Regulation of Sox6 by Cyclin Dependent Kinase 5 in Brain

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Abstract

Cyclin dependent kinase 5 (Cdk5) is a proline-directed Ser/Thr kinase involved in various biological functions during normal brain development and neurodegeneration. In brain, Cdk5 activity is specific to post-mitotic neurons, due to neuronal specific expression of its activator p35. The biological functions of Cdk5 have been ascribed to its cytoplasmic substrates, however not much is known in nucleus. Here, we show that nuclear transcription factor Sox6 is a direct nuclear target of Cdk5. Sox6 is expressed in Tuj1 positive neurons, suggesting that Sox6 is expressed in differentiating neurons. The expression of Sox6 is high in mitotic nuclei during embryonic day 12 (E12) and gradually decreases during development into adult. On the other hand, Cdk5 expression gradually increases during its development. We show that Sox6 is expressed in mitotic nuclei in embryonic day 12 (E12) and in migrating neurons of E16. Sox6 is phosphorylated in vivo. Sox6 was detected by phospho-Ser/Thr and phospho-Ser/Thr-Pro and MPM-2 (Mitotic protein #2) antibodies in brain. Furthermore, calf intestinal alkaline phosphatase (CIAP) digestion resulted in faster migration of Sox6 band. The GST-Sox6 was phosphorylated by Cdk5/p35. The mass spectrometry analysis revealed that Sox6 is phosphorylated at T119PER motif. We show that Sox6 steady state levels are regulated by Cdk5. Cdk5 knockout mice die in utero and Sox6 protein expression is remarkably high in Cdk5−/− brain, however, there is no change in mRNA expression, suggesting a post-translational regulation of Sox6 by Cdk5. Transfection of primary cortical neurons with WT Cdk5 reduced Sox6 levels, while dominant negative (DN) Cdk5 and p35 increased Sox6 levels. Thus, our results indicate that Cdk5 regulates Sox6 steady state protein level that has an important role in brain development and function.

Introduction

When metabolic pathways vital for development go awry, the consequences can be disastrous. Sox [Sry-related high-mobility-group (HMG) box] family of transcription factors play an important role in development and are involved in various physiological processes. Twenty Sox transcription factors (Sox1-20) exist in mice and humans [1,2]. They feature a Sry-related high-mobility-group (HMG) box DNA-binding domain. Sox6 is highly identical to Sox5 and Sox13, forming with them the Sox D subfamily. Sox6 is known to be highly expressed in neuronal cells, chondrocytes, notochord and spermatid cells, and expressed weakly in muscle cells [3,4]. Sox5+− Sox6+− fetuses develop severe skeletal dysplasia [5]. About half die at birth and the others fail to thrive after postnatal day 7 (P7) and die at approximately P14. The cause of death remains unclear. Mice with a chromosomal inversion (p20M8) disrupting Sox6 have the same gross phenotype as Sox6+−/− mice. They were shown to develop cardiac and skeletal myopathy, suggesting that Sox6 might promote myocyte maturation [6]. Interestingly, Sox6 is prominently expressed in mouse glioma [7]. Sox6 is necessary for normal positioning and maturation of medial ganglionic eminences (MGE) derived cortical interneurons. As a consequence, the specific removal of Sox6 from this population results in a severe epileptic encephalopathy [8]. The microRNA-219 (miR-219) is shown to inhibit oligodendrocyte formation through repression of Sox6 [9].

The proline-directed serine/threonine protein kinase Cdk5, together with its neuronal-specific activating cofactors p35 and p39 have been implicated in many normal and pathological processes in the mammalian central nervous system (CNS) [10,11]. Cdk5 is also implicated in numerous complex functions of the adult CNS such as synaptic transmission, synaptic plasticity, and neuronal signaling [12]. We have earlier reported neuronal migration defects in Cdk5−/− brain resulting in developmental abnormalities and early lethality [13–15]. Cdk5 regulates many cellular processes and functions in the brain, and its deregulation via the calpain-dependent cleavage of its activator p35 to p25 has been implicated in neurodegenerative disorders such as Alzheimer’s and Parkinson’s diseases [16,17]. Recently, Cdk5 has been shown to play an important role in regulating transcription factors...
and chromatin remodeling. Transcription factor MEF2 is shown to be regulated by Cdk5 during neurotoxicity induced apoptosis [18]. Furthermore, Cdk5 mediated degradation of MEF2D is known to be mediated by chaperone induced autophagy [19]. Deregluation of histone deacetylase 1 (HDAC1) activity by Cdk5/ p25 induces aberrant cell-cycle activity and double-strand DNA breaks leading to neurotoxicity [20]. The nuclear Cdk5 plays an active role in allowing neurons to remain post mitotic as they mature and that loss of nuclear Cdk5 leads to cell cycle entry [21]. The role of Sox6 in the mammalian brain has not been explored in detail. Since Cdk5 is developmentally regulated [22] and Sox6 has potential Cdk5 phosphorylation sites, it would be of interest to find out whether Sox6 is developmentally regulated by Cdk5.

Here, we report that Sox6 is expressed during early embryonic development and its expression is gradually decreased during its development into adult. Cdk5−/− mice show increased Sox6 expression and inhibition of Cdk5 activity in the primary cortical neurons by DN Cdk5 increases the endogenous Sox6 expression.

**Materials and Methods**

**Chemicals and Antibodies**

Sox6 and Cdk5 antibodies were purchased from Santa Cruz Biotechnology (Santa Cruz, CA, USA). Tuj1 and β-actin antibodies were purchased from Sigma chemical company (St. Louis, Missouri, USA). Inhibitors such as roscovitine were purchased from Calbiochem (EMD chemicals, Gibbstown, NJ, USA).

**Animals**

Cdk5-knockout mice were generated as previously reported [22]. All of the animal procedures were conducted in accordance with the National Institutes of Health guidelines for the care and use of laboratory animals. The animal procedures were approved by National Institute of Neurological Disorders and Stroke (NINDS) Animal Care and Use Committee (ACUC), NIH. This study was carried out in strict accordance with the recommendations in the Guide for the Care and Use of Laboratory Animals of the National Institutes of Health. The protocol was approved by the Committee on the Ethics of Animal Experiments of the Animal Care and Use Committee (ACUC), National Institute Neurological Disorders and Stroke (NINDS), NIH.

**In Vitro Kinase Assays.** In the in vitro kinase assay was performed with either purified recombinant Sox6 protein (0.5 μg) or Sox6 immunoprecipitated from rat brain lysate or from primary cortical cultures. Sox6 protein was incubated with purified Cdk5/p35 complex in a kinase reaction buffer containing γ-32P and cold ATP (50 μM). For some experiments Cdk5/p35 complex was immunoprecipitated from the primary cortical neurons. The kinase reaction was carried out for 30 min at 30°C and terminated by the addition of Laemmli sample buffer. Reaction products were resolved by SDS-PAGE, and 32P-labeled proteins were visualized by autoradiography.

**Culture of Primary Cortical Neurons**

Primary cortical neurons were established from embryonic day-18 (E18) Sprague–Dawley rat embryos (Charles River Labs, NY, USA). An eighteen day timed pregnant rat was euthanized using CO2 and pups were removed, decapitated and the cortex was dissected in Hibernate-E media (Brain Bits LLC, IL, USA). Dissociated cortical neurons were obtained by incubating the cortex in EBSS containing 15 units/mL of papain (Worthington Biochemicals, NJ, USA) for 45 min at 37°C before triturating in neurobasal medium containing 20% fetal bovine serum (HyClone, UT, USA), DNase (0.2 mg/mL). Undissociated neurons were removed from the cell suspension by passing the cell suspension through a 40 μm cell strainer (Fisher Scientific, NY, USA). Neurons were centrifuged at 2000 g for 3 min at 20°C and the pellet was resuspended in neurobasal medium supplemented with B27, penicillin (100 U/mL), streptomycin (100 U/mL) and L-glutamine (0.5 mM, Invitrogen, NY, USA). Neurons were then plated at a density of 150 000 cells/mL on circular glass coverslips and 6-well tissue culture dishes, coated with poly-L-lysine (50 μg/mL, Sigma Chemicals, MO, USA), and incubated in a humidified atmosphere containing 5% CO2: 95% O2 at 37°C.

**RNA Isolation and Real Time PCR**

Total RNA was isolated from the WT, Cdk5−/−, p35−/− using TRIzol reagent (Invitrogen) according to the manufacturer’s instructions. Following TURBO DNA-free (Ambion, Austin, TX, USA) digestion of total RNA sample, to remove contaminated genomic DNA, oligo(dT) primed synthesis of cDNA from 1 μg of total RNA was made using Super-Script III reverse transcriptase (Invitrogen). The PCR consisted of 35 cycles of 30 s each at 94°C, 60°C, and 72°C. For detection of Sox6, p35, and Cdk5 mRNA we used real time PCR, and the following reaction mixture was used for these PCR samples: 1×IQ Sybr Green Super Mix (Bio-Rad, Berkeley, California, USA), 100–200 nM of each primer and 1 μl of cDNA. cDNA was amplified and analyzed in triplicate using Opticon Monitor Chromo 4 (Bio-Rad). The following primers were used to amplify and measure the amount of mouse mRNA by real time reverse transcription-PCR: Sox6 S: 5′-CATGTCCAA-GCAGGAGAAGCA-3′; Sox6 AS: 5′-GGGTACCCTCTAGAGT-GGATTTCG-3′, Cdk5 S: 5′-GCC TAA AAA CCG GGA AAC TC-3′ and Cdk5 AS: 5′-CCA TTG CAG CTG TCG AAA TA-3′ [23]. The mRNA levels were standardized by using the following primers to GAPDH: GAPDH S: 5′-AAT GTG TCC GTG TGC GTG CAT CTG A-3′ and GAPDH AS: 5′-GAT GCC TGC TTC ACC ACC TTC T-3′.

**CIAP assay**

The 25 μg of the brain lysate was subjected to 1 unit of calf intestinal alkaline phosphatase (CIAP) and the 25 μg of untreated and treated lysates were run on SDS-PAGE gel. Western blotting was performed with Sox6 antibody.

**Co-immunoprecipitation assay**

For co-immunoprecipitation of endogenous proteins, neurons were lysed in ice-cold buffer containing 50 mM Tris-HCl (pH 7.5), 150 mM NaCl, 10 mM EDTA, 2 mM EGTA, 1% NP-40, supplemented with the phosphatase and protease inhibitors cited above. Cell extracts were clarified by centrifugation and supernatants (500 μg protein) were incubated with anti-Cdk5 (2 μg) overnight at 4°C, followed by the addition of 30 μl of protein A-agarose (GE Healthcare Life Sciences, Uppsala, Sweden), for 2 h at 4°C. Supernatants (50 μg protein) were incubated with anti-Sox6 or anti-Cdk5 or anti-p35 in the presence of 10 μl of protein A-agarose. Immune complexes were extensively washed with lysis buffer and detected by Western blot analysis. NIH ImageJ software was used to analyze and quantify the bands in Western blots.

**Protein estimation and Western Blotting**

Protein estimation was performed according to BCA method. Western blotting was performed as previously described [23,24].
Immunocytochemical staining of primary cortical neurons

Staining of cortical neurons was performed as described earlier [25,26]. Cortical neurons fixed in 4% paraformaldehyde were permeabilized for 20 min in 0.2% Triton-X100, blocked in 10% horse serum in phosphate-buffered saline (PBS) for 1 h, and incubated with primary antibodies (diluted in 4% horse serum/phosphate buffered saline 0.05% Tween-20) against primary antibody for overnight at 4°C. After three washes in wash solution (phosphate buffered saline 0.05% Tween-20), the neurons were incubated with anti-rabbit (Alexa 488) and anti-mouse (Alexa 568) secondary antibodies (Molecular Probes, Eugene, OR, USA). Secondary antibodies were diluted in the same buffer as the primary antibodies and incubated for 2 h at room temperature. Cells were washed and mounted on the cover slips and analyzed using a TCS software program on an Axiovert 200 M microscope (Zeiss, Germany).

Immunohistochemical staining of developmental brain sections

Paraffin-embedded embryonic brain sections were prepared for immunostaining through xylene treatment and gradual rehydration with 95–75% ethanol. Sections were blocked (5% goat serum/0.1% Triton-X100) and then incubated with primary antibodies (Sox6 antibodies) overnight at 4°C in blocking solution. The sections were incubated with peroxidase conjugated secondary antibodies for 1 h at room temperature. The sections were stained using 3,3’-diaminobenzidine and counterstained with methyl blue, and slides were coverslipped using Permount (Biomedex, Foster City, CA, USA). Images were captured with a ×40 objective on a Nikon Eclipse E100 microscope using SPOT software (Nikon, Tokyo, Japan).

In vitro phosphorylation of Sox6 by Cdk5 and subsequent mass spectrometry analyses.

Culture (~1 liter) of the BL21 bacteria transformed with pGEX-4T3-Sox6 protein (GST-Sox6) was purified by using the Glutathione Sepharose GSTrap 4B columns (GE Healthcare Bioscience Corp.). The purified GST-Sox6 protein was mixed with 10 ng of active CDK5/p35 (Millipore Corp., Billerica, MA) in a buffer containing 20 mM HEPES (pH 7.4), 1 mM EDTA, 10 mM MgCl₂, 0.2 mM dithiothreitol, and a protease inhibitor cocktail (2 μl) in a final volume of 50 μl. The reaction was started by addition of 100 μM ATP and incubated at 30°C for 30 min. The samples were buffer exchanged into 25 mM NH₄HCO₃ by using 30K Nanosep Centrifugal Devices (Pall Corp., East Hills, NY, USA). They were then dried in the SpeedVac System (ThermoSavant Corp., Holbrook, NY, USA) and were taken up in 8 M urea/0.4 M NH₄HCO₃ for reduction (by dithiothreitol) and alkylation (by iodoacetamide) according to the standard protocol [27]. After diluting the samples into 2 M urea/0.1 M NH₄HCO₃ with water, half of each sample was digested with trypsin and half with chymotrypsin overnight at 37°C. Digested samples were then acidified with trifluoroacetic acid, combined, and cleaned up in the Oasis HLB media (Waters Corp., Milford, MA, USA). Eluted digests were dried in the SpeedVac system and were subjected to enrich the phosphorylated peptides by the TiO₂ chromatography, according to the method of Wu et al. [28]. Phosphopeptide-enriched samples were finally analyzed with liquid chromatography/tandem mass spectroscopy (MS/MS) on an LTQ XL mass spectrometer (Thermo Electron Corp., San Jose, CA, USA) where the instrument was set up to acquire a full survey scan followed by a collision-induced dissociation MS/MS spectrum on each of the top 10 most abundant ions in the survey scan. MS/MS Spectra were searched by using the SEQUEST program (Thermo Electron, Inc.) against a FASTA database, which consists of sequences of the pertinent protein and several decay protein to identify phosphopeptides and specific phosphorylation sites.

**Results**

**Sox6 is expressed in post mitotic neurons**

To examine if Sox6 is expressed in differentiating neurons, cortical neurons prepared from embryonic day 17 (E17) dissociated rat cortex were immunostained for TuJ1, a marker for differentiating neurons and Sox6. Cells expressing the neuron-specific marker TuJ1 (red) were also found to express Sox6 (green). The Sox6 protein was detected in the nucleus in the TuJ1 positive neurons, suggesting that Sox6 is expressed in differentiating neurons (Figure 1).

**Sox6 is developmentally regulated in brain**

To investigate the role of the Sox6 protein during mammalian central nervous system (CNS) development, we first characterized the expression of Sox6 protein in the rodent brain by immunoblotting and immunohistochemistry with the Sox6 antibody. Immunoblotting revealed that the Sox6 is detected in the embryonic brain and its expression gradually decreases during its development into an adult stage (Figure 2A and B). From the immunoblotting analysis, it appears that during development, Sox6 is degraded in the adult stage as Sox6 antibodies recognized a degradation product of the lower molecular weight. On the other hand, Cdk5 levels gradually increased during mouse brain
development. Sox6 could be phosphorylated and degraded by Cdk5 during development from E18 until adult (365 days, 1 year) (Figure 2A, middle panel). These results suggest that Sox6 and Cdk5 activity are inversely related. This also raises an intriguing possibility that Sox6 was phosphorylated and degraded by Cdk5 during development.

Immunohistochemistry analysis revealed that Sox6 expression was prominently detectable at embryonic day 12 (E12) in mitotic nuclei (Figure 2C-a (−ve control); Figure 2C-b (Sox6 antibody). Sox6 was expressed in the subventricular zone (svz) in migrating neurons and in the cortical plate in embryonic day 16.5, which indicates that Sox6 was primarily expressed in differentiating neurons in the cortical plate and in actively dividing neuronal precursor cells that populate the subventricular zone (Figure 2C-c). Sox6 was not detected in the adult brain (data not shown).

**Sox6 is a phosphoprotein**

Cdk5 is developmentally regulated and Cdk5 knock out (KO) mice undergo developmental arrest and death (14). Based on the finding that both Sox6 and Cdk5 are developmentally regulated, we examined whether Sox6 is regulated by Cdk5. The domain structure of the full length Sox6 protein revealed that Sox6 has potential phosphorylation sites for consensus Cdk5 signature motif S/TPXR/K/H towards the N-terminal domain (Figure 3A). The Cdk5 consensus motif in Sox6 S98PHK and T119PER is conserved among human, rat and mice (Figure 3B).

We first examined whether Sox6 was phosphorylated in vivo. The mouse brain lysate prepared from embryonic day 18 (E18) was subjected to calf intestinal alkaline phosphatase (CIAP) treatment. Western blotting was performed using Sox6 antibodies. Sox6 showed faster migration in the CIAP treated brain lysate, suggesting that Sox6 was phosphorylated in vivo (Figure 3C). Next, we immunoprecipitated Sox6 protein from the brain lysate from newborn mouse (PN0) and performed Western blot analysis with phospho-Ser antibody. Sox6 was detected by phospho-Thr and MPM2 (mitotic protein monoclonal 2, phospho-Thr/Pro) antibody (Figure 3D, upper panel). Likewise, Sox6 was detected by phospho-Thr and MPM2 antibody detects the proteins phosphorylated at the Ser/Thr/Tyr residues followed by proline. These results confirm that Sox6 is phosphorylated in vivo.

**Sox6 is in vitro substrate of Cdk5**

Because Sox6 has potential Cdk5 phosphorylation sites, we asked if Sox6 is phosphorylated by Cdk5. Sox6 immunoprecipitated from postnatal day 1 (PN1) rat brain lysate was subjected to in vitro phosphorylation with commercially obtained Cdk5/p35. Sox6 was phosphorylated by Cdk5/p35 (Figure 3E). The Sox6 phosphorylation by Cdk5/p35 was inhibited by the Cdk5 inhibitor roscovitine, suggesting specific phosphorylation of Sox6 by Cdk5 (Figure 3E). We then examined the phosphorylation of recombinant Sox6 by Cdk5/p35. We first made GST-Sox6 recombinant construct in pGEX vector, transfected it to express GST-Sox6 and purified it. Next, we determined whether Sox6 was directly phosphorylated by Cdk5/p35. GST-Sox6 was phosphorylated in vitro by Cdk5/p35 (Fig. 3F). Sox6 phosphorylated by Cdk5/p35 is subjected to mass spectrometry to determine the phosphorylation sites by Cdk5/p35. The recombinant Sox6 protein was phosphorylated by Cdk5/p35 (Figure 3G) and phosphorylation sites were identified by MALDI-TOF mass spectrometry. The potential sites identified by mass spectrometry correspond to proline directed Thr residue T119PER (Fig. 3G). The TPER motif is a consensus Cdk5 phosphorylation sequence. Figure 3G shows the MALDI MS spectra of Sox6 phosphorylation motif T*PER. These data suggest that Sox6 is a phospho-protein and Sox6 is phosphorylated by Cdk5.

**Sox6 associates with the Cdk5 and its Activator p35**

Sox6 was tested for its ability to co-immunoprecipitate with the Cdk5/p35 complex. Lysates were obtained from wild-type mouse postnatal brain day 0 (PN0) and were subjected to immunoprecipitation (IP) with Sox6 antisera, Cdk5 antisera, or control rabbit serum. Sox6 was associated with Cdk5 and p35 as revealed by the

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**Figure 2. Sox6 is developmentally regulated in brain.** A) Developmental expression of Sox6 (upper panel) and Cdk5 (middle panel) in embryonic brain (E18), postnatal days 0, 7, 14, 21, 60 and 365 days. The lower panel shows the β-actin staining as loading control. B) Densitometry analysis of Sox6 obtained from Figure 2A. C) Sections of mouse brain during embryonic development were fixed and immunostained with Sox6-specific antibodies from embryonic day 12, E12 (b) and E16 (c). Cb, Negative control shows the section of rat brain (E12) that were fixed and immunostained with Sox6 antibodies preabsorbed with Sox6 protein. Immunostaining with pre-immune serum as well as with no primary antibody showed no signal. Cb, The E12 rat brain section was fixed and immunostained with Sox6 specific antibodies. Sox6 staining is observed in the mitotic nuclei of E12 rat brain. Cc, Sox6 staining in the E16 brain. The neocortex-cortical plate and the subventricular zone-neuroepithelium (svz) are shown in E16. doi:10.1371/journal.pone.0089310.g002
IP of Sox6 and immunoblot analysis with anti-Cdk5 and anti-p35 antibodies (Figure 4A). Likewise, Sox6 was detected in the IP of Cdk5, but not control sera (Figure 4B). These results suggest that Sox6 was present in a complex with Cdk5 and p35.

Sox6 co-localizes with Cdk5 and its activator p35

To examine the cellular localization of Sox6 with Cdk5/p35, immunocytochemical assays were performed on dissociated E18 rat cortical neurons cultured for 6 days. A rhodamine-conjugated secondary antibody was used to show endogenous Cdk5 and p35 expression, and FITC conjugated secondary antibody was used for endogenous Sox6 (Fig. 4C). Figure 4C shows the co-localization of Sox6 (green) with Cdk5 (red). Sox6 was expressed in the nucleus whereas Cdk5 was mainly expressed in the cell bodies, axons and dendrites. We also observed the expression of Cdk5 in the nucleus. We noticed the co-localization of Sox6 with Cdk5 in the nucleus. Figure 4D shows the co-localization of Sox6 with p35. The p35 was prominently expressed in the nucleus, cell bodies and in the axons. In the overlay image, co-localization Sox6 can be seen in the nucleus with the Cdk5 and p35. These results suggest that endogenous Sox6 colocalizes with endogenous Cdk5 and p35 in the nucleus.

Cdk5 Regulates Sox6

In order to determine the effect of Cdk5 deficiency on Sox6 expression, we analyzed its expression in E17 Cdk5\(^{−/−}\) mice brain. We first examined the mRNA expression of Sox6 in WT and Cdk5\(^{−/−}\) brain by quantitative PCR analysis. The expression of Cdk5 was absent in Cdk5\(^{−/−}\) brain as compared to WT (Figure 5A). However, Sox6 mRNA levels remained the same in the WT and Cdk5\(^{−/−}\) brain (Figure 5B). We then examined the protein expression of Sox6 in Cdk5 WT and Cdk5\(^{−/−}\) brain lysate. The protein levels of Sox6 significantly was increased by 4-fold in Cdk5\(^{−/−}\) brains as compared to WT (Figure 5C and D). These results suggest that Sox6 expression is regulated by Cdk5.
Inhibition of Cdk5 activity by DN Cdk5 and roscovitine increases the Sox6 expression in primary cortical neurons

Next we wanted to ask if Sox6 expression is altered by the Cdk5 activity in cortical neurons. Towards this, we transfected the 5DIC neurons with Cdk5 and DN Cdk5 vectors (Figure 6A and B). Transfection of cortical neurons with WT Cdk5 reduced the endogenous Sox6 expression, whereas the transfection with DN Cdk5 increased the Sox6 expression (Figure 6A). The transfected Cdk5 (His-Cdk5) had a slightly lower mobility due to His tag. Endogenous Cdk5 was evident in all 3-lanes (lanes, 1–3) while the transfected Cdk5 (lane 2) and transfected DN Cdk5 (lane 3) showed two bands. The dominant negative Cdk5 has the same molecular weight as the WT Cdk5 except for a mutation at a single residue in the Cdk5 (D144N). The DN Cdk5 reduces the Cdk5 activity; however, Cdk5 levels remain the same, suggesting that Cdk5 activity regulate the steady state Sox6 expression. Furthermore, treatment of cortical neurons with Cdk5 inhibitor roscovitine resulted in the increase of Sox6 expression (Figure 6B). Both dominant negative Cdk5 as well as roscovitine reduced the Cdk5 activity suggesting that Cdk5 activity, regulate the steady state Sox6 expression.

Discussion

In the current study, we provide evidence that Sox6, a developmentally regulated key transcription factor, is a phosphorylation substrate of Cdk5/p35. The co-immunoprecipitation analysis reveals a high-affinity association between Sox6 and Cdk5/p35. More importantly, Sox6 expression is developmentally regulated; it is detected in mitotic nuclei in early embryonic (E12) cortex and in migrating neurons of sub ventricular zone (svz) and cortical plate in E16. In support of this finding, Sox6 is shown to be necessary for cortical interneuron migration [8]. Sox6 expression is abrogated in the adult mouse brain. Our finding that Sox6 expression is significantly high in Cdk5−/− brain suggests that mis-expression of Sox6 is regulated by Cdk5.

A number of cytoplasmic proteins have been identified as substrates of Cdk5. Cdk5 plays a multifunctional role in neurons including neurite outgrowth, cytoskeleton assembly, and synaptic transmission by phosphorylating these diverse cytoplasmic targets [12,29]. Deregulation of Cdk5 activity mediates the hyperphosphorylation of the major cytoplasmic component of fibrillary tangle, tau, triggered by Aβ and aberrant NF-M/H phosphorylation, suggesting that Cdk5 is involved in neuronal apoptosis via a cytoplasmic mechanism [30,31].

On the other hand, Sox6 is a transcription factor present in the nucleus. Genome-wide association studies suggest that Sox6 influences both obesity and osteoporosis [32]. Sox6 is an important enhancer of definitive erythropoiesis [33]. Pertaining to the brain, Sox6 is essential for cortical interneuron development. Sox6 is shown to be important for normal positioning and maturation of MGE-derived interneurons, and that the specific deletion of Sox6
results in epileptic encephalopathy [8]. Recently, it was found that Sox6 is expressed and required in MGE-derived cortical interneurons, and additionally it plays an independent role in pallial/subpallial patterning [34]. Recent work by the same group has also identified Sox5, a close homolog of Sox6, as being required for the specification of deep layer pyramidal neurons in layer V and VI of the cortex [3]. miRNA 219 and miR-338 have been shown to inhibit Sox6 and block oligodendrocyte differentiation [36]. Sox6-deficient mice are anemic due to impaired red cell maturation and show inappropriate globin gene expression in definitive erythrocytes.

In contrast to classic Cdks whose activity is regulated during cell cycle by cyclins, Cdk5 is inactive in cell cycle [29], instead, its activation requires association of Cdk5 with its neuron-enriched regulators p35 or p39 [12]. In vivo Cdk5 activity is tightly regulated. Complete lack of Cdk5 is clearly destructive to central nervous system, with deletion of either p35 or Cdk5 resulting in profound neurological defects [37,38] and Cdk5−/− mice die in utero. Cdk5 hyperactivity is toxic to cells, particularly neurons, leading to neuronal apoptosis under either physiological or...
pathological conditions [39,40]. Neurototoxicity deregulates Cdk5 activity either through generating a more stable proteolytic product of p35, p25, or by stabilizing Cdk5/p25 complex. Transgenic p25 mice display neuronal loss and cognitive impairment [41]. Furthermore, Cdk5/p25 deregulates HDAC1 during neurotoxicity [20].

Cdk5 has been proposed to play a key role in neurodegenerative diseases primarily through a cytoplastic mechanism. Both Cdk5 and p35 are enriched in the cytoplasm of neurons where Cdk5 phosphorylates various substrates including tau, a major component of neurofibrillary tangles associated with neurodegeneration [31]. Interestingly, several studies including this, show that Cdk5 and p35 are localized in the nucleus [42]. Sox6 appears to have multiple functions in the brain. Sox6 is expressed more in an early embryonic stage and gradually reduces during the development in to an adult. In conclusion, this study highlights the role of Cdk5 in nucleus regulating developmental transcriptional factor Sox6.

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Author Contributions
Conceived and designed the experiments: PR. Performed the experiments: PR EU HJ. Analyzed the data: PR EU. Contributed reagents/materials/analysis tools: AK. Wrote the paper: PR.

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