Zipper-interacting protein kinase interacts with human cell division cycle 14A phosphatase

WEI WU1*, HAIYING HU2*, ZI YE1, MANCHEONG LEONG3, MIN HE1, QIN LI3, RENMING HU1 and SHUO ZHANG1

1Department of Endocrinology and Metabolism, Institute of Endocrinology and Diabetology at Fudan University, Huashan Hospital, Fudan University, Shanghai 200040; 2Department of Endocrinology and Metabolism, International Healthcare Center, Second Affiliated Hospital of Zhejiang University, Hangzhou, Zhejiang 310009; 3Department of Endocrinology, City University of Macau, Macau 000853; 4Department of Endocrinology, Shanghai Tenth People’s Hospital, School of Medicine, Tongji University, Shanghai 200000, P.R. China

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Abstract. Zipper-interacting protein kinase (ZIPK) is a novel serine/threonine protein kinase and a member of a large family of protein kinases, known as the death-associated protein kinases. However, the function of ZIPK has yet to be fully elucidated, as few physiological substrates have currently been identified. In the present study, a yeast two-hybrid screen was used and the human cell division cycle 14A (HsCdc14A) phosphatase was identified as a novel ZIPK binding protein. To the best of our knowledge, this is the first study to report the interaction between these proteins. The interaction between ZIPK and HsCdc14A was confirmed by in vitro experiments. In addition, ZIPK-mediated phosphorylation was shown to activate the phosphatase activity of HsCdc14A. These findings indicated that ZIPK may also be involved in the regulation of the cell cycle in human cells, by interacting with HsCdc14A.

Introduction

Zipper-interacting protein kinase (ZIPK) is a novel serine/threonine protein kinase, which was initially cloned and identified in 1998 (1,2). ZIPK is a member of the large family of death-associated protein kinases and has been associated with the regulation of numerous cellular processes, including cell death (1), cell motility (3) and mitotic processes (4), as well as smooth muscle contraction (5,6). In addition, ZIPK interacts with the signal transducer and activator of transcription 3 (STAT3), a latent cytoplasmic transcription factor that plays a role in cell growth and apoptosis. ZIPK has been shown to phosphorylate STAT3 on Ser-727, thus enhancing the transcriptional activity of STAT3 (7). Vetterkind et al (8) have demonstrated that prostate apoptosis response-4 (Par-4), which is characterized mainly as a proapoptotic protein, targets ZIPK to the cytoskeleton in nonmuscle cells, leading to apoptosis. By contrast, in smooth muscle cells, Par-4 supports contractility by targeting ZIPK to the cytoskeleton (9). A previous study, conducted by the authors of the present study, demonstrated that incubation of smooth-muscle cells with glucose induced a time- and dose-dependent increase in the protein expression levels of ZIPK (unpublished data). These findings indicate that ZIPK plays a key role in cellular function.

However, only a limited number of studies have focused on the effects of ZIPK on cell death, motility and mitotic processes. To date, the interacting partners of ZIPK, particularly those associated with the cell cycle, have been rarely identified and the regulatory network of ZIPK is not fully understood. The aim of the present study was to investigate the interaction of ZIPK with the human cell division cycle 14A (HsCdc14A) phosphatase in vitro, and identify whether ZIPK plays a role in the cell cycle regulation.

Materials and methods

Yeast two-hybrid assay. The study was approved by the ethics committee of Huashan Hospital, Fudan University (Shanghai, China). Yeast two-hybrid interaction screening was performed as described in previous studies (10-12). Full-length HsCdc14A was introduced into the GAL4 DNA (Clontech Laboratories, Inc., Mountain View, CA, USA) binding domain as a bait. Its interaction partner was a human testis cDNA library (Clontech Laboratories, Inc.) integrated with GAL4 transactivation domain. The bait plasmids and the cDNA library plasmids were used to transform a yeast strain (AH109; Clontech Laboratories, Inc.), containing HIS3 and LacZ reporter genes, using a lithium acetate method (13). The
transformed samples were spread onto plates with a synthetic defined (SD)/Leu/Trp/His medium. Subsequently they were spread onto a SD/-Leu/-Trp/-His/Ade/X-a-Gal medium for further selection. Positive clones (those with diameter >3 mm) were co-transformed into the AH109 yeast with bait plasmids. Following extraction of plasmids from yeast, the plasmids were used to transform Escherichia coli (Tianz, Inc., Beijing, China). The interacting protein was verified by sequencing the plasmid following extraction from Escherichia coli.

Glutathione S-transferase (GST) pull-down assay. HsCdc14A cDNA [full length, amino acid (aa) 1-623; deletion mutant, aa1-348N terminal; or aa349-623C terminal] was fused with a 6-His tag (pET-28a; Tianz, Inc., Beijing, China). ZIPK cDNA [full length, aa1-454] was cloned into a pGEX-5X-3 vector (Amersham Biosciences, Piscataway, NJ, USA). Using a previously described method (11), these proteins were expressed in Escherichia coli BL21 (DE3) cells (Tianz Inc.), and purified by Ni²⁺ nitrilotriacetic acid (for aa1-348; Sigma-Aldrich, St. Louis, MO, USA) or glutathione beads (for aa349-623; Sigma-Aldrich). An in vitro pull-down assay was performed using purified GST-fused ZIPK and His-tagged HsCdc14A with phosphate-buffered saline (PBS), containing 0.1% Triton X-100 (Sangon Biotech, Shanghai, China) for 4 h at 4°C. The beads were washed three times with PBS containing 1% Triton X-100, and then with PBS alone. The beads were then boiled in SDS-PAGE sample buffer for 5 min and used for western blot analysis.

Co-immunoprecipitation. Human embryonic kidney (HEK) 293T cells from the American Type Culture Collection (Manassas, VA, USA) were grown to 50-60% confluence in Dulbecco’s modified Eagle’s medium (HyClone, Logan, UT, USA), supplemented with 10% fetal bovine serum (HyClone), at 37°C in an atmosphere containing 5% CO₂. The cells were then co-transfected with green fluorescent protein (GFP)-HsCdc14A (full length or deletion mutant aa1-348N-terminal or aa349-623 C-terminal) and FLAG-ZIPK using the standard CaCl₂ method (14), with a GFP-transfected plasmid as a control, 36 h later, the cells were collected and lysed in lysis buffer (50 mM HEPES, pH 7.2; 150 mM NaCl; 2 mM ethylene glycol tetraacetic acid; and 0.1% Triton X-100), together with protease inhibitor mixture (Aprotinin, Bestatin and Leupeptin Pepstatin A; Sigma-Aldrich). The lysate was then clarified using centrifugation at 16,000 x g for 10 min at 4°C. The cell lysate was incubated with anti-FLAG antibody conjugated to agarose beads (Sigma-Aldrich) for 4 h at 4°C, and then washed five times with lysis buffer. The proteins were boiled for 5 min and used for subsequent western blot analysis.

Western blotting analysis. Following SDS-PAGE using a 12% gel, proteins were transferred onto polyvinylidene difluoride membranes, which had been obtained from Millipore Corporation (Billerica, MA, USA). The membranes were incubated with primary antibodies overnight at 4°C with gentle shaking (Qite Analytical Instrument Co. Ltd., Shanghai, China). Membranes were then incubated with secondary antibodies at room temperature for 2 h. Immunoreactive signals were detected using an enhanced chemiluminescence kit (Pierce Chemical, IL, USA) and visualized by autoradiography on Kodak BioMAX film (Kodak, Rochester, NY, USA). The following antibodies were used: Mouse monoclonal anti-His antibody (1:1,000; Cell Signaling Technology, Inc., Danvers, MA, USA); rabbit polyclonal anti-GFP antibody (1:1,000; Cell Signaling Technology, Inc); mouse monoclonal anti-FLAG antibody (1:2,000; Sigma-Aldrich); and anti-rabbit (1:2,000) and anti-mouse (1:2,000) horseradish peroxidase-conjugated secondary antibodies (Cell Signaling Technology, Inc.).

Cell cycle analysis. A cell cycle analysis was conducted as previously described (15). Briefly, HEK293T cells were grown in six-well plates with density ~1x10⁶ cells per well and transfected with the following plasmids: GFP, GFP-HsCdc14A, GFP-HsCdc14A N-terminus, GFP-HsCdc14A C-terminus, GFP-ZIPK, GFP-HsCdc14A+ and GFP-ZIPK. 72 h later, cells were collected and washed with ice cold PBS. Next, they were fixed with 70% ethanol for ≥1 h. The fixed cells were then washed and stained with propidium iodide (PI; Invitrogen Life Technologies, Carlsbad, CA, USA) for 1 h at room temperature. Subsequently, the PI-stained nuclei were analyzed using the BD FACSCalibur™ system (BD Biosciences, Franklin Lakes, NJ, USA).

Apoptosis assay. Annexin V-fluorescein isothiocyanate (FITC)/PI staining using an Apoptosis Detection Kit (Invitrogen, Life Technologies) was used in order to detect apoptotic cells. HEK293T cells were collected at 72 h post-transfection with the plasmids listed above. Cells were then trypsinized and collected by centrifugation at 300 x g for 5 min. Cells were washed once with PBS, resuspended in 1X binding buffer and
stained with Annexin V-FITC for 15 min. The cell nuclei were then counter-stained with PI in order to detect necrosis. Flow cytometric analysis was performed to analyze the percentage of apoptotic cells, using the BDFACSCalibur™ system.

Statistical analysis. All experiments were performed at least in triplicate. The results are expressed as the mean ± standard deviation, and the data were analyzed using Student’s t-test to detect statistically significant differences among the groups. P<0.05 was considered to indicate a statistically significant difference. Statistical analyses was performed using SPSS Version 11 (IBM SPSS, Armonk, NY, USA).

Results

Identification of HsCdc14A as a ZIPK-interacting protein. Nucleotide sequencing was performed in previous experiments using full length HsCdc14A cDNA as a bait in order to screen through a human testis cDNA library. ZIPK was identified as a potential binding protein. Subsequent assays validated that HsCdc14A can bind to ZIPK (Fig. 1). To the best of our knowledge, the present study reported for the first time a potential interaction between ZIPK and HsCdc14A.

ZIPK and HsCdc14A interactions. Subsequent experiments were conducted to verify the interaction between ZIPK and HsCdc14A. In a GST pull-down assay, GST-ZIPK was found to pull down HsCdc14A, as compared with GST alone (Fig. 2A). Various deletion mutations of HsCdc14A were constructed to map the interaction fragments of the two proteins. Two deletions of HsCdc14A were constructed (Fig. 1A). The N-terminus (aa 1-348) consisted of the nuclear localization signal (NLS) and dual-specificity phosphatase (DSP) domains, while the C-terminus (aa 349-623) contained the nuclear export signal (NES) domain. The results of the subsequent GST pull-down experiment indicated that ZIPK interacted directly with the N-terminus of HsCdc14A, but not with the C-terminus of HsCdc14A (Fig. 2A). These results were consistent with the important function of the DSP domain of HsCdc14A.

In order to verify the interaction between ZIPK and HsCdc14A in mammalian cells, HEK 293T cells were transfected with GFP-tagged HsCdc14A or a control plasmid,
along with FLAG-tagged ZIPK. The cell lysates were immunoprecipitated with anti-FLAG antibodies. Subsequently, the obtained immunoprecipitates were analyzed by western blotting using an anti-GFP antibody, in order to detect the presence of bound HsCdc14A. FLAG-tagged ZIPK was found to co-precipitate with GFP-tagged HsCdc14A, but not with GFP alone (Fig. 2B).

The regions responsible for the interaction between ZIPK and HsCdc14A were also determined. A series of GFP-tagged HsCdc14A (full length; GFP-HsCdc14A N-terminus or C-terminus) were co-transfected into HEK 293T cells with FLAG-tagged ZIPK. Co-immunoprecipitation experiments indicated that ZIPK interacted strongly with the HsCdc14A N-terminus, containing the NLS and DSP domains (Fig. 2C).

Co-transfected ZIPK and HsCdc14A plasmids interfere with mitotic progression and cause apoptosis. Due to the identification of the physical interaction between ZIPK and HsCdc14A, the present study aimed to determine the functional relevance of the ZIPK-HsCdc14A interaction in mitotic progression. HEK 293T cells were separately transfected with various plasmids. As shown in Fig. 3A, the cell cycle assay demonstrated that the G2/M phase cell population was significantly increased in cells that were co-transfected with ZIPK and HsCdc14A plasmids, when compared with the cells that were transfected with HsCdc14A alone. These results were similar to those obtained following transfection with the HsCdc14A N-terminus (the phosphatase domain and its activity may be inhibited by the C-terminus). The results indicated that the mitotic progression was arrested in cells co-transfected with ZIPK and HsCdc14A. In addition, the percentage of apoptotic cells following transfection with the various plasmids was determined. A considerable increase was observed in the number of Annexin V-positive cells upon co-transfection with ZIPK and HsCdc14A, compared with the cells transfected with HsCdc14A alone (Fig. 3B). The percentage of apoptotic cells was similar to the cells transfected with the HsCdc14A N-terminus. These results indicated that apoptosis was increased in the cells co-transfected with ZIPK and HsCdc14A. Therefore, the control of HsCdc14A phosphatase activity by ZIPK is hypothesized to play an important role in mitotic progression and cell apoptosis.

Discussion

ZIPK was initially identified as a Ser/Thr kinase that binds activating transcription factor 4 (ATF4). ATF4 is a member of the activating transcription factor and cyclic adenosine monophosphate-responsive element-binding protein family of transcription factors (1). ZIPK aggregates through its C-terminal leucine zipper structure, thereby becoming an active enzyme. A previous study identified that the ectopic expression of ZIPK in NIH-3T3 murine fibroblast cells induced apoptosis (1). By contrast, the kinase-inactive ZIPK K42A mutant was not found to induce apoptosis, indicating that the catalytic activity of ZIPK stimulates cell apoptosis (1). Numerous kinases that mediate cell growth are known; however, only a few protein kinases associated with apoptosis have been identified, besides ZIPK. Previous studies demonstrated that ZIPK participates in the regulation and possibly the coordination of mitosis and cytokinesis, by interacting with the proapoptotic protein, Par-4, and the CDC5 protein (4,16). In addition, ZIPK is a centromere-specific histone kinase that may play a role in the labeling of centromere-specific chromatins for subsequent mitotic processes (17). Furthermore, previous studies have indicated that ZIPK, as a regulator of myosin phosphatase, may play a pivotal role in the regulation of cell motility, reorganization of actin filaments and control of smooth muscle contractions in smooth muscle cells (4,6,7). However, only a few interacting partners of ZIPK have been identified and the regulatory networks of ZIPK remain to be elucidated.

The present study demonstrated that ZIPK interacted with the HsCdc14A protein in vitro, playing an important role in the regulation of the cell cycle. The interaction was shown to involve the highly conserved N-terminus of HsCdc14A.
indicating that ZIPK may be involved in the cell cycle regulation.

Cdc14 is a protein phosphatase conserved between yeast and humans (10). Genetic analyses have suggested that the yeast Cdc14 plays a pleiotropic role during the cell cycle, regulating DNA replication and the exit from mitosis by dephosphorylating cyclin-dependent kinase (Cdk) targets (18-20). Mammalian cells express two functional homologs of the yeast Cdc14, known as HsCdc14A and HsCdc14B (21). These two proteins remain poorly understood; however, recent evidence has indicated that they play an isoform-specific role in centrosome separation/maturation and spindle stability, with the possibility of additional role in the mitotic exit and cytokinesis (21). The majority of previous studies have investigated HsCdc14A, which was demonstrated to interact with interphase centrosomes and regulate the centrosome duplication cycle (22,23). In addition, HsCdc14A is located at the central spindle during anaphase, where it appears to be involved in the spatial regulation of the Aurora B kinase, a key regulator of chromosome segregation and cytokinesis (24). HsCdc14A may also regulate p53 and Cdk1/cyclin B; thus, dysregulation of HsCdc14A may play an important role in carcinogenesis (25). Besides p53, a previous study revealed that HsCdc14A can dephosphorylate the products of Cdk, including hCdcl1 and cyclin E (26). However, the mechanism through which the HsCdc14A function is integrated in mitotic regulation, the substrates involved in chromosome segregation and whether HsCdc14A plays an active role in mitosis remain unclear.

HsCdc14A contains an NLS motif, a DSP domain and an NES motif. Furthermore, HsCdc14 exhibits an inhibitory self-association, in which the C-terminal domain binds and inhibits the phosphatase domain located at the N-terminus (27). Once the inhibition of C-terminus is released, the activity of the phosphatase domain, located at the N-terminus, is increased. The present study demonstrated that ZIPK interacted with the N-terminus of HsCdc14A, which contains the NLS and DSP domains. In addition, the effect of co-transfection of HEK 293T cells with ZIPK and HsCdc14A was similar to the effect of the cells transfected with the HsCdc14A N-terminus; therefore, ZIPK-mediated phosphorylation may activate the phosphatase activity of HsCdc14A. Furthermore, the results of the present study indicated that ZIPK may affect the cell cycle by interacting with the N-terminus of HsCdc14A. Notably, the effect of ZIPK and HsCdc14A overexpression on cell apoptosis was similar to the effect of ZIPK alone, but more evident than the effect of HsCdc14A alone. Apoptosis was not affected significantly more by the increased phosphatase activity of HsCdc14A alone with ZIPK compared with cells transfected with HsCdc14A alone. Therefore, these two proteins may be associated with the same apoptotic pathway. Future experiments are required to investigate this hypothesis.

In conclusion, the results of the present study demonstrated that ZIPK may interact with the HsCdc14A protein. These findings indicated that ZIPK may also be involved in the regulation of the cell cycle. Further research regarding the regulation of HsCdc14A by ZIPK is required to provide valuable insight on the effect of ZIPK on the cell cycle. ZIPK may be a potential target candidate for the treatment of diseases associated with cell proliferation.

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