eukaryotic cell cycle is primarily divided into two basic phases, that is, mitosis and interphase. Interphase consists of three different phases: two gap phases (G1 and G2) and an S (synthesis) phase, during which genetic material is duplicated. In the G2 phase, cells assemble the cytoplasmic materials necessary for mitosis (the M phase), a continuous process comprised of five different stages: prophase, prometaphase, metaphase, anaphase, and...
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NAGK plays a role during prophase NEB by interacting with Lis1 and NudE1 on prophase nuclear envelopes suggest that NAGK-Lis1 complexes and NudE1/NudEL1 influence PNEI formation. Nocodazole disrupts MT fibers and delays prophase NEB, which results in an accumulation of Lis1-NudE1/NudEL1-dynein complex at the prophase nuclear envelope (Cockell et al., 2004; Coquelle et al., 2002; Hebbar et al., 2008; Smith et al., 2000). In the present study, MT associated NAGK expression collapsed in HEK293T cells treated with nocodazole and concentrated around centrosome and on nuclear envelope. Colocalization of latter NAGK punctae with Lis1 and NudE1 on nuclear envelopes occurred, confirming an interaction between NAGK and dynein-Lis1-NudE1 complex during prophase NEB.

In addition, we found NAGK-dynein complexes interacted with Lis1 and NudE1 on metaphase chromosomes, and that NAGK or NAGK-dynein complex colocalized with CENP-B (a KT marker protein) in metaphase chromosomes. Dynein links tubulin to kinetochores (KTs) and adopts a force generating role during metaphase. Dynein is recruited to KTds during (pro)metaphase in two distinct ways. The first involves RZZ (Rod, Zwilch, and Zw10) complex in KTds, which recruits dynein complex (Karess, 2005) via an interaction between Zw10 and the p50 subunit of dynactin (Starr et al., 1998). The second involves CENP-F, Lis1, and NudE1/NudEL1 (Vergnolle and Taylor, 2007) whereby Lis1 and NudE1/NudEL1 work together with dynein complex to link MT fibers to KTds (Liang et al., 2007; Mesngon et al., 2006; Stehman et al., 2007). In addition, Raaijmakers et al. (2013) found that dynein intermediate chain 2, DYNLRB1 and TCTEX1, all contribute to recruitment of the dynein complex to the KTds. Therefore, our observation of an interaction between NAGK and dynein-Lis1-NudE1/NudEL1 complex at metaphase KTds suggests that NAGK plays a role in the dynein-mediated connection of MTs and KTds (Fig. 9B).

In the present study, NAGK-dynein and NAGK-Lis1 complexes were observed at the astral part of the mitotic spindles. Human mitotic spindle consists of different types of microtubules: those that emanates from centrosomes to interact with KTds are known as K-fibers; those that connect to cell cortex are termed astral MTs, and those that form antiparallel overlap between opposing
In the present study, knockdown of NAGK by shRNA delayed cell division, and it has been previously reported native dynein requires the presence of many adaptors for the correct localization and activation of the dynein complex during mitosis, for example, dynein regulators, such as, Lis1 and NudE1/NudEL1, are required to generate sufficient force (Raaijmakers et al., 2013, Tanenbaum et al., 2008). For spindle pole focusing, the p150Glued subunit of dynactin binds with MTs (Waterman-Storer et al., 1995). Furthermore, Lis1, NudE1/NudEL1, numerous dynein-binding adaptor proteins like ZW10 (Starr et al., 1998; Whyte et al., 2008), hSpindly (Gassmann et al., 2008; 2010; Griffis et al., 2007), NuMA (Fant et al., 2004), CENP-F (Vergnolle and Taylor, 2007) as well as dynactin, contribute to the targeting of dynein complex to KTs. In addition, dynein is known to be recruited to the nuclear envelope during G2/prophase in a BICD2-dependent manner (Raaijmakers et al., 2012; Splinter et al., 2010). In view of the many adaptors of dynein required during mitosis, we propose NAGK might be another important adaptor that enables dynein complex to function efficiently, especially during prophase NEB and metaphase chromosome separation. Further molecular dissection of the roles of NAGK-binding complexes in mitosis is required.

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