Neuronal Cyclin-Dependent Kinase 5 Activity Is Critical for Survival

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Cyclin-dependent kinase 5 (Cdk5) null mice exhibit a unique phenotype characterized by perinatal mortality, disrupted cerebral cortical layering attributable to abnormal neuronal migration, lack of cerebellar foliation, and chromatolytic changes of neurons in the brainstem and the spinal cord. Because Cdk5 is expressed in both neurons and astrocytes, it has been unclear whether this phenotype is primarily attributable to defects in neurons or in astrocytes. Herein we report reconstitution of Cdk5 expression in neurons in Cdk5 null mice and its effect on the null phenotype. Unlike the Cdk5 null mice, the reconstituted Cdk5 null mice that express the Cdk5 transgene under the p35 promoter (TgKO mice) were viable and fertile. Because Cdk5 expression is mainly limited to neurons in these mice and rescues the defects in the nervous system of the Cdk5 null phenotype, it clearly demonstrates that Cdk5 activity is necessary for normal development and survival of p35-expressing neurons.

Key words: Cdk5; cerebrum; cerebellum; neuron; astrocyte; phosphorylation; neurodegeneration; transgenic mice

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Cyclin-dependent kinase 5 (Cdk5) is a member of the Cdk family of serine/threonine kinases and is so named because of its sequence homology to other Cdns (Hellmich et al., 1992; Lew et al., 1992; Meyerson et al., 1992). Unlike other Cdns that are involved in cell cycle control, Cdk5 is mainly involved in phosphorylation of target proteins in postmitotic neurons (Shetty et al., 1993). It phosphorylates cytoskeletal components such as high-molecular weight neurofilament protein (NF-H) and microtubule-associated proteins (MAP) tau and MAP1B (Mandelkow et al., 1992; Koba-yashi et al., 1993) and is implicated in regulation of neuronal migration, neurite outgrowth (Nikolic et al., 1996), and axon patterning (Connell-Crowley et al., 2000).

The activity of Cdk5 is regulated in two ways, by its binding with neuron-specific activator proteins p35, p25, and p39, and by phosphorylation (Sharma et al., 1999; Zukerberg et al., 2000). The activator proteins p35 and p39 are noncyclin proteins (Lew et al., 1994; Tsai et al., 1994; Tang et al., 1995), with p25 being a proteolyzed fragment of p35 (Lew et al., 1994). Cdk5 is ubiquitously expressed and is most abundant in the nervous system (Hellmich et al., 1992; Tsai et al., 1993), with p25 being a proteolyzed fragment of p35 (Lew et al., 1994). Cdk5 is ubiquitously expressed and is most abundant in the nervous system (Hellmich et al., 1992; Tsai et al., 1993), with p25 being a proteolyzed fragment of p35 (Lew et al., 1994). Cdk5 is ubiquitously expressed and is most abundant in the nervous system (Hellmich et al., 1992; Tsai et al., 1993), with p25 being a proteolyzed fragment of p35 (Lew et al., 1994). Cdk5 is ubiquitously expressed and is most abundant in the nervous system (Hellmich et al., 1992; Tsai et al., 1993), with p25 being a proteolyzed fragment of p35 (Lew et al., 1994). Cdk5 is ubiquitously expressed and is most abundant in the nervous system (Hellmich et al., 1992; Tsai et al., 1993), with p25 being a proteolyzed fragment of p35 (Lew et al., 1994).
Expression and purification of the recombinant p35. Glutathione S-transferase (GST)-p35 and GST-p25 in pGEX4T-2 were constructed by a PCR method using the oligonucleotides 5'-GAGATCCATGGG-CACG-GTCTCTG-3' (p35) and 5'-GTCCGGATCCCGCCCCCGCGG-3' (p25) as forward primers, 5'-GTATGAAATCTGGATCACG-GTCTG-3' as a reverse primer, and DNA from 6X His-tagged p35 construct (a gift from L. H. Tsal, Harvard School of Medicine, Boston, MA) as a template. The PCR products were digested with BamHI and EcoRI, and the resulting fragments were cloned into BamHI-EcoRI cut pGEX 4T-2. GST-p35 and GST-p25 fusion proteins were expressed and purified as described previously (Pant et al., 1997).

Preparation of the tissue extracts. The cerebrum, cerebellum, spinal cord, heart, liver, kidney, testis, and muscle from the age-matched wild-type (WT) and the transgenic mice were removed surgically, frozen in dry ice, and stored at −80°C. Frozen tissues were further processed as described previously to obtain protein extracts, except for the addition of microcystine LR 2 μM to the homogenization buffer used in these extractions (Veerrana et al., 1996). Protein estimation was performed using the BCA method as described by the manufacturer (Pierce, Rockford, IL).

Electrophoresis and Western blot analysis. Ten to 15 μg of protein was loaded per lane for SDS-PAGE using Novex (San Diego, CA) 10–20% gradient gels. Electrophoretose and immunoblotting were performed as described previously (Shetty et al., 1995) using rabbit polyclonal anti-Cdk5 C-terminal antibody C-8 (Santa Cruz Biotechnology, Santa Cruz, CA) and rabbit polyclonal anti-p35/C-terminal antibody C-19 (Santa Cruz Biotechnology). The slices were developed by the ECL method as described by the manufacturer (Amersham Pharmacia Bio-tech, Piscataway, NJ). Alternatively membranes were developed using alkaline phosphatase (AP)-based 5-brom-4-chlor-indolyl-phosphate/nitroblue-tetrazolium-chloride (BCIP/NBT) single reagent (Kirkegaard & Perry Laboratories, Gaithersburg, MD).

Immunoprecipitation and kinase assays. Immunoprecipitation of Cdk5 and the kinase assays using the immunoprecipitates from the tissues of TgCd5, TgKO, and wild-type mice were performed as described previously using C-8 rabbit polyclonal antibody, which specifically reacts with Cdk5 protein (Veerrana et al., 1996; Pant et al., 1997). Evaluation of the kinase activity in the immunoprecipitates obtained from the Cdk5 overexpression mice and the corresponding controls were performed by the addition of 7.5 μg of bacterially expressed GST-p35 or GST-p25 activator fusion proteins (Pant et al., 1997). The expressed proteins used in these experiments were partially purified, soluble, and active when tested with recombinant Cdk5 (N. D. Amin and H. Pant, unpublished observations).

Tissue preparation for in situ hybridization, histology, and immunohistochemistry. A minimum of three transgenic and two wild-type mice were used at each time point, for at least 3 months. For paraffin sections of the mouse brains, adult mice were perfused transcardially with 0.1 M PBS, pH 7.4 and 4% paraformaldehyde (PFA) in PBS. The brain and spinal cord were removed and immersion-fixed in 4% PFA in PBS for 24 hr at 4°C. The embryos were removed by hysterotomy from their dams, decapitated, and immersion-fixed in 4% PFA in PBS for 24 hr at 4°C. After fixation, the tissues were dehydrated and embedded in paraffin. Six micrometer paraffin sections were cut and stained with hematoxylin and eosin and Nissl stains by standard methods. For thionine staining, two sets of TgKO mice and one set of wild-type control mouse at the ages of 2, 3, 5, and 7 months were perfused transcardially with wash solution (0.8% NaCl, 0.4% dextrose, 0.8% sucrose, 0.023% CaCl2, and 0.034% sodium cacodylate). The mice were then transcardially perfused with fixative (4% PFA, 4% dextrose, and 1.4% sodium cacodylate) and immersed in the same fixative for 3–5 d. Frozen cryostat coronal serial sections with slice thickness of 30 μm covering the entire brain were made for each brain and stained with thionine.

In situ hybridization. Digoxigenin (DIG)-labeled antisense and sense riboprobes were generated by in vitro transcription of pbSSK plasmid DNA containing a Cdk5 cDNA insert using T7 and T3 RNA polymerase to generate cRNA probes. Sections were deparaffinized and hydrated through a series of ethanol grades and incubated in xylene. Sections were washed in Tris-buffered saline I (TBS I) 100 mM Tris Cl and 150 mM NaCl, pH 7.5), the sections were treated with 10 μg/ml proteinase K in 50 mM Tris Cl, pH 7.5, at 37°C for 30 min and then washed in TBS I for three times for 5 min each. The slides were immersed in 0.1 M triethanolamine buffer.
with 0.25% acetic anhydride at room temperature for 10 min to prevent nonspecific probe binding and washed in 2× SSC for 5 min twice. Prehybridization was done by applying 100 μl of prehybridization solution (50% formamide, 4× SSC, 0.1% SDS, 1× Denhardt’s solution, and 400 μg/ml denatured ssDNA) for 1 hr at 45°C, and subsequently sections were hybridized with 30 μl of prehybridization solution with 5–10 ng/μl of either sense or antisense probes. The slides were heated at 65°C for 5 min to denature the target, and hybridization was performed at 45°C for 18 hr in a moist chamber. After three 10 min washes in 2× SSC to remove excess probe, the sections were treated with 100 μl of 40 μg/ml RNome A in 500 mM NaCl and 1 mM EDTA, pH 8.0, at 37°C for 30 min to remove unhybridized probe. Stringent washes were performed with 2× SSC for 30 min at 50°C followed by 0.2× SSC for 30 min at 60°C. After a wash in TBS I, sections were blocked for 30 min with 5% normal goat serum in 500 mM NaCl and 1 mM EDTA, pH 8.0, at 37°C for 30 min to remove unhybridized probe. Stringent washes were performed with three washes in TBS-T, sections were immersed in TBS II (100 mM Tris-Cl and 150 mM NaCl, pH 9.5) for 5 min at room temperature to remove excess probe, the sections were treated with 100 μl of anti-DIG-AP conjugate (Roche Molecular Biochemical, Indianapolis, IN) solution (1.5 U/ml in TBS I) was applied for 1 hr. After three washes in TBS-T, sections were immersed in TBS II (100 mM Tris-Cl and 150 mM NaCl, pH 9.5) for 5 min at room temperature to activate the alkaline phosphatase, and colorimetric detection was performed with BCP/NBT (Roche Molecular Biochemicals) at room temperature for 3 hr. The slides were washed in a stream of distilled water and mounted (Kadkol et al., 1999).

**Immunofluorescent staining of cells.** Mixed cultures of neurons and astrocytes were prepared from embryonic day 17.5 (E17.5) embryos as described previously (Vicario-Abejon et al., 1998). Cells were fixed with 4% PFA for 15 min at room temperature. After blocking for 30 min in PBS with 1% Triton X-100 (PBST) and 5% normal goat serum, cultures were incubated with the primary antibody [monoclonal anti-Cdk5 (Covance, Princeton, NJ), monoclonal anti-GFAP (ICN, Costa Mesa, CA), and polyclonal anti-Cdk5 (Santa Cruz Biotechnology)] for 4 hr at room temperature. After two washes in PBST, the secondary antibody [rhodamine-coupled-anti-mouse (Jackson ImmunoResearch, West Grove, PA) and biotinylated anti-rabbit (Vector Laboratories, Burlingame, CA)] was applied at a 100-fold dilution at room temperature. One hour later, cultures were washed with PBST and incubated with avidin-coupled FITC (Vector Laboratories) diluted 50-fold in PBS. Cells were mounted in 70% glycerol in PBS with 2% 1,4 diazabicyclo-(2,2,2)-octane (Sigma, St. Louis, MO) after two washes in PBS.

**Immunohistochemistry.** The following primary antibodies were used in the immunohistochemical study. Rabbit polyclonal anti-Cdk5 C-terminal antibody (C-8) was used at a dilution of 1:100. Rabbit polyclonal anti-cow GFAP antibody (Dako, Carpentaria, CA) was used at a dilution of 1:500. Mouse monoclonal antibody SM-31 (Sternbergh Monoclonals, Lutherville, MD) was used at a dilution of 1:1000. After deparaffinization, the slides were washed in PBS for 5 min twice and immersed in 90% methanol containing 0.3% H2O2, for 30 min. After washing in distilled water, the slides were incubated in PBST for 5 min. The slides were incubated with blocking solution (normal goat serum diluted with PBS for anti-rabbit antibodies and normal horse serum diluted with PBS for anti-mouse antibodies) (Vectastain Elite ABC kit; Vector Laboratories) for 30 min and then incubated in primary antibody diluted with the blocking solution at 4°C overnight. After three washes in PBST, they were incubated in diluted biotinylated secondary antibody solution (PBS containing normal goat serum and anti-rabbit-IgG antibody or PBS containing normal horse serum and anti-mouse-IgG antibody) (Vectastain Elite ABC kit) for 30 min. Subsequent color development with the ABC and DAB reagents (Vector Laboratories) was performed according to the instructions of the manufacturer.

**RESULTS**

**Generation and analysis of transgenic mice overexpressing Cdk5 driven by p35 promoter (TgCdk5 mice)**

We first confirmed the genotypes of mice generated by the F2 cross. Figure 1B shows the genotype analysis of tail DNA from the WT and TgCdk5 mice by Southern blot analysis, in which the entire transgene was used as a probe. Additional bands derived from the transgene as well as the band derived from endogenous Cdk5 were revealed in the TgCdk5 mouse. Expression of mRNA and protein in the brain was confirmed by Northern and immunoblot analysis, respectively. There was a twofold to fourfold overexpression of Cdk5 mRNA and protein in the brains of TgCdk5 mice over WT mice (Fig. 1C,D). The overexpression of Cdk5 in TgCdk5 did not influence the expression levels of p35 and p25, as shown in Figure 1D.

**The Cdk5 transgene is functional**

The level of Cdk5 kinase activity was determined in immunoprecipitates of Cdk5 from TgCdk5 and WT mouse brains. An unexpected reduction in Cdk5 kinase activity was observed in Cdk5 immunoprecipitates from TgCdk5 mice overexpressing Cdk5 (Fig. 2A). To rule out a functionally inactive Cdk5 transgene product as a cause for the reduction in kinase activity in TgCdk5 mice, the in vitro kinase assays were performed after the addition of the bacterially expressed p35 protein to the kinase assay mixture containing Cdk5 immunoprecipitates from TgCdk5 and WT mice. There was a marked increase in Cdk5 activity upon addition of exogenous p35 to the kinase assay mixtures derived from both the TgCdk5 and WT mice, indicating that the product of the Cdk5 transgene was functional (Fig. 2B,C). Addition of bacterially expressed p35 to the kinase assay mixture of WT mouse showed a 2.5-fold increase in kinase activity over the corresponding control without addition of exogenous p35 (Fig. 2B). A similar experiment performed using an immunoprecipitate from the TgCdk5 mouse revealed a 47-fold enhancement in the kinase activity upon addition of exogenous p35 (Fig. 2C). Thus, exogenous addition of p35 enhances Cdk5 activity in vitro, indicating that the transgenic Cdk5 protein is functional. Experiments performed by the exogenous addition of p25, an activator of Cdk5 derived from proteolytic cleavage of p35, yielded similar results (data not shown).

**Generation and analysis of transgenic mice in which Cdk5 is expressed only in tissues that express p35 (TgKO mice)**

To determine whether expression of Cdk5 in p35-expressing areas was sufficient for reversing lethality of Cdk5 null mice, we
crossed the TgCdk5 line to the Cdk5 knock-out line to generate mice lacking endogenous Cdk5 but with the Cdk5 transgene under the control of the p35 promoter. Of the F1 generation, the mice with a Cdk5 \(1/2\) and transgene \(1/2\) genotype were identified by Southern blot analysis. These mice were intercrossed and, of the F2 generation, the mice with the genotype of endogenous Cdk5 \(2/2\) and transgene \(1/1\) (TgKO mice) were identified by Southern blot analysis. Figure 3A shows additional bands derived from the transgene and the mutant Cdk5 allele, and lack of endogenous Cdk5-derived band in TgKO mice. The TgKO mice were born with normal sex ratio. There was no difference in the number of the live newborn mice from that of the wild-type mice. They did not exhibit any developmental abnormalities. Analysis of mRNA from the TgKO mice showed a marked increase of Cdk5 in the cerebrum, cerebellum, spinal cord (Fig. 3B), and testis (data not shown). The heart, liver, kidney, ovary, and muscle from TgKO mice showed trace amounts of Cdk5 mRNA (data not shown). These findings are compatible with the fact that, in the adult mice, p35 is expressed selectively in neurons (Lew et al., 1994; Tsai et al., 1994; Matsushita et al., 1996; Tomizawa et al., 1996; Delalle et al., 1997; Zheng et al., 1998) and testis (T. Tanaka and A. B. Kulkarni, unpublished observations) and at extremely low levels in other tissues. Cdk5 protein was seen in regions in which p35 is normally expressed, the heart showed moderate expression, and the testis revealed a high level of Cdk5 protein expression (data not shown). Cdk5 mRNA levels corresponded to Cdk5 protein levels in all the organs analyzed except the heart, in which moderate levels of Cdk5 protein were present despite low Cdk5 mRNA levels, which is suggestive of either faster mRNA degradation or slower protein degradation. Although Cdk5 protein levels were higher in TgKO compared with WT mice (Fig. 3C), the kinase activity was \(40-60\%\) lower in TgKO compared with WT in cerebrum, cerebellum, and spinal cord (Fig. 3D). No activity was detectable in the other tissues analyzed, including liver, heart, kidney, and muscle. However, a negligible level of activity was observed in the testis (data not shown).

**Spatial expression pattern of Cdk5 in the TgKO mouse brain**

Because Cdk5 expression is under the control of a p35 promoter in TgKO mice, Cdk5 expression is expected only in p35-expressing regions. Therefore, we performed in situ hybridization using 5-month-old TgKO mice and corresponding WT mice to analyze whether the pattern of Cdk5 transgene expression in TgKO mice was similar to that of the wild-type p35 expression pattern. Figure 4 shows the results of in situ hybridization with Cdk5 riboprobes. The antisense Cdk5
pressed in astrocytes derived from WT mice (Fig. 5). However, the level of Cdk5 expression was higher in the neurons derived from TgKO mice compared with the neurons derived from the WT mice (Fig. 5, compare A, G).

Morphological analysis of brain development in TgKO mice

When the brains of Cdk5 null mice are analyzed perinatally, they reveal abnormal cortical layering, lack of cerebellar foliation, and ballooning of neurons in brainstem and spinal cord (Ohshima et al., 1996b). We analyzed the brains of E18.5 embryos from TgKO mice and WT mice. In contrast to the Cdk5 null embryos (Fig. 6C,F), age-matched TgKO mice that express Cdk5 in p35-expressing regions show normal brain morphology with normal cortical layering and cerebellar foliation (Fig. 6A,D). There were no macroscopic differences in the brain between age-matched TgKO and WT adult mice. Nissl-stained coronal sections of the cerebral cortex from TgKO adult mouse showed a well-defined cortical layering pattern (Fig. 6G), which was similar to that of WT mouse (Fig. 6H). The hippocampal formation of the TgKO adult mouse was well organized, and the pyramidal cell layer and granule cell layer were clearly defined (data not shown). The cerebellum from the TgKO adult mouse had normal foliation (Fig. 6I), which was identical to that of WT (Fig. 6J). The Purkinje cell layer, molecular layer, and granular layer of the cerebellum from the TgKO mouse was clearly defined and well organized. Thionine-stained coronal serial frozen sections obtained from two sets of TgKO mice and one set of WT mouse at the ages of 2, 3, 5, and 7 months were examined. Coronal serial sections from each of the above aged brains revealed no abnormalities in cerebral and cerebellar cortical layering pattern in TgKO mouse. By immunohistochemical analysis, cerebral cortical neurons, cerebellar Purkinje cells, and granule cells expressed Cdk5 when analyzed with anti-Cdk5 antibody (data not shown), which was identical to that of WT mice. Unlike Cdk5 null mice, TgKO mice did not display ballooned neurons in spinal cord and brainstem (data not shown). Thus, all of these data indicate that the TgKO mice are similar to the WT mice in morphological parameters analyzed in the CNS.

Aberrant phosphorylation of the cytoskeleton in Cdk5 null mice is corrected in TgKO mice

Cdk5 is involved in cytoskeletal protein phosphorylation and may lead to NF-tangle formation, a critical change associated with neurodegenerative diseases (Mandelkow et al., 1992; Baumann et al., 1993; Nakamura et al., 1997; Julien and Mushynski, 1998; Bajaj et al., 1999). A comparative study of the phosphorylation status of cytoskeletal elements shown previously to be phosphorylated by Cdk5 was performed with SM1-31 antibody. SM1-31 specifically reacts with phospho-epitopes of high-molecular weight NF, MAP, and tau proteins. Cdk5 null mice showed densely stained neuronal cell bodies in the brainstem, indicating a hyperphosphorylated status of the cytoskeleton resembling that seen in neurodegenerative disorders (Ohshima et al., 1996b). In contrast, TgKO mice (Fig. 7A) did not reveal such changes but exhibited axonal staining patterns similar to the WT (Fig. 7B). Thus, restoration of Cdk5 expression corrected the aberrant phosphorylation of cytoskeletal elements in the soma of brainstem neurons in TgKO mice.

DISCUSSION

Targeted disruption of Cdk5 in mice leads to embryonic lethality and associated defects, such as abnormal cortical layer-
ing, lack of cerebellar foliation, and ballooning of neurons in brainstem and spinal cord (Ohshima et al., 1996b). Ballooning of neurons may result from hyperphosphorylation of cytoskeletal proteins in cell bodies, reminiscent of neurodegenerative disorders. Although neuronal death in vital brainstem regions may be responsible for lethality in Cdk5 null mice, the absence of Cdk5 and its activity in other tissues such as muscle and heart may also contribute to embryonic lethality. In this study, we sought to determine whether limited re-expression of Cdk5 in the nervous system was sufficient to reverse the abnormalities and lethality. Because expression of p35 is predominantly restricted to the nervous system, transgenic mice were gener-

Figure 5. Mixed neuronal–astrocytic cultures were prepared from E17.5 WT and TgKO mice embryos as described by Vicario-Abejon et al. (1998), with minor modifications. After fixation with 4% PFA, double immunocytochemistry was performed on the cultures with either GFAP (an astrocytic marker) and Cdk5 or TuJ1 (a neuronal marker) and Cdk5. Panels on the left (FITC, green) show the staining pattern of Cdk5 with the corresponding GFAP or TuJ1 (rhodamine, red) staining to the right. WT mice show Cdk5 staining in neurons as well as astrocytes (A, B, E, F), whereas there is no Cdk5 seen in the astrocytes of TgKO mice (C, D, G, H). In addition, the levels of transgenic Cdk5 in TgKO neurons is higher than that seen in neurons derived from WT mice (compare E, G). Scale bar (in B), 100 μm.
ated in which a Cdk5 transgene was expressed under the control of a p35 promoter.

**Cdk5 kinase activity in the brain from the TgCdk5 mice is lower than that of WT mice**

Cdk5 activity was analyzed in whole-brain lysates from TgCdk5 mice that overexpress Cdk5. Unexpectedly, TgCdk5 lysates showed a markedly lower Cdk5 kinase activity compared with wild-type controls. Northern and Western blot analysis showed robust expression of Cdk5 mRNA and protein, ruling out lack of expression as a cause of decreased Cdk5 activity. To determine whether a relative deficiency of activator protein p35 caused the decreased Cdk5 activity, we added bacterially expressed p35 to the kinase assay mixture. Addition of p35 augmented Cdk5 activity in TgCdk5 mice by 47-fold compared with a 2.5-fold enhancement in wild-type mice. Hence, the lower level of Cdk5 activity in TgCdk5 mice is indeed attributable to relative deficiency of p35. The other possibility is that the higher level of transgenic Cdk5 protein may interfere with the transcription of p35 and reduce its production. This could lead to a decreased level of Cdk5 activity observed in TgCdk5 mice compared with the WT. Indeed, this is not true, because the Western blot analysis of p35 in TgCdk5 mice revealed no difference in the p35 protein levels compared with the WT mice. Together, our results confirm the active nature of the Cdk5 protein in TgCdk5 mice. These initial results were the basis for the generation of TgKO mice lacking endogenous Cdk5 but expressing transgenic Cdk5 under the control of p35 promoter.
Cdk5 is restricted to p35 regions in TgKO mice

Restricted expression of Cdk5 in the regions in which p35 is expressed helped us to understand the importance of Cdk5 in peripheral areas that do not express p35. Because astrocytes lack p35 (Tsai et al., 1994; Honjyo et al., 1999), our observation that astrocytes from TgKO mice lack Cdk5 is consistent with the expected results. TgKO mice clearly showed the Cdk5 mRNA transcript in the nervous system and testis. The Northern blot analysis also showed an additional band despite high-stringency wash conditions indicative of the fact that the Cdk5 cDNA probe hybridized with a closely related sequence. This transcript might represent an alternatively spliced Cdk5 or a transcript closely related to Cdk5. Analysis of Cdk5 kinase activity in cerebrum, cerebellum, and spinal cord of TgKO mice revealed a 60% reduction in kinase activity compared with the WT controls. This reduction in kinase activity is similar to that of TgCdk5 mice that show a decreased kinase activity, probably because of an autoinhibitory phenomenon caused by excess concentration of the Cdk5, competing for a limited amount of the activator protein p35. The absence of Cdk5 kinase activity in the peripheral tissues analyzed such as heart, in which Cdk5 is expressed, could be attributable to the absence of the activator p35. The low level of Cdk5 activity observed in the testis despite the presence of abundant quantity of both the Cdk5 and p35 proteins in TgKO mice strongly argues for the presence of an unknown inhibitor of Cdk5.

Cdk5 is a key molecule critical for survival and neuronal development

Cdk5 null mice are embryonic lethal, but p35 null mice survive with abnormal brain development (Chae et al., 1997; Kwon and Tsai, 1998). p35 and other activators such as p39, an isoform of p35 encoded on a separate gene, influence Cdk5 activity (Tang et al., 1995; Zheng et al., 1998). The lack of embryonic lethality of the p35 null mice suggests that p35 is not essential for embryonic survival, although p35 null mice still display neuronal migration defects. This is possibly attributable to the redundancy in the activator system, such as p39 expression. The level of Cdk5 kinase activity in the cerebral cortex and cerebellum of the p35 null mice is ~5 and 20%, respectively, compared with the WT mice (Ohshima and Kulkarni, unpublished observations). This difference in the level of Cdk5 kinase activity in cerebrum and cerebellum is consistent with the level of p39 in the adult mice (Zheng et al., 1998). This might represent a minimal level of Cdk5 activity in the brain that may be necessary for survival but yet might be insufficient for normal neuronal migration because migration defects are seen in p35 null mice. The survival of p35 null mice also suggests that p39 by itself is capable of supporting the survival and may not be involved in neuronal migration defects observed in the p35 null mice. It is possible that disruption of both activators p35 and p39 would lead to lethality and phenotype similar to Cdk5 null mice because it may result in total loss of kinase activity. Our results with TgKO mice clearly support the indispensability of Cdk5 for survival. Our results also make it clear that the expression of Cdk5 in p35-expressing regions is sufficient for neuronal development and survival.

Cdk5 outside of the p35-expressing regions is not critical for survival

TgKO mice do not express Cdk5 in the liver, kidney, and ovary, among the organs analyzed, whereas WT mice have Cdk5 expression in such regions. On the other hand, the astrocytes in TgKO mice do not express Cdk5, whereas astrocytes from WT mice do express Cdk5. This suggests that neuronal expression of Cdk5 is sufficient for survival. Cdk5 outside of the p35-expressing regions does not seem to play a critical role for survival, although it may have more local actions in peripheral tissues. Although Cdk5 is expressed abundantly in the testis, the fact that the castrated animals survive rules out possible importance of its role in testis for survival. The heart of the Cdk5 null mice is normal. However, the functional significance of Cdk5 expression in the heart is yet unknown. Thus, these data indicate that Cdk5 expression in other tissues is either dispensable or has redundant functions with other Cdk5s.

Cdk5 plays a key role in modulation of cellular signals

Although a clear pathway delineating the upstream and downstream effectors of Cdk5 is yet to be uncovered, mounting evidence suggests that Cdk5 is a key molecule in mediating important signals involved in developmental and functional regulation of neurons. Integrins augment p35 expression, leading to enhanced Cdk5 activity facilitating neurite outgrowth (Pigino et al., 1997; Paglini et al., 1998). Brain-derived neurotrophic factor enhances Cdk5 activity leading to enhanced NF-H phosphorylation during synapse formation in cortical neurons (Tokuoka et al., 2000). Calcium-activated protease calpain cleaves p35 into p25 leading to augmented Cdk5 activity. This phenomenon has been implicated in apoptosis of neurons in culture (Kusakawa et al., 2000). Cdk5 is also reported to be involved in signaling in dopaminergic neurons, in which it phosphorylates DARPP-32 (dopamine and cAMP-regulated phospho-protein; relative molecular mass of 32,000) leading to an inhibition of PKA activity (Bibb et al., 1999). Interestingly, Cdk5 also downregulates the activity of protein phosphatase-I by phosphorylating inhibitor-I at serine 67 leading to its activation (Huang and Paudel, 2000). Cadherin-mediated cell adhesion is influenced by Cdk5 (Kwon et al., 2000), because the aggregation of cortical neurons is higher in the p35 null mice compared with the WT mice because of the absence of Cdk5–p35–β catenin complex. In muscle, Cdk5 is required for the activity of transcription factors responsible for the synthesis of muscle-specific proteins. Transfection of a dominant negative Cdk5 in myoblasts leads to failure of their development into myotubes.
myotubules (Lazarro et al., 1997). Together with our findings, Cdk5 plays a central role in mediating important physiological signals and it is irreplaceable in neurons.

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